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CENTERS FOR DISEASE CONTROL AND PREVENTION NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

convenes

MEETING 54

ADVISORY BOARD ON RADIATION AND WORKER HEALTH

DAY TWO

The verbatim transcript of the 54th

Meeting of the Advisory Board on Radiation and

Worker Health held at the Crowne Plaza Tampa East,

Tampa, Florida on Apr. 8, 2008.

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TRANSCRIPT LEGEND

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- -- "*" denotes a spelling based on phonetics, without reference available.
- -- (inaudible) / (unintelligible) signifies speaker failure, usually failure to use a microphone.

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PROCEEDINGS

(8:45 a.m.)

WELCOME AND OPENING COMMENTS

DR. PAUL ZIEMER, CHAIR

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DR. CHRISTINE BRANCHE, DFO

DR. BRANCHE: Good morning. We're starting the second day of the Advisory Board on Radiation and Worker Health meeting 54. I'm Christine Branche. I have the pleasure of being your Designated Federal Official, and we are beginning today. I need to make certain -- Mr. Robert Presley, are you on the line? MR. PRESLEY: Right here. DR. BRANCHE: Do you have the same telephone number that I gave you yesterday in case you somehow lose contact? MR. PRESLEY: 813/623-6363. DR. BRANCHE: Great, and we're still in the Cypress Room. Thank you so much; glad you're aboard. DR. ZIEMER: Thank you very much. I want to remind everyone again to please register your attendance with us this morning.

registration books are in the corridor.

again, the copies of the agenda and other materials are on the table in the back of the room.

On today's agenda I just want to point out that the third item, which is really the second business item, the NIOSH program update, we -we moved forward and covered yesterday. will likely move something else up, probably the -- if -- if it's okay with Jeff, and you checked with him, from Department of Labor, we'll probably move the Labor update forward to this morning so that we utilize the time effectively.

AREA IV, SANTA SUSANA FIELD LABORATORY SEC PETITION

But we'll begin this morning with the Santa Susana Field Laboratory SEC petition. presentation on their evaluation report will be given by Stuart Hinnefeld, and then we'll have an opportunity for the petitioners. just check and see if LaVonne Klea is on the line.

MS. KLEA: Yes, I -- yes.

DR. ZIEMER: Good morning.

MS. KLEA: Good morning.

DR. ZIEMER: And after the presentation by Mr.

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1 Hinnefeld, we'll have an opportunity for you, 2 LaVonne, also to make comment. 3 MS. KLEA: Thank you. DR. ZIEMER: Thank you. 5 DR. BRANCHE: While Mr. Hinnefeld makes his way 6 forward, for those of you who are on the 7 telephone, if you would please mute your 8 phones. If you do not have a mute button, then 9 please use star-6. And when -- for example, 10 Ms. Klea, when you're ready to speak, you can 11 use that same star-6 to unmute your phones. 12 Again, please mute your phones while the 13 discussion here at the board room is going on. 14 Thank you. 15 **UNIDENTIFIED:** Okay. Star-6? 16 DR. BRANCHE: That's right. 17 DR. ZIEMER: Yes. Stuart Hinnefeld. 18 MR. HINNEFELD: Thank you, Dr. Ziemer. 19 morning, everyone. I'm here today to present 20 the results of the evaluation report on Area IV 21 of the Santa Susana Field Laboratory, which is 22 located just a little ways outside Los Angeles 23 in California. And matter of fact, it's 24 located in the Simi Hills in Ventura County, and it was -- it's divided into four

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administrative and operational areas called, conveniently, Areas I through IV. And the DOE operations that are covered by the Energy Employees Occupational Illness Compensation Program are in Area IV only, so it's Area IV of the Santa Susana Field Laboratory is the covered facility.

That Area was established in 1953, and nuclear operations began in 1955 under the name of Atomics, International. There was also rocket testing operations at the same location under the name of Rocketdyne.

Those -- there were two engineering centers that were group-- sub-groups of Atomics

Engineering (sic), Liquid Metals Engineering

Center and the Energy Technology Engineering

Center. Those were involved in research and development of liquid met-- liquid metals technology and -- and nuclear technologies.

There was a merger -- Atomics International merged with Rocketdyne in 1984 as part of Rockwell International. Today it's owned by Boeing. There's been sort of this conglomeration of company names associated with the -- with the site over the course of its

operation.

The nuclear operation programs at the site operated from 1955 to 1980. They involved development of operation of some ten test reactors and a number of operation -- operation of a number of critical -- criticality test facilities, which are kind of similar to a reactor, I guess.

Nuclear support operations operated from 1956 to the present, includes reactor fuel manufacturing, disassembly of used reactor fuels and rods, production of radioactive sources, research on fuel reprocessing, preparation of waste for disposal, operation -- and the operation of particle accelerators.

And there were also non-nuclear programs that operated in the Area from 1966 to 1998.

The site is still there today in the middle of a, I guess, fairly contentious environmental remediation effort, so it is still there today.

And I believe the workers today are still covered under the program.

The history of petition-related activities for the site -- in June 22nd of 2007 we received the petition. It's our petition number 93. In

October 2007 we issued our professional judgment paper that petition qualified for evaluation based on limited internal monitoring data for the pre-1965 period.

And I might say here very briefly that the petition petitioned for 1955 to the present. That was the petitioned -- period of time in the petition. We qualified the petition up through 1965 on the basis of limited or lack of internal monitoring data. So the entire petition period that -- up through the present was not evaluated. The evaluation then focused on the qualification period, which goes through 1965.

And we announced that the petition qualified for evaluation on October 27th.

We evaluated the petition using the guidelines in 42 CFR 83.13, submitted a summary of the findings in the petition evaluation report -- report to the Board and to the petitioners, and that evaluation report was issued on February 15th of this year. This, as it says, is an 83.13 petition. The petition was received from a member of the public. It was not originated by NIOSH in our dose reconstruction efforts.

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The petitioner-proposed class definition is: All employees who worked in all the areas of the laboratory from 1955 to the present, including the post-1987 remediation period. The class we evaluated was: All employees of DOE and its predecessor agencies, contractors and subcontractors who worked in Area IV from January 1st, 1955 through December 31st, 1965. And our reco-- we are today recommending the addition of a class that's provided here: All employees of the DOE, its predecessor agencies, DOE contractors and subcontractors who were monitored while working in any area of the Santa Suna -- Santa Susana Field Laboratory Area IV for a number of work days aggregating at least 250 days from January 1st, 1955 to December 31st, 1958, or in combination with work days within another SEC. The basis that we qualified the petition for evaluation on was absence of internal monitoring data and a way to do internal dose reconstruction. There's a clear lack of that data up through 1958, or up until late 1958. And there was a health study published by UCLA that described the internal monitoring data

prior to about 1963 as being relatively scarce. So based on that and the fact that the petitioner's employment continued through 1965, we decided to qualify the petition up through 1965 and evaluate that period.

The -- we came to discover during our investigation that the relatively scarce internal monitoring data up through 1963 was due to the far smaller amount of radiological work up through 1963. And as the work ramped up from that period forward, the bioassay of course would ramp up also as more employees would be engaged in it.

The source of available information that were used in our evaluation report are the Technical Information Bulletins and the site profile that's been prepared; the case files and the individual claims in our NIOSH database, which we often call NOCTS; the NIOSH site research database; documentation and affidavits provided by the petitioner; interviews with former Area IV employees; the CEDR database, that's the Comprehensive Epidemiologic Data Resource database, there have been epidemiological studies done for site; and some scientific

publications as well.

A brief summary of the availability -- general availability of dosimetry data. This is the claims in our tracking system, and this data is up to date as of January 9, 2008 when it was compiled for some purpose, and that data was used on this slide. Classes (sic) which have employment during the class definition period that was the evaluated class, that from 1955 to 1965, there are 158 cases in that class as of January 9th. There are 81 of those dose reconstructions have been completed; 36 of those cases contained internal dosimetry and 65 contained external dosimetry.

No claim had internal data before August of 1958. One claim had data beginning in September of 1958, so it kind of -- the bioassay program (unintelligible) started there in late 1958.

And there were quite a large number of Area IV employees who were not radiological workers.

As I said, there were other activities in that Area in addition to the radiological work.

The internal monitoring data that is available is the -- we have urine bioassay available from

1959 to 1966 for this variety of radionuclides, two different uranium analytical methods and a variety of other radionuclides -- plutonium, thorium, mixed fission products, there are some gross alpha and gross beta results, and there are some results from polonium-210, strontium-90 and tritium.

Urine samples were collected based on job assignment that required exposure to radioactive materials, and there are more than 100,000 internal dose data points collected from more than 300 individuals who were monitored for internal exposure at Area IV. Those of course are not all claimants. External monitoring data is available for all years of site operation. The external dosimetry was assigned based on job or exposure potential.

Beta/gamma exposure was measured from 1954 to 1962 with a two-element film dosimeter; from '63 to '66 is a multi-element dosimeter from a commercial vendor; a pocket or pencil dosimeter -- I'm talking now about the evaluation period and then goes a little beyond. Pocket or pencil dosimeters were used for non-routine

1 work, but they were not used to record the dose 2 of record. 3 Neutron doses were record-- are available from 1954 to 1966 using NTA film. 5 And there are 4,665 individuals enrolled in the 6 external dosimetry program for which data is 7 available. Again, those are not all claimants. 8 The petitioner identified several bases in the 9 petition for -- for a class addition: the lack 10 of internal monitoring program, contamination 11 that was found in the sodium disposal facility 12 and the lack of records of material sent there. This was not considered radiological activity, 13 14 but it was found to be contaminated after some 15 period of disposal there. Inadequate air 16 monitoring, a specific reactor incident at the 17 sodium reactor experiment in which a large 18 portion of the reactor fuel failed -- or 19 significant -- not a large portion, a 20 significant fraction of -- of it failed. 21 Uranium fires that occurred, tritium plume in 22 the groundwater and inadequate radiation 23 badges. 24 In our evaluation we identified issues as well. 25 We looked at the lack -- there appears to be a

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lack of internal monitoring data from 19-before 1959. We were concerned about possibly missing records, and the reason we were concerned about possibly missing records because we -- some workers -- when we would receive the response from the site, we asked for the radiation exposure history, we would receive -- some people would get a response of "no record" and other people we would get a response that would include a radiation exposure record that was completely blank. So we -- you know, when you see a -- when you first get a response and you -- and they respond "no record," you say well, this person probably was not a radiation worker. And then you get a response that has a radiation worker that has nothing on it and you say well, now what does this mean? Because normally I would say this person wasn't a radiation worker. what it turns out is there were restricted areas in Area IV, meaning -- restricted being a term I believe that used to be used to denote contam -- a radiological control aspect. And if you went into the restricted area, whether you were a radiation worker and worked around

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radiation or not, because inside the restricted areas there were places -- not every site inside a restricted area were you subject to radiation exposure. If you went in there, you got a radiation exposure record, and so you got a pink card or blue card filled out and that was placed in your folder. If you were -- if you went in there and you were not a radiation worker, then nothing was ever written on your exposure record. So workers who entered the restricted area but were not radiation workers got an exposure card. Workers who never went into the restricted area did not get one, and so that's why we got the two categories of record responses of either no response (sic) or a blank exposure record.

We were also concerned about monitoring of emergency personnel who may have to enter a restricted area for an emergency.

So addressing each of these concerns -- I guess maybe I should speak quickly, I'm going to -- I'm tak-- I'm taking a lot of time up here.

The lack of internal monitoring data -- based on our evaluation, we found that there was no established routine bioassay program before

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August of 1958, and there was exposure potential to uranium and fission products. have insufficient source term information to bound the dose. And we do have bioassay data in term-- form of urine data after 1958, which we believe we can use to bound the dose. Petitioner raised a concern about sodium disposal burn pit and the lack of records. burn pit was used to react sodium and an alloy -- potassium -- a sodium-potassium alloy, which I've heard commonly referred NaK -- referred to They react it with water as a means of disposal. Some of that material apparently was contaminated with fission products because it was coolant from the sodium reactor coolant. It was not expected or intended to be used (unintelligible) radioactive waste disposal. But given that we have a robust bioassay data set for the people who were radiologically exposed, we are confident that use of that data set will allow us to bound people's doses. And we also have actually for both internal and

Petitioner raised a concern of lack of air monitoring at the -- at the Area IV. Our

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internal dose evaluation relies primarily on the urine bioassay estimates or the internal monitoring database in order to do that. is some air sampling data available, gross alpha and gross beta. To be honest, I think the urine bioassay or the internal monitoring data would be our pathway to do these doses. And again, since we have a fairly robust data set for that, we're confident we can provide a bounding estimate for the doses. Reactor -- the petitioner raised a concern about the sodium reactor incident, including -that there was a release of core gases after the SRE cladding failure in 1959. And that clad-- well, cladding failure or fuel melt, probably some of both, I would guess -incident resulted in release of gaseous fission products to the hold-up tanks, and then there were hold-up tanks over this so it was followed by a controlled release to the atmosphere. And again, since we have bioassay data and a robust bioassay data set, we believe we can analyze these doses -- or bound these doses. There is -- since it's a sodium reactor, of course, it wasn't open to the atmosphere. It

was -- had a helium cover gas and that cover gas then was -- there was some venting of that to the hold-up tanks.

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Exposure from workers in fires, like sodium metal and uranium fires. There were quite a number of fires and incidents like that. tend to be very well documented at Area IV, since they're very well documented with oftentimes radiological readings associated with that documentation. That, in combination with the bioassay data, we believe that we have sufficient data that we can bound these doses. A petitioner concern was that the groundwater used for drinking at the site and tri-- tritium was later found to be in sampling wells. groundwater was exclusive water supply from 1948 to 1964. All but one of the wells for the Santa Susana Field Laboratory were in Areas I through III, not in Area IV. Since 2000 all the water supply has been from off-site. the groundwater supply wells were less than 1,000 picocuries per liter. That sounds like a high number, but that's the detection number on the analysis, so that was not detected in those well samples at 1,000 picocuries per liter.

There's a sampling well, not a groundwater supply well but a groundwater sampling well, near the reactor site that was never used for drinking water where there is measureable tritium on the order of 3,000 picocuries per liter. And because of some knowledge of the amount of tritium in the groundwater, we can assume that tritium made it into the drinking water and provide a bounding dose in that instance.

Petitioner raised a concern about the -- that the inadequate radiation badges -- that was taken from the tiger team report, which I believe those were written in the '90s, if I'm not mistaken. What the tiger team report actually commented on was the fact that the dosimetry system at Santa Susana was not DOELAP accredited. DOELAP accreditation is actually kind of a moot issue for the evaluation period, which goes through 1965, since DOELAP accreditation didn't exist until about 1986. And even when it did come into existence, it provided for smaller sites to seek an exemption as long as they used what we used to call NVLAP accredited -- DOELAP is Department of Energy

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Laboratory Accreditation Program. NVLAP is
National Voluntary Laboratory Accreditation
Program. NRC-regulated and State-regulated
entities typically used NVLAP, where as DOELAP
-- Department of Energy wanted to do their own
thing so they invented DOELAP, which was very
similar.

Concern we ran across about possible missing records, I think I covered this already. Workers who worked outside a restricted area had no dose record. Areas (sic) who went inside a restricted area but had a blank record, and they were not radiation workers, those people had a record but it was blank. Our concern about firemen from Areas I through III who might be called on to respond to emergencies in restricted areas, we found that the Area IV firemen were monitored. is some -- there was at least one incident of apparently missing dosimetry file, someone who engaged and seems like is in fact monitored, but we didn't have a file for him. still feel that because of this, because firemen are included in our monitored population, that our coworker data will be

sufficiently bounding for this situation as
well.

Oops, I (unintelligible) two -- hit the button twice.

Now we included on the O drive a number of sample dose reconstructions to illustrate our ability -- these are hypothetical. These are not actual data -- actual cases. But if we had a hypothetical reactor operator who was male, he worked there during the period of the sodium reactor -- the sodium reactor experiment incident, we would do -- we could do a dose reconstruction based on the -- his internal monitoring and external monitoring, provided the employment starts in 1959 or later. So these are just some of the assumptions we made for this hypothetical person.

The person -- we would expect to have routine monitoring for uranium and fission products.

We said well, how about acute uranium intake in 1965 based on his bioassay record, and so the external dosimetry throughout for all the types of external radiation. Based on this information -- which it may in fact be real -- probably an amalgam of data taken from several

different files, it wouldn't be one person's 1 2 file, if you -- when we do intakes, we always 3 assume the solubility class that provides the most favorable outcome for the claimant if we 4 5 don't know for sure what the solubility class 6 So depending upon which you choose in this case, some organs -- it'll be -- you know, 7 8 S will be more favorable for some organs, M 9 will be more favorable for others, depending 10 upon how you -- which you choose, you'll have 11 these two different intake regimes of a fairly 12 large -- quite large type S intake for the 13 acute intake, on top of a quite small chronic 14 intake over the course of employment. 15 If you had -- if your assumption is type M, the 16 acute intake is a little more moderate, but 17 your chronic intake is quite a lot larger over 18 the entire time of the employment. 19 Strontium-90 is most favorable as type F and 20 the bioassay would provide a chronic intake of 21 -- of that nature. 22 And then the external -- external doses were 23 thrown in here. I doubt that this is real data 24 'cause I doubt that anybody really got the same

dose every year, but this was thrown in for --

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1 to show that we would have those measurements. 2 The outcome of a case like this for various 3 organs giving the organ dose and the probability of causation is provided here in 5 the next table. Up to -- and that's with a 6 1990 cancer diagnosis. 7 Sample dose reconstruction number two is the 8 hypothetical again, hypothetical fireman, again 9 male, tasked with removal or burning of sodium 10 reactor components, et cetera. These are his 11 demographics that would relate to how the case 12 works out eventually. Presumable an Area IV 13 fire-- Area IV fireman would have his own bio--14 his own data, so we would be able to make the -15 - use his data with -- to -- in order to do the 16 dose reconstruction. We would expect someone 17 like this would probably have acute intakes, 18 more so than chronic intakes, so we could go 19 through and do the dose calculation. 20 And you will arrive at doses and probability of 21 causation (unintelligible) provided in this 22 next -- again, these are strictly hypothetical 23 cases. 24 Dose recon-- sample dose reconstruction number

three is a technician, doesn't handle

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1 radioactive materials but drinking the 2 groundwater. So you can understand -- we can go through and do this. There would be an 3 intake -- realistically, I would believe probably the tritium would be included in 5 6 everybody's, as I'm looking through these now. 7 So these again just kind of show magnitude of 8 exposures on certain of these scenarios. 9 In our evaluation process to determine -- as we 10 evaluate a Special Exposure Cohort, a two-11 pronged test -- of course you've all seen this 12 before, is it feasible to estimate the 13 radiation -- level of radiation doses, and is 14 there a reasonable likelihood that such 15 radiation dose may have endangered the health 16 of members of the class. 17 We've de-- NIOSH has determined it's not 18 feasible to complete dose reconstructions with 19 sufficient accuracy for the time period -- the 20 period of limited internal dos-- internal 21 bioassay data, that's from 1955 to 1958, and 22 that the health of the employees covered may 23 have been endangered. 24 The evidence reviewed indicates workers in the 25 class received chronic internal and external

exposure from reactor operations, fuel production and other support and research activities at Area IV of Santa Susana Field Laboratory sufficient to potentially be harmful to them. We did not recognize a particular incident that would indicate that they were subject -- they were likely to be harmed from just being present.

Our recommended class definition is: All employees of the DOE, its predecessor agencies and the DOE contractors and subcontractors who were monitored while working in area (sic) area of the Santa Susana Field Laboratory Area IV for a number of work days aggregating at least 250 days from January 1st, 1955 to December 31st, 1958, or in combination with work days within the parameters established for one or more -- one or more other classes of employees within the SEC.

The summary -- the brief summary of our findings is for -- feasibility findings for internal doses from all radionuclides from 1955 to 1958, there's no data, we don't believe it's feasible to reconstruct those doses. From '58 -- from '59 to the present, we believe that it

1	is feasible. For external, for all the years,
2	we believe the dose reconstruction is feasible.
3	That ends my presentation. I know the
4	petitioner wants to speak. I think the
5	petitioner has even made some comments since
6	our evaluation report was was presented.
7	DR. ZIEMER: Stu, before the petitioner comes
8	with a presentation, a couple of quick
9	questions. Could you clarify your class
10	definition does not include the "should have
11	been monitored" category, so you're confident
12	that anyone included was monitored.
13	MR. HINNEFELD: It's our belief today that they
14	con that they conscientiously monitored the
15	radiation workers.
16	DR. ZIEMER: And slide 21 where you indicated
17	concern for non-monitored workers
18	MR. HINNEFELD: Which one is this, slide 21?
19	DR. ZIEMER: Slide 21.
20	MR. HINNEFELD: I'm not going to sure I'm
21	going to
22	DR. ZIEMER: No dose records for some non-
23	monitored workers. So what does that mean in
24	this issue
25	MR. HINNEFELD: Well, that that speaks to

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our --

DR. ZIEMER: The ones that were outside the controlled area?

MR. HINNEFELD: There were unmonitored workers outside the restricted area, and there were unmonitored workers in the restricted area. What -- our concern about possibly lost records was that when we would receive a response that said "no record" and we -- that -- we would normally assume that that meant this person wasn't a radiation worker. But in this instance, not only did we receive responses that said "no record," we received responses that said -- that had an exposure record that was blank. An exposure record that's blank indicates they were -- they were non-radiation workers. Our concern was that the records of the first group where we got no response (sic), those records had been lost and so we would not know what their exposure history was. That was our concern.

Our concern was allayed by our investigation, which revealed that in fact there were two -- these two groups of non-monitored workers.

DR. ZIEMER: Thank you. Any other questions

1 before we hear from the petitioner? Yes, Brad 2 Clawson. 3 MR. CLAWSON: Stu, who did the bioassay for these people? Was it done in-house or was it 5 done by a contractor? 6 MR. HINNEFELD: In most cases it was done by a 7 contractor. There -- there are a number -- a 8 variety of companies who did it. 9 MR. CLAWSON: Okay. I -- I's just trying to 10 picture what you were drawing up there, though. 11 You've got a secured area that the people that 12 go into this are supposed to be monitored, but 13 we have some that aren't monitored. So I'm 14 trying to figure -- you know, they should have been monitored. What controlled them from 15 16 going into the rad areas? 17 MR. HINNEFELD: I think -- I think what it is 18 is that the restricted area was probably the 19 area that was used by the nuclear operations. 20 Remember, there were nuclear operations and 21 rocket operations. And the restricted area, 22 meaning restricted as a radiological control 23 term --24 MR. CLAWSON: Right. 25 MR. HINNEFELD: -- was where the radiological

1 operations occurred, not the rocket operations. 2 Within there, there would be office buildings, 3 there would be other -- you know, other aspects 4 that were apart from the radioactive material, 5 so there would be areas where there was really 6 no potential for exposure, but that wa-- it was 7 on that part of the plant. I believe that 8 would be the situation. MR. CLAWSON: Did -- did I -- now I can't 9 10 remember, so did we have air sampling data? 11 MR. HINNEFELD: There is some air sampling data 12 for -- gross alpha and gross beta air sampling data --13 14 MR. CLAWSON: Just --15 MR. HINNEFELD: -- starts relatively early. 16 MR. CLAWSON: -- just air, or surrounding 17 areas, or were they personal air --18 MR. HINNEFELD: I don't know where it was 19 collected. 20 MR. CLAWSON: Okay. 21 DR. ZIEMER: Okay. Let's then hear from 22 LaVonne Klea. LaVonne, you still on the line? 23 MS. KLEA: Yes, this is LaVonne. 24 DR. ZIEMER: Please proceed. 25 MS. KLEA: I do have some comments on what you

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said, but I'll -- I'll read what I've written
here. I hope you've all read my comments,
which have been forwarded twice.

I have no evidence that the site contractor had good monitoring data after 1958. This is contrary for all the evidence that I have. Ιn 1994 Rockwell was asked for their employees' monitoring records for a UCLA worker study on radiation and chemical exposure. They stated that very little in the way of records exist. They said that you cannot invent records. NIOSH themselves have used estimates of external environmental doses from 1952 to 1974 and calculated doses from 1975 to 1999. this mean that they have no records from '52 to My dose was estimated from Portsmouth. The contractor used Landauer film badges. were not DOE DOELAP approved. [Name redacted] pointed out certain questionable practices. The first is that data obtained by dosimeters is normalized to a 1000 feet altitude, by using an adjustment factor equal to 15 mr per 1,000 feet elevation difference to obtain site averages. Two nationally renowned experts had never heard of this practice.

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Second, the contractor did not have a comparison study of the dosimeters placed by the State of California versus DOE. According to the DOE there was no procedure or technical basis for operation of the internal dosimety program. Urinalysis was used as the bioassay technique for insoluble cobalt-60. There was no technical analysis for the suitability. And what about super S plutonium? It was highly insoluble. DOE and the site contractor had a long history of giving low priority to environmental safety and health. The site contractor was basically allowed to monitor themselves with almost no oversight from the San Francisco office of DOE. They had no dedicated staff for DOE compliance and staff time was used on NASA and DOD contracts. Radiological protection personnel were not trained and qualified by DOE. The study of air flow patterns at ETEC facilities requiring air sampling was not done and did not meet DOE performance standards for the internal dosimetry program. Often doses were not added to the records because it was manpower intensive. Radioactive particulate monitoring

did not conform to DOE, EPA and CFR requirements. No swipe tests were used for handling packages. ETEC is at 1000 foot elevation from the valley floor. Last week EPA declared that the site qualifies for Superfund eligibility. Radioactive and chemical pollution has flowed down the mountain on all sides, contaminating community drinking water, children's camps, state parkland and new construction projects. It is estimated that contaminants will fill the Rose Bowl 55 times. Contrary to the site profile, well water was given to the employees until 1985, not 1965, that was contaminated with TCE and 1,2 dichlorethylene. The contractor saved 50,000 dollars a year. We have a tritium plume in Area IV of 119,000 pci per liter discovered in 2004. And every year the measurements go up, suggesting that there are sources and also a water supply well in that area, yet Rockwell never tested for tritium and the tritium wasn't Rockwell is a convicted felon. They are convicted felons for illegally burning waste without permit and killing workers. In 1996 we had an FBI raid, grand jury conviction and a record fine of 60 million dollars. Bladder cancer is very high among the workers, probably the highest percentage of the 22 cancers. Also bladder cancer is 50 percent higher in the population closest to the site, suggesting chronic internal exposures.

I also mention that the site profile is flawed. According to EPA, SNAP 8DR operated in Building 59 from 1962 to 1964, shut down and restarted from January 1969 to December '69. Yet the site profile states that it was in operation from 1968 to 1969. The site profile basically was written by the Boeing Company. One large notebook was used and the Boeing consultant was the company's own expert witness who has been fighting workers compensation claims for years, and now he is involved intimately with the NIOSH program. This is an extreme conflict of interest.

And to the Department of Labor, I see no change in the corporate culture of fighting workers compensation claims, contrary to what the law promised, that the corporations would be instructed to stop fighting claims and assist

the workers. We had a UCLA Worker Death Study for which NIOSH had a representative on the board, yet in evaluation of my petition the BOICE study was quoted. The BOICE study was an in-house, corporate paid study, another conflict of interest.

I have a question for the Board on the fairness of the NIOSH program for women. BEIR VII states that women have a 50 percent greater risk for solid tumors than men. Shouldn't the dose reconstruction project be adjusted accordingly? I thank you for listening. I will not give up. I request that my petition be investigated for the whole period from 1955 to the present because most of the data I have was written in late '80s, early '90s, and there looks like corrections to the problem. Thank you.

DR. ZIEMER: Thank you very much, LaVonne. I'd like to point out, Board members, just to remind you that LaVonne distributed to us her rebuttal to the SEC petition evaluation report -- her rebuttal is dated February 6th, 2008. I think, Christine, you distributed this in early April. Is that correct? Right.

1 DR. BRANCHE: Yes, 'cause she added something 2 to it -- she said she sent it twi-- Ms. Klea 3 said that she sent it twice, and so on the second occasion, at which time I was then the DFO -- she sent it a second time and that's 5 6 when I sent it to all of you. 7 MS. KLEA: I sent two different sets of 8 comments and then one I just read I just wrote 9 yesterday, so if you'd like, I can forward that 10 also to the (unintelligible). 11 DR. BRANCHE: If you -- actually that will be 12 very helpful if you could please send that to 13 Ms. Breyer and then we will make certain that 14 it is entered into the record as well as 15 distributed to the Board. Thank you. 16 MS. KLEA: Thank you. 17 DR. ZIEMER: Board members, do any of you have 18 questions for LaVonne, either on her previous 19 materials or on her comments today? 20 (No responses) 21 Okay, thank you very much. Then let me open 22 this for discussion on any of the issues 23 related to either the NIOSH presentation or 24 LaVonne's points, or other issues that any of 25 you wish to raise or any comments you wish to

1 make. 2 MS. KLEA: Oh, sir, could I just --3 DR. ZIEMER: Yes? 4 MS. KLEA: -- (unintelligible) one more 5 statement? 6 DR. ZIEMER: Sure. 7 MS. KLEA: Okay. I have read (unintelligible) 8 documents on the reactor (unintelligible) that 9 we had two reactors that for sure opened their 10 doors to -- to neutralize the radiation 11 exposure inside, and that would have been 12 Building 24, which (unintelligible) Building 28 13 they call the swimming pool reactor and it 14 operated 1961 until 1972 and it was 15 (unintelligible) every day. They ran 16 (unintelligible) reactor (unintelligible) open 17 doors (unintelligible) vent the room. 18 (unintelligible) reactor (unintelligible) which 19 ran from 1956 to 1966. I have an old 20 (unintelligible) very (unintelligible) that 21 they had a large (unintelligible) wearing 22 protective clothes. They had no monitoring in 23 that building and (unintelligible) 1959 and in 24 that report that they have the doors were

(unintelligible) to this (unintelligible)

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1 contaminated air, so I know for sure that it 2 happened twice where (unintelligible) and not 3 captured. It was released by (unintelligible) 4 the door. Thank you. 5 DR. ZIEMER: Thank you. I believe -- and Stu, perhaps you could clarify this -- I assume 6 NIOSH has seen the -- the points that she had 7 8 raised in her letter. I believe you're saying 9 that in -- in spite of those issues, you still 10 believe that you can bound the dose for the 11 later periods. The early period is not in 12 question. MR. HINNEFELD: Yeah, the earlier -- the early 13 14 period is not in question, and today we didn't 15 see anything that would cause us to pull back 16 and amend our evaluation report. 17 DR. ZIEMER: Okay, looking to see if there's 18 any other comments or --19 DR. MELIUS: I -- I have a --20 DR. ZIEMER: -- yes, Dr. Melius. 21 DR. MELIUS: -- just a factual question, and 22 that is has SC&A reviewed either the site 23 profile -- I shouldn't say either. Has SC&A 24 reviewed the site profile? I'm trying to get a 25 -- a handle on where we are in terms of looking

1 at the site. 2 DR. ZIEMER: John? 3 DR. MAURO: Yes, at the last meeting the Board 4 authorized us to proceed with a site profile 5 review, which is underway as we speak. DR. MELIUS: Okay. 6 So that's -- that's in process. 7 DR. ZIEMER: 8 DR. MELIUS: Right. 9 DR. ZIEMER: Right. And John, what was the 10 expected delivery date on that? 11 DR. MAURO: In general our site profile reviews 12 require about a four-month period, and we are 13 only -- we started last month, so three months. 14 Right. And so to the extent that DR. ZIEMER: 15 that may impact on the Board's action here 16 today, take that into consideration. 17 It would be appropriate to take some action. 18 Your poten -- I'll remind you of the 19 possibilities here. The board may -- may move 20 to recommend addition of this class. 21 point out that doing so would not preclude 22 taking additional actions later -- wouldn't 23 require it, but it wouldn't preclude it. 24 You could -- you could move to postpone action 25 until the site profile is received, although

the site profile doesn't directly address
necessarily SEC issues, but it may include
them.

Or the other action would be to not approve the recommendation from -- from NIOSH to add the class.

DR. MELIUS: Can I have a --

DR. ZIEMER: Dr. Melius.

DR. MELIUS: -- another question then for -for NIOSH, so this is the -- the -- the way you've written up the SEC evaluation report, it's -- it's -- you make it appear that in 1958 there was suddenly a full monitoring program there, and at least our past experience has been that usually those monitoring programs are phased in over time -- I mean before they capture all the workers or all the work areas and -- and so forth. And I just can't tell from your report, and I don't have access to the site profile to tell if that's really true or -- or is it -- if it was phased in, then I -- it just raises some questions about time periods involved and sort of how I think we should proceed, but maybe it -- maybe in this case it did --

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1 MR. HINNEFELD: Well, I can --2 DR. MELIUS: -- go from zero to full. 3 MR. HINNEFELD: Our position -- it started in late 1958 and our position is by 1959 --5 DR. MELIUS: Yeah. -- is when we say we can start. 6 MR. HINNEFELD: 7 They had -- the people who were exposed were 8 appropriately monitored, and if you get a 9 bioassay sample not on January 1st but on March 10 31st for someone, you can do a pretty good job 11 of estimating his exposure from -- from January 12 31st, so our -- our view is that by January it 13 was sufficient. Recognize that radiological 14 operations ramped up as years went on and 15 additional monitoring then came on line, too. 16 Okay. John Poston. DR. ZIEMER: 17 DR. POSTON: Well, it seems to me that if we 18 ask SC&A to look at this that we should give 19 them a chance to finish their work, and 20 therefore I'd make a motion that we take no action at this time and -- upon receipt of the 21 22 SC&A review. 23 DR. ZIEMER: Okay, there's a motion to postpone 24 action until an opportunity to see the SC&A --25 and that'll be a site profile review. Is there

1 a second to that motion? 2 UNIDENTIFIED: (Inaudible) 3 DR. ZIEMER: And seconded. Discussion on the motion? 4 5 DR. MELIUS: Yeah, I agree with most of it. quess my only question would be would -- should 6 7 we have them focus just on this area and then 8 the -- on particular issues related to the SEC 9 initially so that we could expedite it, to the 10 extent that it can be, and I -- I just don't 11 know enough about where they are with their 12 work or -- to know if that's going to make any 13 difference or not. 14 DR. POSTON: Well, my motion was just to delay, 15 it wasn't implied that other things couldn't be 16 done and so forth. It was just to -- not to 17 take action --18 DR. ZIEMER: Actually --19 DR. POSTON: -- at this time. 20 DR. ZIEMER: -- let me just suggest the 21 following. If the motion carried, then when we 22 do our -- our other Board work, which will 23 include SC&A tasking, we can specifically 24 address how to task this particular one. 25 DR. MELIUS: That -- that's fine with me then.

1 Yeah, that was my only concern, I -- reason I'm 2 a little reluctant to -- we take any action on 3 the first part, the approved ar -- is this 4 question of -- of what happens on the margins 5 of the -- the -- in terms of years and so forth 'cause again in the past we've -- we've often, 6 7 you know, had questions about what years to put 8 in there in terms of when is there adequate 9 data in order to be able to estimate that --10 again, it -- it -- there may very well turn out 11 to be what NIOSH said, but I'd like to have 12 some more information, some -- before we make 13 that decision. 14 DR. ZIEMER: Any other comments, pro or con? 15 We're debating the motion to delay. 16 appear to be no other comments. Are you ready 17 to vote? 18 All in favor, aye? 19 (Affirmative responses) 20 Mr. Presley's on the line. DR. BRANCHE: 21 MR. PRESLEY: Bob Presley, aye. 22 DR. ZIEMER: Bob Presley votes aye. Any 23 opposed? 24 (No responses) 25 Any abstentions?

(No responses)

The motion carries and we will delay action or delay a recommendation on this particular petition pending the completion of the work by SC&A on the site profile. And again, we'll have the opportunity, if we wish, to focus that in some way as well.

(Pause)

DEPARTMENT OF LABOR UPDATE

Okay, without objection then, we'll proceed to the presentation by the Department of Labor, and Jeff Kotsch is here this morning to present that. Jeff, welcome.

MR. KOTSCH: Good morning. This is a status report from the Department of Labor for activities related to the Energy Employees Occupational Illness Compensation Program.

Just as background, there are two portions to this program. There's Part B, which became effective on July 1st, 2001, and at the bottom are the dates of the slides. There are a couple of differences through this -- through the series of the slides, but this one is March 25th, 2008. And because of that difference, then I think the dates that maybe Larry had in

his, some of the numbers that we share or that
are of similar activities may be a little bit
different.

But anyway, as of March 25th the Department has
-- or there have been 61,234 cases filed with
the Department of Labor. That encompasses

bigger because survivor cases may have one or

more claimants involved in them. Of those,

40,025 are cancer cases, and we have referred

The number of claims is always

26,766 cases to NIOSH.

89,282 claims.

The other half of the program, the Part E program, became effective on October 28th, 2004. This was formerly the Part D program administered by the Department of Energy. And on that side of the program we've had 51,164 cases filed, which includes 70,992 claimants -- or claims. And at the beginning of that program over 25,000 cases were transferred from DOE.

As far as compensation as of, again, March 25th, the Department has put out in compensation \$3.6 billion total; \$2.3 of that is Part B, breaking down into \$1.8 billion for cancer claims, \$282 million for RECA, and the

1 remainder of that would be for silicosis and 2 beryllium. \$1 billion has been paid as part of 3 the Part E program and \$206 million as part of medical benefits. 5 Just quickly under the Part B benefits 6 overview, who's eligible, it's Department of 7 Energy and its contractors and subcontractors, atomic weapons employers, beryllium vendors; 8 9 uranium miners, millers, ore transporters who 10 worked at facilities covered by Section 5 of 11 RECA -- that's the Radiation Exposure 12 Compensation Act, that program's actually 13 administered by the Department of Justice and 14 we supplement it; and certain family members of deceased workers. 15 16 And quickly, the claim categories for Part B 17 are cancer, chronic beryllium disease or CBD, 18 beryllium sensitivity, chronic silicosis --19 which is primarily the miners/millers, and the 20 RECA Section 5 cases. 21 And who -- who eventually becomes -- or who is 22 potentially covered that is compensable are 23 workers or claims that are determined that the 24 covered employee was a member of the SEC and 25 was diagnosed with one of the specified

1 cancers; or it is determined through a dose 2 reconstruction conducted by NIOSH that the 3 covered employee's cancer was at least as likely as not, that's greater than 50 percent, 5 caused by exposure to ionizing radiation. 6 So the Part B cancer case status as of, again -7 - this is a little different date, March 20th -8 - there have been 40,000 -- about 40,000 cases 9 having 61,549 claims; 32,000 of those have had 10 final decisions, that's about 80 percent; 1,800 11 have recommended but no final, that means that 12 their case is with our Final Adjudication 13 Branch for another look -- I mean that's the 14 part of the process of -- of turning a 15 recommended into a final decision; about 4,500 16 are at NIOSH, and about 1,700 are pending 17 initial DOE (sic) -- an -- an initial DOE (sic) 18 -- that is an initial recommended. 19 This is the breakdown for the Part B cancer 20 cases as far as final decisions. On the left 21 side you see that there's 12,559 of final 22 decisions were approved. On the right, the red 23 bar is 19,470 denied, and the breakdowns go 24 across from left to right -- non-covered 25 employment -- reasons for denial, non-covered

1 employment; 11,735 with a POC less than 50; 2 about 3,000 with insufficient medical evidence; 3 about 1,100 with non-covered conditions and 365 with ineligible survivor. 5 Just a quick one on the Special Exposure 6 Cohorts. Of course there's the statutory ones 7 that were in the Act -- the three diffusion 8 plants, certain nuclear tests -- and then the 9 new SEC class designations that have been 10 recommended by the Board and passed by the HHS 11 Secretary. Then there's specified cancers, 12 causation is presumed but no dose 13 reconstruction, and then the HHS recommends the 14 SEC designation as -- if it -- after 30 days 15 with Congress. 16 As far as new SEC-related cases, we've had 17 1,565 cases withdrawn from NIOSH. This number 18 is as of March 20th. 1,421 have had final 19 decisions issued, that's 92 percent; 45 have 20 recommended but no final decisions. We have 20 21 cases pending and we have 69 cases that are 22 closed. 23 As far as our NIOSH referral case status, we 24 show now, as of March 25th, 26,760 cases 25 referred to NIOSH; 18,645 have been returned

from NIOSH. Of those, 16,000 -- about 16,500 have had dose reconstructions, 19 are being reworked for return, 2,077 have been withdrawn with no dose reconstruction.

Then the other portion of that is 8,115 are currently at NIOSH; 4,628 of those are initial or originally referrals, 3,487 are reworks or returns.

The NIOSH dose reconstruction case status numbers, we have -- we're showing as of March 25th 16,549 cases with dose reconstructions.

That's -- and of those, 14,261 have a final decision. That's about 86 percent. 1,952 have a final -- I'm sorry -- have a recommended but no final decision, and 336 are pending with -- at Labor with a recommended decision. I'm sorry, are pending a recommended decision by Labor.

Now as far as NIOSH case-related compensation, as of March 20th \$956 million has been paid out in compensation to 900 -- I'm sorry, 9,908 payees in 6,405 cases. \$779 million has been paid on dose reconstructed cases to 7,364 payees, which is 5,213 cases. And the other \$177 million has been added for SEC cases.

That includes payments to 2,544 payees in 1,192 cases.

Paid cases under the Act, there's 20 -- have been -- again, this is March 25th, 28,613 paid Part B and E cases; 19,777 of those are Part B cases. The breakdown there is about 12,367 for cancer payees; 5,600 -- little over 5,600 for RECA case payees; about 1,760 for other Part B payees. Those are, again, the silicosis/beryllium. And there have been 8,836 Part E cases. That's the toxic side of the program -- toxic chemical exposure.

Just an update quickly on some of the SECs that have been in front of the Board during the meeting or -- or is scheduled. For Texas City, which is an AWE, Texas City Chemicals, we show 84 cases. It's only affected by Part B of the program. There's no toxic -- there's no Part E applications for AWEs. We show two NIOSH dose reconstructions. There have been 14 final B decisions and no compensation.

For the SAM labs at Columbia, we're showing 42 cases, one NIOSH dose reconstruction. We've had ten final Part Bs, nine Part B approvals, six Part E approvals, and have paid \$2 million

1 in compensation. 2 For Horizons we show five cases, Part B only --3 again, this is an AWE -- and no dose 4 reconstructions and no compensation. 5 For Area IV at Santa Susana Field Laboratory we 6 show 729 cases. This is covered under both 7 parts B and E because it's a DOE facility. 8 We're showing 132 NIOSH dose reconstructions, 9 155 final decisions for Part B, 44 Part B 10 approvals, 46 Part E approvals, and total 11 compensation for both parts of \$9 million. 12 And for Kellex-Pierpont we show seven cases and 13 no dose reconstructions or approvals, or 14 compensation. 15 NUMEC Parks Township in Pennsylvania, we show 16 143 cases. This is an AWE so this is only Part 17 B cases, ten dose constructions, 29 final Part 18 Bs, 15 approvals -- Part Es are not applicable 19 -- and total compensation under Part B of \$1 20 million. 21 Pinellas we show 1,137 cases, 300 dose 22 reconstructions, 367 final decisions for Part B 23 under the Department of Labor, 70 Part B 24 approvals, 86 Part E approvals -- again, for 25 toxic chemicals -- for a total compensation for

1	both parts of the program, Parts B and E, of
2	\$12 million.
3	At this point I wanted to ask Gen I can give
4	a quick update on Linde Ceramics, either now or
5	later I guess when you do your update.
6	DR. ROESSLER: I think we need to have it at
7	at some point.
8	MR. KOTSCH: You can have it now
9	DR. ZIEMER: Why don't you do it when you do
10	your report on
11	MR. KOTSCH: We can do it later. Do you want
12	to do it later?
13	DR. ROESSLER: I'll get or you can do it
14	your part when I do the report.
15	MR. KOTSCH: Right, but I mean you will be
16	doing an update on that, Linde Ceramics.
17	DR. ROESSLER: Yes, tomorrow.
18	MR. KOTSCH: We'll do it then.
19	DR. ROESSLER: Okay.
20	MR. KOTSCH: So are there any questions?
21	DR. ZIEMER: Thank you. Jeff, can you give us
22	some idea of what the rate of claim numbers
23	of claims coming in nowadays on Part B? I
24	assume you're tracking that. Is it you
25	know, has it leveled out, is it going up, going

1 down? 2 MR. KOTSCH: It -- actually Larry might have a 3 better -- have a feel, too, but I haven't 4 looked at the numbers recently. It seems that 5 it leveled out. There's -- occasionally 6 there's little rises in it, but we're at a 7 semi-steady state situation as far as 8 additional claims. 9 DR. ZIEMER: Well, what they see at NIOSH has 10 been impacted by their ability to put things 11 back out the door. They were ahead of you for 12 a while, but I wasn't sure whether what comes 13 in to NIOSH is that typically reflected --14 MR. KOTSCH: Yeah --15 DR. ZIEMER: -- what -- what you have coming in 16 or has yours been affected by budget in terms 17 of your --18 MR. KOTSCH: No --19 **DR. ZIEMER:** -- turnover ability? 20 MR. KOTSCH: -- I don't think it's that, but we 21 have -- we saw a dip for a while, and then it 22 started coming up again. Some of it comes off 23 of -- or responds to when we have town hall 24 meetings and other outreach types of things. 25 There's sometimes small increases in the input

1	or the you know, the new cases.
2	DR. ZIEMER: But it's more would it be
3	steady state
4	MR. KOTSCH: I think we're Larry, do you
5	agree we're kind of 'cause you kind of see
6	most of the the baseline, obviously.
7	MR. ELLIOTT: We see about 200 a month come at
8	us. I caution you when you look at the numbers
9	Jeff has provided on the total number of B
10	claims that they receive, we don't see all of
11	those.
12	DR. ZIEMER: No, understood, right.
13	MR. ELLIOTT: So you can't look at my numbers
14	and reflect upon those numbers given by DOL
15	today because they're the include I think
16	I understand this. They include other claims
17	that we don't see
18	MR. KOTSCH: Yeah,
19	MR. ELLIOTT: but we're seeing at NIOSH
20	about on average, about 200 a month.
21	DR. ZIEMER: And that's been pretty steady now
22	for a while.
23	MR. ELLIOTT: Yeah. It may go up to 200, 225,
24	but it'll dip down next month to 170, so
25	MR. KOTSCH: Yeah, I think for right now that's

1 a -- that's a reason-- that's the number I would think in my mind, 200 to 250 probably 2 3 toward the low end as an average. 4 DR. ZIEMER: Dr. Melius. 5 Yeah, and this may -- question, I 6 don't know who best can answer, but I'm trying 7 to get a handle on the reworks, returns to --8 to NIOSH and so forth. Seems that those have 9 gone up dramatically and they don't appear to be getting caught up with very quickly, either, 10 11 and I'm just trying to understand what some of 12 the issues are there. I missed Larry's 13 presentation yesterday so... 14 I'll let Larry com-- comment as MR. KOTSCH: 15 far as the workload for him. I mean I think 16 the -- the cause of those over the past few 17 quarters has been the -- the release of the 18 Program Evaluation Reports. 19 DR. ZIEMER: Yeah, the PERs has impacted on 20 I think Larry discussed that a bit 21 yesterday. 22 MR. ELLIOTT: Yeah, if you look in my slides 23 from yesterday there are a couple of slides 24 that show -- speak to reworks and the Program 25 Evaluation Reviews. The spike that we see in

1 the one bar graph for the last four quarters 2 are really due to super S and other PERs that 3 cover a large range of sites. You'll also see 4 in a later slide in that presentation that we 5 have returned a large number of late. dealing with a case load of reworks around 6 7 3,000 and some. We've returned quite a number 8 just recently, another -- with an evaluation 9 letter saying whether we need to -- we don't 10 need to do a rework or we do need to do a 11 rework, so... 12 DR. MELIUS: Yeah, it also looks as if you -some of the older reworks have never been 13 14 returned. At least you have them broken down 15 here by quarter, so for example, in the third 16 quarter of 2004 you received 113 and have 17 returned 42. Now is --18 MR. ELLIOTT: Well, you know, the reworks --19 you're looking at the bar graph. 20 Bar graph, yeah. DR. MELIUS: 21 MR. ELLIOTT: There are some reworks that --22 that we are -- we have not been -- we have not 23 been able to return quickly. We're working 24 through that. I think you'll see our pace pick 25 up very soon on that. But --

1	DR. MELIUS: I mean is is the date the date
2	that they're received or the date that they
3	were originally I mean I done? I mean
4	I'm just trying to understand this is the
5	first time I think at least I recall seeing
6	this graph. Maybe I've
7	MR. ELLIOTT: Well, it's been there every time.
8	DR. MELIUS: Oh, it's been there? I apologize.
9	MR. ELLIOTT: But it the date well, when
10	we receive them, that when we receive them
11	back from DOL, that's when we lo start
12	counting
13	DR. MELIUS: Okay.
14	MR. ELLIOTT: time on ourselves.
15	DR. MELIUS: Okay.
16	MR. ELLIOTT: The date we turn them back over
17	to DOL is when we stop our clock.
18	DR. MELIUS: Yeah, 'cause 'cause in that
19	case, some of these would be four years old.
20	MR. ELLIOTT: No, I don't believe they're four
21	years old.
22	DR. MELIUS: Okay, I see, this is a cumulative,
23	not a
24	MR. ELLIOTT: It's a cumulative graph
25	DR. MELIUS: Okay, okay, I understand. Okay.

1 MR. ELLIOTT: -- and the trend that we're 2 watching there is this late blip of the PER 3 activity, but they're not -- I don't believe 4 that we've got any rework that's over four 5 years old. I don't believe we've got any rework right now that's over a year old. 6 DR. MELIUS: Okay. Maybe you could report on 7 8 that next time, get some data -- just out of 9 curiosity. Thanks. 10 DR. ZIEMER: Other questions for Jeff? 11 (No responses) 12 Okay. Thank you very much, Jeff. We 13 appreciate the update. 14 DEPARTMENT OF ENERGY UPDATE 15 DR. BRANCHE: We have time to catch DOE. 16 DR. ZIEMER: We're pleased also to have Dr. 17 Worthington here with us today, and if you're 18 agreeable, we'll proceed with your 19 presentation. Are we set to go? Yeah. 20 For those of you participating by DR. BRANCHE: 21 phone, if you could please mute your line. 22 you do not have a mute button, then please use 23 star-6 to mute the phone. And then when you 24 are ready to speak, use that same star-6 to 25 unmute your phone. Thank you.

(Pause)

DR. WORTHINGTON: Good morning. Can you all hear me okay? Good. It's always a pleasure to be here and appear before the Board and to meet with many of the counterparts and in some cases to interface with some of the actual workers. This morning I wanted to give you an update -- it really is an update. We don't have any major changes in the program. We're committed to the things we've talked about over the last sessions, and I want to tell you where we are with those.

And before I get started, I probably should start with something that may be of concern to some of you on the phone and to some of you that are actually here today. We have had some -- some funding constraints over the year for the DOE program in terms of being able to deliver the services that we need to deliver. We've worked hard with our counterparts and we've revisited a number of things, and we're pleased to tell you today that we are fairly confident that all the things that we need to deliver this year, that we have the funds to do that and we're working with the sites and the

1 various organizations to make sure that we're 2 being very efficient and effective in delivering those services. 3 And if I can find the right button here, we'll 5 go to the next slide. Okay. Again, as I said, 6 this was really intended to be an update and to 7 talk about where we are with the things that we 8 need to do for this program. As I've mentioned 9 in the past, we have three major 10 responsibilities here, and one -- the first one 11 is the individual claims. And as I go through the various slides today, that certainly is the 12 13 biggest part of our program and we're committed 14 to -- to doing those efforts. 15 We want to provide support to Department of 16 Labor and to NIOSH and the Advisory Board, and to their contractors, to do a number of things, 17 18 including research, 'cause in some cases it's 19 not a very simple activity to be able to deliver the documents. We want to do research, 20 21 retrieval and to provide the various records 22 from the various DOE sites. 23 We want to research issues related to the 24 EEOICPA covered facilities or time frame 25 designation, as appropriate.

The DOE activities -- as I mentioned on the earlier slide, 90 percent of our activities are focused on the individual claims. And we -- you've seen these numbers before and I'll just mention them again to put sort of in perspective the magnitude of the work that we -- we have before us. And that is that typically we do about 8,000 a year employment verifications. Dose documentation for NIOSH, about 5,000 a year. And our DARs, we have about 9,000 a year. So those are the big things and we remain, you know, committed to -- to working on those areas.

Total number of requests. The total number of requests that we had in 2006 -- and what we wanted to do, as we've done in some of these previous meetings, is just to revisit the previous years, get some idea of where we're going, and hopefully be able to make some good and accurate predictions for the future. So in 2006 we had over -- almost 17,000 requests. The total number for 2007 was nearly 22,000. So as you can see, we had certainly an increase -- what we view as a significant increase from 2006/2007, more than 32 percent increase in

that area.

I want to go to the next slide, slide number five. It's a graph, and I think you've seen that graph at some -- some previous presentation. I want to just -- just talk about it just a little bit. And as you can see, we've experienced an upward trend overall in claims, although for some months we had things that went up and went down. But you can see there's certainly been somewhat of an increasing trend in terms of the number of things that have been requested.

The next slide, I think we've had some similar ones before. I want to talk a little bit about that. Here we wanted to kind of depict what we had over nearly the last year, and we looked at that slide and we looked at the data and tried to determine what it's actually telling us.

And we believe that although individual claims have been down so far this year, the number of large-scale record -- research projects are up significantly from last year, and we expect that to be the case for the rest of the year.

But again, we are looking backwards at the kind of requests that we've received were the things

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that we have on our plate right now. And to the best we can, some projections of the -- of things that will come, and then making sure that we work with all the individual sites and with the organizations so that we can deliver those products and deliver those services. I want to talk a little bit about the SECs. They're certainly very important to all of us, and our current research in support of the SEC activities, you see that we have a number of them here -- Fernald, Hanford, Mound, Nevada, Savannah River and Pantex -- and we expect that all of these efforts will be significant, both in the volume of records gathered and the complexity. We've talked about I think at some of these previous meetings the complexity of finding the documents. We have a legacy of many different types of document collection and retrieval processes, and we have to sort of search all of them to be able to come up with the documents. So it remains to be complex. We have not yet been able to consolidate that legacy into one system at the various sites, but we're working on that and trying to be able to deliver what we believe to be a quality

service in a timely manner. And timely certainly constantly being redefined as we find ways to be -- to do this better.

DOE activities, I want to -- I think we've -we've had a slide similar to this before. We
want to talk a little bit about our efforts.
Certainly we are the funding source for these
large and complex activities, but we fund and
coordinate the large scale record retrieval
activities, and there a number of them going on
and many of you are affected by -- you know, by
the work there.

In terms of Department of Labor, they have developed a good process in the site exposure matrix. I think it brings together a lot of information that certainly should be able to facilitate things. And then we have over -- we've completed over 20 in FY07. And those again a large, complex activities that require quite a bit of -- of interfacing.

We want to continue our work with the Board and with their contractor, and we've had technical reviews of NIOSH site profile documents -- I think we've had six over the past year.

And our Special Exposure Cohort, we had six

large projects and they're active right now and we're working on those.

Again, continuing discussions on the things that we're working on with respect now to NIOSH, NIOSH data capture activities. Again, those things can be extensive. These activities are ongoing and NIOSH has been working with up to ten DOE sites in a single month, so that's quite a bit of juggling and coordinating and facilitating for us, but we're working with NIOSH to be able to do that even better.

In terms of special cohorts, again, it's a number of things that we do, both research, record retrieval, various activities, and there are six that are active and current at the moment.

The next slide deals with DOE responsibility for research and maintaining the covered facility database. We have 343 covered facilities, and we recently updated the Dow Madison and the Chapman Valve facilities.

A little bit about the DOE record retrieval activities that are going on. I've listed three of them here on the slide. One is GE

Ohio and then the Westinghouse Atomic and the Stauffer Metals. The Stauffer Metals is not a direct result of anything that NIOSH or the Board is doing. We're doing some work -- routine research to ensure that the covered facilities period and descriptions are accurate. We are researching the covered period for both, as I mentioned, GE and Westinghouse, and that work is ongoing. We have no specific -- because we've not completed that, no specific updates, but just to remind everybody that that work continues.

A little bit -- and I think we talked about

A little bit -- and I think we talked about this before and we're very proud of our DOE Office of Legacy Management. They certainly have a large number of professionals -- certified records managers and senior staff with security clearances, and they're formally trained in requirements of the National Archives. And a lot of the work that we do, we're looking for very old documents and we have to reach back to our archives, and these individuals are certainly experts in that area. They're readily available to us and they have a good understanding of the DOE process, and it's

easy for us to get to them and to help them to
facilitate things. And they have been
supporting our office and we're certainly very
pleased with that and we hope that you're

6 having with them.

A little bit about some initiatives that we have that are ongoing and that we continue to work them and try to make them a mature part of our processes. We think it's very important to have a single point accountability when we can do that, and that you know and everyone will know where to go to get information and to help resolve their problems. And we've named a POC within our office -- Greg Lewis is in the back of the room, he's with me, very active, very important person on this program, and he's coordinating with all the records requests from the Advisory Board and their contractor and with Department of Labor in trying to address any concerns in a timely manner.

benefiting from that interaction that we're

We've been looking at various ways to be able to communicate and to understand and to make our process smoother. We've been holding conference calls with members of NIOSH and

1 their contractors to ensure that these groups 2 are getting the information and support they 3 need from the DOE sites. And I -- again, I talked about our support from Legacy 5 Management. In terms of the initiatives, as I said, we've 6 been closely working with the DOL -- Department 7 8 of Labor federal POCs and their contractors, 9 again on the site exposure matrix project. We 10 think that's important and we want to make sure 11 that things can run smooth there. 12 Each site undertook a comprehensive review and 13 updated their records search procedures. 14 asked them to do that and to find ways to do it 15 better. And as a result of this effort, a 16 number of sites took steps to improve their 17 data-gathering methods and sources. 18 that's good. We're -- all of us are in the 19 mode of continuous improvement in being able to 20 deliver services much better. 21 And of course we think that the training 22 sessions have been good and that we all learned 23 from all of the organizations that participated 24 in the training sessions. 25 We are, as I said, committed to making

improvements, to continuous improvement. One of them is we're committed to providing site experts to participate and contribute to the Advisory Board working groups and conference calls. I believe that we had a request recently from the Rocky Flats staff to participate on the Advisory Board, and I think that that was good. We would offer to do this at request by the Board or NIOSH at any time. We certainly have the experts. They've been working on things. We want to make them available, you know, whenever we can to work through the various issues.

And again, we've been looking for a way to streamline our processes. We're looking for a way that we can gather information and come up with what we would call a draft project plan that would kind of drive the activities and inform the sites about what might be coming up. And we certainly recognize that initially this might slow the process down, but we believe that in a very short period of time it will certainly expedite things, that people will certainly be more aware of what is expected, what kind of documents and the time frames that

might be needed to deliver those documents. It also would provide, we believe, the best possible opportunity for us to minimize any, you know, overlaps or duplications that we would need and so we think that -- just bear with us as we work through this. We believe it certainly is the right way to go.

At this point I am available -- and again, Greg Lewis is in the back, he's available -- for any questions that you might have about our process. I want to just reiterate what I said in the very beginning and that is that DOE is committed to delivering these services and working with the various organizations. We have what we believe identified the funds within our existing program to be able to fund these efforts and to not have any significant delays in getting the materials to the various organizations.

I thank you for your attention and I welcome questions and discussion at this time.

DR. ZIEMER: Thank you very much, Dr.

Worthington. Let me start off by asking you if
you would elaborate on what you referred to as
the covered facilities database. What's --

1 just -- could you describe briefly the kinds of 2 things that are in that? I assume it's a 3 database that you are building as you retrieve records for these programs for Labor and NIOSH. 5 Is that correct? DR. WORTHINGTON: That's correct. I'll start 6 7 talking and then Greg is going to walk up to the mike and is going to provide some 8 9 additional clarification on this because he's 10 worked quite a bit with that. So Greq, if you 11 want to go on. 12 I think there's about 358 MR. LEWIS: Sure. 13 facilities in there and this is --14 DR. ZIEMER: A little closer to the mike, Greg. There's about 358 covered 15 MR. LEWIS: 16 facilities in there and it was originally 17 developed about four or five years ago, but 18 we've been constantly updating it ever since at 19 -- you know, based on questions from DOL and 20 NIOSH, or whoever's raised issues. So at this 21 point we have it developed but, you know, as 22 different questions are raised -- arise we, you 23 know, have been making changes and doing 24 further research. You know, some of the --25 like we've just recently made changes to

Chapman Valve and to Dow Chemical based on, you know, activities and research for the Board.

DR. WORTHINGTON: We look for every opportunity to make sure that that list is actually accurate, so at any point that we've generated new information and there's consensus and final decisions have been made, we go back and revisit that list to make sure that it's accurate.

DR. ZIEMER: Okay, very good. If I might ask one question on budgeting, there was some indication earlier this year, or maybe toward the end of last year, that because of the continuing resolution situation you pretty much had to focus on primarily the records retrieval and then secondarily the other issues. Is that pretty much corrected now for --

DR. WORTHINGTON: Yes.

DR. ZIEMER: -- from your point of view?

DR. WORTHINGTON: Yes, that is pretty much corrected. As you indicated, the continuing resolution certainly offered some unique challenges to us, and we were at a point that we had to focus first -- because we did have limited funds, we had to focus -- and the way

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that we had to issue those funds, a little bit, a little bit, a little bit, so we certainly had to focus on the individual claims. This is a very recent accomplishment in terms of being able to work with the sites in determining what requests were actually at the site and to look forward to things that might come, and to look at the funding that we had within our organization. And we believe that we have addressed that concern and we will be able to not have any significant delays in any of the services that we have to deliver. And we are probably in the -- and I believe Greg's correct -- in the next round of sending money to the sites, which I believe will be in the May time frame. I'm not sure that we've mis-- then we would -- based on what we have on the plate and what they're expected, provide additional funds and work with them throughout the course of the year. But we are monitoring things very carefully to make sure that we don't have to again, you know, ask for, you know, delays because we -- of funds. But we're in -- in pretty good shape at this point.

MR. LEWIS: Yeah, that's exactly right. I mean

1 we're allocating funds based on the need and 2 it's obviously claims-driven, as well as driven 3 by different research efforts and projects, you 4 know, for SEC research, et cetera. That can --5 that can constitute a significant effort so we have to make sure that we have the funds in the 6 7 right place, depending on the need at the 8 various sites. 9 DR. ZIEMER: So I -- and I think now you've 10 probably answered my final question. That has 11 to do with whether the sites themselves have 12 funding or you are funding the sites for the 13 work. 14 DR. WORTHINGTON: We are --DR. ZIEMER: 15 It sounded like (unintelligible). 16 DR. WORTHINGTON: We are funding the sites for 17 the work. We are monitoring very carefully 18 their requests and what they have already --19 DR. ZIEMER: That is, it's --DR. WORTHINGTON: -- and making sure --20 21 DR. ZIEMER: -- out of your budget and not in 22 their budget request. They -- they -- do they 23 -- they may tell you what -- what they need, 24 but you fund it out of your office. 25 DR. WORTHINGTON: And we provide the funds to

1 the sites to be able to do that. 2 DR. ZIEMER: Thank you. 3 DR. WORTHINGTON: Yes, it is an HSS-funded 4 activity. DR. ZIEMER: Okay, I think we have questions 5 6 here. Dr. Melius and Josie Beach. 7 DR. MELIUS: Josie was first. 8 DR. ZIEMER: Josie, go ahead. 9 MS. BEACH: I was wondering if you could give 10 us an update on the medical records retrieval 11 for Los Alamos. 12 DR. WORTHINGTON: It certainly still is a work 13 in progress. A week ago -- maybe a week or ten 14 days ago -- we had a -- what we viewed as a 15 very good face-to-face meeting with the 16 hospital staff and the corporate organization 17 and our organizations to talk through next 18 steps. We believe we have a path forward for 19 actually cleaning up the records and packaging 20 the records, and then relocating those records to either our natural -- one of our archives or 21 22 to a space at the Laboratory yet to be -- to be 23 determined. We understand their schedule in 24 terms of when they need to have us out of the

warehouse, so we believe we have a path

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1 forward. But there is some specific details 2 that have to be worked in terms of the overall 3 cost associated with that and how that cost 4 would be provided, and then to finalize the 5 actual plan and the contractor that we would 6 use to be able to -- again, to clean up the 7 records, sort them, and then to, you know, have 8 them repackaged and in another location that 9 then would be easy for people to retrieve the records that they were -- needed for any claims 10 11 or whatever. 12 MS. BEACH: Thank you. 13 DR. ZIEMER: Dr. Melius. 14 DR. MELIUS: Yeah, could you give an update on 15 the Hanford situation, please? I note that 16 we've been waiting I think at least six months 17 to a year for records and significantly holding 18 up any progress on that site. 19 DR. WORTHINGTON: I'm not sure that I 20 understand the question. Is the question --21 DR. MELIUS: Well, when will we have access to 22 the records that have been requested at 23 Hanford? 24 DR. WORTHINGTON: Do you understand the actual

question, Greg, in terms of --

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MR. LEWIS: Yeah, I mean there's -- you're -you're in the process of research based on the SEC determination and so we're -- we've been trying to help facilitate the records-gathering process. And again, as -- as Pat mentioned, we did have some funding issue due to the continuing resolution late last year and early this year. We have worked past those and, you know, at that point, to -- hopefully to make the process more efficient and streamline it, we had requested that the groups involved in the research prepare one single consolidated plan that would let us know how, from front to back, they were, you know, planning on obtaining the records and what -- you know, what types of steps that they would need to identify what records they needed and then look at them and then -- and then gather them and keeping in mind security issues and -- and personnel and things like that. So they have -

DR. WORTHINGTON: I think --

MR. LEWIS: -- put together such a plan and then the -- the only -- we are working through some classification and security issues right

1 now, making sure that the right steps are in 2 place to facilitate the data-gathering within, 3 you know, our -- our limits with, you know, classification and security. 5 DR. WORTHINGTON: I think I understand your 6 question now. Thanks, Greq, for the 7 clarification. As I mentioned, we did have 8 those funding constraints. We worked through 9 that. We've -- bringing all the parties 10 together to come up with a plan on what is 11 needed so we can move forward, and we are 12 addressing those security concerns that we have so -- and I admit, we have had delays because 13 14 of all of those things, but we have a good path forward and believe we can move out on those 15 16 things. 17 DR. MELIUS: But when will that path forward 18 deliver some records, I guess is my question. 19 DR. WORTHINGTON: The question, Greg, is when will that path forward deliver some records? 20 21 It's my understanding --22 Is there a time table that --DR. ZIEMER: 23 DR. WORTHINGTON: That we are --DR. ZIEMER: -- has been established? 24 25 DR. WORTHINGTON: -- working on the records as

1 we speak, and --2 MR. LEWIS: Yeah, we believe we have a meeting 3 at the end of this week with our Hanford people 4 and our headquarters classification folks to, 5 you know, finalize our path forward. And after that, we should be moving forward with the plan 6 7 that was put together by the NIOSH and SC&A 8 team. So you know, as soon as next week we 9 should be able to start the data-gathering 10 process. 11 DR. MELIUS: Okay. 12 DR. ZIEMER: Okay, thank you. DR. WORTHINGTON: Are you -- is that --13 14 DR. MELIUS: I just -- I'll believe it when I 15 see it, so -- I'm not going to ask any more 16 questions now. 17 DR. WORTHINGTON: But we are committed to -- to 18 moving out on this. As I said, we've overcome 19 a number of hurdles, and so we -- we want to 20 move forward, and we understand the importance 21 of doing that. 22 DR. ZIEMER: Right. You -- you recognize we --23 we face a little frustration here. We know you 24 have the same frustrations, the funding drives 25 a lot of this. But in turn, our clientele also

1 get frustrated 'cause they think we're not 2 doing our job in getting documents reviewed and 3 so on. So it's a kind of a domino --4 DR. WORTHINGTON: And it certainly has --5 DR. ZIEMER: -- effect all the way 6 (unintelligible). 7 DR. WORTHINGTON: -- been longer than any of us 8 would have liked, but in terms of looking for 9 the funds --10 DR. ZIEMER: We'll appreciate whatever can be 11 done to expedite certainly that, and I'm sure 12 there will be others, particularly the big 13 complex sites. I'm not sure -- Savannah River 14 may face the same thing as we get into that further, too. 15 16 Okay. Let's see if we have any other questions 17 for Dr. Worthington today -- or the DOE in 18 general. 19 (No responses) 20 Okay, thank you again. 21 DR. WORTHINGTON: Thank you again. 22 DR. ZIEMER: We appreciate not only the update, 23 but participation of you and your staff in the 24 program and your attendance at the meetings as 25 well.

1 DR. WORTHINGTON: It's always a pleasure. 2 Thank you. 3 DR. ZIEMER: I think we'll go ahead and take 4 our break now. Let's take a 15-minute break 5 and then we'll resume. 6 (Whereupon, a recess was taken from 10:15 a.m. 7 to 10:40 a.m.) 8 DR. ZIEMER: We are ready to resume our 9 deliberations. First a comment on phones --10 Christine. 11 DR. BRANCHE: Good morning. If everyone 12 participating by phone could please mute your phones. And if you do not have a mute button, 13 14 please use star-6. And when you're ready to 15 speak, you can use that same star-6 to unmute 16 your phones. By muting your phones you're 17 helping us maintain the quality of our court 18 reporting. Thank you so much. 19 This -- this is Bob. MR. PRESLEY: 20 DR. BRANCHE: Thank you. 21 DR. ZIEMER: We have about 20 minutes before 22 our Kellex presentation and we want to keep 23 that as a time certain because of participants 24 who will join us by phone, so I'm going to use 25 the 20 minutes to begin some of our working

time.

WORK GROUP REPORTS

I'd like to just take a few of the workgroup reports, and I'll ask Dr. Branche just to go down the list. We'll take them in the usual order. Workgroup chairmen, when it's your turn you can give us an update report. If there's been no action since your last meeting, you can so report. So let's go through them -- but hang on just a moment.

(Pause)

Just workgroups. Just workgroups, not the subcommittee.

DR. BRANCHE: Because we already have time allocated for Blockson and Chapman Valve, I'm going to skip over those and go to Fernald.

MR. CLAWSON: Yeah, with Fernald workgroup, we met with NIOSH in Cincinnati about a week and a half ago. We've still got some outstanding issues, but we're working through the process.

There -- many white papers have been provided to us and so forth and we're just -- we're proceeding forward as we speak. But we met a week and a half ago and we're waiting for some information back and then we'll set up the next

1 workgroup and proceed on. 2 DR. BRANCHE: Los Alamos -- Los Alamos National 3 Laboratory, site profile and Special Exposure 4 Cohort, Mr. Griffon chair. MR. GRIFFON: Yeah, I don't really have a 5 6 report, I -- I -- again, I'd like to ask the 7 status from NIOSH's side. We've been waiting 8 on an updated site profile. We've sort of held 9 off on our review until we got an update on that, and I don't know where NIOSH stands on 10 11 that right at this point or... Or maybe we can 12 get that tomorrow if someone's not here. 13 anyway, the -- the LANL workgroup's been on 14 hold, but I think we need to get back to the --15 and the -- the question really is the second 16 time period. We've already addressed one time 17 period, but I think we need a follow-up on the 18 second time period and we're waiting on updates 19 to the site profile, I believe, so... 20 DR. ZIEMER: Okay. Jim, are you giving a --21 DR. NETON: (Off microphone) (Unintelligible) 22 MR. GRIFFON: Yeah, Stu -- Stu --23 DR. ZIEMER: Okay, a brief pow-wow here and 24 then we'll get an update -- a quick status 25 report, perhaps, from Stu or someone.

1 DR. NETON: Unfortunately we have no one here 2 right now that can answer that question but 3 we'll -- we'll research it and get back to you 4 shortly. 5 Well, we -- we understand that, DR. ZIEMER: 6 from the workgroup's point of view, they're --7 they're simply awaiting that for the next step. 8 MR. GRIFFON: Yeah, and I -- I will follow up 9 on -- I'll (unintelligible) with the NIOSH 10 folks and see when we can -- as soon as we can, 11 we're going to schedule a workgroup meeting on this, though. And we'll let all -- all the 12 13 interested parties know about it, so... 14 DR. BRANCHE: I'm going to skip over Linde 15 because I don't see Jeff Kotsch and he was 16 going to supply some information. 17 Mound, Ms. Beach, chair. 18 MS. BEACH: Yes, we were -- we held our first 19 workgroup meeting. Mou-- SC&A and NIOSH were 20 able to go through the matrix and clarify some 21 of the -- some of the concerns with the matrix. 22 We have not scheduled another meeting but we 23 hope to do that shortly. 24 DR. BRANCHE: Nevada Test Site profile, Mr. 25 Presley chair.

1	MR. PRESLEY: This is Robert Presley. We are
2	in trying to set up our next meeting, which
3	will be sometime around the 9th through the
4	21st of May. NIOSH sent contractor to NTS
5	to pick up some data that was needed for our
6	final closure for comment 11, and
7	(unintelligible) that information comes back
8	and they get it into a final form, we will be
9	ready to meet and hopefully (unintelligible)
10	that's at our next meeting in St. Louis.
11	DR. BRANCHE: Thank you. We had an extensive
12	discussion about procedures yesterday.
13	Rocky Flats site profile and Special Exposure
14	Cohort SEC petition, Mr. Griffon, chair.
15	DR. ZIEMER: I I told Mark that may be a
16	little more extensive and we'll
17	DR. BRANCHE: Hold off?
18	DR. ZIEMER: delay till tomorrow on that
19	one, yeah.
20	DR. BRANCHE: All right. Dr. Melius is out.
21	Savannah River Test (sic) Site profile, Mr.
22	Griffon, chair.
23	MR. GRIFFON: Yeah, at this point on Savannah
24	River we just have not had time to re to
25	schedule a follow-up workgroup meeting, so

1 that's another one that's been on hold a little 2 bit. No update at this point. 3 DR. BRANCHE: We're going to hold off on the 4 subcommittee as well, Mark? 5 MR. GRIFFON: (Off microphone) (Unintelligible) DR. BRANCHE: Worker outreach, Mr. Gibson, 6 7 chair. 8 MR. GIBSON: We haven't had any significant 9 activities recently. We're still -- NIOSH is 10 in the process of modifying a procedure on 11 worker outreach, and also their database where they track comments. So we're waiting on that. 12 13 DR. ZIEMER: Could I ask a question here? And 14 incidentally, Board members, if you have 15 questions as we go, that'll be fine. 16 Mike, some of your -- you and some of your 17 committee did attend some outreach meetings, 18 did you not, in the last month or so? 19 just give us an update on that. 20 Yeah. We've -- different members MR. GIBSON: 21 of the group have attended different types of 22 meetings. As you know, NIOSH holds different 23 type of outreach meetings. I think as I 24 mentioned the last meeting, one I came to down 25 here in Tampa back in February -- late February

was a worker outreach on SEC process, and
Laurie Breyer and Denise did a real fine job at
expl-- I think explaining to the claimants
about the SEC process, the steps to go through
and seemed to be real well received and if
there's any of the other workgroup members want
to talk about meetings they've attended and how
they felt they went...

DR. ZIEMER: Wanda Munn, hang on just a minute. Again, remind folks on the phone, please mute your phone. We're getting some back talk and background conversations.

Ms. Munn.

MS. MUNN: I spent an interesting three days at Argonne East with our contractor's team looking through the extensive records that they have there, interviewing some of the workers and talking to the medical personnel on that site - which of course has such an extensive history. It goes all the way back, literally, to CP-1. So it was an extremely informative and I think most productive visit from our workgroup's point of view.

DR. ZIEMER: Thank you.

DR. BRANCHE: That's actually the end of the

1 list. I'm sorry, Dr. Neton has some 2 information. 3 DR. NETON: Yeah, we have an update already on Los Alamos site profile. 5 MR. RUTHERFORD: I apologize, I stepped out of 6 the room for one minute and there you go. 7 We actually had a schedule for our contractor -8 - for ORAU to provide a draft document in March 9 for us that addressed the feasibility of post-10 '75, and the ultimately what would -- we would 11 use that document to update the site profile. 12 We've reviewed that document. They're in comment resolution with that document. We are 13 14 looking at adding a little more to that document, and we do have a schedule for 15 completion of that. I don't have the schedule 16 17 with me right now, but as soon as that is 18 available we will provide that to the 19 workgroup, and I expect that to be completed 20 within the next cou-- within the next month or 21 month and a half, I would suspect. Okay? 22 Is Mr. Kotsch from the Department DR. BRANCHE: 23 of Labor in the room? 24 UNIDENTIFIED: He's not. 25 DR. BRANCHE: Okay.

1	DR. ROESSLER: I could do
2	DR. ZIEMER: Do your part and then
3	DR. BRANCHE: Okay, so Linde
4	DR. ZIEMER: We we can get the statistics
5	afterwards.
6	DR. BRANCHE: All right. Linde, Dr. Roessler,
7	chair.
8	DR. ROESSLER: I had planned to put this
9	together tonight so this is off the top of my
10	head, but we expect soon to have completed the
11	site profile review. We started this with
12	SC&A's help and had our first meeting in March
13	'07. We had 22 issues to deal with. By
14	November '07 we had reduced it to six issues,
15	and by January '08 we had just one remaining
16	issue to look at.
17	This has to do with burlap bags on-site. The
18	bags were used to deliver ore to the site.
19	They were then emptied and apparently stored.
20	These empty bags, though, would have had some
21	residual radioactivity in them.
22	The issue came up because of a worker who
23	recalled that some of the bags were in a
24	certain location at a certain time. So the
25	assignment to NIOSH and ORAU at our last

workgroup meeting was for them to model the situation to be able to calculate doses to persons who might have been on or near the bags.

The white paper that ORAU or NIOSH was working on came to all of us, and all the Board members received it, I think last week. Joe Guido completed that. So now we're waiting for SC&A to take a look at it and if they feel that this handles this issue, then we will have the site profile completed.

So then I assume the next step will be for NIOSH to eval-- well, for -- the next step then will probably depend on what Jeff has to tell us. So as far as I can see, I guess the bottom line is that we hope to have the site profile review completed.

DR. ZIEMER: We -- we didn't -- we will have a report on Chapman Valve later this afternoon.

We will have a report on Sandia Livermore later this afternoon, as we will for Hanford on the Hanford -- and those are part of the SEC petition updates. But for -- let me report, since I'm part of the Hanford workgroup and Dr. Melius is the chair, and he can update that

1 further if he wishes when he returns, but as 2 was already indicated during the DOE 3 discussions with Dr. Worthington, the Hanford workgroup basically is awaiting some documents 5 from DOE. So -- thus that workgroup has not met since our last meeting, so basically the 6 7 only thing to report from the workgroup is that 8 they are awaiting those documents for -- for 9 further action. 10 DR. BRANCHE: This might need -- thank you, Dr. 11 This information might need to be Ziemer. 12 repeated when we get to that time this 13 afternoon because there -- some of the people 14 who plan to participate I believe were not only 15 members of the petitioner and other workers, 16 but also members of Congress. 17 DR. ZIEMER: Yes, thank you. 18 MR. GRIFFON: Paul, can I ask, just -- just 19 from the subcommittee standpoint, I -- I can do 20 the update --21 DR. ZIEMER: This is the subcommittee on dose 22 reconstruction --23 MR. GRIFFON: Dose reconstruction. 24 DR. ZIEMER: -- which we -- always is part of 25 our workgroup review, but they're -- yeah.

1 MR. GRIFFON: And I -- I wou-- I'm just going 2 to lay -- I mean we can certainly do the 3 primary update tomorrow, but I -- I look in our provided materials and I don't see the 4 5 subcommittee information so I think maybe that 6 I need to get that to the Board members. 7 mean the -- we should be able to move on a 8 tenth set of cases, and I'm assuming that Stu 9 Hinnefeld updated -- we -- at the last 10 subcommittee meeting we -- we went through a 11 list of -- of possible cases and we gave that 12 to Stu, as is our normal process. And then Stu was going to provide more detailed information, 13 14 and I don't see that matrices (sic) in our --15 in these handouts, so I'm wondering if we ever 16 got those. 17 DR. BRANCHE: No, he --18 MR. GRIFFON: Did Stu step out of the room 19 again? 20 UNIDENTIFIED: Yes, he did. 21 MR. GRIFFON: Okay. 22 DR. ZIEMER: Well, in the meantime --23 MR. GRIFFON: It's like he's avoiding me, huh? 24 DR. ZIEMER: In the meantime, Jeff is back --25 is he back?

1 DR. BRANCHE: Yes, he is. 2 DR. ZIEMER: Jeff, we just had a brief report 3 on -- from Dr. Roessler on Linde. Are you in a 4 position to give us those statistics, that you 5 referred to in your report, on the Linde site? 6 MR. KOTSCH: Yeah. I mean they weren't really 7 statistics, and I have to apologize, I forgot 8 to mention also that Labor is here both in the 9 form of the Jacksonville Office and our 10 Resource Center from Savannah River on the 11 other side of the building over by where the 12 NIOSH (unintelligible) are. 13 DR. ZIEMER: Get a little closer to the mike, 14 Jeff. 15 MR. KOTSCH: Sure. 16 DR. ZIEMER: Or raise it up a little bit there. 17 MR. KOTSCH: Now for Linde I was just going to 18 update you -- I think during the phone call --19 the telephone meeting of the Board in February 20 we discussed -- or I presented the rationale or 21 the background for the change in site 22 designation for Linde Ceramics where it went 23 from strictly AWE to four of the buildings

becoming DOE facilities, and then I think

Building 14 remaining as an AWE. And at that

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1 point I -- I -- the discussion was what happens 2 for those -- we were going to continue to 3 review employees that were -- worked strictly 4 within the residual contamination period. 5 so that was -- that was just the essence of the update that I -- Friday I had a meeting with 6 7 our -- our legal staff and they noted that for 8 the four buildings -- Buildings 30, 31, 37, 38 9 -- that were switched from AWE to DOE 10 designation, that -- those four buildings, 11 based on further review of the 2004 amendments 12 to the Act, that workers in those buildings who worked only during the residual period --13 14 residual radiation period are also eligible for 15 Part B's benefits as atomic weapons employees, 16 even though they have changed -- they, we --17 even though we, Labor, have changed the status 18 of the buildings as a DOE facility. 19 So I think the issue was, when we last 20 discussed it was what happened to those people 21 who were solely employed during the residual 22 period, so now they will be covered under Part 23 В. 24 DR. ZIEMER: Okay, thank you. 25 MR. KOTSCH: Okay?

1 DR. ZIEMER: Any questions on that? 2 Roessler. 3 DR. ROESSLER: I quess I expected maybe you 4 were going to say something about the SEC 5 petitions for Linde. I understand that the 6 petitioners have moved forward on that? 7 MR. KOTSCH: For --8 DR. ROESSLER: Perhaps that's still in 9 progress. 10 MR. KOTSCH: I -- I'm not aware. 11 MR. RUTHERFORD: I can --12 MR. KOTSCH: Okay. 13 MR. RUTHERFORD: I don't know that Jeff can 14 answer the SEC -- we do have SEC petitions for 15 Linde Ceramics that we are in the process right 16 now in the qualification phase for the 17 operational years that were past the al-- the 18 SEC that we've already designated, and for the 19 residual period. Now this does affect that 20 petition because we had -- we were not 21 operating under that same, you know, knowledge 22 that Jeff just gave us, so we're going to have 23 to go back and look at that for that residual 24 period. 25 DR. ZIEMER: Okay, thank you. Further comment? Jeff.

MR. KOTSCH: I'm sorry, I just needed to add a disclaimer that I remembered my legal people mentioned to me on Friday concerning that -- the statement I made about the coverage at Linde. That only applies -- I mean their evaluation only applies to, at the current time, the Linde Ceramics. You know, the way they interpret that site.

DR. ZIEMER: Okay. Thank you for clarifying that.

KELLEX/PIERPONT SEC PETITION

Okay, we're going to now move to the discussion of the Kellex-Pierpont SEC petition. Dr. Glover from NIOSH is going to present the NIOSH evaluation report. I wanted to check first to see if [name redacted] is on the line. Or [name redacted].

(No responses)

My notes indicate that [name redacted] or [name redacted] may wish to be on the line.

DR. BRANCHE: Given that we've asked people to do star-6, maybe you need to dial star-6 in order for us to hear you if you're speaking.

DR. ZIEMER: Or put -- take your mute button

1 [names redacted], are either of you on 2 the line? 3 (No responses) 4 If you are, we're not hearing you. 5 DR. BRANCHE: She's going to -- our public health advisor's going to call. 6 7 DR. ZIEMER: We'll --8 Mr. Presley, are you on the line? DR. BRANCHE: 9 (No responses) 10 Is anybody on the line? 11 DR. ZIEMER: Have we -- have we lost -- does it 12 show whether people are on --13 MR. PRESLEY: (Unintelligible) 14 DR. ZIEMER: Oh, Pres-- okay, Robert, you're on 15 the line, okay. 16 MR. PRESLEY: (Unintelligible) a tremendous 17 amount of static (unintelligible) that's just 18 got on there. 19 DR. ZIEMER: Okay. Well, we'll ask again if 20 [names redacted] are on the line. We have 21 somebody trying to reach them right now. We'll 22 wait just a moment, give them the opportunity, 23 'cause they may want to hear the presentation 24 as well, so we'll wait just a moment. 25 (Pause)

1 DR. BRANCHE: Dr. Ziemer, go ahead and get 2 started. 3 DR. ZIEMER: Okay. 4 UNIDENTIFIED: Hello? 5 Hello. Did -- we'll check again. DR. ZIEMER: 6 Did [names redacted] -- are you on the line? 7 (No responses) 8 Apparently not, but we're going -- then I've 9 been given the signal that it's okay to go 10 I quess there was some uncertainty as 11 to whether they actually wanted to be present, 12 but anyway, here we are with the petition for 13 Kellex. Sam Glover. 14 Thank you, Dr. Ziemer. Can you DR. GLOVER: 15 hear me okay? All right. 16 So I did forget to put -- this is the -- Number 17 0100, the hundredth SEC petition, so 18 (unintelligible). This is the Kellex-Pierpont 19 facility for the Special Exposure Cohort 20 petition. This is an 83.14. We evaluated this 21 petition in accordance with the 83.14 rule. 22 The petition was submitted by a claimant whose 23 dose reconstruction could not be completed by 24 NIOSH due to lack of dosimetry-related 25 information. The claimant was employed at

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Kellex-Pierpont from 1943 to 1946. Keeping with the 83.14, NIOSH -- determination that it is unable to complete a dose reconstruction for any EEOICPA claimant is qualified basis for submitting an SEC petition, and currently there are four claims at -- when we submitted -- when we prepared this SEC petition analysis at NIOSH with Kellex-Pierpont employment during this class period. From 1943 to 1953 Kellex-Pierpont was classified as an Atomic Weapons Employer facility. It was first established by the M.W. Kellogg Company in 1943 to design and construct the K-25 plant. It's approximately 43 acres, with about 20 buildings. Radioactive work was conducted in only one of those facilities. 1951 Kellex-Pierpont merged with Vitro, which then become the Vitro Corporation of America. If you look through the records you will note that it's often referred to as just Kellex. Pierpont is actually part of a mini-ma-- a mini-mall that was added later on, so it's often discussed as the Kellex-Pierpont property in the later time frame, but in -- if you look at the actual early documentation you'll see it

1 often referred to as just Kellex. 2 One more background, they conducted design, 3 engineering and research on diverse radiological programs including gaseous 5 diffusion pilot studies, solvent extraction of 6 uranium from reactor wastes associated with the 7 Hanford processes, and also solvent extraction 8 of valuable components from low-grade wastes. 9 Radiological operations were completed by 1952, 10 and the facility was demolished in 1953. 11 Data capture efforts involved searches at the 12 Germantown offices, multiple visits to the National Archive and Records Administration 13 14 facilities in Atlanta and Kansas City; the 15 Fernald legal database/OpenNet/Nuclear 16 Regulatory Commission Agency Wide Documents 17 Access and Management System, the ADAMS system; 18 also DOE Office of Scientific & Technical 19 Information, OSTI. 20 Furthermore, inquiries were made with the State 21 of New Jersey. Kellex is a company that no 22 longer exists so they obviously could not be 23 contacted to request additional records. 24 relevant -- all records relevant to the Kellex-25 Pierpont petition have been uploaded to the

SRDB.

The radiological operations were conducted in what was known as Building 11, also known as the Kellex-Pierpont -- Kellex Laboratory. The initial mission of the -- of Kellex was to develop the barrier technology for gaseous diffusion. Numerous documents provide shipment evidence of UF-6 canisters to the site. Other doc-- other documents establish operations using ores, ores residues and pilot projects on mixed fission products conducted at the facility.

Information related to the radiation exposure period, internal sources of exposure include significant uranium research conducted on-site, and possible enrichment activities associated with the K-25 pilot studies. Research included AEC-funded research on uranium ore and metals, K-65 and Q-11 residues, with enhanced thorium, radium and radon levels. There was thorium work and fission product operations. And also we had ore and ore byproducts, uranium, possible enriched uranium, and other PUREX type wastes associated with Hanford.

External sources of exposure would be the beta

1 and photon sources, primarily from the uranium 2 and thorium progeny. 3 For data, none of the four claims have bioassay 4 data in the files. From our broad scope 5 searches we have identified 25 uranium 6 urinalysis records for a few individuals in the 7 1950 to '51 period, so nothing predating that 8 for a facility that started in 1943. 9 a few radon breath samples from 1951 for a 10 single employee. 11 For external monitoring data, badging results 12 are available, at least in part, from the 1948 13 through 1953 operations. One of the four 14 current claimants has external dosimetry information in the file. 15 16 Some workplace monitoring data is available. 17 There are some health physics reports in the 18 1950s discussing positive smear readings and 19 locations. There are some evidence of air 20 sampling, primarily for radon. And mostly 21 these were general area samples. Again, these 22 were limited to the 1951 and forward time 23 frame. 24 Feasibility of internal dose reconstruction, 25 NIOSH has obtained bioassay results for only a

handful of claimants or individuals in two very small time frames. Based on the diverse scope of source terms, coupled with the lack of operational data, NIOSH has determined that internal dose cannot be reconstructed.

Lack of information regarding source term location and usage leads NIOSH to conclude (sic) all employees at the Kellex-Pierpont facility in the SEC class definition.

Obviously this requires a health endangerment determination.

The evidence reviewed in this evaluation indicates that some workers in the class may have accumulated chronic radiation exposures through intakes of radionuclides and direct exposure to radioactive materials.

Consequently, NIOSH is specifying that health

may have been endangered, with the parameters - for those workers covered by this evaluation
who were employed for a number of work days
aggregating at least 250 work days within the
parameters established for this class, or in
combination with work days within the
parameters established for one or more other
classes of employees in the SEC.

1 NIOSH's proposed class is all AWE employees who 2 worked at the Kellex-Pierpont facility in 3 Jersey City, New Jersey from January 1, 1943 through December 31st, 1953 for a number of 4 5 work days aggregating at least 250 work days 6 occurring either solely under this employment 7 or in combination with work days within the 8 parameters established for one or more classes 9 of employees in the SEC. 10 As a final kind of summary, the period January 11 1 for -- to -- January 1st, 1943 through 12 December 31, 1953, NIOSH finds that it cannot 13 estimate radiation doses -- radiation doses 14 cannot be reconstructed for compensation 15 purposes. The feasibility is no; the health 16 endangerment is yes. 17 Thank you, Sam. Now as I DR. ZIEMER: 18 understand it, there's 20 buildings, only one 19 of which involved work with radioactive 20 materials. But the -- the class definition 21 covers everyone who worked, regardless of the 22 20 buildings. Is that correct? 23 DR. GLOVER: It's -- there's really no way to -24 25 DR. ZIEMER: We don't know --

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1	DR. GLOVER: (unintelligible) with class
2	titles and and it's very difficult to tell.
3	DR. ZIEMER: So there's no records, is what
4	you're saying, to indicate that that they
5	would be restricted from entering that building
6	if they were assigned to a different building.
7	Is that
8	DR. GLOVER: That's correct.
9	DR. ZIEMER: Thank you. Wanda Munn.
10	MS. MUNN: I'm prepared to move that we accept
11	the NIOSH recommendation for this SEC.
12	MR. CLAWSON: Second it.
13	MR. PRESLEY: This is Bob Presley. I second.
14	DR. ZIEMER: Mr. Presley has seconded it. I
15	think that that Mr. Griffon is prepared to
16	read a formal form of that motion, if that's
17	agreeable to Wanda Munn as a friendly
18	amendment.
19	MS. MUNN: I was sure someone would have it.
20	DR. ZIEMER: Here is the motion.
21	MR. GRIFFON: Yeah, Jim actually handed this to
22	me, so this is the motion, the same format that
23	we're all used to.
24	The Board recommends that the following letter
25	be transmitted to the Secretary of DHHS within

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21 days. Should the Chair become aware of any issue that in his judgment would preclude the transmittal of this letter within the time period, the Board requests that he promptly informs the Board of the delay and the reasons for this delay, and that he immediately works with NIOSH to schedule an emergency meeting of the Board to discuss this issue.

The Advisory Board on Radiation and Worker Health, the Board, has evaluated SEC Petition 00100 concerning workers at the Kellex-Pierpont facility in Jersey City, New Jersey under the statutory requirements established by EEOICPA and incorporated into 42 CFR Section 83.13 and 42 CFR Section 83.14. The Board respectfully recommends Special Exposure Cohort, SEC, status be accorded to all AWE employees who worked at the Kellex-Pierpont facility in Jersey City, New Jersey from January 1st, 1943 through December 31st, 1953 for a number of work days aggregating at least 250 work days, occurring either solely under this employment or in combination with work days within the parameters established for one or more other classes of employees in the SEC.

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1 The Board notes that although NIOSH found that 2 they were unable to completely reconstruct 3 radiation doses from -- for these employees, they believe that they are able to reconstruct portions of the external radiation doses and the occupational medical dose. 6 7 The recommendation is based on the following 8 The Kellex-Pierpont facility was 9 involved in early research and development work 10 for the manufacture of atomic weapons. 11 was unable to locate sufficient monitoring data 12 or information on radiological operations at 13 these laboratories in order to be able to 14 complete accurate individual dose 15 reconstructions involving internal exposures 16 throughout the time period the facility 17 operated. The Board concurs with this 18 conclusion. 19 NIOSH determined that the health -- that health 20 may be -- may have been endangered for the 21 workers exposed to radiation at the Kellex-22 Pierpont facility in the Jersey City, New 23 Jersey during the time period in question. 24 Board concurs with this determination. 25 Enclosed is the supporting documentation from

1 the recent Advisory Board meeting held in 2 Tampa, Florida where this Special Exposure 3 Cohort class was discussed. If any of the 4 items are unavailable at this time, they will 5 be -- they will follow shortly. 6 Sam, could you clarify one thing. DR. ZIEMER: 7 The way we have it here, it only refers to the 8 internal doses not being reconstructed. You --9 your slides didn't give us the -- the usual 10 chart that show-- are you saying -- the 11 implication here is that external can. 12 there -- there are some -- you said there were 13 some external monitoring but not complete, it 14 appears. 15 This is part of the 83.14 16 difference in how we usually present the end of 17 that about what we can and cannot do. 18 believe we can reconstruct, at least in part, 19 the external doses. We have the -- a number of 20 records (unintelligible) --21 So it's sufficient just to say DR. ZIEMER: 22 internal (unintelligible) anyway. You may be 23 able to do external. 24 DR. GLOVER: Yes. 25 DR. ZIEMER: Thank you. So that is the motion

1	that's before us. Any discussion?
2	(No responses)
3	If not, we'll vote by roll call and we will
4	also get the vote later from Mr. (sic) Lockey
5	and Dr. Melius. Here's the roll call.
6	DR. BRANCHE: Ms. Beach?
7	MS. BEACH: Yes.
8	DR. BRANCHE: Mr. Clawson?
9	MR. CLAWSON: Yes.
10	DR. BRANCHE: Mr. Gibson?
11	MR. GIBSON: Yes.
12	DR. BRANCHE: Mr. Griffon?
13	MR. GRIFFON: Yes.
14	DR. BRANCHE: Ms. Munn?
15	MS. MUNN: Yes.
16	DR. BRANCHE: Mr. Presley?
17	MR. PRESLEY: Yes.
18	DR. BRANCHE: Dr. Poston?
19	DR. POSTON: Yes.
20	DR. BRANCHE: Dr. Roessler?
21	DR. ROESSLER: Yes.
22	DR. BRANCHE: Dr. Ziemer?
23	DR. ZIEMER: Yes.
24	DR. BRANCHE: Oh, excuse me, Mr. Schofield
25	forgive me. Mr. Schofield?

1 MR. SCHOFIELD: Yes. 2 DR. BRANCHE: Please forgive me. And I'll get 3 the votes from the other two gentlemen. 4 DR. ZIEMER: Okay. The motion does carry. 5 Board members, you will have an opportunity to 6 see a written version of this tomorrow 7 afternoon before the Board meeting ends, make 8 sure that everybody's comfortable with the 9 wording. 10 DR. POSTON: It was such an eloquent motion. 11 MS. MUNN: It was. 12 NUMEC PARKS SEC PETITION We are a little ahead of schedule 13 DR. ZIEMER: 14 on NUMEC. However, I note that the NUMEC 15 petitioner was undecided as to whether she 16 would be on the phone. Do we know -- oh, we 17 have someone here. 18 DR. BRANCHE: No, no, she's the -- she's the 19 NIOSH staffer. 20 DR. ZIEMER: Oh, she's NIOSH staff, right. is -- I'm looking for -- do we know whether 21 22 [name redacted] will be on the phone? 23 DR. BRANCHE: I think you can't say her name 24 until we know -- you can't say her name until

we know she's going to speak.

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1	DR. ZIEMER: I didn't say her name. That was
2	just a pseudo name, it's
3	MS. BREYER: She wasn't sure that she'd be able
4	to to listen in at this time. I usually
5	tell them to call in about half an hour earlier
6	earlier than the scheduled time in case the
7	agenda gets moved up a little, so it's very
8	likely that she just wasn't wasn't able to
9	(unintelligible)
10	DR. ZIEMER: Jane Doe, are you on the phone?
11	(No responses)
12	Is there a petitioner from NUMEC on NUMEC
13	Parks on the phone?
14	(No responses)
15	Okay. This says will try to listen, but
16	probably won't comment. So
17	DR. BRANCHE: And and as we've heard from
18	Ms. Breyer, she encourages the petitioners to
19	call about 30 min at least 30 minutes before
20	
21	MS. BREYER: Right, I usually tell them
22	DR. BRANCHE: the scheduled time.
23	MS. BREYER: about a half an hour. It may
24	be off either way, earlier or later.
25	DR. ZIEMER: Sure.

1 MS. BREYER: (Off microphone) And I don't have 2 a number to contact her (unintelligible) she 3 may be unavailable (unintelligible). 4 DR. ZIEMER: Okay. Well, Dr. Hughes is with us 5 anyway. Dr. Hughes will present for NIOSH. 6 Thank you. 7 DR. HUGHES: Thank you, Dr. Ziemer and the 8 I'm going to present on behalf of NIOSH 9 the SEC petition evaluation for the --10 DR. BRANCHE: Dr. Hughes, would you please 11 speak up? 12 DR. HUGHES: Okay, I'll -- I'll try. 13 DR. ZIEMER: Or get closer to the microphone. 14 DR. HUGHES: Can you hear me better? 15 Okay, how's this? 16 MR. GRIFFON: Better. 17 DR. HUGHES: Okay. Okay, I'm going to present 18 the NIOSH SEC evaluation for the Nuclear 19 Materials and Equipment Corporation -- or 20 short, NUMEC -- Parks Township plant. 21 a petition that was submitted to NIOSH under 22 83.14 for a petitioner whose dose -- dose could 23 not be reconstructed with the available data. 24 The petition evaluation also considered a class 25 of workers similar to the petitioner.

This is a slide you've seen before, the evaluation process, the two-step process which consists of the -- first the feasibility determination, followed by the health endangerment determination.

A little bit of background. The NUMEC Parks
Township plant is located in Parks Township
near Leechburg, Pennsylvania, which is about 30
miles northeast of Pittsburgh. It is a sister
facility to the NUMEC Apollo facility, which
was also evaluated by NIOSH and I believe was
presented last year, in October, to the Board.
This plant had its first license granted in
1961, March of 1961, and it is -- the covered
time period is actually 1957 to 1980. However,
we found information that there was no
radioactive material on-site before June of
1960.

The radiological operations relevant to the class consisted of production of plutonium-containing nuclear fuels and experimental fuels, the recovering of plutonium from scrap, the production of highly enriched uranium nuclear fuels, and the processing of depleted uranium. In addition, the production of

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uranium metal alloys, also the production of thorium experimental fuels and encapsulation of thorium fuels; the production of alpha, beta and neutron sources such as plutoniumberyllium, polonium-beryllium or americiumberyllium sources, in addition to thermal sources; and the production of gamma sources such as iridium-192 and cobalt-60 sources. The exposure potential to the class is obviously as a result of the operations that were conducted at the plant such as plutonium from fuel fabrication and scrap recovery, uranium from the machining of depleted uranium and highly enriched uranium production, exposure potential to thorium from the fuel production operations; and exposure to polonium, plutonium, americium, cobalt, iridium and cesium from source production. NIOSH looked into acquiring all available information to determine the feasibility of dose reconstruction, and the data capture attempts included formal requests to the former operator of the site, which is BWXT; requests to the Nuclear Regulatory Commission; a data search at the Office of Scientific and

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Technical Information, data requests to the DOE and also information was collected through worker outreach meeting and interviews with former employees.

There is monitoring data available. Internal monitoring data is available in the form of urine and fecal bioassay for plutonium, americium and uranium. Also some workers had occasional whole body counts for uranium and plutonium, starting in 1968 through 1985. urine bioassay is available from 19-- starting in 1960 to 1976; fecal bioassay was conducted from 1966 to '76. The whole body count it appears were given to employees who had exposure potential or potential uptakes. are very limited bioassay data for thorium available. All data also appear -- or are unclear whether or not they are for the Parks Township facility or the Apollo facility. As I mentioned earlier, they were sister facilities. They shared the same management. They also shared health and safety, so if you look at a given health and safety record it is not always clear which facility these are actually pertaining to. The process information that's

available for the Parks Township plant for the thorium operation is insufficient for source term determination. Additional urine sample is available in form of mixed -- in form of urine data from mixed fission products, which is also very limited.

There's also no bioassay or air monitoring data for radionuclides from source production, and there's also insufficient process information available to determine the source term.

And lastly, NUMEC used the contractor Controls for Environmental Pollution, or CEP, as a bioassay contractor starting in 1976 to 1993.

In 1994 both DOE and NRC advised contractors and licensees that the analytical results provided them by that company should be considered suspect because there were some implications of data falsifications. And for that reason, NIOSH has concluded not to use any CEP data for dose reconstruction.

There is limited air sampling available at the

site, only for uranium and plutonium which started in 1961, that consists of general air sample data and breathing zone sample data.

There was in general a large variation in

sampling frequency, which it is unclear whether this large variation in the data that we have is a result of a change in radiological risk or if there's a -- if there are gaps in the

External monitoring data is available starting in 1961 through 1980, and it -- it appears that all employees who had exposure potential were required to wear -- be monitored for external

This is the petition overview. NIOSH was unable to obtain sufficient information to complete the dose reconstruction for an existing claim. And on March 10, 2008 a claimant was notified that the dose reconstruction could not be completed and the claimant was provided with a copy of the Special Exposure Cohort petition Form A. the petition was submitted to NIOSH on March

The feasibility conclusion is that NIOSH lacks sufficient monitoring, process or source information from thorium and source production operation to estimate internal radiation doses to NUMEC Parks Township employees for the

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period of June 1st, 1960 through December 31st, 1980. NIOSH does believe that it has sufficient information to estimate the internal doses from uranium and plutonium from 1960 to 1976, and occupational external exposures, including the medical exposures, for that same period. And NIOSH will use individual personal monitoring data, with exception to the CEP data, for partial dose reconstruction, as appropriate.

NIOSH has determined that it is not feasible to estimate with sufficient accuracy external or internal radiation doses, and that the health of the covered employees may have been The evidence indicates that endangered. workers in the class may have accumulated intakes of uranium, plutonium, thorium and other radionuclides during the covered period. This is the summary slide. Again, dose reconstruction is believed to be feasible for uranium and plutonium only up to 1976, and dose reconstruction is not feasible for any of the other radionuclides on-site. reconstruction is believed to be feasible for external exposures, and including occupational

medical X-rays.

Therefore the NIOSH proposed class definition is all employees who worked at the Nuclear Materials and Equipment Corporation plant in Parks Township, Pennsylvania from June 1st, 1960 through December 31st, 1980 for a number of work days aggregating at least 250 work days occurring either solely under this employment or in combination with work days within the parameters established for one or more other classes of employees in the SEC.

And the recommendation is that feasibility is

no and health endangerment, yes.

Thank you.

DR. ZIEMER: Okay, thank you. Let's open this for questions. Let -- let me begin. And again, for clarification -- and I'm looking at

again, for clarification -- and I'm looking at the feasibility chart which is toward the end there, feasible to construct uranium and plutonium. What -- what was the status on americium, was or was not feasible on americium? I thought they -- I thought it's -- you said that they did bioassay for americium.

DR. HUGHES: Ye-- well, yes.

DR. ZIEMER: Is that included in the --

1 DR. HUGHES: It's included with the --2 DR. ZIEMER: Okay, that's included. 3 DR. HUGHES: Yes. 4 DR. ZIEMER: On the other nuclides, for those 5 for whom they did bioassay, were they simply 6 not looking at anything -- or -- I'm just 7 wondering why they wouldn't have done other 8 nuclides if they were doing bioassay. 9 they simply doing -- was this an alpha process 10 or -- can you clarify that? What -- what would 11 they -- what were the other nuclides that they 12 -- that would be in this category, other than 13 uranium, plutonium, americium? 14 DR. HUGHES: Thorium -- they did not do --15 DR. ZIEMER: So thorium would be the main one 16 here. 17 DR. HUGHES: Yes. 18 DR. ZIEMER: They simply weren't looking for 19 it, or were they -- what was --20 MR. RUTHERFORD: At least -- this is LaVon 21 Rutherford. At least for -- from my knowledge 22 with Apollo, when we looked at Apollo, it 23 appeared that for these smaller -- or through 24 the operations that were more -- on a smaller 25 scale, that they were not looking for those

1 isotopes when they were doing actual bioassay 2 monitoring, and the whole body count. Because 3 if you look at the actual sheets, and Dr. 4 Hughes can correct me if I'm wrong, they're 5 very specific on what they -- and it's anno--6 annotated what they're looking for. 7 DR. ZIEMER: Do you know in the bioassay here -8 - was it nuclide-specific versus gross alpha, 9 gross beta, or --10 DR. HUGHES: Yes, it was nuclide -- well, for 11 the biggest -- for the largest part, it was 12 nuclide-specific. 13 DR. ZIEMER: Okay. 14 DR. HUGHES: I do believe that the uranium and 15 plutonium was a very large portion of the 16 production and the other -- the thorium 17 production were relatively smaller programs of 18 the site, but -- so... 19 DR. ZIEMER: I -- I'm -- I'm really trying to 20 get a feel for whether or not -- if someone had 21 a positive bioassay and -- would they likely 22 have missed the thorium, even if it was there? 23 That's what I'm -- I'm not clear on. 24 DR. NETON: I'm not sure what you mean by 25 missed it. If it was specific for uranium, it

1	wouldn't show up in the uranium analysis
2	DR. ZIEMER: Well
3	DR. NETON: obviously, but and
4	DR. ZIEMER: if there were thorium there and
5	they're they're just not looking for
6	anything else, is what you're saying.
7	DR. NETON: Right, if they (unintelligible)
8	DR. ZIEMER: Other words, are they doing alpha
9	spectroscopy or what what are they doing
10	here?
11	DR. NETON: It seems to me I think Dr.
12	Hughes knows better, but she would
13	(unintelligible)
14	DR. ZIEMER: Oh, was it a chemical separation?
15	DR. NETON: It was a uranium chemical
16	radiochemical separation.
17	DR. ZIEMER: So they were separating out
18	DR. NETON: Yeah.
19	DR. ZIEMER: specifically for uranium.
20	Okay.
21	DR. NETON: Usually if you're going to go to
22	the trouble to digest a sample, it's easy to
23	pull off the uranium and then electrodeposit it
24	or something like that.
25	DR. ZIEMER: Okay, thanks. Other questions or

1	comments?
2	MR. GRIFFON: I I think
3	DR. ZIEMER: Mark.
4	MR. GRIFFON: this this is similar to the
5	other NUMEC site, but I just and I I
6	notice the language, it says all workers, right
7	so that would include the question of
8	administrative folks or guards or any workers
9	on the site okay. 'Cause there was that
10	issue at the other right.
11	DR. ZIEMER: It would be appropriate to have a
12	motion on this one if the group is prepared to
13	make such a motion.
14	DR. POSTON: Mark?
15	MR. GRIFFON: I don't have that complicated
16	detailed motion, but
17	DR. ZIEMER: I think
18	MR. GRIFFON: The author is not here.
19	DR. ZIEMER: if you wish to make the motion,
20	we can have it in simple form and our our
21	agenda is catching up with our ability to get -
22	-
23	MR. GRIFFON: Right.
24	DR. ZIEMER: the words together. Wanda
25	Munn.

1 MS. MUNN: I move we accept the recommendation 2 for -- that NIOSH has made for this particular 3 SEC class. 4 DR. POSTON: Second. 5 DR. ZIEMER: And seconded. Is there further discussion on this one? 6 7 MR. GRIFFON: The only -- the only thing I 8 wanted to ask was -- I -- I -- I think that in 9 the last one -- and this is not -- well, it's 10 sort of around the motion, but the -- in the --11 in the other NUMEC site I think we considered -12 - or we asked the workgroup on the 250-day issue to consider the NUMEC Apollo, and I think 13 14 we should probably put Parks in there with that, you know, just -- I -- I think the 15 16 petitioner mentioned both when they had spoken 17 with me before, so I'm not sure it's going to 18 fall in-- you know, at least let it be 19 considered by the 250-day workgroup whether 20 they could have had exposures in -- in a 21 smaller -- shorter time frame that could affect 22 that 250-day criteria. So -- but that's -- I 23 don't think that --24 DR. ZIEMER: That's a separate issue. 25 MR. GRIFFON: -- affects the motion -- right,

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              separate issue, but -- yeah.
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              DR. BRANCHE: Do you want to include it or not?
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              DR. ZIEMER: No, not --
4
              MR. GRIFFON: Not in the motion, no, no, no.
5
              DR. ZIEMER: -- not in the motion, no. No.
              Are you ready to vote on the motion? And again
6
7
              we'll have -- the formal words are available
8
              for you tomorrow.
              Okay, we'll do it by roll call.
9
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              DR. BRANCHE: Okay. Ms. Beach?
11
              MS. BEACH: Yes.
12
              DR. BRANCHE: Mr. Clawson?
              MR. CLAWSON: Yes.
13
14
              DR. BRANCHE: Mr. Gibson?
              MR. GIBSON: Yes.
15
16
              DR. BRANCHE: Mr. Griffon?
17
              MR. GRIFFON: Yes.
18
              DR. BRANCHE: I'll get a vote from Dr. Lockey.
19
              Ms. Munn?
20
              MS. MUNN:
                        Yes.
21
              DR. BRANCHE: Mr. Presley?
22
              MR. PRESLEY: Yes.
23
              DR. BRANCHE: Dr. Poston?
24
              DR. POSTON: Yes.
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              DR. BRANCHE: Dr. Roessler?
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1 DR. ROESSLER: Yes.

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DR. BRANCHE: Mr. Schofield?

MR. SCHOFIELD: Yes.

DR. BRANCHE: Dr. Ziemer?

DR. ZIEMER: Yes. Then I declare that the motion has carried and we will prepare a formal recommendation to the Secretary in accordance with the formal wording that will come in the motion.

NINTH SET OF CASES FOR DOSE RECONSTRUCTION REVIEWS

We have a little bit of time before lunch and I'm going to take care of the -- a part of a subcommittee item. The Chair had the task of assigning workgroups for the ninth set of reviews, and I have done that and I wanted to give you those assignments, and then we will give you a hard copy of this before you leave the meeting. But I'm going to give you the assignments verbally so that they are in the record, and you can jot these down as we go. This is the ninth set of cases for dose reconstruction review. On the selection ID number, and the selection ID number is not at all related to the NIOSH number -- case number, so I simply point that out. It is a Board

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The -- on all of these the first digits are 2008-01, which represents January of this year, which was the final date at which we made the actual selection of those cases. then the ID number that I will use here, the last three digits, all following the 2008-01 --I'm not going to repeat the 2008-01 each time. So I will give you the case number and I will give you the facility, and then I will give you the review team. Then I will make this available to you in writing and I will make it available to John Mauro and SC&A because they will be working with the individual teams to review those dose reconstructions. So -- and there's I think 40 of these, so I'll go through the list.

MS. BEACH: Is this from the ninth set, Paul?

DR. ZIEMER: Ninth set.

MS. BEACH: Thank you.

DR. ZIEMER: Ninth set. Case 125 from Feed
Materials Production Center, otherwise known as
Fernald. Team one -- and I'm using the same
team numbers as we used last time, so -- but
I'll give you the names as well. Team one is
Poston and Presley.

1	The next case is 135 at Argonne East, and Los
2	Alamos and U. of California, it's a person that
3	worked at three facilities. This is for team
4	two, Roessler/Lockey.
5	Next, ID is 183, Ashland Oil, team three,
6	Griffon and Clawson.
7	Case 184, Vitro Manufacturing, team four,
8	Ziemer/Gibson.
9	Next, 198, Y-12 and K-25, Oak Ridge, team five,
10	Melius/Schofield.
11	Next is 418, Herring Hall, Marvin Safe Company,
12	that's team six, Munn/Beach.
13	Case 1 case 434, Los Alamos National Lab.
14	This will be team two, Roessler/Lockey.
15	DR. ROESSLER: What was the number again?
16	DR. POSTON: 434.
17	DR. ZIEMER: 434. Maybe I should read the team
18	first and then
19	DR. ROESSLER: That would help, yeah.
20	DR. ZIEMER: Okay. The next will be assigned
21	to team one, Poston/Presley. It's case 442 at
22	Fernald.
23	Next is team three, Griffon/Clawson, case 454,
24	Bridgeport Brass.
25	Next is team four, Ziemer/Gibson, case 461,

1	Paducah.
2	Parenthetically I will say normally I'm
3	going in order unless there's a conflict, and
4	then I'm switching teams. That's why they're
5	not all completely in order.
6	Next is team six, Munn/Beach, case 464,
7	Fernald.
8	Next, team five, Melius/Schofield, case 465, K-
9	- Oak Ridge K-25 and Hanford.
10	Next is team one, Poston/Presley, case 477,
11	Downey facility and some others as well.
12	Next is team two, Roessler/Lockey, case 490,
13	Weldon Spring plant.
14	Next, case or team three, Griffon/Clawson,
15	case 491, Hanford.
16	Next, team four, Ziemer/Gibson, case 492,
17	Hanford and PNNL, Pacific Northwest National
18	Lab.
19	Next, team five, Melius/Schofield, case 521,
20	Huntington Pilot Plant.
21	Team six, Munn/Beach, case 523, Y-12 plant.
22	Next, team two, Roessler/Lockey, case 533,
23	Lawrence Livermore.
24	Team one, Poston/Presley, case 537, Brookhaven
25	National Lab and Idaho National Lab.

1	Next, team three, Griffon/Clawson, case 461,
2	Clarksville facility and Pantex.
3	Team four, Ziemer/Gibson, case 565, Savannah
4	River Site.
5	Team five, Melius/Schofield for case 568,
6	Savannah River Site.
7	And then team six, Munn/Beach, case 571, Linde
8	Ceramics.
9	Team one, Poston/Presley, case 583, Idaho
10	National Lab.
11	Team two, Roessler/Lockey, case 584,
12	Albuquerque Operations Office and Los Alamos.
13	Then team five, Melius/Schofield, case 585,
14	Medina facility and Pacific Proving Ground.
15	Team three, Griffon/Clawson, case 588, Oak
16	Ridge X-10.
17	Team four, Ziemer/Gibson, case 614, Hanford,
18	Nevada Test Site, Los Alamos and Pacific
19	Northwest National Lab.
20	Team six, Munn/Beach, case 639, Y-12 plant.
21	Next, team two, Roessler/Lockey, case 648, Y-12
22	plant.
23	Next, Poston/Presley, team one, case 652,
24	Savannah River Site.
25	Team three, Griffon/Clawson, case 653, Y-12

1	plant.
2	Team four, Ziemer/Gibson, case 664, Nevada Test
3	Site.
4	Team six, Munn/Beach, case 672, Idaho National
5	Lab, Y-12, K-25 and X-10.
6	Okay, we're getting down to the final page
7	here.
8	Team five, Melius/Schofield, case 677, Grand
9	Junction Operations Office and DeSoto facility
10	and Hanford.
11	Then team one, Poston/Presley, case 69 679
12	again, 679, Hanford.
13	Team two, Roessler/Lockey, case 681, Idaho
14	National Lab.
15	Team three, Griffon/Clawson, case 690, General
16	Steel.
17	Team four, Ziemer/Gibson, case 697, Hooker
18	Electrochemical.
19	That completes the list. We've got six teams,
20	40 cases, so each of you has six or seven
21	cases. This is a double whammy review. Okay?
22	John, I'll give you a copy of this for SC&A so
23	that we're ready to go on that.
24	Any questions on those assignments? I've
25	checked this these assignments with counsel

1 and they have cleared this as far as conflicts 2 of interest for all Board members. 3 MR. GRIFFON: Can I just ask --4 DR. ZIEMER: Yeah, Mark Griffon, question? 5 MR. GRIFFON: Just -- Stu Hinnefeld, I see him 6 in the room now. Stu, I might ask about the 7 tenth set. We're going to consider those 8 tomorrow in the subcommittee meeting, and the 9 tenth set -- we had a subcommittee -- or in the 10 regular meeting. 11 DR. ZIEMER: In the report. 12 MR. GRIFFON: And -- and we had a subcommittee 13 meeting recently. We went through a first tier 14 review of the tenth set. We selected some, and 15 I think you produced a expanded matrix on those 16 17 MR. HINNEFELD: No -- no, I haven't. 18 MR. GRIFFON: Oh, you have not. 19 MR. HINNEFELD: No, we're not prepared to 20 actually make --21 MR. GRIFFON: So you're not prepared to do it 22 here, okay. 23 MR. HINNEFELD: What -- what happens or what I 24 have done is I have culled out cases that had 25 actually been selected for the ninth set.

1 MR. GRIFFON: I -- I --2 MR. HINNEFELD: Because recall that, unlike our 3 other preselection list --4 MR. GRIFFON: Yes. 5 MR. HINNEFELD: -- the ninth set had not been 6 selected when that preselection list was run, 7 and so they had not been culled out. So after 8 the selection of the -- the preselection of the 9 tenth, there were some 54 cases that the 10 subcommittee preselected. I went through and 11 culled out 11 -- I guess there were 56. 12 culled out 11 that had -- that had actually 13 been selected in the ninth --14 MR. GRIFFON: Right, and --15 MR. HINNEFELD: -- part of the ninth --16 MR. GRIFFON: -- we saw that e-mail, but --17 okay. 18 MR. HINNEFELD: So the remaining 45 then we are 19 compiling to fill out the rest of the matrix. 20 MR. GRIFFON: Okay. 21 MR. HINNEFELD: The rest of the matrix requires 22 us to enter in the way in which the internal 23 dosimetry was done --24 MR. GRIFFON: Yeah. 25 MR. HINNEFELD: -- the way in which the

1 external dosimetry was done and whether 2 neutrons were present before and after --3 MR. GRIFFON: Okay, I just wanted to clarify --4 I thought we would have that for this meeting, 5 but may -- for the phone call meeting I guess 6 we'll do that. 7 MR. HINNEFELD: For the phone call meeting 8 we'll have it. We didn't have enough time to 9 do it by now. 10 MR. GRIFFON: All right, all right, thank you. 11 TRACKING STATUS OF TRANSCRIPTS AND MINUTES 12 DR. ZIEMER: Thank you. I think we have a 13 little time for a couple of things. 14 Branche, I'm wondering if it would be useful to 15 cover the tracking status --16 DR. BRANCHE: Yes, that'd --17 DR. ZIEMER: -- now? 18 **DR. BRANCHE:** -- be great. 19 DR. ZIEMER: We're -- I've skipped ahead to some of the Board working time for tomorrow 20 21 afternoon because these are things that, if we get done, we might be able to leave a little 22 23 bit early. But we'll take care of pieces of 24 this. This is -- bullet under Board working

time called tracking status of transcripts and

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minutes, so we get an update on where we are on minutes and transcripts.

DR. BRANCHE: Okay.

DR. ZIEMER: Do we have this in our packet?

DR. BRANCHE: No, you don't have it in your package this time because I'm happy to report that as it concerns the Board meeting transcripts, both the face-to-face meetings and the telephone meetings, we are completely up to date. Everything is on the web site. We are at the steady state for those issues.

I -- my compliments to the staff at NIOSH and as well as our contract court reporter for getting this information to us in a timely fashion.

Now as it concerns the subcommittees -- sorry, the subcommittee meetings and the workgroup meetings, we are still behind, but I don't think we've ever made a promise to you as to when those would be coming. We have tried -- I actually put a little bit of a moratorium on requests that many of you had been making for workgroup meeting transcripts until we got ourselves back into -- into a smooth delivery of our Board meeting transcripts. Our

constituents -- and you all, appropriately -- have demanded that that backlog be cleaned up and that has been done.

Ray has been giving us information -- the transcripts from our -- from our workgroup meetings and the subcommittee meetings, and he is catching up. As well he's catching up on the minutes.

As it concerns the minutes, we've been working with the Federal Advisory Committee Act staff at the Centers for Disease Control and Prevention because they're undergoing a review from I guess the HHS Federal Advisory Committee Act office, and they were concerned about the fact that our minutes were lagging. I've put forth an argument that our transcript should serve as fulfilling that need, and so I'm -- so far I seem to be winning that argument, but I will appreciate any good vibes you can give me on that score.

Again, our -- the time line that Dr. Wade and other staff at NIOSH outlined for you all at a previous meeting about the time frame that we need to be able to get the transcript for the Board meeting produced, redacted and posted

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seems to be exactly the amount of time we need. We have -- we would have great difficulty in producing that sooner. But we -- given that time line, we seem to be honoring it now and we are playing catch-up with our other information. I'm probably talking too long.

DR. ZIEMER: No, that's fine.

DR. BRANCHE: Any questions? I wish Dr. Melius were here to hear me say all that.

DR. ZIEMER: So you'll be working on the workgroup meetings as -- transcripts to bring those up. The minutes themselves -- Board members, you've probably noticed that we haven't had to approve any minutes lately, and the reason for that is we haven't had minutes to approve, working hard on the transcripts. And we've found in practice many of the folks who are involved in pursuing petitions -- that is, petitioners themselves -- prefer the transcripts rather than the minutes because the transcripts more accurately reflect -- or at least include everything that was covered, as opposed to the minutes, which may be condensed and may summarize what occurred rather than giving it verbatim.

One of the issues is, and what Christine was referring to, is that in the past the folks who operate or who set forth the rules for federal advisory boards have required that I sign off that the minutes are a true reflection of what occurred. Since we are -- and -- and we're behind on those.

Since the transcripts are what we're focusing on, we're -- what she's trying to do is get them to agree that we can assert that the transcripts fully and correctly reflect what occurred at the meeting. And if -- if we have that, then it's not clear to me that we have to necessarily approve the minutes. Do we still approve the minutes?

DR. BRANCHE: Well, I would -- I would say that in -- in my -- my being assertive about the fact that the minutes (sic) serve our needs and the needs of our constituents, and if the federal --

DR. ZIEMER: Minutes or the transcript?

DR. BRANCHE: Sorry -- forgive me, thank you, the transcript. If the Federal Advisory

Committee Act staff agree with me/us, then it would require that Dr. Ziemer in his capacity

1 as the Chair of this Advisory Board, and Mr. 2 Griffon in his capacity as the chair of the 3 sub-- subcommittee, essentially sign off on the 4 transcript. And if that is the case, then I 5 think we would need to go through an opportunity for each Board member to receive 6 7 those in advance and then -- whether by e-mail 8 or another mechanism that we work out -- agree 9 that the transcript is something that you all 10 accept, and then we would have -- I would have 11 them sign off on those and -- and potentially 12 forego the minutes. 13 DR. ZIEMER: So we may not even have minutes to 14 deal with in the future. 15 The other thing I think on transcripts -- I'm 16 not sure we're ever in a position to say that 17 what the transcript says is not what I've said 18 because that's a --19 MR. GRIFFON: Right. 20 DR. ZIEMER: -- an official court reporter type 21 of thing. Ray is --22 DR. BRANCHE: That's all you, Ray. 23 DR. ZIEMER: Similar to a court proceeding. 24 -- I think you might not like what you said or 25 how you said it, but -- but it's going to be

1 there for the record. I don't think we're 2 going to go back and -- and edit the 3 transcript, unless there's a spelling error or 4 something like that. 5 DR. BRANCHE: But of course that -- if we take 6 the step that -- and -- and because we do have 7 a certified court reporter, it would certainly 8 expedite our signing off on a lot of the 9 documents that the Federal Advisory Committee 10 Act office would -- would require. So I'm 11 quite hopeful that they would see the wisdom of 12 this approach. And if you don't see the wisdom 13 of this approach, this is now your opportunity 14 to tell me. 15 DR. ZIEMER: Now would you rather have minutes 16 than transcripts? I think our petitioners have 17 been relying on the transcripts rather than the 18 minutes. 19 DR. BRANCHE: Again, I don't know that we'll be able to forego minutes, but we will be able to 20 21 have -- sign off on the transcripts, which 22 might be able to forego needing a signature on 23 the minutes, so... 24 I have some updates for the other information 25 if you'd like --

1 MR. GRIFFON: I was just going to ask about --2 before you move on -- the moratorium on the 3 other minutes -- or the other transcripts. I -4 - I know --5 DR. ZIEMER: Well, there's not a moratorium --Well, moratorium on producing 6 MR. GRIFFON: 7 them for workgroup members or --8 DR. BRANCHE: I -- I was -- I was actually 9 asking people, when they would come first to 10 Lew and now me -- actually nobody has asked me 11 for --12 MR. GRIFFON: In a while. 13 DR. BRANCHE: -- an expedited transcript from a 14 workgroup or subcommittee meeting, and I 15 appreciate the sensitivity that you all have 16 shown because I think -- actually did cry a lot 17 at our last Board meeting about how we were in 18 -- in a bit of a -- a flurry in trying to do 19 that. I'm not asking for floodgates, either, 20 but if you do need an expedited transcript for 21 your workgroup meeting or your subcommittee 22 meeting, I would ask that you simply come to me 23 and I'll let you know where we are in the 24 production cycle. 25 MR. GRIFFON: Okay, I was just going to say, in

1 -- or ask a question, I guess. In the past I -2 - I've asked -- especially for the Rocky Flats 3 process, we had several meetings very close to 4 each other and it was useful to have the 5 previous minutes and Ray --DR. BRANCHE: The minutes or the transcript? 6 7 MR. GRIFFON: -- transcripts, and Ray produced 8 them very quickly in a -- in a raw form, a -- a 9 draft form that we wouldn't circulate, but we 10 had them there for our information. 11 don't know when you say -- if -- can we still 12 get those kind of minutes if we need them and -13 14 Transcripts? DR. BRANCHE: 15 MR. GRIFFON: Yeah, transcripts, I'm sorry --16 in a pinch if we need those kind of things to 17 facilitate the workgroup process, can we get 18 those draft transcripts? 19 DR. BRANCHE: Very reasonable question. think producing the draft -- it's easier to 20 21 promise when we've gotten past the window for when -- for when Ray needs to produce the 22 23 transcript for our Board meeting. So for 24 example, there was a request for a workgroup 25 meeting from November, but the person requested

it just this past week. And so my honest reply was if Mr. Green can produce it without it 3 interrupting the schedule -- which again, is already tight -- for getting out the transcript for the Board meeting, then that wouldn't be a problem. So I would say the same to any 6 7 workgroup chair or subcommittee chair in your 8 case, Mr. Griffon, that as long as the timing 9 of your request is not going to jeopar -- and I 10 always check with Mr. Green first -- is not going to jeopardize the -- the time line for 12 our producing the Board minute -- I'm sorry, 13 transcript, you've got me in -- the Board 14 transcripts, then I think we can try to honor And in -- and in those I use a first come-15 16 first served approach. Okay?

UPDATE ON BOARD'S CONTRACTOR

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Some of you have been asking questions about the -- the Board's contractor --

MR. CLAWSON: Yes.

DR. BRANCHE: -- and I can give you an update on that. I appreciate Dr. Wade, Dr. Neton and the staff in the Centers for Disease Control's Office on Procurement and Grants. We expect that the first solicitation -- or rather the

solicitation announcement will come out in the middle of this month, April 2008, and that the full request for proposals will be announced at the very beginning of May. And we're expecting to be able to select certainly someone before any deadlines are appro-- approach. But those are the two dates that you need to have in mind. Mid-- mid-April for the solicitation announcement, which essentially just gives people a heads-up -- rather potential applicants a heads-up. And then the full-fledged request for proposals would be the first few days of May.

DR. ZIEMER: Mark?

MR. GRIFFON: This may be just my -- you know, 'cause we have so much documentation going around this Board, but have we -- has the Board seen a final -- or are we entitled to see a final copy of the RFP? I know -- I know you asked for comments on certain sections from me, and I appreciate that and I -- I did send them in, but I -- I don't know that I've seen the final form of it, and are we -- do we get one last read-through or -- or -- I don't know.

DR. ZIEMER: Well, let me ask if David Staudt

1 happens to be on the phone. Do we know if he's 2 going to -- or anyone from procurement. I 3 think --4 DR. BRANCHE: Talk about this tomorrow 5 afternoon, though. 6 MR. GRIFFON: Okay, we can talk -- yeah. 7 DR. BRANCHE: He was going to be available. 8 might be on the line. 9 We'll try to get that answer by DR. ZIEMER: 10 tomorrow. I -- I believe that David had told 11 us before we would see a final copy of that, 12 but let's -- we'll get it clarified before --13 MR. GRIFFON: I -- I would like to if we can, 14 yeah. I think that'd be useful. 15 DR. BRANCHE: I will get an answer to you as to 16 when we would be able to get that to you. I 17 know that by e-mail you and Mr. Claws-- or 18 actually Mr. Clawson copied you on his request. 19 You all had questions about some very elaborate 20 language that had to do with conflict of 21 interest, and the conflict of interest language 22 that the Board apparently labored through on 23 the last announcement -- apparently it has been 24 undisturbed. 25 MR. GRIFFON: Remained, right, right, yeah.

DR. BRANCHE: So that remains completely unaltered. I believe Dr. Wade gave you all information to review -- the main sections that -- again, my understanding is that the Board members labored over in a previous version of this when the announcement was done three years ago, and that language, to my understanding, has been left undisturbed. But the staff and the -- at NIOSH and Procurement and Grants have been working on the specific parameters that are date-sensitive. But I'll work to get a copy of that to you.

DR. ZIEMER: Thank you. Any other questions for Dr. Branche on those administrative matters?

(No responses)

Okay. Now I want to point out that after lunch today the Board is going to undergo ethics training so that we will be even more ethical in the second half of our meeting after that. Actually it -- we're -- we're required every year to participate in what is called ethics training. Is that the right word?

DR. BRANCHE: That is correct, that's correct.

DR. ZIEMER: I think it's ethics training. And

that is really an administrative session of the Board. It's -- if you'll notice on the agenda, it says for Board members only. We recognize that none of the members of the public nor the federal staff people, nor our contractors, need ethics training and so you do not need to come back early from lunch. In fact, I -- I've been told the attorneys don't want you to come back early. Our session will become unethical if you're here, for some reason. Well, in any -- any --

MR. GRIFFON: It's not -- it's not a closed
session, though, is it? No --

DR. ZIEMER: Well, let's say it's not officially a closed session, as defined, but I'm told that it is considered to be for Board members only. It sort of sounds closed to me, but I -- I think if -- probably if someone's out there and they really feel they need this badly, we might let them in the door. I don't know. We'll see what -- we'll see what -- MS. HOMOKI-TITUS: Dr. Ziemer, this is Liz Homoki-Titus. (Unintelligible) administrative session under FACA, so therefore it is not open to the public. So even if someone really badly

1	wanted to come in, they would not be permitted
2	to.
3	DR. ZIEMER: Oh, okay, even if they wanted to
4	badly. Okay. That that was word from
5	counsel. This is considered an administrative
6	session of the Board, which is although not
7	closed, Mark, it's not open to the public.
8	Okay.
9	DR. BRANCHE: I have an administrative note on
10	that piece.
11	MR. GRIFFON: I'll have to look that one up.
12	DR. ZIEMER: Okay. Anyway, take I'm simply
13	telling folks have a leisurely lunch
14	DR. POSTON: Paul
15	MR. PRESLEY: Hey, Paul Paul?
16	DR. ZIEMER: Yes, yes, Mr. Presley.
17	MR. PRESLEY: I got something from counsel that
18	says it's going to start at 1:00 o'clock. Is
19	it going to start at 1:00 o'clock or 1:30?
20	DR. ZIEMER: 1:30.
21	MR. PRESLEY: Thank you.
22	DR. BRANCHE: Oh, and and and Mr.
23	Presley, you will note that I sent you an e-
24	mail message yesterday with a separate phone
25	number that I would like you to dial in to for

1	that administrative session. Did you receive
2	my e-mail?
3	MR. PRESLEY: I've got it right here.
4	DR. BRANCHE: Okay, thank you.
5	DR. ZIEMER: So the open or the regular
6	session will begin at 2:30, and at 2:30
7	since we have already covered the Department of
8	Energy and Labor update, I'm proposing that we
9	consider the SEC petition update, so I'll ask
10	LaVon to be prepared for that.
11	So thank you very much. We're recessed until
12	the appropriate time.
13	(Whereupon, a recess was taken from 12:00 p.m.
14	to 1:30 p.m.)
15	(Whereupon, the meeting reconvened in
16	Administrative Session, transcript of which is
17	not included as part of the public document.)
18	(2:53 p.m.)
19	DR. BRANCHE: Mr. Presley, are you on the line?
20	MR. PRESLEY: I sure am, Christine.
21	DR. BRANCHE: Thank you. For those of you who
22	have joined the re-established telephone line,
23	we ask that you mute your phones. If you do
24	not have a mute button, then please use star-6
25	to mute the telephone line. And when you are

ready to speak, please use that same star-6 to unmute your phones. Thank you. I heard all that noise, so thank you very much for muting your phones.

Dr. Ziemer.

SEC PETITION UPDATE

DR. ZIEMER: Okay, we're going to proceed now.

We have a number of SEC petitions that we're
going to have updates on, beginning with

Hanford, then Sandia Livermore, and then
Chapman Valve, and then we also will add Mound
to that list. Well, we'll start with Hanford -

DR. BRANCHE: Oh, I thought we were going to start with LaVon Rutherford from tomorrow morning.

DR. ZIEMER: Oh -- wait a minute, I'm -- I'm sorry, I'm ahead of myself on the schedule. Yes, those come at 3:45, and I was just looking at the schedule as it was and realizing it's not 3:45. We're -- we're picking up tomorrow morning's part of the SEC updates, and that is everything but the ones I just named, let's put it that way.

MR. RUTHERFORD: Are you ready for me?

1 DR. ZIEMER: I'm ready for you. I think I'm 2 ready for you. 3 MR. RUTHERFORD: All right. Thank you, Dr. 4 Ziemer. I'm going to give the update to the 5 SEC petitions. There will be a little overlap between some of the SEC petition discussion 6 7 that you probably already heard and that you're 8 going to hear later on. 9 We provide this update to the Board and -- to 10 allow them a little preparation for future 11 workgroup meetings and for future Board 12 meetings, and so they can get a little look 13 ahead. 14 At this time we've -- as of March 26th we had 15 108 petitions. I don't know if we've received 16 any since I've been in the office or not. 17 We've quali -- or we have four petitions that 18 are in the qualification phase. We have 56 19 petitions that we have -- we are -- we have 20 qualified for evaluation. Of those 56, six of 21 those are in the evaluation process and 50 of 22 them have -- we have completed our evaluation. 23 We had 48 petitions that have not qualif -- that 24 did not qualify for evaluation. 25 Now I'm going to go over the petitions that are

with the Board at this time and are awaiting recommendation from the Board.

Chapman Valve, the evaluation report was approved and sent to the Boa-- Advisory Board and petitioners on August 31st, 2006. We presented our evaluation at the September 2006 Advisory Board meeting. At that time the Advisory Board established a workgroup to review the evaluation. The workgroup presented its findings at the May 2007 Advisory Board meeting and a decision at that time was made to postpone a recommendation until the petitioner had received all the documents and had had a chance to review those documents.

The Advisory Board voted 6 to 6 on a motion to deny adding a class to the SEC at the July 2007 meeting. In light of the vote, the Advisory Board determined they -- they needed more -- or needed to receive a response from the Department of Labor and Department of Energy concerning potential covered work at the Dean Street facility. Prior to the October 2007 Board meeting Department of Labor provided a response to the Advisory Board's questions about the Dean Street facility. DO-- the

Department of Energy provided an update during the November 2007 Advisory Board conference call. At that time they had not completed their investigation.

The Department of Energy presented their findings at the January 2008 Advisory Board meeting that the Dean Street facility should be included as a covered facility, but they indicated that they had no information that there was any additional radiological activities. NIOSH also indicated at that meeting that they would revise the Chapman Valve evaluation report based on DOE's finding, but also indicated they anticipated there would be no changes in our feasibility determination based on these findings.

We issued our revised evaluation report at the February 2000 -- February 5th -- we issued our evaluation report February 5th. At the February 2008 Advisory Board conference call the Board asked SC&A to do a focused review of the new information provided by Department of Energy, and asked that the information be available prior to the April Board meeting. SC&A provided a report to the workgroup on

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March 12th, and status -- current status is NIOSH plans to present our revised evalua -actually the revisions to our evaluation report at this meeting. I think that's later today. Blockson Chemical, the evaluation report was initially approved and sent to the Advisory Board in September 2006. We presented our evaluation report at the December 2006 Advisory Board meeting. However, we determined that we had not addressed all covered exposures and we withdrew that evaluation report. The Advisory Board established a workgroup to review the evaluation report at its December 2006 meeting. NIOSH issued a revised evaluation report on July 3rd, 2007, and we presented that revised evaluation report at the July 2007 Advisory Board meeting. The workgroup met in Cincinnati on August 28th and a public meeting was held on September 12, 2007 to explain changes made to the dose reconstruction technical approach. was the work-- the workgroup held a conference call on November 2nd, 2007, and then at the January 2008 Advisory Board meeting Dr. Melius indicated he wanted to review the pedigree of the bioassay data and he wanted to discuss the

1 radon model with Mark Griffon. 2 Current status is the petition and evaluation 3 report are with the workgroup. An update will be provided at this meeting. 5 Feed Materials Production Center, Brad Clawson 6 has already provided an update. I'll try to be 7 really brief here. The evaluation report was 8 approved and sent to the petitioners on 9 November 3rd, 2006 and we presented our 10 evaluation report at the February 2007 Advisory 11 Board meeting. The Advisory Board established 12 a workgroup at that meeting and -- and May SC&A provided a draft review of the evaluation 13 14 report to the workgroup, petitioners, Advisory 15 Board and NIOSH. And the workgroup has met on 16 a number of occasions, and the current status 17 is that workgroup is still reviewing the Feed 18 Materials Production Center evaluation. 19 Bethlehem Steel, the evaluation report was 20 approved and sent to the Advisory Board and 21 petitioners on February 27, 2007, and NIOSH 22 presented our evaluation report at the May 2007 23 Advisory Board meeting. At the time, the 24 Advisory Board determined that it needed 25 further information before making a

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recommendation on the SEC petition. Advisory Board tabled the discussion of Bethlehem Steel SEC evaluation report until the workgroup that is looking at the use of surrogate data reports back to the Board. the petition and the evaluation report are with the Advisory Board for recommendation, and I believe an update is scheduled for tomorrow. Sandia National Lab Livermore, we appro-- the evaluation report was approved and sent to the Advisory Board on March 29th of 2007. On April 25th, 2007 -- that was just before the May meeting -- we received new information from the petitioner. However, we presented the evaluation report at the May 2007 Advisory Board meeting and discussed the new information provided by the petitioner. The Advisory Board asked NIOSH to provide an

The Advisory Board asked NIOSH to provide an update that addressed the new information. We issued an addendum to the evaluation report and presented that addendum at the October 2007 Advisory Board meeting. The Advisory Board tabled the vote at the October meeting until information provided from the petitioner could be reviewed by the Board. NIOSH informed the

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Advisory Board during its conference call in November that all information had been made available to the Advisory Board, and the Advisory Board indicated they needed more time to review the information.

The status -- an update is -- is scheduled for this meeting.

Hanford Part 2, the evaluation report was approved and sent to the Advisory Board and the petitioners on September 11, 2007. presented our evaluation report at the October meeting. The Advisory Board established -sent the report to the already-established Hanford workgroup that was working on the site profile review. The Advisory Board's contractor, SC&A, issued a white paper questioning whether additional buildings should be included in the proposed class definition. And we reviewed that white paper and in March we issued a revised evaluation report with a modified class definition, and I believe that update is scheduled for later on today as well. The petition and evaluation report are with the workgroup, and SC&A and the Board workgroup and NIOSH will provide an update at this meeting.

1 Nevada Test Site, the evaluation report was 2 approved and sent to the Advisory Board and the 3 petitioners in September of '07. We presented our evaluation report at the January 2008 5 meeting, and the Advisory Board sent the report to the contractor and the NTS Board workgroup. 6 7 That workgroup had already been established. 8 The petition and evaluation report are with the 9 Advisory Board and SC&A for review. 10 Mound plant, evaluation report was approved and 11 sent to the Advisory Board in December. 12 presented our evaluation at the January meeting and the Advisory Board concurred with our 13 14 recommendation to add a class for the early 15 years, but sent the report to their contractor 16 for review and established a Mound workgroup. 17 The Mound workgroup met on April 1st, and the 18 status is the current -- the petition and 19 evaluation report are with the workgroup and 20 SC&A for review. 21 Texas City Chemical, Dr. Neton presented our 22 evaluation report on -- that was -- was 23 approved on January 18th and Dr. Neton 24 presented that evaluation report at this 25 meeting. And the vote and recommendation has

1 been postponed until a future meeting. 2 NUMEC Parks, Dr. Hughes presented our 3 evaluation report that was approved -- it was approved on February 14 and Dr. Hughes presented our evaluation report earlier today, and the Board concurred with our recommendation 6 7 to add a class for NUMEC Parks. 8 Santa Susana Area IV, the evaluation report was 9 approved and sent to the Advisory Board and 10 petitioners on February 15th. Stu Hinnefeld 11 presented our evaluation this morning, and the vote has been delayed until SC&A completes 12 13 their review of the site profile. 14 SAM Laboratories, the evaluation report was 15 approved and sent to the Board on February 16 19th, and we presented our evaluation of the 17 SAM Laboratory yesterday's -- during this 18 meeting. The Board concurred with our 19 recommendation. 20 Kellex-Pierpont, the evaluation report was 21 approved and sent on February 27th and Dr. 22 Glover presented our evaluation report earlier 23 today, and the Board concurred with our recommendation. 24 25 Horizons, Inc., the evaluation report was

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approved and sent to the Board and petitioners on March 14th. I presented that evaluation report yesterday, and the Board concurred with our recommendation.

Now I'm going to go to SEC petitions that are currently -- that are in the evaluation process, give you a little update where we are with each one of those.

Pantex is a petition that we've had for quite The petition was initially not some time. qualified. The petitioner requested an Administrative Review of the qualification and the Administrative Review Panel recommended that we qualify the petition. We -- we have been in the evaluation for -- for some period We have run into some difficulties with data capture efforts and also some site reviews that have taken place. We are working to complete the evaluation report in June. However, I believe that the evaluation report is going to be approved at the later part of June and will be -- we'll have limited chance of getting that report to the Board in time to

Spencer Chemical, this petition was actually

present for the June meeting.

delayed, at the petitioner's request, for four months while the petitioner reviewed additional documentation. We anticipate this report will be completed in June and presented at the June Board meeting.

Westinghouse Atomic Power Development, we have
-- during our review of the documentation for
Westinghouse Atomic Power Development we -- we
uncovered some concerns with covered activities
at the facility and we have been corresponding
with Department of Energy and Department of
Labor concerning this. However, we expect to
have some answer for the June meeting with the
-- concerning Westinghouse.

We also, if you remember, had some issues with the early Y-12 with administering the class. Back in our earlier SEC we had defined a class that -- at Y-12 for uranium enrichment operation and other radiological activities, and administering that class has been a little difficult based on our class definition. We are working with the Department of Labor to resolve this issue. We've actually sent a letter to the Department of Labor outlining an approach to allow us that we would not have to

-- to complete a SEC evaluation report, that we can resolve this without that. And so current status is we are working with Department of Labor and we should have a update for the June

meeting.

Massachusetts Institute of Technology, we presented at the beginning of the meeting -- this meeting that we -- we had issued our evaluation report and we pulled back that evaluation report after it was recognized that there were other prime contractors or other contractors at the Hood facility and we needed to review additional documentation. We anticipate that we will have this evaluation completed or with a path forward at the -- and we will make that presentation at the June meeting.

Dow Chemical, we have been working with the former -- with Dow Chemical to retrieve additional documentation based on the new covered per-- or new designation that Department of Energy had fin-- its -- had -- Department of Energy had indicated that the -- had made a determination that the thorium alloy at Dow Chemical was actually part of the

1 weapons program or it could have been part of 2 the weapons program and had determined that it 3 would be a covered activity. We had not evaluated that in our initial evaluation, so we 5 have gone back now, requested additional 6 documentation through the State of Illinois and 7 through Dow Chemical on the thorium operations, 8 and we plan to issue a revised evaluation at 9 the June 2008 meeting. 10 Savannah River Site is currently in the 11 evaluation phase, and we anticipate issuing a -12 - a report and presenting that report -- we --13 issuing the report in August 2008. 14 And currently there are six sites that are in 15 various stages of the 83.14 SEC process, and a 16 -- we are working to have a few of those 17 available for the June meeting. 18 That's it. Ouestions? 19 Thank you, Sam (sic), that's very DR. ZIEMER: 20 helpful. I think Larry Elliott has a comment 21 and then we'll (unintelligible). 22 Thank you. I just want to make MR. ELLIOTT: 23 one addition to the Y-12 and that is, beyond 24 what Bomber said -- or LaVon said about us 25 working with DOL, we are also working with

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individual claimants that come forward to us or to Denise Brock, the NIOSH ombudsman, who have in their hands a situation where DOL has not found them to be eliqible for this class. And so we're -- we're working on their behalf to go back to DOL and talk to DOL about why, and so there is that component going on here, too,

I assume that once NIOSH and Labor agree to that definition, you will come back to us and clarify it to us so that we can be assured that what we think we voted on is -- is properly covered in the definition.

Denise may have an additional comment here on

MS. BROCK: I do. I just wanted to report that we've actually had probably about three or four cases that have actually been overturned that have went back to the Department of Labor and have some very pleased claimants for that, so that -- that's very exciting. So hopefully, as Larry said and as LaVon said, we'll get the rest of that taken care of.

Thank you. LaVon, you had a

1 MR. RUTHERFORD: Well, my only comment was --2 on that as well was actually to say that we 3 have a pretty good approach. We actually have reviewed all the claims that fit into that pre-5 1947 period, and we've laid out -- we've looked 6 at their interviews, we've looked at all their 7 documentation, and we've identified claims that 8 we felt that should have fit into the class and 9 made that -- made those available in a 10 spreadsheet to the Department of Labor and 11 we're working with them, as Denise had 12 mentioned. 13 DR. ZIEMER: Jim. 14 DR. MELIUS: Yeah, how many petitions do you 15 have that are under evaluation now for whether 16 they qualify? 17 MR. RUTHERFORD: We have four. 18 DR. MELIUS: Four, okay. 19 MR. RUTHERFORD: We have two at the Linde 20 facility and we have recently received Los 21 Alamos National Lab and -- I thought I had them 22 Those are three of them, anyway. 23 DR. MELIUS: That's not bad. 24 DR. ZIEMER: You have -- in your last slide you

mention the six sites --

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1	MR. RUTHERFORD: Yeah, we
2	DR. ZIEMER: under consideration. That's
3	MR. RUTHERFORD: Yeah, those are actually
4	you don't those are not considered a
5	petition until we get the Form A back from the
6	
7	DR. ZIEMER: That's in addition to these four.
8	MR. RUTHERFORD: Yes, yes, that's in addition
9	to those four.
10	DR. ZIEMER: Okay, thank you. Other comments
11	or questions?
12	(No responses)
13	Okay, thank you. That's very helpful.
14	MR. RUTHERFORD: All right.
15	DR. ZIEMER: We have a we have a little time
16	before we actually take up these next ones
17	have to be fairly time certain, so I want to
18	see if we have some items that we can handle in
19	the meantime.
20	DR. BRANCHE: There was going to be a report
21	is John Mauro in the room? There was going to
22	be a report by SC&A on the budget issues. That
23	was going to be part of the Board working time.
24	I don't know if you want to take
25	DR. ZIEMER: Let me let me

1 DR. BRANCHE: -- or not. 2 DR. ZIEMER: -- ask -- hang on a minute. 3 (Pause) 4 DISCUSSION OF SC&A BUDGET ISSUES 5 Is John Mauro in the assembly? John, you were 6 going to report on SC&A budget issues. 7 this be something you're prepared to do now if 8 we --9 DR. MAURO: Sure, yeah. 10 DR. ZIEMER: -- pick that up? This is part of 11 the --12 DR. MAURO: Let me get my notebook, I have some 13 (unintelligible) --14 DR. ZIEMER: -- Board working time. 15 DR. BRANCHE: For individuals who joined the 16 telephone line, if you could please mute your 17 phones. If you do not have a mute button, 18 please use star-6 to mute your phone, and when 19 you're ready to speak you can use that same 20 star-6. But we do need you to mute your phones 21 now. Thank you. 22 DR. ZIEMER: For members of the assembly to --23 if you're trying to track where we are, 'cause 24 we've had to jump around here a little bit on

the agenda, the part of the agenda called SEC

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Petition Status Updates scheduled for 3:45, we're holding that at that time -- we're keeping it at that time because of individuals who will be joining us, either -- well, mainly by phone, some in person. But we want to keep that at a -- basically a fixed time. So what we're doing now, we're picking up pieces of what's called Board Working Time, which is on your agenda for tomorrow afternoon. For example, we covered the tracking status of transcripts and minutes. We talked a little bit about the status of next year's Board contractor support. Now we're going to have a brief report from the Board contractor, SC&A, which deals with some budgetary issues. basically want to bring the Board up to date on some issues relating to this year's budget. John Mauro will fill us in on that.

DR. MAURO: I guess it was about a week or so ago when I informed the working group, the Task III working group on dealing with procedures, that -- and that was -- we were about to run out of money on Task III, and to -- up -- that was -- that was about a month in advance, and we had a special conference call with Dr.

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Ziemer, yourself and David Staudt and Lew Wade and Christine, and we discussed what to do about that, strategies. And I did co-- come up with one strategy that might work to help resolve that problem.

In effect, if you -- we would step back a bit, our -- our -- we have a system of keeping track of how much more work we have to do, how much money do we have, do we have enough money to finish all the work. The work -- the contract will end in the end of September. Bottom line is -- I'm stepping a little back from -- from this Task III issue, but Task III is sort of like a subset of this issue. In effect, our budget for this contract was about \$13 million. We project, using this earned value system that I keep track of on a monthly basis, that we are probably going to run short by about \$1.2 million and about -- in effect, the cost of -our projection, using our system, says we're probably going to be ten percent un-- overbudget, if -- if, now there's a big "if" here -- keep in mind what an earned value system is. It's a system that you start in the beginning of a project to score -- to -- let's say I

think that this is the rate at which we will proceed in terms of spending money and accomplishing things. In developing that system I made certain assumptions about how much money we're going to need to do various tasks.

That brings me to Task III. It turns out we ended up spending more money on Task III than we anticipated, and there's a variety of reasons for that that I -- no need to go into that now, but -- and one way to help resolve the problem goes to Task I. Please bear with me.

In Task I we currently have ten site profiles that we've completed -- the review of the site profiles that we've completed, and they're sitting on a shelf, but we have not activated a workgroup and so no work is -- we're really not in a closeout process. Now what I do -- what I did is for every time we deliver a report, a site profile review report, I put in the bank \$61,000. That's basically saying I'm going to hold \$61,000 available for the -- when the day comes when we open up -- when we have a working group and we start the closeout process. This

is based on historical experience on what does it cost to go through the whole closeout process for all the issues on one of our site profile reviews.

So in effect right now we have ten site profile reviews -- you can read them if you'd like the list of them; they are INL, LANL, X-10, Pinellas, Paducah, K-65, Lawrence Livermore National Laboratory, Pantex, Portsmouth and Argonne National Laboratory West. All of those are sitting and waiting for a working group. Which means there's \$610,000 sitting in the bank waiting to take that on.

Now one of the things that -- and they've been sitting there for -- for some time. But in the interim, from when we wrote those reports to today, of course we've learned a lot. We've enga-- we've had a closeout process with many procedures, closeout process for many site profiles, many SEC petitions. Where I'm heading is that I suspect if SC&A were to go back and look at the findings contained in the ten site profile reviews as a group, just collect them all and come and report back to the Board, or to a working group of the Board,

and summarize -- and sort of do a -- some triage, would -- let's say it turns out there are ten findings per site profile, just for discussion. We don't need 100 findings associated with these ten site profile reviews. I strongly suspect that some fraction of that, maybe 25 percent, are issues that have already been resolved in another venue because we've seen them before. Can't say that for certain -- I'm using this as an example. There may be other issues that we're currently resolving and dealing with actively, either as part of procedure review process or we've -- or as part of one of the other site profile reviews that are currently active.

Where I'm headed is -- and this is a bit speculative, though -- in theory, let's assume that 50 percent of the issues imbedded in those ten site profile reviews can be -- say are -- are already well in hand, we're dealing with them. The way I look at it is this: That means we have \$610,000 sitting in the bank when perhaps we really only need \$300,000. It frees up \$300,000 -- this is all hypothetical now. I haven't done any of this yet, it's a concept.

It frees up \$300,000, which I believe could be money that's sitting there to be used, one, to help out on Task III, to help keep the closeout process going 'cause, you know, we're through with all the procedure reviews but we're not through with the closeout, so that has to continue.

Also, as I understand from watching these proceedings, there may be a number of new SEC petition support work. Those resources could be made avail-- right now we're -- we're okay on Task V, which is the SEC. But if we start to add in a number of additional focused reviews on some of the new SEC petitions, it may turn out that we -- we could use those resources and -- in helping to relieve the pressure on Task Order V.

So I guess what I'm saying is I'm bringing before the Board an idea that -- I guess there are a couple of things that could be done -- add additional resources to the contract, or shift some resources between tasks. And I think one place where I think it's a very good possibility that we could be able to shift some resources is regarding this Task I matter. But

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it would require, I would say, SC&A to take a look at these ten site profiles and then go through this triage process. We would do it initially within SC&A and -- and perhaps make some type of matrix, sorting; bring that before perhaps a special workgroup that might be formed to help with this process; or of course, bring it before the Board and -- and tell our story. Say out of these 100 issues, so to speak, here's how many we believe are being handled or have been handled, but -- and here's what's remaining. And in that way we can get a sense of how much resources we could break free from Task I closeout and make it available for other tasks that could use the resources. DR. ZIEMER: Thank you, John. Let me add a couple comments to that and then we'll open The immediate issue of this for discussion. being short of funds has been handled between Dr. Branche and David Staudt and -- and some of the funds have been shifted from one task to another to allow Task III work to continue. Further, a fair amount -- and I don't know how much, but a fair amount of what's been carried out under Task III might -- one might arque is

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more generally administrative work; that is the development of the -- of the database tool that was described to the Board yesterday by Kathy Behling. One might argue it's not exclusively a Task III item. One might argue that it is either administrative or it perhaps could be spread over other tasks since it has potential applicability to others as well.

In any event, the -- the immediate problem has been handled. But going forward, as John has indicated, he has put funds in reserve to handle clear work that has to be done by our contractor, and by us as we have workgroups available, to resolve the findings in those completed site profiles. So what John has described here to us now is a sort of streamlining process that supposes that there are common issues in the remaining site profiles -- perhaps issues like the neutron dosimetry issue which seems to have reoccurred in several past site profiles, which have been largely addressed. Maybe they'd have to be further tweaked for a given site, but for which we don't need the full funding to address yet another time. So I think that's kind of the

1 thrust of -- of what John is suggesting. And I 2 think what we're looking for is some feedback 3 from the Board for -- basically this -- this is 4 the last year for that contract. We don't 5 know, going further, whether the new contractor will be SC&A or some other entity, but we have 6 7 to think about completing this year and 8 completing those tasks. And it's important 9 that SC&A be able to complete what they've been 10 assigned. They've been assigned to close out 11 those site profiles. That's an obligation to 12 them and also to us because they can't complete 13 that without Board input and without exchange 14 with NIOSH on -- on issues as well. 15 So with that as background, I think -- Dr. 16 Melius, you have a comment or a suggestion? 17 DR. MELIUS: No, I have first a question --18 actually a couple of questions. Fir-- first of 19 all, when we're talking about a year left on 20 the contract, what are -- what are you 21 specifically talking about, when --DR. ZIEMER: Well, I believe it's the fiscal 22 23 year of the government, so it would go to 24 October. Is that not correct? 25 DR. MAURO: Yes, October 1st.

1 DR. ZIEMER: By end of September. 2 DR. MAURO: October 1st would be the 3 termination date. 4 DR. ZIEMER: And actually I -- I think what 5 would happen, as I understand from David Staudt -- for example, if there were a new contractor, 6 7 it would still be possible for funds -- a no-8 cost extension to carry forward to allow 9 closeout with the present contractor. 10 DR. MELIUS: Uh-huh. 11 Or even if it's the same DR. ZIEMER: 12 contractor, you can carry some funds forward. 13 So we are still looking at sort of this year as 14 a package, either way. 15 DR. MELIUS: Uh-huh. Okay. But so -- so we're 16 talking about six months. 17 DR. BRANCHE: Yes. 18 DR. MELIUS: Yeah, yeah -- I mean roughly --19 roughly. 20 DR. ZIEMER: And I might observe, and I think 21 you realize that closing out ten site profiles 22 in six months, even if we had all -- even if we 23 had to use all the money, would be a formidable 24 task for this Board in terms of our workgroup 25 activities.

1 DR. MELIUS: Yeah. Secondly, has NIOSH 2 responded to the SC&A review of any of these 3 site profiles, or how many have they responded 4 to? 5 Those ten, no response. words --6 DR. MELIUS: No, no --7 8 DR. MAURO: -- these are ten site profile 9 reviews, the ones I just read, that were 10 delivered but the -- to the Board, but there 11 has not been a workgroup formed or the process 12 started to close out those issues. 13 DR. ZIEMER: I think we should recognize that 14 the reality is, as we prioritize things, 15 usually the closeout process is triggered by 16 the Board saying we're ready to move ahead. I 17 understand that they're there. 18 information's there for NIOSH to look at. 19 20 DR. MELIUS: Yeah --21 DR. ZIEMER: -- unless -- unless we're ready to 22 go with it, it -- well, you understand --23 DR. MELIUS: Yeah, yeah, I'm -- I'm not 24 faulting -- trying to fi-- fault NIOSH for not 25 having done something. I'm just trying to

1 think if we're going to try to do this process, 2 one, does NIOSH have the time to do it in six 3 months, and --4 DR. ZIEMER: Well, they would have --5 DR. MELIUS: -- secondly --6 DR. ZIEMER: -- the same issue as the Board; 7 it's a formidable task. 8 DR. MELIUS: And they have some resource 9 issues, too, at least in the sort term, with 10 the renewal of the ORAU or whatever's going on 11 with the contract and so forth, so -- the new contract. So I mean I think -- I'm not sure 12 13 how far we're going to get with those six --14 DR. ZIEMER: Well --15 DR. MELIUS: -- though I think we can think of 16 a way of starting, but I think what John was 17 talking about, I -- I just don't see us being 18 feasible to do in that -- that time period and 19 20 DR. ZIEMER: Let me add one other comment and 21 then we'll hear from Larry. A concern I have -22 - let -- let's assume, for example, that -- one 23 scenario is we have a new contractor. It would 24 be very difficult for us to be closing out 25 those -- those documents with a contractor for

1 whom it -- it's not their findings. 2 DR. MELIUS: Yeah. 3 DR. ZIEMER: So -- okay. 4 MR. ELLIOTT: Larry Elliott. I didn't take it 5 that you were finding fault with us, Dr. 6 Melius. When you first made your comment, I 7 thought well, is he talking about a departure 8 from process here, because certainly I could 9 say that NIOSH could pick up one or two of 10 these reviews and examine them and tell the 11 Board where -- you know, where we're at on 12 them. 13 DR. MELIUS: Yeah. 14 MR. ELLIOTT: We can do that. DR. MELIUS: 15 Yeah. 16 MR. ELLIOTT: It's going to -- you know, it's 17 going to take time and resources to do that --18 DR. MELIUS: Yeah. 19 MR. ELLIOTT: -- but that's -- that would be a 20 departure from process and so -- I'm not 21 advocating one way or the other, but I'm saying 22 that's something that could be looked at as 23 well. We could work with SC&A on a technical 24 level and react. 25 Here's the other part that I would like the

Board to think about --1 2 DR. MELIUS: Yeah. 3 MR. ELLIOTT: -- and as each one of these 4 reviews are sitting on the shelf, they are also 5 on our web site, and claimants hold these up to 6 DOL and say look at this review on this site 7 profile, and there are deficiencies noted in 8 this. And so the Final Adjudication Branch of 9 DOL turns that back to us to answer. All 10 right? 11 DR. MELIUS: Uh-huh. 12 MR. ELLIOTT: And in some cases when we get 13 those back, we pend them. 14 DR. MELIUS: Uh-huh. MR. ELLIOTT: We hold those claims. 15 And so 16 those claimants are now further frustrated --17 DR. MELIUS: Uh-huh. 18 MR. ELLIOTT: -- because they're back at NIOSH 19 and they're not getting any answer. 20 DR. MELIUS: Uh-huh. MR. ELLIOTT: And we can't move forward because 21 22 we don't have closure on a set of issues. 23 other cases we are able to provide a definitive 24 disposition of the claim without going to the 25 point of closure.

DR. MELIUS: I guess a third question then is that -- say we -- okay, we have the procurement for the -- the new review contract that's going to go in place. Should SC&A be the -- successful in that, then would this money roll over into the new contract? How does that work? Or -- or would it -- would it roll over or would the activity be allowed to continue, I guess, it'd sort of be melded into the new contract if -- if SC&A were successful and decides to apply and wins and all that stuff.

DR. BRANCHE: Okay. You asked a couple of questions, let me try -- at them all.

DR. MELIUS: Yeah, yeah.

DR. BRANCHE: How -- how a new Board contractor is selected in this next cycle, to a degree, is a bit divorced from the current set of activities. And that's what I think Dr. Ziemer was saying. You've got professional opinions that you've sought from your current Board contractor, and those are pending further action from this Board.

Now, he also mentioned that in our discussion with David Staudt, the procurement and grants officer for this -- for the Board from CDC,

1 there is an opportunity for SC&A to be in a no-2 cost extension situation to close out those --3 the information from their current contract if SC&A is not selected on the next round. 5 DR. MELIUS: Yeah, but I think there's -- Dr. 6 Ziemer mentioned that possibility, but I -- the 7 possibility I was -- what happens if SC&A is 8 selected in the next round? 9 DR. BRANCHE: Then --10 DR. MELIUS: The -- then, you know --11 DR. ZIEMER: I think Lew has some comments. 12 ahead, Lew, you can help us on this 'cause 13 you've been involved with David on procurement, 14 but I believe those funds can still roll 15 forward, can they not? 16 DR. WADE: Right, they can in -- in two ways. 17 I mean theoretically, if SC&A was to secure the 18 next contract, you could have two contracts 19 running concurrently. You could have the 20 existing contract with the money in it, or it 21 might be prudent for the government to in some 22 way combine those two --23 DR. MELIUS: Uh-huh. 24 DR. WADE: -- but those --25 DR. ZIEMER: One thing we do know, that the new

1 contract -- let's say it is a different entity 2 -- does not contain money for closing out the 3 old contractor's work. DR. MELIUS: Yeah, no, I -- that's 4 5 (unintelligible). 6 DR. ZIEMER: And the deliverables under the contract were -- for example, the site profiles 7 are deliverables under the contract, so S&CA 8 9 has met that. But then we have the other task 10 which involved the resolution of those. 11 MR. ELLIOTT: You have to think of it this way. 12 The money that has been awarded under this current contract goes to SC&A --13 14 DR. MELIUS: Yeah. 15 MR. ELLIOTT: -- to finish the task. 16 DR. BRANCHE: Right. 17 DR. MELIUS: No, no --18 DR. BRANCHE: And you've heard from John Mauro 19 on a coup-- this is Christine. You've heard 20 from John Mauro on a couple of occasions that 21 they are holding money on reserve to be able to 22 close out those reports. And I would just say 23 it's just prudent bookkeeping for the cycle of 24 -- of assignments to be completed in the ti--25 as close to the time frame as possible.

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DR. MELIUS: Uh-huh.

DR. BRANCHE: -- and I think Mr. Elliott's given you additional reason why, for other reasons as it concerns the claimants, that there are some reason-- that there's -- there are good reasons to be able to close those out.

DR. MELIUS: Uh-huh.

DR. ZIEMER: Now -- oh, go ahead, Jim.

DR. MELIUS: I would guess then sort of my opinion on how we should do here is, one, we -the Board does need to work out a process for closing those out. I'm not sure a single workgroup is the -- is the ri-- best way and most efficient way, and I'm not sure I'd wish it on my other Board members or -- or they would wish it on me, I guess, given the potential scope of that. But I -- but I do think we -- we're obliged to come up with a mechanism and a schedule for figuring out how to resolve that within the -- the available resources. I'm just very skeptical that we -that the way John suggested would -- would actually be workable in a -- in that period of time, given what it would take, not only from

1 the Board members but also from NIOSH, to be able to -- to devote in terms of resources, 2 3 time and -- time and effort. 4 DR. ZIEMER: Well, let -- let me suggest a 5 couple of things, perhaps to help focus our thinking. There's two parts to this. 6 One is a 7 resource issue for, in a sense, moving money 8 out of the site profile part to make it 9 available -- 'cause that's -- that's John's 10 bank right now. 11 DR. MELIUS: Yeah. 12 And if the -- if the Board thinks DR. ZIEMER: 13 something along the lines that John has 14 described, maybe some variation of that or 15 maybe that exactly, is a useful thing, we could 16 get them underway on that, as a mat-- as an 17 effort to free up funds, for example, to cover 18 Task III. 19 The other part of it is, we need to be thinking 20 about the schedule itself for the closeout. 21 DR. MELIUS: Uh-huh. 22 DR. ZIEMER: Whether or not we go to 23 streamlining, I think we all recognize closing 24 out ten site profiles, most of which are pretty 25 sizeable facilities -- they're not -- not the

little small guys, so to speak -- that a sixmonth turnaround is just not feasible.

DR. MELIUS: Uh-huh.

DR. ZIEMER: I would like to ask the question can we even think about a -- an 18-month turnaround, which is -- you know, for carrying money forward, if we're going much beyond that, that's a problem, but -- but I -- I guess I'd like the Board and SC&A to start to think seriously, maybe -- well, we -- we can't postpone thinking about this and say well, let's be thinking about it and we'll start to take action in three months or six months.

DR. MELIUS: No.

DR. ZIEMER: We need to -- to start looking and say okay, what site profiles are we going to start working on sort of right away off the shelf, and -- and get some kind of a schedule on those. And perhaps how can that process be streamlined to free up the resources, so there's two parts to that. And per-- I -- I would suggest, maybe when we come to our work session tomorrow, that we come -- I don't want to invent this --

DR. MELIUS: Yeah.

1 DR. ZIEMER: -- on the spot. 2 DR. MELIUS: So you'll be --3 DR. ZIEMER: But I don't want us to say yeah, 4 let me cogitate for the next three months, or 5 by October 1st I'm going to have a solution, because it's -- it's pressed upon us. 6 7 - in essence was thrust upon us by -- perhaps 8 this was a good thing for that particular 9 budget to call attention to what was going on. 10 DR. MELIUS: Yeah. Can -- you know, I -- thank 11 you for letting us procrastinate at least until tomorrow 'cause -- but two pieces of 12 information I think would be useful to have by 13 14 -- before we meet tomorrow. One is I would 15 certainly like to have a fuller understanding 16 of -- of what is the amounts of money left --17 funding left in the different tasks. 18 you said you were able to do a short-term take 19 -- take care of this, but I just need to 20 understand what --21 DR. ZIEMER: Yeah. 22 DR. MELIUS: -- what -- what happens and what 23 was --24 DR. ZIEMER: Well, we have the monthly roll-ups 25 and we can come up with that very easily.

1 fact, I have it on my computer, but we'll --2 we'll have that in our work session tomorrow --3 DR. MELIUS: Tomorrow, it -- yeah --4 DR. ZIEMER: -- on where we are on each task, 5 and John can give it to us by percent. 6 -- we want to be around 50 percent of the task 7 8 DR. MELIUS: Right. 9 DR. ZIEMER: -- and this one was getting up 10 toward 90 percent --11 DR. MELIUS: Yeah -- no. DR. ZIEMER: -- and of course, as John said, 12 13 he's put some money away, so he's been a 14 prudent guard of -- of some of that money as well, so --15 16 DR. MAURO: There is a -- there's \$3 million 17 left in this project for SC&A to use, in 18 theory, over the next six months. So there are 19 a lot of resources, but a large fraction of 20 that is in the bank because we're moving -- we 21 -- we've completed the vast majority of our 22 deliverables, our procedures, our site profile 23 reviews. We only really have two site profile 24 reviews that are be-- in process right now, 25 Santa Susana and Weldon Springs. We've

1 completed 26 site profile reviews. So I mean -2 - think of it like this. There are -- of all -3 - of all the work we do, we deliver our work 4 product as a draft and then we move into the 5 closeout. Well, in effect, what I -- what I'm 6 saying is that we have substantial funds, but 7 we also have a substantial amount of work that 8 has to be done to close out all of these produ-9 - work products, so -- and I certainly have all 10 the -- every -- all the information you might 11 need, how much resources are left in each one 12 of the tasks, and how to use those resources. DR. ZIEMER: And also it's -- it's probably --13 14 as we think about this we may need to think 15 about, for example, whether or not it's prudent 16 to assign more profiles when we have all this 17 backlog to resolve. 18 Right. DR. MELIUS: 19 DR. ZIEMER: What good does it do to put 20 another one on the shelf at this point, you 21 see. 22 DR. MELIUS: And -- and then can we get a list 23 of the -- these site profiles that are in limbo, so I think we --24

DR. ZIEMER: Right, I think you read my --

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It's a rare

1 2 DR. MELIUS: -- a written -- a written list

would be --

Wanda?

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DR. ZIEMER: We'll get the list for -- right.

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The budgetary concerns that we

express here often fail to take into

budget plan on a regular basis.

consideration what we as a Board do to the

occasion that we do not postpone something that

we are doing -- case in point, this very

meeting -- with a request that our contractor

perform a, quote, focused review or a broader

review of a point or other points. Now these

items have not been factored into our budget

process, and we continually ask our contractor

-- this is not an obscure case. We do this

almost every meeting. We're asking them to add

So when we find something more to the process.

ourself in a position where we're squeezed in

terms of where we want the budget to be and

where our contractor wants the budget to be, it

would seem wise for us to be very conscious at

each step of our own process that we're

creating a portion of the problem that we're

attempting to overcome every time we say let's

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postpone this for six months and get a focused review in the meantime. We're adding to the issue.

DR. ZIEMER: Now keep in mind that the budget -- or the tasks and the budgets do in fact specify certain monies for certain numbers of focused reviews. And Lew has been helpful, and Christine now, in making sure that in fact when we do that -- and they touch base with David Staudt to make sure it's within the framework of the larger task. So yes, we do add to that workload, but in general we've kept within the framework of -- of the annual big picture. what has happened in this particular case -- I think in the procedures review, as we've developed -- I think particularly the -- the new instrument for -- for data sorting and so on, I believe that's taken more resources than we had originally thought it would, and the product is great and probably well worth it, but it has brought us to this -- this -- or at least focused on this issue, so -- Lew, you had another comment -- or Jim, you --

DR. MELIUS: No, I -- you gave my comment for me. Thank you.

1 DR. ZIEMER: I took the words out of your 2 mouth. 3 DR. MELIUS: Out of... Out of the mouths of babes. 4 DR. ZIEMER: 5 DR. MELIUS: Go that far with... DR. ZIEMER: 6 Let's see -- well, we'll return to 7 this tomorrow during our work session. have a time certain item that -- or items that 8 9 are before us here, and that is the -- the SEC 10 petition updates dealing with Hanford, Sandia and Chapman, and we will also have -11 SEC PETITION STATUS UPDATES: 12 HANFORD 13 DR. BRANCHE: And we have -- we're also going 14 to have one thing back on Mound. 15 DR. ZIEMER: One Mound item, okay. So let us 16 begin with Hanford, and actually LaVon 17 mentioned all of these in his summary, and now 18 we will have the specifics on Hanford. 19 also with regard to Hanford -- just checking my 20 list here -- Mary Ann Carrico, Rosemary Hoyt 21 and [name redacted] I think are going to be 22 with us. Are either or all of you on the 23 phone? 24 MS. HOYT: This is Rosemary and I am on.

DR. ZIEMER: Okay, Rosemary. And Mary Ann, are

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1	you on?
2	MS. CARRICO: Yes, I am.
3	DR. ZIEMER: Very good. [name redacted]?
4	(No responses)
5	Mary Ann or Rosemary, do you know if [name
6	redacted] is going to be with us?
7	UNIDENTIFIED: No, I have not heard from him.
8	DR. BRANCHE: Laurie, have you heard from him?
9	DR. ZIEMER: [name redacted]?
10	MS. BREYER: I mean I I mean I haven't
11	called him today. I think he was a maybe, so
12	he said yeah, he'd probably be on so I have him
13	on there as a yes, but
14	DR. ZIEMER: But we should proceed?
15	MS. BREYER: I can try to give him a call
16	yeah, I'd proceed and I'll call him.
17	DR. ZIEMER: Okay, we're going to proceed, so
18	let's hear first from from Sam Glover.
19	DR. GLOVER: Bomber already gave a good update
20	so he stole my thunder. All right, so we're
21	going to step back just a little bit and
22	because this was a two-part review, I thought
23	I'd just remind everybody how we got to this
24	where we are.
25	So this is a the Hanford update for the

1 Special Exposure Cohort petition, Part 2. 2 just a little bit of background. We had three 3 Hanford petitions qualify. One November 9th, 4 2006, which was all production workers in the 5 100 and 300 areas in the very earliest years of 6 Hanford. We had another one, SEC-57 -- that 7 was SEC-50, the first one. SEC-57 was November 8 -- qualified on November 21st, 2006, which was 9 all employees in all facilities in areas of 10 Hanford Reservation from 1942 though December 11 31, 1990. And then we had a third qualifying 12 on February 28, 2007, which was for all roving 13 maintenance, carpenters and apprentice 14 carpenters that worked in the 100, 200, 300 and 15 400 areas of Hanford from April 25th, 1967 16 through February 1, 1971, and that was SEC-78. 17 So just a brief reminder, these three petitions 18 were merged into a single petition and 19 evaluated under SEC-57. 20 UNIDENTIFIED: (Unintelligible conversation) 21 DR. GLOVER: We split them into two periods 22 because there was some... 23 UNIDENTIFIED: (Unintelligible conversation) 24 DR. BRANCHE: Excuse me. 25 DR. GLOVER: Yes.

DR. BRANCHE: The pers-- the people who are on the phone, if you could please mute your phones. If you do not have a mute button, then please use star-6. Thank you.

DR. GLOVER: So one of the major reasons that we split that was because Part 1 was evaluating the DuPont years, during the DuPont contract time, from 1942 through September 1, 1946. And then Part 2 was from September 1, 1946 through 1990. This presentation reports the conclusions of the evaluation for Part 2, and that first re-- that report was issued September 9th, 2007 -- I'm sorry, the evaluation on May 2007 and presented to the Advisory Board in July of 2007.

Just a brief reminder, the summary of the class added under Part 1, employees of the Department of Energy -- this just summarizes -- this was added October 12th, 2007, employees of the Department of Energy, its predecessor agencies or DOE contractors or subcontractors who were monitored, or should have been monitored, for internal radiological exposures while working at the Hanford Engineering Works in the 300

area fuel fabrication and research facilities
from October 1, 1943 through August 31st, 1946;
and the 200 area plutonium separation
facilities from November 1, 1944 through August
31st, 1946; or the 100 B, D and F reactor areas
from September 1, 1944 through August 31st,
1946 for a number of work days aggregating at
least 250 work days or in combination with work
days within the parameters established for one
or more other classes of employees in the
Special Exposure Cohort.

So that brings us to the second evaluated class, which we evaluated all employees in all facilities and areas of the Hanford Nuclear Reservation from September 1, 1946 through December 31st, 1990. This was presented to the Advisory Board in September of 2007. As part of this report, NIOSH's original class recommendation was as follows: All employees of the Department of Energy, its predecessor agencies and DOE contractors or subcontractors who were monitored, or should have been monitored, for, one, internal thorium radiological exposures from September 1, 1946 through December 31st, 1959 in the 300 area

facilities: the metal fabrication building, 313; the reactor fuel manufacturing pilot plant, 306; the 300 area maintenance shops, 3722; or the radiochemistry laboratory, 3706; or internal americium radiological exposures from January 1, 1949 through December 31st, 1968 in the following areas: the isolation building, 231-Z; the waste treatment facility, 242-Z; and the plutonium finishing plant, 234-5Z while working at the Hanford Nuclear Reservation -- the standard language regarding the 250 days.

So that brings us to the update. So as LaVon mentioned, SC&A has had -- issued several white papers since we had our report, and NIOSH has continued to evaluate these param-- these class -- they issued a report on americium, thorium and uranium, and they discussed primarily where operations were conducted outside of the areas that we -- they limited the scope of their evaluation to the time frame that we had proposed. And they put forth that there may be other facilities that these were -- would be a concern.

So NIOSH has continued to research these and

1 other topics. I will admit, as we have 2 discussed previously, that progress has been 3 hindered by inability to access DOE data. We have had several workgroup calls and 5 meetings. We had a workgroup call on March 6 6th, 2008 and what I would like to put forward 7 is that, based on the additional research, 8 NIOSH proposed to revise the class definition 9 and reissue the evaluation report for Part 2 of 10 SEC-57. The proposed changes to the class will 11 allow DOL to effectively administer the 12 proposed class. We also followed this up with a working group 13 14 call. These follow-up discussions were held between SC&A and NIOSH. The matrix was updated 15 and discussed. We had some initial 16 17 prioritization to the matrix items and kind of 18 worked out what will be -- how we're going to 19 proceed. 20 Finally, NIOSH issued a revised evaluation 21 report on March 31st, 2008, of which I believe 22 everyone was provided a copy. 23 So as part of that, NIOSH updated the proposed 24 Hanford class, and the language will now -- now 25 reads: All employees of the Department of

1 Energy, DOE or its predecessor agencies and DOE 2 contractors or subcontractors who worked from 3 September 1, 1946 through December 31st, 1961 in the 300 area; and from January 1, 1949 5 through December 31st, 1968 in the 200 area at 6 the Hanford Nuclear Reservation for a number of work days aggregating at least 250 work days 7 8 occurring either solely under this employment 9 or in combination with work days within the parameters established for one or more other 10 11 classes of employment (sic) in the SEC. 12 As part of that, we'll restate the health endangerment, that NIOSH has determined that it 13 14 is not feasible to complete dose 15 reconstructions with sufficient accuracy for 16 1949 through 1968 period in the 200 area for 17 hazards associated with americium, nor for the 18 1946 through 1961 period in the 300 area for 19 hazards associated with thorium. 20 NIOSH finds that the health of employees 21 covered may have been endangered from chronic 22 exposures from production and research 23 activities in these areas. 24 (Pause)

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So then a summary of our standard summary of

1 feasibility slides. You note that for -- it is 2 not feasible for thorium or americium during 3 these time periods. For the other materials, 4 we still retain that being feasible for 5 plutonium, fission products, tritium, polonium, 6 iodine, ambient environmental, and that --7 believe we can reconstruct external doses in 8 this time period. 9 With that, Dr. Ziemer, I conclude the update. 10 DR. ZIEMER: Thank you, Sam. Let's see if --11 before we hear from the petitioners, let's see 12 if we have any questions from the Board members on Sam's report. 13 14 (No responses) 15 If not, I'd like to ask Mary Ann or Rosemary, 16 do either of you have comments for the Board 17 today? 18 DR. BRANCHE: We may have some speakers from 19 the audience as well. 20 MS. HOYT: Yes, this is Rosemary and I would 21 like (break in transmission) (unintelligible) -22 23 DR. BRANCHE: If she could speak up. 24 MS. HOYT: -- worked so hard any (break in 25 transmission) (unintelligible) --

1 DR. ZIEMER: Mary Ann, you are breaking up, 2 we're having a hard time hearing. Let's ask if 3 -- oooh. 4 MS. HOYT: Hello? 5 Yeah, try it again. We're having DR. ZIEMER: trouble -- your phone seems to be breaking up 6 7 as you speak. We're hearing just clipped vowel 8 sounds. Maybe move back a little bit from the 9 mouthpiece and try again. MS. HOYT: Okay, I took it off speaker phone. 10 11 Is that better? 12 That's much better. Thank you. DR. ZIEMER: 13 MS. HOYT: I would also like to 14 (unintelligible). I've had difficulty with 15 (unintelligible) getting his (unintelligible). 16 I'd like to just thank everybody who has worked 17 on this and (unintelligible) also to Dr. Melius 18 (unintelligible). 19 There are a few things that I would like to go 20 over on the (unintelligible) petitioner 21 requested (unintelligible) basis in NIOSH-22 proposed class (unintelligible), and this is a 23 quote. The SEC-00057 petitioner 24 (unintelligible) exists for several individual 25 workers listed in the petition. NIOSH found

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monitoring data information (unintelligible)
petitioner did not support submission date
qualifying the petition. However, during the
qualifying process NIOSH identified
(unintelligible) monitoring for
(unintelligible) NIOSH qualified SEC-00057 on
this date, end quote.

There are two misrepresentations in the above statement. The first is petition 00057 cites far more than (unintelligible) monitoring records. We responded (unintelligible) request (unintelligible) 2006. This letter became a supplement to (unintelligible) petition. letter (unintelligible) were not monitored or (unintelligible) monitored, falsification of records, (unintelligible) records, underreported neutron doses, (unintelligible) not accurate and in adequate. Bioassay records did not exist or were lost or destroyed. section (unintelligible) point four of the original petition, we included the Hanford site profile (unintelligible) requesting that NIOSH qualify the petition based on their findings. When Laurie Ishak, now Breyer, called to tell us the petition had qualified for evaluation,

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she told us it was (unintelligible) based on the SC&A report. Finding two in the (unintelligible) of that report specifically addressed (unintelligible). I request that (unintelligible) be corrected to accurately state the facts.

4.3 of the (unintelligible) report, facility employees and experts. There were two groups interviewed. The minutes (unintelligible) really something that gets to me. The minutes of the March meeting are (unintelligible) on the OCAS web site, but (unintelligible) minutes are not. The (unintelligible) minutes were specifically for the class covered by (unintelligible) petition 00057 Part 2. (unintelligible) of this information is worthless. It's outrageous that these minutes cannot be (unintelligible) in a timely manner. External monitoring in general (unintelligible) not be consistently applied (unintelligible) stated. I state that if, quote, consistently applied, end quote, equates consistently absent, this might be a true statement. In the comments to the Advisory Board for October 4th, 2007 we stated worker outreach

1 meetings (unintelligible) the worker exposure A 2 was not monitored for all employees, B 3 (unintelligible) bucket at the end of the 4 shift, C employees were transported 5 (unintelligible) without monitoring devices, D 6 monitoring devices were worn under layers of protective clothing and not on areas of the 7 8 body being exposed, and we question that 9 (unintelligible) monitoring (unintelligible). 10 Somebody's dog (unintelligible) --11 DR. ZIEMER: I gather that's not your dog. 12 someone on the line has a dog barking, please 13 mute your phone or mute your dog, whichever 14 works better. 15 MS. HOYT: Okay, (unintelligible) statement 16 (unintelligible) affidavit (unintelligible) 17 petition and then (unintelligible) out of the 18 ER it says potential unreported neutron dose 19 (unintelligible) distribution (unintelligible) 20 August 27th, 1997 (unintelligible). We did not 21 submit that letter. The letter 22 (unintelligible) stated was in a file 23 (unintelligible) former worker who assisted us 24 and submitted an affidavit for the petition. 25 (Unintelligible) only record in our response

1	(unintelligible). We also claim there is a
2	conflict of interest (unintelligible). This is
3	(unintelligible) the Board's (unintelligible)
4	in the past. We would appreciate
5	(unintelligible) being fully addressed.
6	(Unintelligible) quote (unintelligible)
7	submitted by SEC-00057 petition regarding
8	(unintelligible) records, location of records,
9	(unintelligible) condition of individual are
10	you there?
11	DR. BRANCHE: Yes.
12	DR. ZIEMER: Yes, go ahead.
13	MS. HOYT: Hello?
14	DR. BRANCHE: Yes, we're here.
15	DR. ZIEMER: We're still here.
16	MS. HOYT: Okay. It it sounded
17	(unintelligible). But (unintelligible) workers
18	incidents and exposures. The affidavit also
19	covered falsification of records, coercion to
20	falsify records, loss or destruction of
21	bioassay records and lack of cooperation from
22	the FO from the FOIA office, DOE
23	(unintelligible). We would appreciate 4.7
24	being corrected to accurately reflect the
25	facts.

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On 7.1, pedigree of the Hanford data, the Hanford pedigree has little credibility or reliability. (Unintelligible) is not accurate, (unintelligible) does not (unintelligible) favorable and the TBD is not accurate and is incomplete. Out of the -- a quote out of 7.1, current and past Hanford workers have access to their records at any time upon request, end This is not true. The FOIA process is burdensome, unfriendly, inaccurate, not (unintelligible). The FOIA officers are uncooperative. Personal exposure records are not accurate. Worker outreach meeting speakers that were rad techs confirmed that their own exposure records were not accurate. (unintelligible) issues in, quote, official use only, quote, issues limiting access to records. I think that this is (unintelligible). 7.4, evaluation of (unintelligible) for SEC-(Unintelligible) accurately states 00057-2. the fact (unintelligible), (unintelligible) were not monitored or consistently monitored, falsification of records, coercion to falsify records (unintelligible) accurate and inaccurate bioassay records did not exist

1 (unintelligible) employees. 2 On behalf of my sister and all the former and 3 current workers of Hanford, we would appreciate 4 your resolution of these (unintelligible). 5 Thank you very much. 6 DR. ZIEMER: Thank you, Mary Ann (sic). 7 like to ask a question for clarification on the -- of the manuscripts that you were having 8 9 difficulty -- I think you mentioned the June 10 minutes, was that correct, or was it the 11 transcripts? 12 MS. HOYT: The June minutes. They're all posted. According --13 DR. BRANCHE: 14 DR. ZIEMER: Are they the transcripts or the minutes? 15 16 DR. BRANCHE: Everything we have -- Ms. Hoyt, 17 this is Christine Branche. According to my 18 records, everything that we have for all 19 meetings in June of 2007 have been posted. 20 UNIDENTIFIED: (Off microphone) 21 (Unintelligible) worker outreach --22 MS. HOYT: Well, I (unintelligible) to --23 DR. BRANCHE: Now if you're talking about a 24 worker outreach meeting, that's different. Ιs 25 that what you're talking about, a worker

1 outreach meeting? 2 MS. HOYT: Okay. Now that's -- that -- that's 3 something that the NIOSH staff handles and not 4 the Board. 5 MS. HOYT: Yes, and I received a reply back and 6 it said that it was not on their web site at 7 this time. It's from --8 DR. ZIEMER: We don't handle those. 9 MS. HOYT: -- update Ms. Hoyt, at this time I 10 have not been provided with final 11 (unintelligible) the meeting you are referring 12 to in the e-mail below. Please note that if 13 the meeting was a worker outreach meeting, 14 information about the meeting is not posted 15 until the final minutes are approved and 16 available for public distribution. 17 DR. BRANCHE: This has nothing to do with --18 MS. HOYT: Which brings up another question, 19 and that is why are the worker outreach 20 meetings being redacted now when they were not 21 redacted in the past? Why is this process so 22 burdensome that from June to now they cannot be 23 posted? 24 DR. ZIEMER: Okay, we're going to try to track that down. The Board is not involved with the 25

1 worker outreach minutes, but we're going to try 2 to find out -- also on the redaction here --3 Larry Elliott has a comment here that was --I don't -- I don't know which 4 MR. ELLIOTT: 5 worker outreach effort we're speaking about and I don't know who sponsored it. If it was -- if 6 7 it was a meeting in --8 DR. ZIEMER: Might it have been Labor? 9 MR. ELLIOTT: We did sponsor it? Okay. Well, 10 if we sponsored it, then there should be a set 11 of minutes that are being created for that -that meeting. The minutes of these worker 12 13 outreach meetings that NIOSH has sponsored and 14 held, whether it be a town hall type meeting, a 15 -- a focused panel group meeting or individual 16 interviews, those things have always gone 17 through Privacy Act review before we post them, 18 before we share them, so... 19 DR. ZIEMER: And keep in mind that those are 20 not -- those aren't part of the same group 21 covered by the Board's policy on that 22 redaction, probably, or -- is there -- is the 23 redaction policy different than --24 MR. ELLIOTT: The Board's activities are 25 covered under FACA.

1 DR. ZIEMER: Right. 2 MR. ELLIOTT: The program's activities are 3 covered under the Privacy Act in FOIA. Which is different. 4 DR. ZIEMER: 5 MR. ELLIOTT: Yes. 6 DR. ZIEMER: Yeah, so the redaction policy of 7 the agency on those -- on those matters is 8 different than the Board's, which is under the 9 Federal Advisory Act issue -- or laws, so there 10 is a difference in the redaction there. 11 MS. HOYT: So the worker outreach meetings were 12 not redacted. (Unintelligible) the site will 13 be that former worker outreach meetings 14 (unintelligible) meetings (unintelligible) Mr. 15 (unintelligible) who was a worker who has (unintelligible) and I would like to have the 16 17 redaction policy for worker outreach meetings 18 (unintelligible) because they are 19 (unintelligible). 20 MR. ELLIOTT: Well, these are -- these are 21 meetings that, when minutes are captured -- as 22 they have always been -- will have to go 23 through Privacy Act review before they are 24 released. Yes, there will be names in these

minutes. Names of government employees are not

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1 Names of individuals who worked at a 2 site and are representing themselves as having 3 worked at that site may not be redacted, or may 4 be redacted, given the particular context of 5 how they -- of what they had to say. 6 may expect to see minutes continue to be 7 redacted before they are shared and publicly 8 distributed. You may expect to see holes in 9 those where certain people's names or personal 10 identifiable information that is sensitive has 11 been struck out. I am sorry for that, but that is the way we have to live. 12 13 DR. ZIEMER: Right, and that is different from 14 the Board meetings and minutes. 15 Okay, Board members, any other --16 MS. HOYT: (Unintelligible) Elliott and I 17 should discuss this further in a different --18 DR. ZIEMER: Sure, sure, right. 19 MS. HOYT: -- (unintelligible) if I could have 20 Mr. Elliott's phone number (unintelligible) e-21 mail to me I would appreciate it. 22 Yeah, he -- he has your number and DR. ZIEMER: 23 will call you then, Mary Ann. 24 MS. HOYT: This is Rosemary. 25 DR. ZIEMER: Oh, Rosemary, okay. I'm sorry.

1 MS. HOYT: Thank you. 2 DR. ZIEMER: Okay, any other comments from 3 either of the petitioners? MS. CARRICO: This is Mary Ann Carrico. 5 like to comment. 6 DR. ZIEMER: Okav. 7 MS. CARRICO: I would like to say that we were 8 relieved that NIOSH and SC&A came to agreement 9 on (unintelligible) areas rather than specific 10 buildings within the areas of Hanford. 11 Also on the white paper prepared by SC&A 12 (unintelligible) issue for the proposed Hanford petition to the special cohort, there's a 13 14 specified roving workers. This includes construction workers, instrument technicians 15 16 and maintenance workers. These people would 17 generally perform work in various parts of the 18 They'd be required to go into a variety 19 of buildings. There were several 20 (unintelligible) workers who were not mentioned 21 in the white paper. The security emergency 22 response people, the transportation 23 (unintelligible) to name a few; there may be 24 others. I (unintelligible) in the roving

workers. We question how NIOSH

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1 (unintelligible) worker. 2 Also, my other question is, are the other 3 findings on the matrix (unintelligible) for 4 exclusion in the SEC. Thank you. 5 DR. ZIEMER: Okay, we will ask that those 6 questions be looked at. I don't know if we 7 have the answers to those at the moment. 8 MR. ELLIOTT: It's all workers. 9 DR. ZIEMER: It's all workers, so it's not a --10 any ro-- any of the roving workers. 11 The definition is who worked in DR. MELIUS: 12 those areas --13 DR. ZIEMER: Yeah, so --14 DR. MELIUS: -- so it's not -- you know. 15 DR. ZIEMER: The naming of some of them does 16 not -- is not limiting; it's an example, more 17 or --18 DR. MELIUS: Right. 19 DR. ZIEMER: Yeah, okay. 20 DR. MELIUS: Can I just comment on --21 DR. ZIEMER: Dr. Melius has --22 DR. MELIUS: -- the last question? 23 DR. ZIEMER: -- a comment for you. 24 DR. MELIUS: Yeah, just to say that -- I think 25 as we all know, we're early in the review of

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this SEC evaluation report, and so the many other issues that are in the matrix, most of them are still open and we're trying to figure out a -- sort of a way forward, what's the most efficient way forward. We have been stymied by the fact that both NIOSH and SCA have very limited, if any, access to records from the site at the moment. It's been over six months now, have not had access, and this is causing a I think very significant delay in the work that -- that both NIOSH needs to do to complete some of the work to both respond to SC&A's comments, as well as some other work they've already planned in their evaluation report, and certainly makes it essentially impossible for SC&A to review any of the -- the work in the NIOSH -- any more of the work in the NIOSH evaluation report. We've talked -- as part of the working group, as well as some techni-technical call to try to, you know, make -develop a way that -- that we can go forward on some of these issues, but it's -- it's very limited until DOE resolves the issue of access to records at this site. As I said, it's been over six months and I -- I don't know, Larry,

1 if you have any further information. All we 2 heard today from DOE was that there may be an 3 update or some plan or something, but I don't know -- I haven't -- yet to hear a schedule for 4 5 access. 6 I don't -- I don't know if -- Sam MR. ELLIOTT: 7 has been working as the NIOSH point of contact 8 to coordinate with SC&A on what information 9 needs we have and prioritize those so we can 10 put them in front of DOE. I think that has 11 happened. Right, Sam? 12 DR. GLOVER: That is correct. 13 MR. ELLIOTT: And -- and DOE is telling me 14 today that -- that they are going to respond to 15 that prioritized set of requests, so I take 16 that to mean -- and I asked specifically, does 17 that mean the logjam is broken and work can 18 start? Yes, that's what I hear. So we should 19 start knocking on the door and seeing, you 20 know, how far we get with our requests. DR. MELIUS: Okay. 21 22 DR. ZIEMER: Thank you. 23 DR. MELIUS: Thank you. 24 DR. ZIEMER: Any further questions or comments? 25 I was told there might be some Hanford folks

1	here with us in person. Are there any here?
2	(No responses)
3	Apparently not.
4	UNIDENTIFIED: (Unintelligible) Department of
5	Energy en route to Hanford. I'm on the phone.
6	DR. ZIEMER: Okay.
7	MR. ELLIOTT: One of the things that I
8	understood DOE was going to do first is give
9	you guys some instruction on how to do do
10	your own searching and sorting, but you know, I
11	don't know, so we'll see.
12	DR. GLOVER: Do you want any details? Okay.
13	DR. ZIEMER: There there was someone else on
14	the phone, a Hanford person, we didn't catch
15	your name.
16	UNIDENTIFIED: This is Gail (unintelligible),
17	I've been working with Sam on giving them on-
18	line access to some of our finding aids to
19	DR. ZIEMER: Oh, okay. Did you have any
20	comments on that for us? Any any additional
21	comments on this discussion?
22	(No responses)
23	Apparently not. Okay, thank you.
24	DR. GLOVER: It's we are working with DOE
25	we have developed a formal strategy of keyword

searches that will help support the matrix, a closure of these items. Both SC&A and NIOSH have put together a common set of search terms to reduce the duplication of this effort, so we're going to share those resources. We've put that forward to the DOE so they can prioritize and understand a better -- have a better understanding of how much resources they need to put forward. And so pending a meeting I think in the next week which we will begin -- able to have an understanding of what the schedule will be so we can gain access to those records.

DR. ZIEMER: Greg has --

MR. LEWIS: This is Greg Lewis from DOE. I just want to back up what -- what Sam said. You know, we are committed to getting them in there to start looking at the records as soon as possible. We're working with Gail, working with Sam. We believe we have a plan that should be successful. There are a couple of final issues that we'll be meeting on later this week, and we should be able to dive right in as --

DR. ZIEMER: Thank you.

1 MR. LEWIS: -- soon after that as possible. 2 DR. ZIEMER: Further questions? 3 DR. MELIUS: No, I've... 4 DR. ZIEMER: Is there anything further we need 5 to do on Hanford at this time then? 6 DR. MELIUS: Yeah, I need to make a motion. 7 DR. ZIEMER: Yes, go ahead. 8 DR. MELIUS: I'll offer a motion, it's a 9 lengthy motion -- get Ray ready. 10 Board recommends that the following letter be 11 transmitted to the Secretary of Health and Human Services within 21 days. 12 Should the Chair become aware of any issue that in his 13 14 judgment would preclude the transmittal of this 15 letter within that time period, the Board 16 requests that he promptly informs the Board of 17 the delay and the reasons for this delay, that 18 he immediately works with NIOSH to schedule an 19 emergency meeting of the Board to discuss this 20 issue. 21 The Advisory Board on Radiation and Worker 22 Health, the Board, has evaluated SEC Petition 23 00057-2 concerning workers at the Hanford Nuclear Reservation in Richland, Washington 24 25 under the statutory requirements established by

1 EEOICPA, incorporated into 42 CFR Section 2 83.13. The Board respectfully recommends 3 Special Exposure Cohort status be accorded to all employees of the Department of Energy, its 5 predecessor agencies and DOE contractors or 6 subcontractors who worked from, number one, 7 September 1st, 1946 through December 31st, 1961 8 in the 300 area; or two, January 1st, 1949 9 through December 31st, 1968 in the 200 area at 10 the Hanford Nuclear Reservation for a number of 11 work days aggregating at least 250 work days 12 occurring either solely under this employment 13 or in combination with work days within the 14 parameters established for one or more other 15 classes of employees in the SEC. 16 The Board notes that although NIOSH found that 17 they were unable to completely reconstruct 18 radiation doses for these employees, they 19 believe they may be able to reconstruct 20 external doses, and internal doses (other than 21 americium and thorium). 22 This recommendation is based on the following 23 factors: Hanford Reservation facility was 24 involved in development, manufacture of nuclear

weapons; two, NIOSH found there was

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insufficient monitoring data, information on 1 2 radiological operations at these laboratories 3 in order to be able to complete accurate individual dose reconstructions involving 5 internal exposures to thorium in the 300 area 6 and americium in the 200 area of the facility 7 during the time periods in question. The Board 8 concurs with this conclusion. 9 NIOSH determined that health may have been 10 endangered for the workers exposed to radiation 11 in the 200 and 300 areas of the Hanford Nuclear 12 Reservation during the time periods in 13 question. The Board concurs with this 14 determination. 15 Enclosed is rec-- supporting documentation from 16 the recent Advisory Board meeting held in 17 Tampa, Florida where this Special Exposure 18 Cohort class was discussed. If any of these 19 items are unavailable at this time, they will 20 follow shortly. 21 DR. ZIEMER: You heard the motion. Is there a 22 second? 23 MR. CLAWSON: Second. 24 DR. ZIEMER: Got a couple of seconds here. 25 discussion?

1 I have a question. This is kind of a devil's 2 advocate type of question. In light of the documents that we don't have from Hanford, what 3 level of confidence does NIOSH have that the 4 5 issues that lead you to -- to recommend this as a class of the SEC would not be resolved in the 6 7 materials that may be forthcoming? Probably a 8 question you can't answer, but it seems to me 9 we have to ask it anyway. 10 DR. GLOVER: The driving point is that there 11 were no bioassay during that early phase for 12 those nuclides. DR. ZIEMER: So -- and we know that for certain 13 14 15 DR. GLOVER: Absolutely. 16 DR. ZIEMER: -- it's not simply that we haven't 17 seen the records. 18 MR. GRIFFON: Through interactions with many 19 different sites and different levels, there is 20 absolutely no bioassay at that facility. 21 DR. ZIEMER: That's what I wanted to hear. 22 DR. MELIUS: Yeah, and I think the issue was 23 the -- the sort of class definition, how do you 24 -- and given, I think -- or you -- what 25 information we did have on operations, the

1	facility and so forth, that that going to
2	the the area definition as opposed to
3	building definition was was probably much
4	more appropriate as a way
5	DR. ZIEMER: Yeah, that that part's all
6	right.
7	DR. MELIUS: yeah, yeah yeah, regardless,
8	yeah.
9	DR. ZIEMER: Just to make sure that
10	DR. MELIUS: Yeah.
11	DR. ZIEMER: there's no doubt that that
12	there were no bioassay.
13	DR. MELIUS: Yeah, there are a number of other
14	issues at the site that that that, as I
15	said, we are stymied until we have access to
16	records and NIOSH I mean just to be able to
17	begin discussions on on some of these
18	issues, and that's why so adamant about the
19	records access issue.
20	DR. ZIEMER: Other comments or discussion on
21	this?
22	DR. POSTON: I have ques
23	DR. ZIEMER: Dr. Poston?
24	DR. POSTON: I have a clarification Jim, you
25	named some specific areas

1 DR. MELIUS: Uh-huh. 2 DR. POSTON: -- in your motion, and I didn't --3 maybe I missed it. I didn't think I heard all 4 the areas included, which was what LaVon -- I 5 thought LaVon was talking about. I mean -- not 6 -- I'm sorry, I've -- did you name all the 7 areas, is that -- just for clarification? 8 DR. MELIUS: It's the same -- they match the 9 definition. 10 DR. POSTON: Since I don't have it to read, I 11 have to ask the question. 12 DR. MELIUS: Yeah, yeah, no, I -- take another look at it tomorrow 'cause it's a little tricky 13 14 to write, given the -- two separate areas, but 15 16 DR. POSTON: So you did -- not --17 DR. MELIUS: Yeah. 18 DR. POSTON: So it's all the workers in all the 19 areas. 20 DR. MELIUS: Yeah. 21 DR. ZIEMER: It's all facilities and areas. DR. MELIUS: Yeah. 22 23 DR. BRANCHE: They're -- Dr. Poston --24 DR. GLOVER: No. 25 DR. MELIUS: No. So --

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1
              DR. GLOVER: It is the 300 area and the 200
2
               area.
3
              DR. MELIUS: Yeah, it's both those areas, yes.
4
              DR. GLOVER: The 100 area -- for -- just for
5
               review, the 100 areas are primarily the reactor
               facilities.
6
              DR. MELIUS: Yeah, yeah.
7
8
              DR. POSTON: Okay.
9
              MS. BEACH: I have a question.
10
              DR. BRANCHE: You need to come to the
11
              microphone.
12
              DR. ZIEMER: Josie's question is --
13
              MS. BEACH: The 200 area is --
14
              DR. ZIEMER: -- as a site expert.
              MS. BEACH: -- actually plural.
15
                                                There's an
16
              east and a west. Does it cover both east and
17
               west?
18
              DR. GLOVER: It -- it -- yes.
19
              DR. MELIUS: Yes.
20
              DR. ZIEMER: Are you ready to vote then?
                                                          Ιt
21
              appears we're ready to vote.
22
              DR. BRANCHE: Mr. Presley, you're on by phone,
23
               so may I get your vote first?
24
              MR. PRESLEY:
                             Yes.
25
              DR. BRANCHE: Yes, I can get your vote, or yes
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1
               is your -- is your decision? Mr. Presley?
2
              MR. PRESLEY:
                             Yes, yes.
3
              DR. BRANCHE: Mr. Clawson?
4
              MR. CLAWSON: Yes.
5
              DR. BRANCHE: Mr. Gibson?
6
              MR. GIBSON: Gibson.
7
              DR. BRANCHE: Mr. Griffon?
8
              MR. GRIFFON: Yes.
9
              DR. BRANCHE: Dr. Melius?
10
              DR. MELIUS: Yes.
11
              DR. BRANCHE: Dr. Poston?
12
              DR. POSTON: Yes.
              DR. BRANCHE: Dr. Roessler?
13
14
              DR. ROESSLER: Yes.
              DR. BRANCHE: Mr. Schofield?
15
16
              MR. SCHOFIELD: Yes.
17
              DR. BRANCHE: Dr. Ziemer?
18
              DR. ZIEMER: Yes.
19
              DR. BRANCHE: And I'll get Dr. Lockey's vote.
20
              DR. ZIEMER: And the record will show that Ms.
21
              Beach is not voting on this.
22
              DR. BRANCHE: Nor is Ms. Munn.
23
              DR. ZIEMER: Nor Ms. Munn.
24
              DR. POSTON: I apologize to Sam for calling him
25
              by the wrong name.
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DR. ZIEMER: I declare that the motion has carried and we will transmit the appropriate recommendation to the Secretary for action.

DR. MELIUS: And can -- can I make one final comment? I would just like to thank Sam and the people -- other staff at NIOSH for sort of -- we're trying to do this -- that -- move this forward sort of incrementally and -- been very good to work with and I think we've -- we've got a process in place that, once we get access to the information, I think will allow us to go through -- it's a large facility with a lot of complicated issues -- I think pretty rapidly and I think between S-- SC&A and NIOSH and the workgroup, I think we've -- able to make good progress.

SEC PETITION STATUS UPDATES:

SANDIA NATIONAL LABORATORY-LIVERMORE

DR. ZIEMER: Thank you. Next we have Sandia National Lab Livermore. It says that -- that Paul Ziemer is making the presentation.

Actually that's not the case. I'm simply declaring that that's where we are on the agenda. The --

DR. BRANCHE: May I -- may I, Dr. Ziemer?

DR. ZIEMER: Yes.

DR. BRANCHE: For those of you participating by -- by phone, I know I sound a bit like a broken record. I would like you to please mute your phone, and if you do not have a mute button, please dial star-6. It's important not only that our court reporter be able to record information correctly when he prepares the transcript, but also understand the background noise that's created when your line is open makes it difficult for other participants by phone to get all of the information that's going on in our Board -- in our room here. So I do ask -- I encourage strongly that you mute your phones. Thank you so much.

DR. ZIEMER: Thank you. The Sandia National Laboratory material was actually presented to us at our last meeting by Sam Glover. Sam, perhaps you would take a moment and just remind the Board of -- of the recommendation and where we were on that. We -- we had a -- an evaluation report and recommendation from NIOSH on Sandia, and we need to see the bottom line, and I believe in -- in LaVon's review he reminded us on -- on that particular one that -

1 - that was tabled so the Board could review the 2 -- I believe it was the addendum. 3 DR. GLOVER: I did not bring the presentation 4 down here yet. If you want the -- the 5 (unintelligible) --DR. ZIEMER: I don't think we need the whole 6 7 presentation, just... 8 DR. BRANCHE: LaVon is bringing up his stuff, 9 too -- there it is. 10 DR. ZIEMER: There it is. Just go to the 11 bottom line on that one. 12 DR. GLOVER: Let's see -- so let me make sure 13 we have the -- we actually had several -- we 14 had the first -- the evaluation was approved, 15 sent to the Board in March of 2007, as -- as 16 LaVon mentioned that -- right before the Board 17 meeting we received additional information and 18 di -- after -- following the presentation on 19 March -- actually the beginning of April at the 20 Board meeting the Board asked us to update the 21 evaluation report. We did so, and there was a 22 change because what -- the real change -- the 23 material that came was that there was potential 24 for direct beam interactions, and so that was 25 the major change in the evaluation report that

we reissued and presented in October 2007.

At that point in time the Board tabled the vote. The petitioner provided -- it's weird to look at two different screens -- provided some additional information regarding I believe a number of items that were to be reviewed by the Board to see if they would make an impact on their evaluation of our -- of the evaluation report. We certainly could -- I -- I didn't bring that down, Dr. Ziemer. I'd be happy to put that forward. I could -- that presentation's previously in my room, depending on what level of detail you would like to see it.

DR. ZIEMER: The -- the additional materials for the -- from the petitioner, as I recall, were distributed to the Board members to look at after that meeting. My understanding is that NIOSH found no reason to change their recommendation on the basis of those materials. Board -- Board members, I ask you now if any of you wish -- or are ready to make a recommendation on this particular petition?

(No responses)

If you are not ready to do that, I'm going to

1 ask you again tomorrow if you are ready to do 2 that. I -- I don't believe there's any point 3 in continuing this for any longer. We've had the material in our hands for a fair amount of 4 5 time. If you need to review it tonight, you can do that, but otherwise we need to take 6 7 action. 8 Give us your bottom line recommendation. The 9 recommendation --10 DR. GLOVER: Yeah, the recommendation was that 11 we could do --12 DR. ZIEMER: -- was that you could reconstruct dose --13 14 DR. GLOVER: Yes, sir. 15 DR. ZIEMER: -- and I believe it was only an 16 external dose issue, it was an X-ray device, as 17 I recall. So that if the Board accepts --18 accepts that recommendation, then there is 19 nothing that would go forward to the Secretary 20 since we would not be recommending a class. 21 MR. ELLIOTT: I might add, for the Board's 22 understanding, if you recall, there are a very 23 small number of individuals involved in this 24 class, a total of three, one of which we have a 25 claim for. And upon revisiting that dose

1	reconstruction, I believe that claim is now at
2	a compensable state.
3	DR. ZIEMER: Okay. Is there anyone that wishes
4	to make a recommendation or make a motion?
5	MS. MUNN: Well
6	DR. MELIUS: We'll do it tomorrow.
7	UNIDENTIFIED: Can we pick it up
8	DR. ZIEMER: We can pick it up tomorrow
9	well, I I've already told you that that's
10	what's going to happen if I have no if no
11	one is moving today, they
12	DR. MELIUS: If we all leave the
13	DR. ZIEMER: If you all I think
14	DR. MELIUS: If we all leave the room
15	DR. ZIEMER: You become motionless, I can tell
16	that. Okay.
17	DR. GLOVER: Dr. Ziemer, I would be happy to
18	quickly review the main points of that tomorrow
19	morn before your
20	DR. ZIEMER: I'll ask you to do that. That may
21	help the Board recollect this particular one.
22	MR. CLAWSON: I I just wanted to see the
23	bottom line. I couldn't
24	DR. ZIEMER: Well, and and keep in mind
25	that, in essence, this petitioner as Larry

1	has indicated this will not affect this
	has indicated this will not affect this
2	petitioner any longer, in any event, either
3	way. It's kind of a moot point issue.
4	Although there are two there are two
5	possible other petitioners.
6	MR. ELLIOTT: Don't take my statement as to say
7	that this won't affect the other two
8	DR. ZIEMER: No, no, I
9	MR. ELLIOTT: because if one of those other
10	two are both
11	DR. ZIEMER: said this particular
12	petitioner, but there are two potential other
13	ones.
14	MR. RUTHERFORD: There one I'm sorry.
15	There are two two people that are
16	potentially in the class. However, they are
17	not claimants
18	DR. ZIEMER: Right.
19	MR. RUTHERFORD: at this time. I'm just
20	making sure
21	DR. ZIEMER: Right, they are not claimants.
22	[name redacted], are you on the line? I I
23	didn't give his last name, did I?
24	DR. BRANCHE: No, I'm trying to keep you from
25	saying it.

1 DR. ZIEMER: Is there anyone on the line named 2 [name redacted]? Are there any petitioners... 3 In the future, don't give me a list with people's names on it -- give me a redacted 4 5 list. 6 Apparently there's no one on the line from 7 Lawrence (sic) Livermore. Thank you. 8 becoming motionless myself. 9 SEC PETITION STATUS UPDATES: 10 CHAPMAN VALVE 11 We're going to move on to Chapman Valve, for 12 which we have petitions from unknown people. 13 DR. BRANCHE: And we have people in the room. 14 DR. ZIEMER: We have some people in the room, 15 but I'm not even going to tell you who they 16 are. 17 DR. BRANCHE: I think that might be smart. No 18 one fr-- Dr. Ziemer, no one from Chapman 19 Valve's name can be mentioned because it's --20 DR. ZIEMER: Unless they wish to mention it. 21 DR. BRANCHE: -- unless they wish to mention 22 their own names if they are on the phone. 23 DR. ZIEMER: Are there any folks from Chapman 24 Valve on the phone or any --

Yes.

UNIDENTIFIED:

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1 DR. ZIEMER: -- Congressional people on the --2 from -- representing Chapman? 3 MR. CLAWSON: Yes. 4 MS. BLOCK: Yeah, this is Sharon Block from 5 Senator Kennedy's office. DR. ZIEMER: 6 Okay. 7 MS. REALE: Maryanne Reale, petitioner. 8 DR. ZIEMER: Very good. We will have a report 9 from the working group chair, Dr. Poston, on 10 Chapman Valve, and then we will hear from those 11 on the phone who wish to make comment. 12 DR. POSTON: Well, Mr. Chairman, I don't have 13 much of a report. As you know, there's been 14 some -- addition of the Dean Street facility. There's been a reconsideration of that added to 15 16 the group, and I guess the -- we've had a 17 revision that was -- I'm -- can't find it here 18 -- SEC petition evaluation report was reissued 19 February the 5th of this year. And as far as I 20 can see, it really hasn't changed the -- the 21 conclusions that -- that NIOSH can do the dose 22 reconstruction. And Jim Neton may want to have 23 -- say something. 24 DR. ZIEMER: And the recommendation from the 25 workgroup is?

DR. POSTON: That -- that the petition is still
as -- as it was, that the petition be denied.

DR. ZIEMER: Okay. Does NIOSH have any
additional comments on this facility? Jim
Neton.

DR. NETON: Okay. Thank you, Dr. Ziemer. I'd just like to have a -- I have a few slides so I'll be somewhat brief. But I'd just like to address the changes that were made to the evaluation report that Dr. Poston alluded to that are based on the Department of Energy's research into additional activities that may have occurred at Chapman Valve.

If you recall, NIOSH requested the Department of Energy to review the information surrounding Chapman Valve and to look at that definition to see if there were any additional work activities or sources of radiation-related work activities that occurred at Chapman Valve. And this was in direct response to some statements made in a site expert interview that SC&A captured during their worker outreach activities that indicated that there may have been activities off-site -- that is, at Dean Street. In fact, there was some speculation

that maybe those activities could have involved additional exposure to radioactive materials.

We sent -- we requested this from Department of Energy, and they provided a response to us in a letter report dated January 7th, 2008. And their conclusion in that report was the Dean Street facility is indeed -- should be covered under the Atomic -- AWE facility, it is now part of the covered facility definition. But they did state in their report that they found no indication of additional radiological activities that occurred at Dean Street after their somewhat I think apparently detailed review of the records.

I'd just like to go a little bit into that review. We did look at the additional information that was provided by Department of Energy. They sent over -- and we have put all this information on the -- on the O drive for the Board to review as well -- about 30 letter documents documenting the manufacture of valves and manifolds for the Y-12 electromagnetic enrichment facility that was being constructed in Oak Ridge. No doubt that Y-12 -- or Chapman Valve manufactured these manifolds for the --

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what's called the racetracks down at -- in Oak Ridge. In addition to these letters there were about 40 engineering drawings of various valves and manifolds, sort of documenting what their configurations were.

In those -- in that information, though, there was no indication of shipment of valves for repair. And if you recall, that was one of the -- the statements made by the subject expert from Chapman Valve, that she had a very de-very real recollection of -- of things being shipped up from Oak Ridge to Chapman Valve, being staged at the main facility and then being transferred over to the Dean Street facility. However, in looking through these records, I was caught by the similarity of some of the test specification documents for valves and manifolds that were included in these 30 essentially letter reports. And that is, as -what I would call test specification for valves and manifolds, and I worked for equipment manufacturers and we would call these factory acceptance tests. That is, when you make a new product, oftentimes in the purchase specifications you'll require that -- you know,

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the purchaser will visit the site, hook up the equipment and test it to make sure it meets the specs. And in fact there were about three or four letters that -- that spoke to this, that each pump and manifold should be tested before shipment down to Oak Ridge, and that is in the presence of the purchaser's representative -if you recall, the expert talked about people coming up to the site -- and using purchaser's test equipment. And this is not little equipment. They speak of these 440 volt valves and pumps and test gauges and all these sort of things. Matter of fact, one of the letters spoke about -- from Chapman Valve telling Stone and Webster they actually needed six complete sets of these things to do their tests around the clock. So there is no doubt there was a lot of equipment shipped -- I'm not sure from where, but to Chapman Valve. Sometimes the shipment was requested from Westinghouse, sometimes the letter originated in Boston -could have come from Oak Ridge; we don't know. At any rate, there were a number of tests conducted on these manifolds, presumably at the Dean Street facility. And also there were

1 specifications about how these valves must be 2 cleaned prior to shipment using these solvents, 3 and also they must be coated with some sort of 4 a drying agent, a desiccant, and that was also 5 in the recollection of the site expert. 6 there's -- there seemed to be a match here 7 between the recollections. 8 At any rate, after looking through this --9 looking at the DOE letter and also the 10 documents provided by the Department of Energy, 11 we saw nothing in their review that indicated 12 there were additional sources of radioactive materials present at Chapman Valve, 13 14 specifically nothing in the 1948 and '49 period 15 -- that is the right period, am I right? 16 have to look and see if it -- what's the 17 covered period for Chapman Valve? That was --18 '46 -- whatever the covered period was for 19 Chapman Valve, I think it's '48 and '49 --20 MR. CLAWSON: '48 through '49. 21 DR. NETON: '48 through -- I'm sorry. 22 nothing in there, in that period specifically. 23 All this information that we talked about with 24 these acceptance testing of the -- of the 25 manifolds and such occurred in the early '40s,

1 1943, 1944 time frame. Although we would say 2 the acceptance requirements for these new 3 products in the '43-'44 time frame are not 4 inconsistent with the site expert's 5 recollections. 6 So to that end, after looking at all this information, we revised the evaluation report 7 8 in February and distributed it to the Board and 9 -- and put it on our web site, and additional 10 text was added to pages 13 and 14 specifically 11 of the report to summarize DOE's findings that 12 the Dean Street facility is now part of the 13 covered facility, and that their conclusion was 14 that there was no -- and we support the conclusion that no additional sources of 15 16 radioactivity were identified. 17 So in essence, nothing changed in our 18 evaluation report other than adding Dean Street 19 to the covered facility, and you'll see that 20 here. The previous definition just listed 21 Building 23 at Chapman Valve. And then if you 22 look at the revised definition, I've 23 highlighted in yellow here "work at the Chapman 24 Valve Manufacturing Company" and now we say 25 "(i.e., Building 23 and the Dean Street

1 Facility) " -- that is the total sum change of 2 the report, other than a summary, like I say on 3 pages 13 and 14, of what the DOE identified. So we still maintain that it's feasible to do 4 5 dose reconstructions at Chapman Valve during the covered time period, and the summary is 6 7 feasibility is possible, yes, and therefore 8 health endangerment's not applicable. And 9 here's the covered periods -- January 1, 1948 10 through December 11th, 1949 -- in addition to 11 this residual period, if you recall, that the 12 petitioner has requested to be evaluated. 13 is January '91 through '93. We still do not 14 have information on the '94-'95 time frame. That's it. 15 16 DR. ZIEMER: Okay, thank you. Questions for 17 Jim? Yes, Jim Melius. 18 DR. MELIUS: Jim, did you talk to the site 19 expert yourself about these -- these letters 20 and --21 DR. NETON: No, I did not. 22 DR. MELIUS: -- go over that -- okay. 23 DR. NETON: No. 24 DR. MELIUS: So it's just based on what information --

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DR. NETON: I'm just suggesting that I looked at -- these factory acceptance tests were in there and they're very similar -- there's no information about contaminated manifolds being shipped up for repair, but there was a lot of information about factory acceptance tests being conducted on the newly-produced manifolds and a lot of test equipment being shipped to Chapman Valve. It was just an observation on my part and I just bring that up as an -- as just that, an observation.

DR. MELIUS: Well -- okay.

DR. ZIEMER: Other questions? Okay, I
understand that Sharon --

MR. CLAWSON: I've got one question.

DR. ZIEMER: Oh, hang on -- Clawson.

MR. CLAWSON: I -- I've still just got the question. You know, we pulled three samples, two of them were natural and we've still got one that's high enriched, so I -- I don't know how you can just cast that one sample off and say well, it didn't. There wasn't anything there -- or, you know, this -- this was kind of food for the fire, repairing manifolds or so forth like that and I -- and I just don't see

1 how we can take and --2 DR. NETON: Well --3 MR. CLAWSON: -- discard that. DR. NETON: I -- I -- well, we could --4 5 MR. CLAWSON: (Unintelligible) 6 DR. NETON: We could talk more about this --7 DR. POSTON: It's not highly enriched. 8 a misstatement. 9 MR. CLAWSON: What is it? 10 DR. POSTON: It was what, less than two percent 11 or about two percent? 12 DR. NETON: It was 2.16 percent enriched, I 13 believe. 14 DR. POSTON: That's not highly enriched. 15 UNIDENTIFIED: (Off microphone) that's not high 16 enriched. 17 MR. CLAWSON: Okay, but --18 DR. NETON: Not high enriched, but it was 19 slightly enriched by their calculation. It is 20 still unknown to us, even though SC&A has done 21 some reviews in their -- their report, whether 22 the sample truly was enriched uranium or not. 23 I think it's SC&A's opinion that they feel it 24 was. I'm not convinced that it was, based on -25 - there's a lot of unknowns of what happened

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here.

I could tell you a few more things that are also inconsistent here. I've spent a lot of time thinking about this. SC&A correctly indicated in their report that the -- the activity that was discovered at the west ramp, the loading ramp on the west end of Building 23 where the sample was found, was I think 120 picocuries per gram of uranium, and that's the one they cited was 2.16 percent enriched. was interesting to me, though, was that the gamma measurement there was 32 micro R per hour, which was well above -- that's three times basically above background. That, to me, is not consistent with 120 picocurie per gram material. That just doesn't make any sense to me.

In addition to that, if you look at the FUSRAP report and the cleanup activities, there's an indication that when Bechtel came in there in 1995, they actually still found some elevated contamination at that spot, but they actually found it to continue underneath the ramp and actually had to jackhammer out part of the concrete to dig under there to pull out the

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rest of the contamination. So I really have no idea as to what -- what that might have been and how it -- when it got there. But it's still, to me, sort of an unknown.

There is evidence that radium was used at the site. One of the worker outreach meetings someone talked about 100 to 200 radioactive radium sources that were used to X-ray materials. I know SC&A has commented that they believe that it couldn't have been radium because they're aware of the fact that the radium and U-235 share the same energy line, but I'm not convinced that that's necessarily the case 'cause you can't tell how they stripped out the contribution from the U-235 in that analysis.

So anyway, there are a lot of unknowns there, you're right. But again, in the contract -- in the SEC period, 1948-'49, we have a complete picture, with a closure report and a 100-page document or so that documents every natural uranium activity that was carried on at that location; no evidence of enriched uranium being processed that would have exposed the workers, in our opinion.

DR. ZIEMER: Dr. Poston?

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DR. POSTON: Mr. Chairman, I apologize for not being as complete as I should have, perhaps. If you read the revised report, there's basically no question about the external doses. Those -- the workers were monitored, lots of badges exchanged on a regular basis, processed by Rochester, so this -- there's no question about the external dose. That's never come up in -- as far as NIOSH is concerned, as far as I -- I can see, they feel that -- very strongly and very -- that they can reconstruct the external doses. The approach that they took in reconstructing the internal doses is based on bioassays, and the assumption that they took is what we call in health physics extremely conservative. They took the highest bioassay that they measured. They calculated an air concentration for the -- for that -- that would result in that bioassay. They assumed that every worker at the site was exposed at that air concentration every work day for about a year and a half, which is the period of time in which the Chapman Valve facility was operating. The point here is that if a -- if -- in these

1 dose calculations and the calculation of POC, 2 if a person doesn't reach 50 percent, they're 3 never going to reach 50 percent, and that's the whole crux of the matter. We understand the 4 5 external dose. We've made what in physics we 6 would call a bounding calculation to see what 7 it would -- would be the maximum, and if the 8 POC doesn't come anywhere close to 50 percent, 9 it will never be 50 percent. 10 DR. ZIEMER: Thank you. I think we have Sharon 11 Block on the line from Senator Kennedy's 12 office. I'm allowed to say both of those 13 names, I'm told. 14 MS. BLOCK: Yes, (unintelligible). 15 DR. ZIEMER: And Sharon, do you have a comment 16 from the Senator? 17 MS. BLOCK: Yes, I do. Can you hear me? 18 DR. ZIEMER: Yes, very well. 19 MS. BLOCK: I think you're aware the Senator 20 sent you a letter last week concerning the --21 this petition and (unintelligible) for dose 22 rate (unintelligible) significant questions 23 about what is known about the (unintelligible) 24 Chapman Valve. And (unintelligible) question 25 (unintelligible) the answer or (unintelligible)

1 determination needs to be (unintelligible) now 2 (unintelligible) this has gone on too long 3 without (unintelligible) any answers 4 (unintelligible) petition for the -- for the 5 Chapman Valve petitioners, that this is not --6 a process this lengthy was not what Congress 7 had in mind when they created this -- this 8 system and he would like to see 9 (unintelligible). 10 DR. ZIEMER: Okay. Thank you. Are there any 11 other folks on the line representing Chapman 12 Valve? 13 (No responses) 14 We have a couple folks here that -- do you wish 15 to speak? 16 Yes, I would. MR. PETERSON: 17 DR. ZIEMER: Yeah, please address the assembly. 18 MR. PETERSON: My name is Carl E. Peterson. 19 [redacted] is a petitioner in the Chapman Valve 20 This is really my first meeting and 21 [redacted] has been handling this, but my eyes 22 were widened today in terms of the whole process and I've -- I have talked to some 23 24 gentlemen and some people about what has 25 transpired about Chapman Valve and I have a

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couple of issues -- not necessarily on the technical issues at this particular point because I have not had a chance to read them all and I'll certainly get a copy of the report. But certainly the dialogue I've heard just then from Mr. Clawson and -- and the assumptions made -- my understanding, the bill was written to give the applicant the benefit of the doubt. That's my understanding when the bill was written in Congress, and that's what is supposed to transpire today. What I just heard was something about manifolds that we say well, you know, I worked and they did this with manifolds, they cleaned manifolds, I don't think they sent manifolds that were radiated, but I didn't hear conclusively that that happened. I didn't -- I didn't really hear back that something was shipped or someone thought it was shipped. That -- that's not That -- that is certainly not justification. enough evidence to say that it didn't happen. The next issue, and I'll be very short in this. I was -- I was going to say something else just about [redacted] and -- you know, she lost her dad. He was 37 years old. She has spent seven

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years because the Department called her and says look, we want to do something for people. And they said okay, we're going to open up your heart again, and for seven years she's been sitting by -- and what I heard today was people making assumptions and people basing those assumptions on -- you know, I understand there was a report by Ferguson. I understand -- they were the contractors making the product at Chapman Valve. Now to me, just in simplistic terms, if I'm a businessman and I have someone making something for me, I don't depend on them to give me the report whether it's right or wrong. I haven't heard any justification -- if that person who could have a liability in relationship to what they're doing at a facility is giving you the report that everything's just fine and dandy and we're cutting up concrete and finding other items, I think just from a basic layman's understanding there seems to be something wrong. There's not conclusive evidence here. We could talk about science, and science can work both ways. I'm an engineer. I understand that. But it seems there's certain items related to this

1 particular facility that we're making a broad 2 assumption at this particular point in time. 3 And I think we need to be more conclusive. And 4 if the benefit of the doubt goes to the 5 claimant, you can't just say that I think the 6 valves weren't contaminated. You -- I don't 7 think you have a right to say that. 8 That's what I have to say at this point. 9 you. 10 DR. ZIEMER: Thank you very much. Any other 11 comments? 12 Okay, Jim. 13 DR. NETON: I'd just like to -- I think I 14 pointed this out but I just want to be clear 15 that the shipment of the valves was in 1943 and 16 1944, which is outside the covered period for 17 Chapman Valve right now. So right now the 18 petition requested an evaluation for the 19 current covered period, which is 1949-1950. 20 think that's an important thing to keep in 21 mind. 22 The valves were -- were not DR. ZIEMER: 23 involved at that time. 24 DR. NETON: The valves were not involved and 25 it's not covered. I mean it may be indeed at

1 some point in time become covered --2 MR. GRIFFON: So you're saying it leaves it 3 open for --4 DR. NETON: Certainly it leaves -- the 5 possibility's open for that period to be opened 6 and valves to be discovered to have been 7 contaminated, if they indeed were. But right 8 now '49 and '50 -- or '48 and '49 is the time 9 period under evaluation. And the Department of 10 Energy, with the Department of Labor 11 collaboration, we've come -- come to that 12 conclusion. 13 MR. PETERSON: If I might? 14 DR. ZIEMER: Yes. 15 MR. PETERSON: Correct me if I'm wrong. 16 let's -- let's go on the assumption the valves 17 were contaminated. Does that mean they were 18 set on the site in '43 and '44, there's no 19 contamination and '45 there's no contamination? 20 I mean I understand there's half-life, full 21 life of these particular elements -- they just 22 don't go away. You have to be realistic here. 23 Because something's under -- not under your 24 domain, that doesn't mean to say that

particular item affects something in 1947. I -

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1	- I think that's should be pretty clear.
2	What's what's the life of the particular
3	item? Is it less than five years? I don't
4	think so.
5	DR. ZIEMER: Thank you. Dr. Melius?
6	DR. MELIUS: Yeah, I believe at our last
7	meeting we asked SC&A to review some issues.
8	I'd sort of like to have a chance to hear from
9	them.
10	DR. ZIEMER: Okay.
11	DR. MELIUS: If that's okay.
12	DR. ZIEMER: Who who is
13	DR. BRANCHE: There's Arjun.
14	DR. ZIEMER: reporting for SC&A?
15	DR. BRANCHE: There's Dr. Makhijani.
16	DR. ZIEMER: Okay. Okay, Arjun Makhijani.
17	DR. MAKHIJANI: Thank you, Dr. Ziemer. We had
18	five conclusions in our report, and I can just
19	go over them very quickly if you'd like.
20	Chapman Valve manufactured manifolds on a large
21	scale and tested them prior to shipment to Oak
22	Ridge. This would be during the Manhattan
23	Project, and we agreed with the DOE and NIOSH
24	finding there. So those were the first two
25	conclusions.

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We looked at the documents posted on the O drive and did not find evidence of returns of manifolds from Y-12 to Chapman Valve for repair and testing. And there was no available evidence in the reviewed documentation that manifolds were returned during the 1948-'49 period covered by the NIOSH evaluation report. That was the third conclusion.

We -- the fourth conclusion was about this M-31 sample which was on -- just on the inside of a building on the north side of the west ramp. In view of -- we examined the -- the measurement pretty closely and by -- by a number of different methods. We looked at the measurement techniques that were used at the time and concluded that it's reasonably certain that M-31 sample, that sample in question, was an enriched uranium sample. It would be very difficult to conclude otherwise because (unintelligible) a lot of things about the -about measuring samples and measuring uranium samples into question because the measurement methods were specified in considerable detail, and we had several people at SC&A look at this before arriving at this conclusion because we

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realized that obviously this would be a -- a significant matter for you to consider. We -specifically, we looked at the NIOSH work. We looked at the Oak Ridge '90-'92/9092* report, and we looked at the measurement protocols, which was a 1987 Oak Ridge document. Finally was the question of the -- ques-question of what might have been done at The -- the '90-'92/9092* Oak Chapman Valve. Ridge report describes a wider range of activities that was done at AWEs in the context we're referring to Chapman Valve. It doesn't actually say that -- that additional activities were done at Chapman Valve. I recognize that. And the -- the evidence that is available about the enriched uranium sample, the only evidence is from the site expert interview, who had said that material was returned for repair during the Manhattan Project. The -- the only thing about the enriched uranium sample that isn't consistent with that is that it's on the inside of the building rather than on the outside of the building, and the activities described were the manifolds were returned and then transferred from train to truck at the Chapman

Valve main facility and taken by truck to the Dean Street facility. And given the size of the manifolds, this -- this would have -- this would have obviously happened on the outside, although it is a surmise. The person -- the site expert wasn't actually present at the main facility, but -- but this -- I think it would be a very reasonable surmise. The -- let me read the rest of it, since this is a sensitive matter. Why don't I just read the fifth conclusion so -- so you can have it all on the record for you in case you haven't had a chance to look at it.

Oak Ridge '90-'92/9092* describes a wider range of storage and other historical activities than are described in NIOSH 2008, but as -- as I have said, this is generally for AWEs. It is possible that enriched uranium sample may have been associated with these other activities; however, there is no evidence of this in the reviewed documentation. The only piece of evidence as to the possible source of the enriched uranium is a site expert interview which described the returns of contaminated manifolds from the electromagnetic separation

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plant at Oak Ridge that was operated during the Manhattan Project and for a short period thereafter. While this does not prove that that was the source or that there was not another source, it is consistent with the available evidence, including the fact that the sample was very close to the entrance ramp and that it is the only sample that was enriched uranium. If manifold returns were -- were the source of the enriched uranium, it would have been deposited prior to the period covered by the evaluation report and the SEC petition. However, the fact that it was on the inside of the building creates some uncertainty since the site expert stated that the main Chapman Valve site was at -- was the location of transfers of the manifolds from train to truck, all of which would have taken place on the outside. The only thing -- the only last thing I'd like to add is that, as we understood the charge given to us by the Board, was to look at the new documentation posted on the O drive and the letters from the DOE, and we stuck to that charter and did not go beyond that, so -- so we did not look for additional documentation or do

1 any additional search. 2 DR. ZIEMER: Okay, thank you. Dr. Poston? 3 DR. POSTON: Arjun, I have two -- a couple of 4 questions. One, could you describe a little 5 bit further what you did in evaluating this one measurement to ascertain that you -- that you 6 7 felt it was correct? I mean you said you had 8 three -- three of your folks look at it very 9 carefully and I'd like to know exactly what you 10 did. 11 DR. MAKHIJANI: Well, Chuck Phillips is also 12 here. We -- well, it's described in detail in 13 the report you have, Dr. Poston. 14 DR. POSTON: Yeah, but I'd like to hear you describe it, please. 15 16 DR. MAKHIJANI: The -- the NIOSH report 17 describes the sample as being enriched uranium 18 of 2.16 percent enrichment, but that there was 19 no U-235 measurement and that there was no 20 uncertainty on the U-235. We found in looking 21 22 DR. POSTON: Wait, wait, I'm confused already. 23 You -- you said --24 DR. MAKHIJANI: I'm just going through the 25 process.

1 DR. POSTON: Yeah, but you said there was no U-2 235 present? 3 DR. MAKHIJANI: No, I'm just saying what --4 DR. POSTON: Did you misstate or... 5 DR. MAKHIJANI: No, I just -- I'm just saying 6 what the NIOSH report said. That was our 7 starting point, the NIOSH report. 8 I'm sorry, I thought you were... DR. POSTON: 9 DR. MAKHIJANI: We then went to the original 10 Oak Ridge document and found that there was 11 additional information about the sample. 12 Uranium-238 was given at 120 picocuries per 13 gram, as Jim has stated. An uncertainty bound 14 was described, which we have cited. Since the 15 enrichment was provided, you can estimate a U-235 concentration, which would be about 17 16 17 Radium was also described picocuries per gram. 18 in this particular sample, and that is a very 19 important point, as less than one picocurie per 20 gram, because then this allowed a comparison of 21 the 186 keV gamma emission from radium --22 almost 186 -- with a similar line from U-235. 23 And given the various concentrations and the 24 relative probability of emission of the 186 keV 25 line, you could conclude that almost none of it

1 was due to radium. And in fact, the radium 2 intensity would be less than one -- the -- the 3 uranium intensity would be more than 280 times 4 the radium intensity. 5 DR. POSTON: I quess I don't understand. 6 mean I would expect to find radium in normal 7 soil samples, whereas I wouldn't expect to find 8 uranium-235, so --9 DR. MAKHIJANI: That's right, that's why this 10 was an enriched uranium sample of artificial 11 provenance. That's -- that's the reason for 12 the conclusion. If you had found radium at --13 on --14 DR. POSTON: Well, how can you tell if you --15 if they have the same -- roughly the same 16 energies? 17 DR. MAKHIJANI: No --18 DR. POSTON: I mean how can you conclude one 19 way or the other? DR. MAKHIJANI: They have the same energy per 20 21 photon. 22 DR. POSTON: Yeah. 23 DR. MAKHIJANI: They're at least 280 photon 24 more from the U-235 than they are from the 25 radium, so you can tell that whatever you're

1 measuring would be --2 DR. POSTON: Wait, wait, wait, the specific 3 activity of radium is much higher than the specific activity of uranium-235. 4 5 DR. MAKHIJANI: Yes, but the number of -- num--6 number of emissions of 186 keV photons is only 7 3.2 percent, whereas it is 54 percent from U-8 235, and U-235 concentration was 17 picocuries 9 per gram, and it is not related to specific 10 activities because everything is in terms of 11 picocuries, so it's comparing radioactivity 12 with radioactivity and nothing to do with 13 weight. 14 DR. POSTON: But you're talking about activity. 15 Right? 16 DR. MAKHIJANI: Exactly. That's why it had 17 nothing to do with specific activity. 18 DR. POSTON: I guess I'm still not clear why 19 this -- you know, what -- what evaluation you 20 made that would make -- lead you to this 21 conclusion, other than --22 DR. MAKHIJANI: I explained it in detail. 23 -- the gamma -- the measurement protocol is 24 described in the Oak Ridge 1987 document. 25 Really, since the specific isotopes are

1 mentioned, there are only two measurement 2 protocols that are reasonable. The measurement 3 protocol that's actually described is -- is gamma spectro-- spectroscopy. That's why we 5 actually compared the intensity of the line --6 whether it was possible that you would confuse 7 what was measured between radium and uranium, 8 and we concluded that you could not confuse it 9 because a radium measurement was actually 10 provided as being less -- well, it was a upper 11 limit as being less than one picocurie per 12 gram. And when you look at the characteristics 13 of the emissions of photons from U-235 and 14 radium and their frequencies, and the described 15 concentrations, you come to the conclusion that 16 you could not mistake this for radium, that it 17 would have been associated with U-235. 18 you do the same with alpha spectroscopy, of 19 course, you would come to the same conclusion. 20 DR. POSTON: Did they use alpha spectroscopy --21 DR. MAKHIJANI: (Off microphone) 22 (Unintelligible) 23 DR. POSTON: -- or just gamma? 24 DR. MAKHIJANI: It's not described. 25 DR. POSTON: Okay, I was just asking why you

1 brought that up. 2 MR. PHILLIPS: The reason we did the 3 calculation from the radium is if you had that amount of radium in it, you would have 5 certainly seen that in the gamma analysis and 6 the lead and bismuth-214 (unintelligible), to 7 that extent. So you could -- you know, you --8 it wasn't reported as that. It was reported as 9 less than one. So if there was sufficient 10 amount of radium in there to have sufficient 11 interference in the uranium-235 (unintelligible) to account for this amount of 12 13 uranium-235 that would be required for this 14 amount of enrichment, you would certainly have detected that. And we -- was that 185 15 16 picocurie (unintelligible)? 17 DR. MAKHIJANI: Sir? 18 MR. PHILLIPS: How much radium (unintelligible) 19 have to have to --20 DR. MAKHIJANI: Well, you have to have 280 21 times more radium (unintelligible). I didn't 22 measure (unintelligible). 23 DR. NETON: Can I speak for a second? This is 24 getting pretty technical, but I think it's 25 important to discuss this. My -- my reading of

the protocol that was used to measure the sam-first of all, I don't think they ever said that
they used the 609 or the 239 keV

(unintelligible), that's not even covered.

What they do say is they recognize the fact
that uranium can -- higher levels of uranium
can interfere with measurements of radium-226.

That's what they say. Presumably when they say
that, that must be talking about the 185 keV
line.

Now what they did say is that if there were higher levels of interference in that 185 keV line, they would quote a detection limit above that peak. So in other words, the higher the U-235 peak that's there, the less ability they would have to measure radium-226 because it would essentially be there with an interfering background. So that's -- that's what they did, so I don't know -- I looked at SC&A's analysis and it makes no sense to me when they say they basically stripped out the 235 peak -- or they -- they assumed that all the 185 keV was primarily due to interference from -- from uranium, and then calculated a detection limit above that. So there's no -- I don't know how

you can make an inference from that about the degree of enrichment. And in fact, the protocol that was used in the manual said that enriched -- enrichment of uranium was established using neutron -- neutron analysis, like measuring the proper fission neutrons in a reactor. That was their standard protocol. I have no idea what the uncertainty is of that measurement, and that's what concerns me quite a bit, how they would have done that and what the total uncertainty is of the -- of the neutron measurements that they -- that they used, by their own method, to calculate the U-235.

DR. POSTON: I have a -- I have another question for Arjun. I'm -- Arjun, I'm 71 years old and my memory sometimes fades, but as you may remember, I attended those meetings at the Chapman Valve and when we talked with the folks, and I was there when we made the interviews and I attended every interview that you attended. My recollection is that we found out about these manifolds from a woman who worked primarily at the Dean Street facility, and her recollection was she was sure that they

came from the Dean Street facility 'cause she always typed the shipping orders. I do not remember her saying that they were contaminated. I do remember a discussion between you and John Mauro and myself that says maybe that explains the enriched uranium sample, maybe they were processing contaminated systems that came from Oak Ridge. But as Jim stated in his -- there's no evidence that those were shipped back to the -- the valve company at all, so I -- I don't recall anyone testifying or stating during the meeting that we had that evening in that question and answer session that these -- these manifolds were

DR. MAKHIJANI: Dr. Poston, the -- the record of that interview of course was part of -- you looked at it, and the interview, we looked at it, and that is part of our original review, and I also attached it for convenience --

that is stated in our report is that the existence of an enriched uranium sample is consistent with what she said, since she had

1 said that the manifolds were returned from the 2 elec-- from -- from Y-12. 3 DR. POSTON: I'm reacting to your statements 4 today when you said --5 DR. MAKHIJANI: No --6 DR. POSTON: -- they were contaminated. Yeah, 7 we -- we did agree, the three of us, that that 8 is a potential pathway and that was a potential 9 way that these were contaminated, that's all. 10 DR. MAKHIJANI: I believe -- I believe -- Dr. 11 Poston, our report is quite carefully written, 12 and what --13 DR. POSTON: I'm sure of that. 14 DR. MAKHIJANI: -- (unintelligible) said --15 yes. Well, I hope that you expect nothing 16 less. The -- what we've said that the 17 existence of the enriched uranium sample is 18 I didn't say consistent with what she said. 19 that she said that it was an enriched uranium This is obviously an inference, and 20 21 because it's only an inference, we also have to 22 leave open the possibility that it didn't come 23 from there, and I believe that we have also done that. I mean I -- I believe -- if you 24 25 would like, I would -- I could read it again,

1 but I have already read it into the record. 2 DR. POSTON: Well, no, I'm -- I'm -- again, 3 Arjun, I'm not talking about what's in the record. I'm talking about what you're saying 5 orally when you stood up there. You -- you 6 emphasized that she was -- contaminated 7 manifolds. There --8 DR. MAKHIJANI: I (unintelligible) --9 DR. POSTON: -- is an inference. You didn't 10 indicate that these were inferences, one way or 11 the other. You indicated that these were --12 DR. MAKHIJANI: Well --DR. POSTON: -- contaminated, and that leaves 13 14 the thought in people's minds that they were 15 contaminated. That's why the gentleman is 16 reacting the way he is. DR. MAKHIJANI: Well, in order to be accurate, 17 18 I actually decided that I was going to just 19 read what we had written, and I read that into 20 the record, and I'd like to say again, I don't 21 know -- I can't remember every single word that 22 I said and we'd certainly have to go back to 23 the record and look at that, but for the 24 record, what is written here I'd like to read 25 again since it seems to have engendered some

1 confusion. Let me read the -- perhaps caused 2 by me, I do not know. 3 The point number five that we wrote about this. It is possible that the enriched uranium sample 5 may have been associated with these other 6 activities. However, there is no evidence of 7 this in the reviewed documentation. 8 piece of evidence as to the possible source of 9 the enriched uranium is the site expert 10 interview which described the return of 11 contaminated manifolds from the electromagnetic 12 separations plant at Oak Ridge that was 13 operated during the Manhattan Project and for a 14 short period thereafter. Now I do see here that I think -- I think what 15 16 we have return -- written is not 100 percent in 17 conformity. You're quite right. 18 DR. ZIEMER: Uh-huh, it says --19 DR. MAKHIJANI: It should have said potentially 20 -- return of manifolds from Y-12 which may have 21 been contaminated. 22 DR. POSTON: Yeah, that was --23 DR. MAKHIJANI: You're quite right. 24 DR. POSTON: -- that was the contention between 25 you and John and me, and had nothing to do with

1 the site expert. 2 DR. MAKHIJANI: I'm -- you're quite right, Dr. 3 Poston. What we will do is we will issue a 4 page change to this report and -- and make that 5 correction so it's clear that it is an 6 inference that that could possibly be the 7 source. But the rest of it is very clear that 8 it may not -- it may not be the source. 9 DR. POSTON: You have made my point. 10 you. 11 DR. MAKHIJANI: No, I did, I'm agreeing. 12 DR. POSTON: Yeah, like I say, you did very... 13 DR. MAKHIJANI: That's (unintelligible) --14 I think the gentleman from Dow --DR. ZIEMER: 15 or from Chapman had an additional comment. 16 MR. PETERSON: (Off microphone) 17 (Unintelligible) 18 DR. ZIEMER: Use the mike, please. 19 MR. PETERSON: (Off microphone) 20 (Unintelligible) actually quite a number of 21 comments. One being a inference or being 22 proactive, it appears to me or what I just 23 heard is we're not sure. That -- I think 24 that's what we heard. Maybe I'm wrong, but 25 there's documents that were presented that say

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one thing and I just heard this gentleman say another thing. And again I would go back to the premise that we have to be proactive. we think it might be possible, then we have to go on the assumption that it did happen. know, we -- we're making -- we're making judgment calls with people's lives and issues that happened based on well, it could have happened. I -- I think -- I think the approach should be if you don't know for sure, you can't act in the negative. You have to act in the positive or do more tests. If in fact we found uranium in the building, you -- I just heard an assumption that the manifolds could not be in the building. I'm a registered architect. could sit here and tell you that I don't care what size they are, they make buildings and they make doors and they move space shuttles in and out of buildings. You cannot make a statement that a particular item that you built cannot be housed in a building, moved out of a building and put on a loading deck. absurd assumption, as far as I'm concerned. You know it and I know it. So we're -- we're compounding our assumptions to come up with a

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conclusion that are based on assumptions that we have no right saying. And I guess -- I quess that's my concern. Unfortunately, this gentleman -- he's restricted as to his investigation because he has to go on information that originally by an agency that we're -- we're supposed to be looking at in terms of were their assumptions correct. And -- and I quess that gives me a very bad feeling that all I heard today was maybe, maybe not, and no real conclusive evidence and we're dealing with a lot of scientists here, a lot of brain power here and I don't hear anything conclusive. I hear people making assumptions. DR. ZIEMER: Let me make two comments, somewhat in reaction to that. Number one, the restrictions on the contractor are set by this Board, not by NIOSH.

MR. PETERSON: Okay, fine.

DR. ZIEMER: We -- we define the task for them, so NIOSH did not restrict them in any way.

Number two, I think Dr. Poston described the process that is used in terms of -- to handle uncertainties. John, I think you described it well, and that is to make the assumptions that

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every worker got the highest exposure found, every work day of their -- of their time. So if there was contamination there, then that -- that -- those kind of assumptions, those worst-case assumptions, are -- are intended to cover that.

Now we -- we all -- we all know that if -- if we make a worst-case assumption, I can always think of something worse than the worst-case assumption. But within reason, not just in Chapman Valve but in many of these cases, the bounding -- the bounding assumption or the bounding calculations made by NIOSH are intended to cover those uncertainties. Actually we have found in -- in many cases the better we know the data, the lower the exposures tend to be. These bounding assumptions are extremely user-friendly, if I They're not 100 percent can use that term. certain, but that -- that is the intent. recognize -- not just here, but in almost every case -- there are uncertainties and that's built into the system. So that -- that -nonetheless, we understand the point you made. We -- we don't know whether these -- these

1 manifolds were inside or outside, but --2 although at this point the -- they -- they are 3 not there during -- during the time interval. The contamination was. John's -- the 4 5 description of his calculation is intended to cover that. Whether it does or not, the Board 6 7 members have to make that determination as to 8 whether that satisfies their concerns or not. 9 MR. PETERSON: I -- I quess the only point I 10 would make --11 DR. BRANCHE: Please use the microphone. 12 MR. PETERSON: Oh. I guess the only point I 13 would make, and correct me if I'm wrong, they 14 were not put in the calculation to -- to reach a level. If -- if the manifolds -- we're --15 16 we're stating as far as coming up with the 17 criteria to gauge the amount of exposure, the 18 manifolds were looked at as being neutral. 19 Correct? 20 UNIDENTIFIED: Yes. 21 MR. PETERSON: They were looked at as not being 22 contaminated. 23 DR. ZIEMER: If the contamination -- in spite 24 of the manifolds, if the contamination is 25 present during that period, if it's

1 contributing -- for example, number one, to 2 external, that's covered. If it's con--3 MR. PETERSON: No, but you -- you're doing a reconstruction. You're doing an assumption. 5 You're creating a value level, and in creating 6 that value level the manifolds are not part of 7 that value level. Is that my understanding? 8 DR. POSTON: Well, fir-- let me see if I can 9 clarify this. First, we have no evidence that 10 they were contaminated or not contaminated. 11 Okay? 12 MR. PETERSON: (Unintelligible) 13 DR. POSTON: Now we had this one person, and 14 Arjun has correctly corrected his statement, 15 saying potentially contaminated because there 16 was a sample taken outside of the building that 17 in 19--18 Inside the building. UNIDENTIFIED: 19 DR. ZIEMER: Inside the building. 20 DR. POSTON: -- inside the building that in 21 1992 or something that showed elevated levels 22 of uranium-235. That -- that -- that was not 23 taken into --24 MR. PETERSON: Account. 25 DR. POSTON: -- account in the worst-case

1 bounding sort of calculation of the dose. MR. PETERSON: We're saying the same thing. 2 3 quess that's my point. That -- that has not 4 been factored into the level. 5 DR. POSTON: But that wa--6 MR. GRIFFON: Right, but that's not it. 7 MR. PETERSON: Well, I -- I don't know, I'm not a scientist or -- I'm -- I'm just saying --8 9 DR. POSTON: Yeah, let --10 MR. PETERSON: -- that's just one item. 11 DR. NETON: I don't want to engage in argument 12 or anything, but I just wanted to point out 13 that it was -- the activity that was found was 14 120 picocuries per gram, a fairly low level of 15 activity, the dose of which would be very minor 16 compared to what we've assigned to these 17 workers based on the measurement of uranium in 18 their urines, which were taken on-site. 19 MR. PETERSON: But it would be added to that. 20 DR. NETON: Well, it -- it depends. I mean we 21 do have contem-- we had contemporaneous urine 22 measurements on the people. If -- if -- worst 23 case scenario is if they were all exposed to 24 enriched uranium, it would essentially double 25 their dose. I mean if -- if we -- if we, for

some reason, had no knowledge that two percent enriched uranium was processed at this site as opposed to natural, which I don't believe is the case, the net difference would be a factor of two difference, approximately, in the dose based on our calculations. So it -- it could be factored in if we did find out that there was indeed enriched uranium. And -- and that certainly would be bounding and, again, we -- we could do that if -- if we did have information to that effect.

DR. ZIEMER: Dr. Melius.

DR. MELIUS: Yeah. I mean -- but isn't the real point about the potentially enriched uranium sample -- hopefully I'm correct on that -- is whether or not we have -- whether or not we are aware of all the activities that have gone on at that site. If there were additional activities, then the question is what were they, what time period, and would they further contribute to dose. And I'll remind you that originally the Dean Street facility was not part of this because DOE had not designated -- in fact, DOE came and tried to tell us that the building had disappeared, et cetera, and lo and

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behold, we found that there were -- you know, operations there that's now part of the facility. So I think the question is do we have -- what do we make of this -- this sample and -- as well as the information from this one person who's reported, and how do we resolve that and has there been sort of due diligence -- you know, again, not on the part of NIOSH but on the part of DOE -- in terms of -- of evaluating the operations that were on the site and -- and designating the site. Also remind you that as we -- looking at other older industrial facilities of this type, general type, I think some of them that we've looked at -- in fact, one we looked at earlier today where we're lacking information we are des -- basically designating the whole site as a -- everybody working there as being part of the SEC because there's so much uncertainty about operations and so forth so -- at those sites that NIOSH is not able to describe those operations in a way that's sufficient to develop appropriate individual dose reconstruction. So I think there's also an

issue of how are we being consistent with --

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with what we're doing in -- in other instances. Now not to say there isn't other information on Chapman that may indicate otherwise, but -- but I think -- you know, some of us are very concerned and suspicious about trying to understand this site better and understand the operations there and -- and trying to understand this -- this sample and, again, it's not the situation were we can go back and recreate or resample or whatever. It's -- we have limited information and I think we're trying to understand it and what its implications are in terms of operations at that facility as opposed to particularly exposures at that facility. Go ahead, Jim.

DR. NETON: Yeah, I just -- just a couple of comments on that. I think this is somewhat different than some of the other facilities we've looked at because indeed the exposure that was discussed in those facilities was part of the certified covered exposure period by the Department of Energy. Right now we have no covered period in 1943 to 1944 to even evaluate. We would do that if the Department of Energy said yes, there were covered

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activities there. They've exercised a lot of due diligence, in my mind. In fact, a lot of things had to happen to make this sample a problem. One is it has to be enriched uranium, which I'm still not convinced it is, that it's -- it's -- delayed neutron measurement to establish enriched uranium, which is what their procedure says, is a fairly uncertain measurement technique. Secondly, there are FUSRAP data in '92 and '97, none of which recovered any evidence of additional enriched uranium samples at the site at all. Third, were manifolds shipped back that were contaminated -- were they shipped back at all, or were they actually factory acceptance test pumps. And if they were shipped back, were the contaminated. A lot of things have to happen for that scenario to play out, and it just seems to me the weight of the evidence right now is not there.

DR. ZIEMER: Any other comments or questions.
Yeah, Mark.

MR. GRIFFON: Yeah, just a -- a call out to Jim on that. When you were up here at the podium before you mentioned that there were -- you

1 mentioned the radium issue or -- or the --2 someone had raised early on in the --3 DR. NETON: Yeah. 4 MR. GRIFFON: -- and you followed up by saying, which -- which -- this is where I stand, is 5 6 there's a lot of unknowns here, even if -- you 7 know, is this an enriched sample, you know, and 8 -- and you said --9 DR. NETON: Yeah, I agree. 10 MR. GRIFFON: -- right up here, there's a lot 11 of unknowns here. That's the question, I 12 think. If it was radium interference, maybe 13 they were working on radium beads, I don't 14 know, but that's another operation that we 15 don't know about, so I quess that's the 16 question, going back to the operations 17 question. 18 DR. NETON: That -- that activity, to my 19 knowledge, was not -- not carried out in 20 Building 23 during 1948-'49 --21 MR. GRIFFON: Well, not to mine, either, based 22 on what we're seeing --23 DR. NETON: -- but remember --24 MR. GRIFFON: -- yeah. 25 DR. NETON: -- this is 1048-'49, Building 23

1 and Dean Street are the covered facilities. 2 MR. GRIFFON: Right. 3 DR. NETON: The other buildings are not part of the class definition. And those radium 4 5 activities were carried out elsewhere, if there were indeed, I mean --6 7 MR. GRIFFON: Oh, but I thought you were 8 referring to those, that they might have been 9 in-- the reason --10 DR. NETON: Well --11 MR. GRIFFON: -- for interference with that 12 enriched uranium sample. 13 DR. NETON: It could have been near the loading 14 dock --15 That's Building 23. MR. GRIFFON: 16 DR. NETON: It could have ended up there 17 because there was a pile -- if you look at 18 pictures in the H. K. Ferguson report, there's 19 a pile of material laying over where -- where 20 that sample may have been taken. In fact, this 21 was not -- this was sort of a dust sample. 22 They had to --23 MR. GRIFFON: Yeah. 24 DR. NETON: -- the way they described it, they 25 almost like kind of scooped it up and got a

1 sample of some contaminated dust, if you will, 2 so -- so I -- I'm not sure, but it's -- the 3 radium sources were asserted to have been used by one of -- one of the site experts. 4 DR. ZIEMER: Okay, any other comments? 5 6 (No responses) 7 Board members, are you at a position where 8 you're prepared to take action on this? We're 9 late in the day. We have -- we need a break 10 before public comment period. We can continue 11 the discussion further or if you've heard as 12 much as you wish, we -- we -- it would be also 13 in order to have a motion, so -- one way or the 14 other. 15 MS. MUNN: (Off microphone) (Unintelligible) 16 DR. POSTON: Is that true? 17 MS. MUNN: Could we prepare the motion for 18 tomorrow morning? 19 DR. ZIEMER: We can have a motion. 20 DR. POSTON: We already had a motion and 21 (unintelligible). 22 DR. ZIEMER: Well, one of the options is to do 23 nothing. I guess that's an option. We've had 24 a motion on this, als-- and we -- we ha-- the 25 vote was split.

1 MR. GRIFFON: Right. 2 DR. ZIEMER: It was a -- we had a 6-6 split on 3 Chapman the last time. The effect of a -- of a 6-6 vote is that there is no recommendation 4 5 sent forward. It has the same effect as a 6 motion to deny the petition. I'm asking if 7 there's a motion -- we've had some additional 8 information that's been looked at and so on. 9 If the Board wishes to make a motion, you're 10 entitled to do that. 11 DR. BRANCHE: You would first need to vote to 12 take the motion off the table. 13 DR. POSTON: It's not on the table. 14 DR. ZIEMER: There's no motion on the table. 15 DR. BRANCHE: No, to put it back on the table. 16 You've tabled this issue --17 DR. POSTON: No, we have not. 18 DR. ZIEMER: No, no --19 DR. POSTON: That was incorrect. 20 DR. ZIEMER: -- the motion has never been 21 tabled, we --22 **DR. BRANCHE:** Okay. 23 MS. HOWELL: I think the split --24 DR. ZIEMER: -- had a vo-- huh? 25 DR. POSTON: The vote --

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               DR. BRANCHE: I -- I gotcha. The split
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               effectively tabled it.
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               MS. HOWELL:
                            Yes.
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               DR. POSTON:
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               MS. HOWELL: You -- you would be voting -- you
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               would be picking that back up because as a
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               split vote there was no Board decision so this
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               issue has never been --
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               DR. BRANCHE: Officially tabled.
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               MS. HOWELL: -- officially determined.
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               DR. ZIEMER: Well, okay, if -- if we're going
               to be -- do it parliamentary-wise, this has not
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               been tabled, but you can always vote to
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               reconsider.
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               MS. HOWELL: Right.
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               DR. ZIEMER: A vote to reconsider would put it
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               back on the table.
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               MS. HOWELL: I'm not saying it was tabled.
                                                            I'm
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               saying you're bringing it back --
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               DR. ZIEMER: Right.
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               MS. HOWELL: -- up for a vote.
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               DR. ZIEMER: We can vote to reconsid--
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               DR. BRANCHE: So the motion as it was is what
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               you're --
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               MS. HOWELL:
                            Yes.
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1	DR. ZIEMER: Yeah. A motion to reconsider
2	would be in order, and and you have to vote
3	to reconsider, and then you have the motion to
4	deal with if you if reconsideration is is
5	approved.
6	DR. MELIUS: Perhaps we should refer this to a
7	workgroup on Roberts' Rules of Order to
8	DR. ZIEMER: Wanda Munn.
9	MS. MUNN: If a vote if if a motion to
10	reconsider is in order, I so move.
11	DR. ZIEMER: Okay, there's a motion to
12	reconsider. Is there a second to the motion to
13	reconsider? If if there is
14	MR. PRESLEY: Bob Presley, and I'll second that
15	motion.
16	DR. ZIEMER: Okay, I hear a hear a second.
17	Let me instruct you, if the motion to
18	reconsider comes before us, then we will be
19	considering the motion that we had before. My
20	recollection is that that was a motion to deny
21	the petition.
22	MS. MUNN: I believe that's correct.
23	DR. POSTON: That's correct.
24	DR. ZIEMER: But what I'm going to suggest to
25	you that if we pass the motion to reconsider

1	UNIDENTIFIED: (Unintelligible)
2	DR. ZIEMER: if we pass the motion to
3	reconsider, then I'm going to suggest that we
4	do the reconsideration tomorrow after we have a
5	chance to confirm the nature of the original
6	motion. Is that a would that be well
7	MR. CLAWSON: That'd be agreeable with me.
8	DR. ZIEMER: Okay. All in favor of
9	reconsidering, say aye.
10	(Affirmative responses)
11	Any opposed?
12	(No responses)
13	Mr. Presley, did you vote?
14	DR. BRANCHE: He seconded the motion.
15	MR. PRESLEY: Bob Presley.
16	DR. ZIEMER: Thank you. So the motion to
17	reconsider has been approved and we will
18	reconsider then is it agreeable with the
19	assembly that we do this tomorrow, in view of
20	both the time and the need to get the wording
21	of the original motion that we handled before?
22	MS. MUNN: Yes, please.
23	DR. ZIEMER: I hear no objections. I think
24	with that I'm going to postpone anything on the
25	Mound issue till the public comment period

1 because it's simply reading a letter into the 2 record. 3 It's now 5:30, this -- this assembly is going 4 to come back together at 7:30 for a public 5 comment period. 6 DR. BRANCHE: Only, strictly. 7 DR. ZIEMER: Strictly public comment at 7:30. 8 We will not be debating any motions at that 9 time. 10 Thank you very much, everyone, for your 11 attention. We will see you all in two hours. 12 (Whereupon, a recess was taken from 5:35 p.m. 13 to 7:30 p.m.) 14 PUBLIC COMMENT 15 DR. ZIEMER: Good evening, everyone. If you 16 would take your seats, we're going to begin the 17 public comment session of our Board meeting 18 this evening. 19 I should tell you in advance that we do not 20 have a large number of individuals who have 21 indicated that they wish to speak. 22 nonetheless, we will hear from several. 23 Before we call on our speakers, I'm going to 24 ask our Designated Federal Official, Dr.

Christine Branche, to give us some ground rules

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as far as the policies are concerned on redaction and related things. And I will also tell you that we have an operating Board rule that the individual comments are limited to ten minutes. So Dr. Branche.

DR. BRANCHE: Thank you. I want to make sure people are -- Mr. Presley, can you hear me?

MR. PRESLEY: Yes, I can.

DR. BRANCHE: Okay, thank you. I just want to make sure the line is open.

Our policy on redaction of Board meeting transcripts are as follows: If a person making a comment gives his or her name, no attempt will be made to redact that name. NIOSH will take responsible steps to ensure that individuals making public comment are aware of the fact that their comments, including their name, if provided, will appear in a transcript of the meeting posted on a public web site. Such reasonable steps include the statement that I'm reading now at the start of this meeting. When people signed up today and yesterday, a printed copy of the -- of this redaction policy was available on display at the table where they signed up. A statement of

1 our redaction policy was appended to the NIOSH 2 web site along with the agenda for this 3 meeting. And also our redaction statement was also included with the Federal Register notice 5 for this meeting. 6 If an individual in making a statement this evening reveals personal information, such as 7 8 medical information about themselves, that 9 information will not usually be redacted. 10 NIOSH Federal -- sorry -- Freedom of 11 Information Act coordinator will, however, 12 review such revelations in accordance with the Freedom of Information Act and the Federal 13 14 Advisory Act -- excuse me -- the Federal 15 Advisory Committee Act and, if deemed 16 appropriate, will redact such information. All 17 disclosures of information concerning third 18 parties will be redacted. 19 And if you'd like to make a statement but not 20 reveal your name, I would ask that you come to 21 see me -- well, now. 22 And for those persons participating by phone, 23 if you could please mute your phone. And if 24 you do not have a mute button, please dial 25 star-6. And then when you are ready to speak,

1 you may still either use your mute button or 2 use that same star-6 to unmute your phone when 3 you're ready to speak. It is important that 4 everyone use the mute button or the star-6 5 feature because the people participating by 6 phone will end up with a very unclear 7 transmission of this portion of our meeting, 8 and they will not be able to hear everything. 9 I'm concerned that I still hear some background 10 noise, so that person, if you could please mute 11 your phone. Oh, there's still some background 12 noise. Somebody has a radio. 13 MR. PRESLEY: Christine? 14 DR. BRANCHE: Yes? 15 MR. PRESLEY: Somebody's -- somebody's (break 16 in transmission) their telephone on a (break in transmission) that's got (break in 17 18 transmission) signal on. 19 Will that -- and here's a case in DR. BRANCHE: 20 point. I only got every other word and only 21 part of those words, Mr. Presley, so I think we 22 just need to go ahead and get started. 23 you for muting. 24 UNIDENTIFIED: It appears that someone put

their phone on hold.

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1	AV TECH: I can I can turn it off and you
2	won't hear, but only when (unintelligible)
3	you're going to have a problem.
4	DR. ZIEMER: The people on the line may not be
5	able to hear hear the presentations since
6	somebody on the line apparently is causing
7	noise through their "hold" system instead of a
8	mute system. So if your phone is on "hold"
9	rather than "mute" although if it's on
10	"hold," you're probably not around to take care
11	of this. I don't know the answer to that,
12	but
13	MR. CLAWSON: We could call and have that
14	disconnected.
15	UNIDENTIFIED: Dr. Branche, I think you can get
16	(break in transmission).
17	DR. ZIEMER: Say it again.
18	DR. BRANCHE: Ms. Bon Ms. Bonsignore, what
19	were you saying? Someone was trying to get my
20	attention?
21	MS. BONSIGNORE: I wasn't speaking.
22	DR. ZIEMER: It was someone else.
23	DR. BRANCHE: Okay, it was someone else.
24	Forgive me.
25	DR. ZIEMER: Okay, we're going to proceed.

1 First of all, we have two Congressional letters 2 that we want to enter into the record. 3 from I believe Senator Kennedy's office and I forget -- the other one was perhaps from the 5 Ohio -- was it a Mound letter? 6 Yeah, the second one's a Mound MR. BROEHM: 7 letter. 8 DR. ZIEMER: Okay. So Jason's going to read 9 those letters into the record for us first. 10 MR. BROEHM: Okay. So the first letter was 11 addressed to Dr. John Howard, Dr. Lewis Wade and Dr. Paul Ziemer, dated April 2nd, 2008. 12 13 This was cosigned by Senator Edward M. Kennedy, 14 Senator John F. Kerry and Representative 15 Richard E. Neal about the Chapman Valve site. 16 And we were going to read this in later, but 17 the discussion kind of got ahead of us. 18 (Reading) Dear Dr. Howard, Dr. Wade and Dr. 19 Ziemer, we're writing to bring your attention 20 to the Special Exposure Cohort petition filed 21 by former employees of the Chapman Valve 22 Manufacturing Company of Indian Orchard, 23 Massachusetts, pursuant to the Energy Employees 24 Occupational Illness Compensation Program Act. 25 The company was involved in the nation's

1 nuclear weapons program in the 1940s, and its 2 employees were exposed to radioactive materials 3 in their work. Cleanup activities took place 4 from 1991 to 1995 under the Formerly Utilized 5 Sites Remedial Action Program of the Department 6 of Energy. 7 The Chapman Valve petition was filed on August 8 18, 2005. It was qualified on November 9th, 9 2005, and was submitted to the Advisory Board 10 on August 8th, 2006, nearly six months after 11 the statutory deadline for completion. 12 Petitioners have been waiting for a 13 determination from the Advisory Board for more 14 than two and a half years. 15 Last month Sanford Cohen and Associates, the 16 Advisory Board's experts, issued a new report 17 on Chapman Valve in which they conclusively 18 determined that enriched uranium was present in 19 the Chapman Valve facility. The report 20 concludes, however, that we are still no closer 21 to determining exactly when and how the uranium 22 came to be at the facility. 23 It is now seven years since Congress enacted 24 EEOICPA. A primary motivation for this 25 legislation was the need to expeditiously

compensate workers and their surviving family members. The statute specifically states that the purpose of the program is to provide, quote, timely compensation, unquote, for these Cold War victims. The Special Exposure Cohort is Congress's acknowledgement that, because these were programs carried out in secret more than 50 years ago, many workers would not have the records they need to prove their claims. The Special Exposure Cohort process was specifically designed to ensure compensation for workers whose records are not available. Congress clearly did not intend them to get trapped in an endless search for records that no longer exist.

Yet the Chapman Valve petitioners have been forced into just such an endless search. Over the past two and a half years their petition has been debated and discussed repeatedly at Advisory Board meetings, without conclusion. The Advisory Board has asked the Department of Energy to go back time and again to review its records, and yet the latest report by Sanford Cohen and Associates makes clear that NIOSH still lacks the information needed to make an

1 accurate dose reconstruction for the Chapman 2 Valve petitioners and their families, the 3 program has obviously failed to fulfill its 4 promise. 5 It is unfair to ask these petitioners to wait 6 any longer while the Department of energy 7 endeavors to pursue even more avenues of possible evidence raised by the latest report. 8 9 We ask the Advisory Board to fulfill its duty 10 and grant the Chapman Valve petition as soon as 11 possible. The hard-working men and women of 12 Chapman Valve have waited too long for the compensation they deserve from our country for 13 14 their sacrifices. 15 Thank you for your consideration of this issue. 16 If you have any questions or additional 17 information to provide, please contact Sharon 18 Block in Senator Kennedy's office at 202-224-19 5441. 20 With respect and appreciation, sincerely, 21 Edward M. Kennedy, John F. Kerry, Richard E. 22 Neal. 23 Then the second letter is addressed to Dr. 24 Ziemer, and really to the whole Board, by 25 Representative Michael R. Turner from Ohio.

This is about Mound.

(Reading) Dear Dr. Ziemer, I am writing in support of the Special Exposure Cohort petition 0090 and 0091 for former Mound employees whose dates of employment are between 1959 and the plant's closure. Granting these petitions in full would assist workers who could be victims of a magnitude of potential exposures to radiation and other toxins used and handled at the Mound facility.

It is my understanding that Mound employees who were employed between the dates of 1949 and 1959 have recently been granted SEC status. I thank the Advisory Board for their efforts in ensuring those workers were granted this special status.

Former Mound workers have dedicated years of service to this facility, and many have incurred health problems as a result of the dangerous nature of their employment.

Additionally, in order to properly assess benefits, workers must attempt to reconstruct old employment records that may no longer exist or are otherwise incomplete. It is my hope the Advisory Board will help alleviate the health

1 issue affecting these workers by recommending 2 them SEC status for workers from 1959 until the 3 facility's closure. 4 Sincerely, Michael R. Turner, Member of 5 Congress. 6 Thank you very much. DR. ZIEMER: Next we'll 7 hear from Donna Hand, who represents the 8 Nuclear Workers of Florida, Pinellas. 9 welcome. 10 Just push that down a little bit, or tilt it 11 down will be fine. 12 MS. HAND: I want to thank you very much for, you know, coming to Florida and giving me the 13 14 opportunity to be educated and where do I get 15 my certificate on 101 Advisory Board meetings? 16 All right? 17 And then as a same token, I want to thank you 18 and as a claimant, as a private citizen and as 19 a worker advocacy for all your time and energy, 20 because I have seen first-hand how much time 21 and energy you have spent on all these claims, 22 and I really, really appreciate that you are 23 really, really looking after the worker, as 24 well as making sure that the law is fulfilled. 25 Myself and a introduction to Pinellas plant,

1 I'm going to -- I don't have enough but we --2 you can share. 3 DR. BRANCHE: Can I help you? I'll hand these 4 out. 5 MS. HAND: Okay. The Pinellas plant, as you 6 know, was a neutron device facility. I got 7 these facts not only from the site description 8 that NIOSH has, as well as the baseline report 9 that you can get from the Department of 10 Environment Protection on the decontamination, 11 but also there's Pinellas flat -- plant facts 12 that you can get when you Google GE ND into the 13 DOE OSTI web site. 14 As you can see from the Pinellas plant pro--15 site profile, there was ion silirators (sic), 16 pneumatic seals, high voltage generation, 17 lightning arresters, special decapacitors, 18 vacuum systems -- now these were glass vacuum 19 systems -- crystal resonators, active and 20 reserve batteries and radioisotopically-powered 21 thermoelectric generators. 22 This facility, especially Building 100, was a 23 warehouse-type facility. That means that it --24 that there were partitions, that the ceiling 25 did not -- you know, and the walls did not go

1 up to the wall -- the ceiling in this 2 warehouse, and therefore all the radionuclides 3 was exposed everywhere. In fact, when they did the decontamination, one of the cement walls 5 that was supposed to have had a lead panel in it did not have the lead panel in it. 6 The only monitoring that was done was a finger, 7 8 a wrist and two whole badges, dosometers (sic). 9 These whole badge dosometers were sensitive to 10 some radionuclides and insensitive to others. 11 From 1954 to 1980 only 27 percent of the 12 employees were monitored. From 1980 to 1996 when they closed, only 14 percent were 13 14 monitored. And again, these monitors were not 15 adequate. Some of the reports that you would 16 find would have zero, some of them just said 17 that they met the minimum detection, that's it. 18 So therefore they weren't really accurate 19 because, as you know and you have done --20 excuse me -- in other facility sites that a 21 zero is not adequate at all on a radiation 22 dosometer (sic). 23 If you go next to the next page, you'll see a 24 list of 28 radionuclides. The list of these 28 25 radionuclides was confirmed by DOE to be

1 2 3 4 5 6 7 8 added to this list. 9 10 11 lists 433. 12 13 14 radiation. 15 Again, that came from the Pinellas plant 16 17 18 19 20 21 22 23 24 25 Also which is not on this list but was

present at the Pinellas plant. Also, because they are -- according to the definition of the 40 CFR (unintelligible) also the Atomic Energy Act, and 20 CFR 1910, there's already been determined a significant health effect, according to the (unintelligible) Supreme Court requirement before these substances could be

Besides the radionuclides are other 733 toxic substances. The sitometrics* that DOL has only

Anyway, your concern is the 28 radionuclides, because you're concerned with ionizing

environment baseline report. I need a podium. Okay, the next page you'll see is a chronicle list of unusual events. These are all the unusual events that happened at the facility, so therefore these are all incidents. though the employee may not have been right there, because this warehouse-type effect and the residual contamination afterwards, they were still exposed to that ionizing radiation.

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confirmed when we received a file from DOL, there was a plutonium fire in 1973 at this facility in Building 200.

Again you'll see the radio-producing equipment. There's a list of a whole bunch of equipment that would have radiation-producing on it as These radio equipment was cleaned by people that were not monitored. They were also worked and maintained on by people, again, that were not monitored. And several -- several reports we've been finding out, when we get the entire file from DOL or DOE, that these people were injured while working on these equipment. If they're injured, they're required to, in your dose reconstruction, to have a wound quidance bulletin and to have run multiple myeloma and/or other ill-defined sites. of the dose reconstructions have attempted to do this.

You'll see a report from the health physics
Pinellas plant. It (unintelligible) tritium,
krypton, radiation generators, neutron
generators and calibration sources. It goes on
to explain the badges, the -- the neutrons, et
cetera, et cetera, and the -- like I said, the

1 finger and the rings.

I have requested, under the Freedom of
Information Act, a copy of the dosometer (sic)
readings from the fingers, wrist and the two
whole bodies from NIOSH, because this is what
they're supposed to use when they determine an
unmonitored -- so far, to date, I have not
received it.

You'll see a dose reconstruction view -- review -- overreview (sic). Some of the lettering or wording of this is very confusing. example, claimant favorable assumptions are in addition to this report, but even under these assumptions NIOSH has determined that further research and analysis will not produce a level of radiation. How do they know, they haven't con-- taken all in relevant factors. haven't tooken (sic) in the injuries. haven't took in neutron doses. And you've already done several, several facilities regarding neutron doses. You've also regarded several industries or sites regarding tritium. And as you know that the tritium radiocobiological (sic) effect when it hits water, where does it go to? In the body.

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is a human area. This was a pretty hot area so people would sweat, so therefore absorption through the skin was very, very high.

Again on the next page, dose estimate, as you can see, they confirm that this was classified waste that this person got injured on, that he may have been exposed to photon, electron and neutron radiation, but they were not considered in the external dose. Why weren't they?

Wasn't this part of your guidelines? They had the potential of the exposure and that's all that is required by the law.

There was an insignificant amount of dose received so therefore they ignored that. It doesn't matter what type of dose or level of dose, these are not standards. These are potential exposures, period. That's what your methods say. That's what your guideline says, and that's what your law says 'cause it says radiation-related illnesses.

And again they said incidental exposures may have included neutron doses, but the only unmonitored dose assigned was photon. Why not the other? Aren't you supposed to include all the ionizing radiation in your external and

your internal? Are you not required to use inhalation, ingestion, absorption and injection, and the injection is all the injuries.

Some of these areas were unconfined radioaction
-- radioactive areas, so then therefore these
workers could be exposed because just walking
by it or what-- and also their cafeteria. In
order to get to the cafeteria, he had to walk
through the machine shop or right by the
machine shop area. The machine shop area was a
high radioactive area because of the oil and et
cetera, et cetera.

No unmonitored neutron doses were assigned again. The internal doses, as you can see, they've -- confirm and they state the rems, and then they go to the uncertainties. Since this was a high uncertainty because the person was unmonitored, he was a janitor that worked in decontamination. He was noted in a file that he was cut on his left wrist with classified radioactive waste but none of these relative facts were taken into account. You can see that the IREP input program over in -- they ran it all constant, and then they put in the

perimeter (sic) one, .129? Point one, when 1 2 your technical information document, June 2002, 3 says this would be a hybrid of 30 to 250 keVs 4 with the perimeters (sic) of being lognormal 5 and one to five. If you're going to do the 6 constant, then you should look at every single 7 area, who had the highest dosometer (sic) 8 reading in that area, and the 95th percentile 9 of that person's reading. So that means for 10 every year and every location that every worker 11 went to, they would have had to calculate that, and that is very time-consuming and I think 12 13 that's why you passed that June 2002 NIOSH 14 technical information document. 15 Again, I enlarged it so you could see, because 16 what the claimants get is that smaller version. 17 You're talking about elderly people. 18 can't read it, so... 19 Again I put into the guidance on wound 20 modeling, even though it addresses plutonium, 21 at the very end it says this can go for all 22 radionuclides. 23 Pinellas plant is one of the ten that's been 24 put on the shelf. It was done in 2006 by SC&A. 25 Certain questions were asked. To this date,

nothing's been addressed by NIOSH or anyone else. Because they have not been addressed, the health physicist for NIOSH will not comment on a draft. Because they have not been addressed, these workers are being denied the law by really getting to all the external, internal and environmental radiation exposures from all ionizing radiation because even an environmental report done by Oak Ridge at this facility, if a claimant sat out on that north porch overlooking the pond, they could get over 2,000 bcqs. Now I'm just a country farmer girl. I don't know what bcqs are, but I know it's a lot for just a person just sitting out there looking at the scenery.

Really in conclusion is that why does NIOSH not include things according to your own guidelines in the dose reconstruction? Why does DOL not abide by your guidelines for probability of causation? 42 CFR 81.21 says all cancers, even precancers, neoplasms unknown or uncertainty shall be considered as malignant neoplasms.

DOL refuses to send over precancers. In fact, DOL has a bulletin, 614, that lists cancers in that that is not radiogenic cancers, no known

Some of

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1 medical significance. That's not what the law 2 says. That is not their authority nor the 3 responsibility. By law, that was given to NIOSH, HHS. They are the ones to determine all 5 the cancers by the probability of causation 6 quidelines. NIOSH was supposed to require to 7 do the dose reconstruction as methods that this Board has determined, and I haven't seen any 8 9 Code of Federal Regulations where this Board 10 has changed any of those methods yet. 11 them I think you have addressed as far as 12 technical bulletins and have through Oak Ridge, 13 but you -- the basic method you have not 14 changed. The only one that I'm still confusing 15 of and I'm still having a lesson on is your Monte Carlo simulation, and I'm sure some of 16 17 you know what I'm talking about, others do not. 18 But what basic line is is that in order to be 19 claimant friendly, your photons are supposed to 20 be acute. We don't care if they got exposed to 21 chronically, they're supposed to be acute. 22 That's claimant friendly. 23 Your neutrons is supposed to be chronic. 24 don't care if they got it at an acute exposure, 25 it's supposed to be chronic because that's

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claimant friendly.

And that's all we're asking is that the guidelines and the dose reconstruction methods be ascertained as per as in the law for all the Pinellas workers. If it cannot be done, then please inform us. If it's just that they're forgetting it, then, you know, they -- that needs to be addressed because this is really negligence of the law.

9

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Thank you.

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DR. ZIEMER: Okay. Thank you very much, Donna.

The next person I have on the list I think is here by phone. That's Antoinette Bonsignore.

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Antoinette, are you on the line?

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MS. BONSIGNORE: Yes. Thank you, Dr. Ziemer.

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speaking tonight on behalf of the former Linde

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workers and their families who, despite

My name is Antoinette Bonsiquore and I'm

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repeated efforts to get basic information from

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the Department of Labor regarding a request for

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appeal for redesignation of the Linde facility

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that eliminated residual radiation workers from

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(unintelligible), the Linde workers have been

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unable to get any response from the Department

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of Labor to requests -- a letter that was

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submitted to Peter Turcic on February 6th requesting information about revisitation of the decision and an opportunity to appeal that decision.

Additionally, I would like to address two issues that are related to the Department of Label -- Labor's decision to redesignate the Linde facility.

First, on February 20 Jeff Kotsch addressed the Advisory Board regarding the redesignation issue at Linde and a request was made by the Board at that time for a written statement from the Department of Labor explaining the reason (unintelligible) the decision. That statement was provided to the Board, but I have been unable to get a copy of the statement 'cause the Department of Labor designated the document for Board use only. My question is, why are the Linde workers being denied access to this document, and what do the Linde workers need to do at this point to get the Department of Labor to respond to their February 6th letter? Finally, I submitted an SEC petition covering the residual radiation time period (unintelligible) Linde last month and the

petition is currently under qualification review. Our ability to pursue this petition is directly affected by the Department of Labor's decision about the redesignation and whether we will be provided with an opportunity to appeal that decision. It is imperative that the Department of Labor provide an answer to the request to appeal the redesignation decision and stop denying the Linde workers access to the basic (unintelligible) that directly affects their rights under the Part B program. Thank you.

DR. ZIEMER: Thank you, Antoinette. And although, as you know, the Department of Labor issues are not directly the responsibility of this Board, I can tell you that Jeff Kotsch, as well as some other Labor colleagues, are with us today and they have heard your concerns, so -- in fact, maybe Jeff is -- is actually approaching the mike so he may have some comments for you. Thank you.

MS. BONSIGNORE: Great, thank you.

MR. KOTSCH: Antoinette, we -- I know Labor ha- is in receipt of your -- what date is it -February 6th, 2008 letter. I know they're

working on a response, I just don't know where it -- it is. I know it's close to the end of the process of getting out.

I did mention to the Board today that on Friday
I was informed that, as far as the residual
period, the decision was made, after review of
the 2004 amendments to the Act, that the
workers in Buildings 30, 31, 37 and 38, which
are the buildings that changed designation from
AWE status to DOE status, who worked -- so
people in those four buildings who worked only
during the residual radiation period are also
eligible for Part B benefits as atomic weapons
employees, even though they changed the status
of those buildings to a DOE facility. So that
-- you'll be seeing that -- that decision in
your letter whenever it -- when it arrives.

MS. BONSIGNORE: I'm -- I'm sorry -- I'm sorry,

Jeff, I -- I didn't quite understand what you

just (unintelligible).

MR. KOTSCH: Well, first of all, let me say that this decision was made -- and only applies to Linde Ceramics, but Buildings 30, 31, 37 and 38, which are the ones that changed status, workers who worked only during the residual

1	radiation period will now be eligible for Part
2	B benefits as AWE employees.
3	MS. BONSIGNORE: Okay. So essentially the
4	decision (unintelligible) coverage has been
5	rescinded?
6	MR. KOTSCH: Yes, for as far as the review
7	for Linde Cer this is only applicable to
8	Linde Ceramics.
9	MS. BONSIGNORE: Right. Okay. I was not aware
10	that that decision had been made.
11	MR. KOTSCH: Oh, that was it's very recent.
12	Like I said, I got it con not going out the
13	door, but in a meeting on late Friday.
14	MS. BONSIGNORE: Okay. And so are has
15	Senator Schumer's office or Senator Clinton's
16	office been advised of this?
17	MR. KOTSCH: They will be. That that
18	information is also in letters that that
19	will be going to them.
20	MS. BONSIGNORE: Okay.
21	DR. ZIEMER: Okay, you may be the first to
22	know, Antoinette, so
23	MR. KOTSCH: I I have to
24	DR. ZIEMER: but you will be hearing this in
25	writing, is my understanding.

1	MR. KOTSCH: Yeah, they as far as I know
2	I mean the letter is is I asked before I
3	left on Friday, which is when I get kind of
4	get instructions on what
5	MS. BONSIGNORE: Okay, well, I I guess tell
6	you I'm not easily shocked, but I'm I'm
7	shocked. Okay. Well, I appreciate that
8	information and I assume that letters to
9	addressed to me and to Senator Schumer and
10	Senator Clinton will be forthcoming at this
11	point.
12	MR. KOTSCH: Yeah. I mean I can't guarantee
13	the the time frame, but I know that they've
14	been drafted and, as far as I know, they're
15	working their way through the through the
16	end of the process.
17	MS. BONSIGNORE: Okay. All right. Well, thank
18	you very much.
19	DR. ZIEMER: Okay. Thank you, Antoinette.
20	MS. BONSIGNORE: Thank you.
21	DR. ZIEMER: Uh-huh. Now I I'm going to ask
22	if there are any individuals in the assembly
23	here who did not have an opportunity to sign up
24	but do wish to make public comment?
25	(No responses)

1	There appear to be none. Are there any
2	individuals on the phone who wish to make
3	public comment?
4	MR. FUNKE: Dr. Zimmer (sic)?
5	DR. ZIEMER: Yes.
6	MR. FUNKE: This is John Funke.
7	DR. ZIEMER: Yes, John, do you have some
8	comments for us?
9	MR. FUNKE: Yes, I do, I
10	DR. ZIEMER: Okay.
11	MR. FUNKE: (unintelligible) on the agenda
12	(unintelligible)
13	DR. ZIEMER: No, that's fine. You're quite
14	welcome to make comments. Remember the ten-
15	minute time limit.
16	MR. FUNKE: I do.
17	DR. ZIEMER: Okay.
18	MR. FUNKE: I sent the Board the entire
19	Board a packet of information
20	DR. ZIEMER: Yes, we have received that.
21	MR. FUNKE: (unintelligible) got it?
22	DR. ZIEMER: Yes.
23	MR. FUNKE: Okay. There were three very
24	important points to that information. One was
25	the the job classification was was post-

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1992 on the site profile. They are no good.
We're not concerned about what happened after
'92 (unintelligible) the test was over. We're
concerned only with the job classifications
when the tests was going on. In 1992 there was
a major change in the job classification at
Nevada Test Site.

Number two, there was a -- in the site profile made a comment that all the radioactive areas were fenced in and posted with (unintelligible). I sent you a document that tells you as late as 1996 there still was no (unintelligible) and no (unintelligible). There was also a question about area (unintelligible) as I have Fred Dunham on the line is going to comment after me. There was a -- it shows this -- that area -- I know you don't want to talk about Area 51, but it was a part of Nevada Test Site, and up until the realignment in 1999 it was part of the Nevada Test Site. All the employees who worked there came and went through the Mercury gate.

the employees who worked there (unintelligible)

paychecks with DOE allocated funds. All the

equipment used over there was part of

(unintelligible). The general manager of Area 51 was [name redacted] (unintelligible), a REECo area manager. We cannot ignore that area any longer. It has to be considered. It was part of Nevada Test Site and it should be -- if nothing else, we're not interested in the secrets that are out there. We're only concerned about the contamination that the people who worked there would have gotten. That has to be considered and can no longer be ignored.

There's another problem that surfaced. NIOSH - recently there's been some claims that's been
reopened on super S plutanium (sic), and a
couple of other items. And NIOSH has been
sending out arbitrarial (sic) denials with a
single form letter. They're all the same.

I've looked at a dozen of them already, they
just change the names on them. They're not
running the programs through an IREP, they're
simply saying the changes on the site profile
doesn't make any difference, therefore it was
changed by original denial. Your claim is
still being denied. I don't see how they can
do this considering the amount of

1 (unintelligible) as we found in the site 2 profile. I don't even want to begin naming 3 them off. I've already named them off to you a 4 dozen times and they're not considering this 5 and this is not allegations. We've provided 6 documentation. I have more coming. And I -- I would like you to address the issues I sent you 7 8 in the packet, if you would, and also I'd like 9 to find out why NIOSH is just sending out these 10 arbitrarial (sic) denial letters to these 11 people when there is a lot more than just the 12 one issue. They are not rerunning the claims like they're supposed to and I'm -- I'm 13 14 wondering whether that's even the -- the -- the 15 procedure that Congress laid out. 16 So I'll (unintelligible) over to Fred Dunham. 17 He's (unintelligible) the next man behind me. 18 He's one of the people that worked over in that 19 area and he's got some concerns. He's got some 20 more information (unintelligible) to you 21 tomorrow. 22 DR. ZIEMER: Okay, thank you. 23 MR. FUNKE: Thank you, Dr. Zimmer (sic). 24 DR. ZIEMER: So Fred Miller, are you there 25 then?

1 MR. FUNKE: Fred Dunham. 2 MR. DUNHAM: My name's Fred Dunham. 3 DR. ZIEMER: Oh. 4 DR. BRANCHE: Could you please say your last 5 name again? 6 DR. ZIEMER: What's your last name again? 7 MR. DUNHAM: Dunham, D-u-n-h-a-m. Thank you. Proceed. 8 DR. ZIEMER: 9 (Pause) 10 Go ahead, Fred. 11 MR. DUNHAM: Okay. I was an employee out at 12 the Nevada Test Site in the area commonly known I came down with chronic 13 as Area 51. 14 obstructive pulmonary disease after being 15 exposed to chemical fumes from the byproducts 16 of open pit burning of material that was used 17 in a classified aircraft. And through the 18 Department of Labor, my claim has been refused 19 on the basis that Area 51 was out of the 20 boundary of the Nevada Test Site. 21 Now on October 23, 1999 President Clinton did a 22 realignment of the property, transferring the 23 ownership and control of the property known as 24 Area 51 to the Department of the Air Force in a 25 trade for a piece of property that the

25

Department of the Air Force had that was contaminated with nuclear material (unintelligible) Department of Energy. So the Department of Labor clearly erred based upon the information on the site alignment that occurred in the 23 October 1999. Like I say, I worked there from 1981 to 1991.

I would like to have some clarification on that. Also they indicated that the contractor that I worked for, EG&G Special Projects, was not a Department of Energy subcontractor. Well, at the time, EG&G Special Projects was a contractor for both the Department of Energy and the Department of Defense. I worked for -part of the things that I did was the rad safe badge exchange. I traveled to and from across the Nevada Test Site, exchanged the dosimeter badges for that location, manned a post at the 700 gate which was clearly out of the bounds of Area 51 directly across from a (unintelligible) So I can't understand how the quard. Department of Labor can suggest that Area 51, between October -- I mean from 1980 to '92 was not part of the Nevada Test Site. The prime contractor for that particular area was

1 Reynolds Electric. All the contracts went --2 for the subcontractors went through Reynolds 3 Electric and the site manager was [name redacted] (unintelligible), as Mr. 5 (unintelligible) said, and he was an employee of Reynolds Electric, who was the prime 6 7 contractor for the Nevada Test Site. And with 8 that, I -- I'd like to have some clarification. 9 Either the President knowingly signed a bogus 10 document or there's a mistake in the Department 11 of Labor when they looked at my claim. 12 DR. ZIEMER: Okay, thank you, Fred. Again, 13 this is an area -- it appeared to me, an area 14 that is outside the jurisdiction of this Board. 15 However, I will -- since there are some 16 Department of Labor folks here with us tonight, 17 I don't know if they're in a position to answer 18 that at all but at least they've heard your 19 comments and will follow up as they may deem 20 necessary. Jeff Kotsch, you probably don't 21 know the answer to that at the moment -- but he 22 has heard your comment, so --23 MR. DUNHAM: I -- I do have the --24 DR. BRANCHE: Who is this? 25 DR. ZIEMER: Is this Fred?

1	MR. DUNHAM: Yeah, this is Fred again. I do
2	have the particular information on the site
3	realignment that occurred in October 23
4	October '99 that I could fax to you
5	DR. BRANCHE: Not to us.
6	DR. ZIEMER: Well, again, this Board is not the
7	one involved with those decisions by Labor. I
8	think you would need to provide that directly
9	to a Labor representative, and I'm going to
10	pause here a minute. Maybe Jeff, can you
11	advise me on
12	MR. DUNHAM: Yeah, I I have done that
13	DR. ZIEMER: Oh, okay. You have done that
14	already?
15	MR. DUNHAM: Yes, and
16	DR. ZIEMER: Okay, that's what you need to do.
17	MR. DUNHAM: I sent them to the lady here and
18	her name is
19	DR. ZIEMER: Well, she
20	MR. DUNHAM: [name redacted], I believe, and
21	
22	DR. ZIEMER: Yes, well, she can channel that to
23	the proper person then. Thank you very much.
24	MR. FUNKE: Dr. Zimmer (sic), one
25	DR. ZIEMER: Yes.

1 MR. FUNKE: -- one comment, please. 2 DR. ZIEMER: Sure. 3 MR. FUNKE: We are concerned with the 4 boundaries at Nevada Test Site during the 5 period of testing --6 DR. ZIEMER: Yes, I understand --7 MR. FUNKE: -- (unintelligible) 1992. 8 DR. ZIEMER: Yes, and -- and that could -- that 9 could impact indeed on the rest of the program. 10 Again, a determination that Labor would have to 11 make on our behalf. 12 MR. FUNKE: Well, one -- one thing I'd like to 13 remind them of, though, like I said, the 14 general manager was (unintelligible), he was 15 the REECo general manager. All the employees 16 worked over there were REECo employees. 17 all came and left the site through the Mercury 18 gate. They all wore Department of Energy 19 All their paychecks was the same as 20 mine. They were all REECo paychecks, and they 21 was all paid for through the appropriations 22 fund of Department of Energy. 23 DR. ZIEMER: Understood. 24 MR. FUNKE: And this cannot be ignored longer. 25 DR. ZIEMER: Okay, thank you.

1 MR. FUNKE: (Unintelligible) 2 DR. ZIEMER: Is there anyone else on the line 3 that wishes to make public comment? 4 (No responses) 5 Again I'll ask, is there anyone else who wishes 6 to make public comment? 7 UNIDENTIFIED: Hello? 8 DR. ZIEMER: Yes? We hear you. Do you wish to 9 make comment? UNIDENTIFIED: Well, yes, I must have gotten in 10 11 the wrong conversation here 'cause I was -- my 12 concern is relative to the Pinellas plant. 13 DR. ZIEMER: Yes, the Pinellas -- this is 14 dealing with the Energy Employees Occupational 15 Illness Act. Pinellas plant is one of the facilities of interest. Do you have a comment 16 17 relative to Pinellas? 18 UNIDENTIFIED: Yes, only (unintelligible) --19 DR. ZIEMER: Give us your name and then we'll 20 hear what you say. 21 MR. FLYNN: All right. Name is Jim Flynn, F-1-22 y-n-n. 23 DR. ZIEMER: Uh-huh. 24 MR. FLYNN: I live out in Oakland, California. 25 DR. ZIEMER: Okay, what is your comment, Jim?

1 MR. FLYNN: I guess it's more of a question, 2 and that is -- I -- I've been diagnosed now 3 with chemical (unintelligible), and as I look 4 back over my career and the only place that I 5 was exposed to that type of thing would have 6 been at that location working -- I spent my 7 time out there, approximately four years -- on 8 the floor. And that would be making sure that 9 (unintelligible) was in control or 10 (unintelligible), that type thing. So is that 11 (unintelligible) doctors explored this thing, 12 who do I contact relative to information 13 regarding (unintelligible) and, you know, that 14 type of thing? 15 I believe you're going to have to DR. ZIEMER: 16 make contact with Labor, too. And if he's in 17 California, can we give him a contact? 18 would he contact in California for the proper 19 information? We have some Labor folks here. 20 UNIDENTIFIED: He can contact --21 UNIDENTIFIED: You have to go to the 22 microphone. 23 DR. BRANCHE: There's a Labor (unintelligible). 24 DR. ZIEMER: We're going to try to get you a 25 name here 'cause you have to go through the

1 Department of Labor on this. 2 MR. FLYNN: Okay. 3 DR. ZIEMER: Okay. 4 MR. MILLER: Yeah, this is David Miller from 5 the Jacksonville office. We would handle a 6 claim for the Pinellas plant, and you would 7 need to contact our office here in 8 Jacksonville. If you could, if you could give 9 us a telephone number, we can call you back and 10 get that information from you personally. 11 Would you mind doing that? 12 Sure, [redacted]. MR. FLYNN: MR. MILLER: Okay, sir, we'll call you first 13 14 thing tomorrow. Is that fine with you? 15 That's fine. Now who would be MR. FLYNN: 16 calling me? 17 MR. MILLER: I'll call you myself. My name is 18 David Miller. 19 MR. FLYNN: Okay, Dave. All right, sir, look 20 forward to the call. 21 MR. MILLER: Thank you. 22 DR. ZIEMER: And thank you, Jim. Hang on. 23 We're going to repeat the phone number to make 24 sure we have it right. It's [redacted]. Ιs 25 that correct?

1	(No responses)
2	He may we may have lost him.
3	UNIDENTIFIED: I got it on tape.
4	DR. ZIEMER: We have it on tape if we need it,
5	very good.
6	Any is there anyone else on the line that
7	wishes to make comment?
8	UNIDENTIFIED: Is there can you make comment
9	on Chapman Valve?
10	DR. ZIEMER: Yes, you may.
11	UNIDENTIFIED: Okay.
12	DR. ZIEMER: Identify yourself and make your
13	comment.
14	MR. DUARTE: My name is Robert Duarte. I'm
15	from Springfield, Massachusetts.
16	DR. ZIEMER: Could you spell your last name?
17	MR. DUARTE: D-u-a-r-t-e.
18	DR. ZIEMER: Thank you.
19	MR. DUARTE: I'm hoping [redacted] in the
20	audience there. I don't know if she is, but
21	she lives in [redacted]. I'm calling
22	concerning [redacted]. We had a appeals two
23	years ago in [redacted] case. [redacted]
24	worked for Chapman Valve as a foreman in
25	Building 23 and he died at the age of 36. He

1 was the second-youngest one to die there. 2 Chapman Valve destroyed all his records and we 3 have to prove that to the people that we went 4 to the -- went to the tax people, we -- all his 5 records prove that he worked there for 12 -- 12 6 -- from '47 to '59 and that the appeals from 7 Washington said they would give us a decision 8 in 90 days. Now we're two years later waiting 9 on this. I've spoken to Mark Rolfes on many 10 occasions with no -- no -- no results. 11 (Unintelligible) got a reading, they said they 12 were going to send it back to Cincinnati so he 13 could get another reading, and we're still 14 waiting on that. And I asked them before the 15 meeting, at the end of the meeting, if he was 16 going to make the final decision and this 17 gentleman, [name redacted] (unintelligible), 18 said that he was. And now, two years later, 19 [redacted] going to be 85 now. She's still 20 waiting, still getting the same results. 21 I understand they're supposed to have a -- a 22 recount somehow or some kind of 23 (unintelligible) again. You know anything 24 about when that's going to be? 25 DR. ZIEMER: We'll give you a timetable here.

1 Chapman Valve will be on our agenda tomorrow 2 morning, --3 DR. BRANCHE: I'd say after 10:00 a.m., Eastern 4 Standard Time. 5 DR. ZIEMER: -- approximately 10:00 a.m. 6 Eastern time. You can -- again, you can call 7 in and listen in if you wish. 8 MR. DUARTE: On that vote, doing a re-vote 9 aqain. They had a 6-6 vote the last time, I 10 quess. 11 DR. ZIEMER: That's correct. 12 MR. DUARTE: Probably nothing transpired again, 13 which they just keep prolonging this thing 14 here, but nobody's getting results from Chapman 15 Valve. You know, you don't just die at 36. 16 The autopsy showed everything he had 17 (unintelligible) -- slow everything down with 18 [redacted], who's going -- like I said, she's 19 going to be 85 and we still haven't got any 20 results from the people there. I'm sure Mark 21 Rolfes may be there, I don't know. 22 DR. ZIEMER: Mark is not here at this meeting, 23 but as I say, we will be discussing Chapman 24 tomorrow morning, approximately 10:00 o'clock, 25 so you're -- you're welcome to join at that

1 time, if you wish. 2 MR. DUARTE: Has there been any other comments 3 from Chapman Valve tonight? I don't know, I 4 didn't get in at the beginning of this meeting, 5 but --6 MR. CLAWSON: Yes. 7 DR. ZIEMER: Yes. There has been. 8 MR. DUARTE: Okay. 9 DR. ZIEMER: Thank you. 10 MR. DUARTE: Very good. 11 DR. ZIEMER: Anyone else on the line that 12 wishes to comment? 13 MS. RYAN: Yes, my name is Darlene Ryan. 14 DR. ZIEMER: Ryan, is it, R-y-a-n? Yes, it is. 15 MS. RYAN: 16 Thank you. Proceed. DR. ZIEMER: 17 MS. RYAN: You've had many calls from my 18 friends and I who are in (unintelligible) and 19 the contentions of most of the people is that 20 they're spending more time and money denying us and having these meetings -- and one time it's 21 22 in Vegas, the next time it's in Tampa or 23 wherever -- that these people, day by day, are 24 dying off and we keep getting the same results. 25 They're not finding any new information.

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They're not coming up with anything that is positive for them but they're still putting us off like they are again tomorrow for the (unintelligible). And I was a little upset when today Dr. Poston said that he was really on (unintelligible) came up to Chapman Valve (unintelligible) without any (unintelligible) I was there and he sat in the back of the room and at one point one of the girls hit me because he was snoozing. Well, I know (unintelligible) be over (unintelligible) we don't feel anybody is really listening to us and listening -- this -- we didn't ask for this. We did not ask you. You came to us. You opened up a part of my heart that I never knew the suffering and what my father went through and how he did it for his country. took that (unintelligible), he probably, like most of them say, didn't even know how dangerous this job was, but he died for his country. He had a long, four-year death from cancer, and six months later my mother was dead, who took care of him. And finally in the back of our heads, we were going on, and you get a letter telling us that you're looking out

1	for us to give us people something you're
2	helping us, when you haven't helped us. You've
3	hurt us. We have many people.
4	(Unintelligible) paper I picked up a woman from
5	Chapman Valve died. Are they just going to
6	(unintelligible) until there won't be anybody
7	left to pay? We're very hurt here and I feel
8	very (unintelligible) when I get a call from
9	someone who may say to me do you have any
10	answers for me, and I say no.
11	DR. ZIEMER: Okay. Thank you, Bev (sic), for
12	those comments. Let me ask if there's anyone
13	else on the line who wishes to comment?
14	(No responses)
15	No further comments?
16	(No responses)
17	Okay. Again, anyone here in the assembly that
18	has comments? Yes
19	MS. HAND: (Off microphone) (Unintelligible)
20	second chance?
21	DR. ZIEMER: You bet.
22	MS. HAND: Thank you.
23	DR. ZIEMER: Now you don't get another full ten
24	minutes now, remember.
25	MS. HAND: What was my remainder of time, did

anybody notice?

DR. ZIEMER: No, you're -- I'm just kidding.
Go ahead.

MS. HAND: My question is, is that the radionuclides, do they change, as far as factors go, to their distribution or their assumptions, depending on their geographic locations, such as in the dry areas where you've got sites that are in the dry area, does that energy distribution change as compared to Florida? (Unintelligible) has no effect on it, but it does have effect on the soluble and insoluble, does it not?

Also, at the Pinellas plant was phosphoric acid. The phosphoric acid, if it came from central Florida, had a higher degree of uranium inside that phosphoric acid. Because it is mixed with acid, it is soluble, so therefore these workers were exposed to insoluble and soluble. There are eight sites in the state of Florida. Pinellas plant is the only ones having claims. Five of these sites are in the phosphate industry. And if you'll look at those, there's only two or three claims. Some of the sites have zero claims -- everything.

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We would request that you have a resource center or tell us how to go about getting a resource center for the state of Florida. These people are elderly. The survivors don't know anything, don't know how the program -and the phone interviews are not working. For example, one of my clients was going to do a update on his employment history. requested that the form be sent, because I'm his authorized representative, to him and me. I went to his house. We went over it. Again we find out that he was a janitor, but he worked as a decontamination area (sic) -- or that he did this, he did that. Okay, in what area did you work? We did this, we did that. We looked in the Pinellas baseline report, oh, this area -- you've -- exposed to this, this and this. You did this, this and this. You did this, this and this. By the time when we came to the interview, we had everything down pat. We knew exactly where you're exposed to beryllium instead of no, don't know, yes, we were exposed to beryllium. Were you exposed to cobalt? Yes, we were exposed to cobalt. Were you exposed to explosives such as boron? Yes,

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we were exposed to boron. We were exposed to all those incidents, besides the personal incidents. We had all that documentated (sic). Also in the form they are not claimant friendly. The NIOSH form that there was required for the people to sign at the end of the interview that I do not know anymore information is very, very intimidated and not claimant friendly at all. And in the hearing officers -- in their final decisions, use it against the claimants. The -- going back to the employment history, you have a frequency that you have to list. It's one to five in the very center, but over in the column it has one to three, five meaning every day, one meaning hardly ever. So if a -- over the phone you're saying answer me one to three, they're not getting the adequate picture because some of these people it happened every day, but -- and I asked the -- the hearing case person that was taking the interview, did you know that there's a clerical error? Yes, we know it's a technical error but we can't change it because DOL did it. It's a technical error that affects not only the claimant's dose

reconstruction, but their entire claim. You know, these are just two issues that are very, very confusing. I called up the resource center as -- and suggested -- I said 'specially since these claims are elderly, whenever you call up and you make the appointment for a phone interview, why don't you send that employment history ahead of time to them so they can look over it, start refreshing their memory and et cetera, et cetera? We can't do that unless they ask for it.

Again, these are issues that pertain to the law, the regulations, which is in your purview, that is in your job responsibility descriptions to help that part, to tell legislative these things are happening, not just the SEC group and everything, but dose reconstruction, the probability of causation and the problems these people are having with this. We want consistency, which means that if the guy next to me got paid for skin cancer, I did the same job, I worked the same way, how come I'm being denied skin cancer? I had the same dose. I had the same distribution. I had the same exposure to ionizing radiation. Why am I being

1 denied? And again, we need a resource center 2 here because we do have eight sites. We do 3 have a lot of claimants that are not even aware 4 of this program. 5 DR. ZIEMER: Thank you very much. And last 6 call for anyone else here who wishes to 7 comment. 8 (No responses) 9 If not, I thank you all for your time this 10 evening. This Board will reconvene in the 11 morning to complete our business, and we will 12 hope to see many of you then. Thank you and 13 good night. 14 (Whereupon, the day's business was concluded at 15 8:35 p.m.) 16

CERTIFICATE OF COURT REPORTER

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STATE OF GEORGIA COUNTY OF FULTON

I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of Apr. 8, 2008; and it is a true and accurate transcript of the testimony captioned herein.

I further certify that I am neither kin nor counsel to any of the parties herein, nor have any interest in the cause named herein.

WITNESS my hand and official seal this the 10th day of May, 2008.

STEVEN RAY GREEN, CCR, CVR-CM, PNSC

CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102