THE U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES PUBLIC HEALTH SERVICE

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

convenes

MEETING 50

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

VOL. II DAY TWO

The verbatim transcript of the 50th

Meeting of the Advisory Board on Radiation and

Worker Health held at the Holiday Inn Select,

Naperville, Illinois, on Oct. 4, 2007.

STEVEN RAY GREEN AND ASSOCIATES NATIONALLY CERTIFIED COURT REPORTERS 404/733-6070

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TRANSCRIPT LEGEND

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PROCEEDINGS

(9:40 a.m.)

WELCOME AND OPENING COMMENTS DR. PAUL ZIEMER, CHAIR

DR. ZIEMER: Well, good morning. I'll officially 1 2 declare the meeting to be back in session, 3 remind you again to register your attendance. 4 We like you to do that each day. Even if you 5 registered yesterday, do so again today so we have a record of who was in attendance at the 6 7 meeting. 8 Again I'd like to remind everyone that there 9 are copies of the agenda and other documents 10 relating to the agenda on the table in the back 11 of this room. 12 The record will show that all of the Board 13 members are here today in attendance. We have 14 a full quorum. 15 Dr. Wade, do you have any opening remarks 16 before we get to the agenda? 17 DR. WADE: No, only to say to those in the 18 audience, all of the papers that I've given out 19 this morning are on the table in the back as 20 well and for your consideration.

HANFORD SEC PETITION

DR. ZIEMER: Thank you very much. We're going to now go to the first agenda item for this morning, which is the Hanford site -- or the Hanford SEC petition and the review by NIOSH.

Also I -- before we get underway with the presentation, I want to double-check -- although it's early in Hanford -- in Richland, I want to see if Ms. Hoyt or Ms. Carrico are on the line. They are representing the petitioners and we'll hear from them after this presentation, but let's see if either of them, or both, are on the line.

(Pause)

Yes, they are. Thank you. Then we will proceed. Dr. Sam Glover -- is Sam going -- no -- yes, Sam is going to make the presentation on the Hanford SEC petition evaluation report. Sam, welcome.

UNIDENTIFIED: (Off microphone)

(Unintelligible)

DR. WADE: Oh, yes, I'm sorry, we do have conflicts. We have two members conflicted; Ms. Munn and Ms. Beach will need to leave the table.

1	(Pause)
2	You can take your time, Wanda, and get your
3	MS. MUNN: That's okay, I can watch Sam from
4	(unintelligible).
5	(Pause)
6	DR. GLOVER: (Off microphone) All right, can
7	you hear me now?
8	I'm Sam Glover (unintelligible) the second
9	you can't?
10	(Pause)
11	Better?
12	DR. ZIEMER: There you go.
13	DR. GLOVER: There we go. Got to get trained
14	on all these new pieces of equipment. So I'm
15	Sam Glover. I'm here to present the second
16	part of the Hanford Special Exposure Cohort
17	petition evaluation 57. Back in July we were
18	at Hanford and presented the first part of
19	this. This will cover the period from the
20	first part covered for the period from 1943 to
21	September 1st, 1946. This will continue from
22	September 1st, '46 through 1990.
23	Very briefly, this was presented previously at
24	the but we're going to talk there were
25	three petitions that were submitted; Petition

1 50, which covered the earliest period, from '44 2 to September 1st, 1946; the Petition 57, which 3 covers the time span from 1942 to December 4 31st, 1990; and a Petition 78, which covered 5 the period from 1967 to 1971 and is fairly specific. They were discussing maintenance 6 7 carpenters and apprentice that worked in 100, 8 200, 300 and 400 areas of Hanford. 9 This is a ma-- this very large chunk of time --10 what we chose to do with this, as we discussed 11 with the Board, was to merge these into a 12 single petition and evaluate them in two time periods, because there were clear splits with 13 14 where the contractor changed in 1946, the 15 DuPont years, and then after the DuPont years. 16 So we presented that first part in July and 17 this report will be the second part, which goes 18 from September 1st, '46 until 1990. 19 This second evaluation report was -- it's -was issued September 9th, 19-- September 9th of 20 21 2007. The previous evaluation report was 22 issued in May 2007 and presented to the Board 23 July 2007. Previously the basis for the SEC 57 was --24 25 petitioners provided information and affidavit

1 statements in support of the belief that 2 accurate dose reconstruction over time is 3 impossible for the Hanford workers in question. 4 They claimed that personal monitoring data gaps 5 exist in several of the individual workers. And finally, however, what qualified the 6 petition, was that during the early time frame 7 8 NIOSH identified some pre-1949 operational 9 periods which no internal exposure monitoring 10 was performed or was reliable. 11 SEC 50 was qualified based on being completely 12 encompassed by the class proposed by SEC 57. 13 SEC 78 was qualified based on construction 14 workers performed work that took place in these 15 contaminated areas and sometimes required 16 respiratory protection. They asserted that no 17 bioassay monitoring was performed for this 18 class, and they provided documentation 19 regarding the potential for missing external 20 dosimetry records. 21 NIOSH evaluated the following class: 22 employees in all facilities and areas of the 23 Hanford Nuclear Reservation from September 1st, 24 1946 through December 31st, 1990 for this 25 specific petition part. We evaluated the first

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pa-- as ER Part 1 -- SEC 57 Part 1, so that's 1943 through this September 1st, 1946 period. So what (unintelligible) look at? There's a tremendous amount of documentation for Hanford. Hanford estimates some 220 million records exist. There are buildings of documentation. So start out -- part of the -- to do techni-to do dose reconstruction, NIOSH assembled a Technical Basis Document. This has been -undergone several revisions. It is currently under review by the Board. And so over the last -- course of the last year, it has been the subject of a great deal of discussion. As is -- the document is a six-part document having an introduction, a description, how to do medical dose, what is the background for environmental dose, internal dosimetry, and external dosimetry. And some of those are fairly recent revisions, being June of 2007. A variety of Technical Information Bulletins that assist with dose reconstruction were also reviewed. These include maximum plausible doses to workers, OTIB-4, at atomic weapons employers; external coworker dosimetry data for Hanford site, OTIB-30; OTIB-39, internal

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coworker dosimetry data; and OTIB-54, fission product acti-- fission and activation product assignment for internal dose related to beta and gross gamma analysis. Again, many of these are fairly recent documents. Additional Technical Information Bulletins include ambient dose reconstruction for DOE sites, X-ray dose reconstruction for Department of Energy sites, and also O-- apparently I missed the -- OTIB-52, parameters to consider for processing claims for construction trade workers. We conducted many outreach meetings and interviews with unions and with the general public. These include -- so interviews include those provided as part of the Sanford Cohen Associates review -- I'm sorry, (unintelligible) keep (unintelligible) this button. For my laser pointer it's got a different one. The worker outreach meeting in 2004, another one with the Atomic Metals Trade Council in January 2004. The -- and additional interviews include -- we had several in 2004. We had a specific -- we had several days where we addressed early workers and some of those went up to 1950. We were out there in March of

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2007 for several days of worker interviews, and we had a number of people who came from the 1943 to 1950 time frame. I didn't specifically put that because we followed that up with meetings on June 18th and June 19th with workers from '50 through that time frame.

However, those workers and their interviews are included. We did -- we -- certainly as part of the ER 1 and understood that those existed as part of this ER 2.

The Site Research Database, we currently have a little over 1,000 documents that have been identified as pertinent. These include historical background on process descriptions, Hanford Engineering Work monthly reports, Hanford Instrument Section reports, incident documentation, epidemiological studies, documentation and affidavits supplied by the petitioner; information submitted as part of the Comprehensive Epidemiological Data Resource, the CEDR database. There's an extensive on-line documentation of the -- what they call the Hanford Declassified Document Retrieval System, the DDRS. This has a little over 130,000 documents available. The U.S.

Department of Energy OpenNet System; other documentation from the Department of Energy includes logbooks, radiation survey logs, monthly reports, special work permits. We conducted a number of special reviews of their records to query their databases to see what other records may assist with dose reconstruction and evaluating this Special -- this SEC.

Other sources of information include the REX, the Hanford Radiological Exposure Database. This is -- we also have -- there was a significant publication put together regarding the early years at Herbert Parker Memorial where they collected a lot of the publications from the very earliest times on the radiation protection practices.

Some -- an overview of what -- the claims that we have to date. As of August 9th the same as what we -- as the evaluation report, we had 2,564 claims. Eighteen -- 1,827 of those had dose reconstructions completed. In cases which included internal dosimetry data, 1,919; cases which contained external dosimetry data were 2,370. In addition to those claims, our

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Computer Assisted Telephone Interviews are performed for each dose reconstruction. And also these claims have been put into a database. Though it isn't described here, we actually have summary data that can be evaluated against these.

Major operations for internal exposure included the fuel fabrication facilities for uranium and thorium operations; reactor operations.

Chemical separations obviously were a very large part of the Hanford process. Plutonium finishing, and what -- this means converting from plutonium in a nitrate form to a finished metal product. And also they had separations of americium in some of those facilities. was a major source in the late '60s for heat source development, including promethium, plutonium-238 and polonium-210. Obviously Hanford had conducted many research and development activities, and I put some of the radionuclides here. Certainly it would not encompass all the nuclides that are part of the R&D, but they're plutonium, americium, neptunium, some mixed fission products -- and one of these days I won't keep pushing the

1 button -- and also mixed activation products. 2 Of course all of those generated significant 3 amount of waste which we deal with today in a 4 number of very -- very, very large tank farms 5 and those of course contain mixed fission products, mixed activation products. Of course 6 7 today the long-lived ones only -- plutonium and americium. 8 9 The in-- the internal monitoring information, 10 the bioassay, the analytes and methods change 11 over time as methods improve and capabilities, 12 and also the needs change. Plutonium includes total plutonium, and then later became isotope-13 14 specific. You have plutonium-238, in addition 15 to the -- for spectrometry you measure the 239 16 and 240 at the same time, so that's why you 17 have the plutonium-239 and 240. 18 Americium-241 bioassay by both whole body 19 counting and urinalysis. 20 Uranium total, and also isotopic for 234, 235 21 and 238. 22 Tritium, fission products; strontium-90, it's 23 specifically -- and there are also other 24 radionuclides including curium, promethium,

carbon-14, neptunium, to name a few.

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This internal monitoring also includes hundreds per month, if not larger, thyroid scans, particularly in the very early years. Whole body methods became available in late 1950s and are -- continue through today. Air sampling was conducted at many of the facilities and locations at Hanford as well.

For plutonium, the potential source of exposure basically started in 1945 at most -- at many of the facilities. Urinalysis didn't start until September of 1946. This is one of the main reasons why the SEC -- that first part was granted. Many changes in the plutonium have -- chemistry have occurred over time and counting methods. And since 1983 the plutonium and 239 -- 238 and 239 and 240 have been reported as separate analytes.

The reason I mention this is if you look at the graph and look -- as if they doubled right here. This is actually a double reporting.

The 238 -- they -- some -- not everyone had both, but there's a large increase in 1983.

Probably a very similar level, but we have around 2,000 analyses per month in the REX database. One of the limitations of the REX

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we're going to see on some of the other graphs is that they have a huge physical paper record. Some of that stuff gets into the REX database - we actually get all the hard copy records. Not everything has been entered into the REX database, so this is an underestimate of how many records exist, particularly with respect to the early years.

Principal bioassay methods would have been urinalysis. You also have, of course, chest counting for Americium-241 in the later years. And for some workers, fecal sampling was done. Americium -- and this may be confusing to some people that oftentimes it is a contaminant of the plutonium matrix. It is not a separate product. Usually it's just -- it's something that in-grows with time after the material is irradiated. It's created by plutonium-241; it's a decay product of plutonium-241. However, back in the late 19-- in the early 1950s, beginning in 1949, to -- to support essentially the nuclear chemistry operations of the U.S., they began to separate americium and this operation continued until 1976 when they had a very large glovebox explosion which ended 1 the operations -- americium operations at
2 Hanford.

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Specific activity of americium-241 is about 55 times higher than plutonium-239. Operations -- for recovery or -- included both the 231-Z and later in the 242-Z of the plutonium finishing plant.

For production rates -- we were unable to (unintelligible) production rates throughout all time. We do have some of the very early ones and some of the later ones. The glovebox explosion that occurred in 1976 had over 100 grams of plutonium on a column. Early years may have been the order of a few grams to ten grams per month, but a lot of the intermediate time frames are hard to tell. Again, a much higher specific activity than plutonium. As I'd mentioned, the highest actinide exposure to a U.S. -- in U.S. history occurred as a result of a column explosion in 1976. We have found that no bioassay program prior to 1964 exists, no urinalysis or chest counting methods. In addition to that, these early years -- basically were done in -- in fume hoods, a lot of the separation products, so

1 there certainly is potential for -- for 2 exposure. 3 The 1964 REX database shows only 41 plutonium -4 - americium-241 bioassay measurements. 5 are for 19 workers, and probably are the baseline for the new 242-Z process. 6 7 methods start in 1968 with the availability of 8 the chest counter. 9 These are the number of bioassay records 10 available in the REX database. You can see in 11 1976 they went from doing a few dozen to 12 somewhere on the order of 800, probably in 13 response to this very large accident. 14 In addition to plutonium and americium, you 15 also have some other actinides that -- curium, 16 we had curium-242 and 242 (sic). They 17 conducted separation of 244 curium in the 325 18 building in the 1970s. You also see some of 19 the heavier actinides, including californium 20 and berkelium. 21 Tritium production occurred beginning in around 1949, historically called P-10. Separation of 22 23 tritium occurred in the 108B facility from 1949 24 to '55, and at the PRTR, the Plutonium Recycle

Test Reactor -- Reactor facility from 1960 to

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1990 -- 1969, I apologize. The early methods at Hanford for tritium in urine are described in about 1949. As with many sites, the tritium dose was stored with the external radiation data as part of the whole body dose. Tritium is a -- essentially acts like water and distributes through your whole body so they used the (unintelligible) stored the results as part of your whole body dose.

The tritium bioassay results, the individual analyses, are not included in our record. We do not get those. We get the dose from the site. Those dose data are used -- it's a fairly straightforward calculation and certainly things have changed with time, but they are modified to -- to use the more current biological models as dose methods.

These slides need to be bigger. They get too big a -- I should have had more graph, but this provides some level of -- the graph (unintelligible) urine Table 5-29 provides some level of what the mean dose was for the various years. Essentially we use a coworker study to assign the tritium dose, and so looking at the -- the data, you see from 1955 to 1960 there's

insufficient data. There wasn't -- there was very little tritium monitoring or tritium dose that went on.

Uranium was started of course since the very beginning of the Hanford facilities. It was used to load the -- the fuel cores, and so they began machining that at the very beginning in 1944. You don't have urinalysis until '47. At around 1948 essentially they -- they determined that it's reliable. Before that they had problems with the chemistry.

Presented a number of -- in addition, just the -- the machining -- the fuel separations and also the separation facilities at a number of different facilities that have uranium isotopes. Later we have -- in addition to bioassay methods, we also have in vivo methods. This provides some level of detail. Again, this is always biased in the low -- to the low because not all the early records are in there. You can see that there's -- beginning around 1948, 1949, 1,000 to 4,000 measurements done. And that continues until the late 1960s. Fission and activation products, this is one of the areas where the REX database clearly does

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not have all the data because beginning in 1949, '49, there were thousands of measurements that were done. They're detailed in the -- in the records. These fission and activation bioassays started in 1946. They were considered reliable in '48. They continued to 1965, at which time they were replaced by whole body counting and specific urinalysis for strontium -- strontium-90. So you see some level beginning -- the REX database beginning to have an accurate or -- some level of numbers at around 1958 showing around 5,000 per year. You see that it drops off as the strontium-90specific analyses take over, and this doesn't include the whole body counts. Promethium was another heat source.

that we have on the order of a few hundred samples per year. This is the primary time that it was -- when it was used. They used very large quantities. Heat source are on the order of kilocurie-type levels. Twenty-nine bio samples were known to have been taken, were -- 29 bioassay samples were known to have been taken following an incident in 1963. These do not show up in the REX, but they were detailed

1 in a -- in a report. It was monitored using 2 urinalysis and some fecal sampling. 3 That's not good. 4 (Pause) 5 There we go, polonium. Didn't want to let go; 6 I hit the wrong button. 7 Okay. In the very beginning for polonium, 8 Hanford irradiated canned bismuth and shipped these to Mound. This has been the -- a 9 10 deliberation with the Board bef -- coming to you 11 guys before. They did not process the 12 material. They irradiated bismuth to create polonium-210 and material was shipped for 13 14 processing to Mound laboratories. Later 15 polonium-210 was evaluated as a heat source in the late 1960s. You will see that the bioassay 16 17 for pluton-- for polonium-210 is indicated. 18 1968 we have several hundred polonium bioassay 19 measurements, and sporadically a few samples 20 per year through 1983. 21 In vivo measurements began in 1959. 22 became essentially routine as one of the major 23 operations. Chest counting was begun in 1967 24 for the uranium workers. Thyroid scans were 25 conducted, as I previously mentioned, for the

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workers in the separation canyons. Hundreds of scans per month were -- were conducted. Thorium work started as early as 1946. They mentioned 150 pounds of thorium was brought on site and was machined and -- straight for use in the reactors. Continued at significant levels until 1970. Beginning in 1960 whole body counting methods were -- were available for -- capable of evaluating thorium-232. records of alpha spectrometry results for thorium are also in the record. These next few slides really -- for background levels to provide some evaluation of what kind of photon and beta -- beta-gamma exposure we have available. I don't want to belabor it too long, but just to show some of the distribution. One of the difference at (unintelligible) plutonium finishing plant you see the less-than-30 keV associated with the plutonium handling. Other separation areas you see the -- the high prod-- you see a lot of the high-energy fission product gamma rays. fuel fabrication facilities we see the uranium spectra.

And briefly just some of the neutron areas -- I

don't want to go through these individually -
they certainly include the reactor areas. Many

of the 200 areas have neutron as -- neutrons

present. And of course the 300 and 400 areas

as well.

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External monitoring information -- this is perhaps a bit strong, but essentially the dosimeters assigned to all workers that entered restricted 100, 200 and 300 areas is the documented practice. Certainly there are -- we recognize that there are workers who were not adequately monitored, including construction workers. That's why we use construction worker methods to evaluation those classes. External monitoring methods include, in the very early years, the pencil ionization chamber. Those were of course one of the -used in the very beginning. Later the film dosimeter from '44 to '72 was used. element dosimeter was used from 1944 to March of 1957. Weekly results were included in the individual's cards, and the MDA -- the minimum detectible activity -- I'm going to (unintelligible) would have been -- it would

have been around 30 millirem. From March of

1 1957 to December 31st, '71 they used a multi-2 element film badge. 3 On January 1st, 1972 the thermoluminescent 4 dosimeter was -- began use. Several variations 5 have been used over time. They used the basic 6 TLD for -- assigned to personnel expected to have a low chance of dose from 1972 to 1988. 7 8 They had a multi-purpose TLD from 1972 to '94 9 that was -- actually had two different designs; 10 a five-chip design from 1972 to '77 and again 11 from '83 to '95, and a four-chip design from 12 1977 to 1983. Since 1995 a commercial 13 Harshaw/Bicron system has been in use. 14 Extremity monitoring began in 194-- began in 15 1945 with a simple ring badge; has changed over 16 time from film to TLD; a variety of filter 17 configurations. Wrist dosimetry is also seen 18 in places where hand and forearm exposures are 19 a concern. '46 to '89 when we review the REX 20 database we have about an average of 530 21 workers per year with extremity monitoring 22 results. This is included with the EEOICPA 23 claim information. 24 Neutron monitoring began in 1944 with use of 25 boron-lined PICs. The neutron -- the NTA film

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was used from 1950 to 1971. There's a significant change in '58. From '50 to '58 a two-element neutron dosimeter was used. It was calibrated with polonium-beryllium source, which is a fairly high-energy source. In 1958 a multi-- two changes occurred. began using a multi-element NTA, and they also -- which had a cadmium and tin filter which allowed the monitoring for thermal neutrons, and they also began calibrating with plutonium fluoride, a source of significant concern on plutonium fluoride operations in the Z plant. TLD began use of course in the -- on the 1st of January, 1972. NIOSH agrees that pre-1972 NTA film neutron dose is likely biased low. Dose reconstruction using neutrons is feasible using claimantfavorable neutron-to-photon dose conversion In addition to numerous weekly, monthly, annual and topical reports, we've also obtained at least -- we've also found at least 250 boxes of survey log sheets that have been

identified showing the measurements that

to be -- those had not been retrieved.

occurred in the facilities are actually going

actually in the process of re-- of retrieving those next week.

1957 analysis for the AEC indicated that approximately 70 percent of the dose at the reactors occurred during shutdown. So essentially at the -- at the reactor facilities there's not a neutron present for the -- a presence during the -- when the reactor's non-operational. So using a neutron-to-photon ratio when a reactor is not on is a very claimant-favorable process.

A couple of graphs just to provide some of the 1945 -- as we explored the -- the record at Hanford we found where they had completely mapped the front face of these reactors where these people were working, showing -- it actually shows the total neutron beta-gamma dose across the entire face of this reactor. Something they were clearly concerned with the -- with the dose. In '45 they did not have TLD dosimetry, but they were doing measurements to evaluate the photon and neutron dosimetry. One thing I did want to mention that those were three separate reactors, the B, D and F reactor.

Now I apologize for the quality of this. This is a scanning image, but this is a 1955 -- and there's a -- this is about a 25- or 30-page document that shows a series of neutron and photon measurements that were conducted at numerous different areas within the -- in this case we have the K East reactor -- showing the slow, intermediate and fast neutron flux and dose, in addition to the gamma rays that were obtained.

Other routes of exposure include occupational medical X-ray. They did receive routine medical X-rays. NIOSH has procedures and records available to evaluate this dose. Environmental dose, records and models exist to evaluate the exposure from environmental releases.

And for unmonitored workers although most process workers were monitored, unmonitored workers' dose from external sources may be estimated using coworker methods.

Some specific petition topics that -- that addressed -- that the -- petitioners' issue were the Hanford workers were inadequately or inconsistently monitored. Radiation exposure -

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- overexposure and radiation doses potentially incurred by members of all classes at Hanford were not monitored or consistently monitored through individual monitoring or area monitoring.

NIOSH evaluated -- evaluation findings include -- they reviewed the concern. It's evident from the records available and the published monitoring practices at Hanford that all moni-that not all workers were monitored. However, large amounts of monitoring data exists for Hanford employees, particularly those employees who had the jobs with highest exposure potentials. Gaps in monitoring records for specific employees can be filled in conservatively using available coworker data. Petitioner issues include Hanford construction workers were not monitored for internal dose, and noted a lack of internal monitoring for construction trades between '67 to '71. Construction workers exposed to outside air releases without respiratory protection and that they had limited access to properly functioning respirators.

NIOSH findings include that the -- in addition

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to the previously discussed evaluation -- that being on the previous page; I'm sorry, I didn't make that clear -- that NIOSH reviewed several individual dose reconstructions associated with construction trades from the '67 to time -- to '71 time frames which -- which do show this lack of monitoring data for some workers. Continuing on that, though, that they can be covered with the conservative assumptions and existing coworker data and the application of the construction trade worker OTIB. we also include that we do not take protection factors into account when assessing personal dose under this radiological dose reconstruction program. It eliminates the need to consider or account for the subsequent performance or failure of personal protective equipment for EEOICPA dose reconstructions. Petitions also express a concern for underrecording of neutron dose, especially in the '57 -- the 1950 to '71 time frame. And we concur, as part of our ongoing review of the Board, that the TLD systems had a technological inadequacy for measuring -- for accurately measuring neutron dose and that -- however, we

do believe that a claimant-favorable assignment 1 2 of neutron dose, based on the application of 3 neutron-to-photon dose ratio supported by Hanford field measurements and other monitoring 5 that can be done, including Attila and Monte Carlo methods, can be used for external --6 7 external neutron dose reconstruction. 8 Based on the absence -- however, the 9 feasibility of internal dose reconstruction, we 10 find the absence of bioassay data for the 11 period prior to 1960 for thorium and the period 12 pre-19-- up to 1968 for americium, NIOSH has 13 concluded internal dose reconstruction is not 14 feasible for those radionuclides in selected 15 facilities. And a health -- as part of this 16 two-pronged test, a health endangerment 17 determination is required, and we find that the 18 workers' health may have been endangered due to 19 exposure to thorium and americium exp... 20 Feasibility of external dose reconstruction is 21 that the recorded external dosimetry photon 22 data are extensive and sufficient for external 23 dose reconstruction, especially when coupled 24 with this -- the coworker data that... 25 Turning -- as summary, we find that dose

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reconstruction is not feasible for thorium in the period 1946 to 1959; for americium the period 1949 through 1968. And we find that all external and all other sources -- that internal dosimetry are re-- that dose reconstruction is feasible.

So the recommended class definition, all employees of the DOE, 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 three -- in the following 300 area facilities: the Metal Fabrication Building (313), the Reactor Fuel Manufacturing Pilot Plant (306), and the 300 Area Maintenance Shop and Radiochemistry Laboratory (306); or, two, 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 for a number of work days aggregating at least 250 work days, or in

1 combination with work days within the 2 parameters established for one or more of the 3 other classes of employees of the SEC, 4 excluding ag-- aggregate work -- excluding 5 aggregate work day requirements. Additional information is available to the 6 7 Board of course at the following location. 8 With that, I'll take any questions. 9 DR. ZIEMER: Thank you, Sam. Before we hear 10 from the petitioners, let me ask if any of the 11 Board members have questions for Sam while he's 12 at the microphone, either for clarification or 13 -- or comments. 14 (No responses) 15 If not, we'll hear from the petitioners --16 DR. MELIUS: I have --17 DR. ZIEMER: Oh, Jim, did you have a comment or 18 question? 19 DR. MELIUS: Yeah, I have a question, and I 20 actually e-mailed this to Sam earlier. I guess 21 I'm waiting for an answer. But in -- in the 22 report you referred -- regarding the neutron-23 photon ratio issue, which has been a issue that 24 we've been concerned about in -- in regard to 25 the Hanford site that NIOSH and/or your

contractors were working on some other methods for estimating that ratio, and I was trying to obtain a schedule for -- for that -- that work, mainly in a practical sense so that we have to be able to schedule the work of the workgroup and our contractor for reviewing this and so I was just wondering if you have an update on that now or if we'll obtain that information later, or if you're unable to estimate it at this point in time, which --

DR. GLOVER: I believe I did. Right now we -next week we will be at Hanford. We've
identified 250 boxes of these -- these survey
reports.

DR. MELIUS: Okay.

DR. GLOVER: We have pulled 50 of those in addition to around 35 other boxes we've -- have actually found (unintelligible) to be the original tritium bioassay results and a number of other different documents. So next week Tim Taulbee's actually traveling to Hanford to begin collection of that data, going through that looking for the neutron survey results as part of that.

We'll have a better idea -- we've actually --

1 then of course -- compiled numerous other 2 reports as part of this, so we want to see how 3 we're -- it's not a final thing, but next week 4 we'll actually have a very good idea of where 5 those are going to go. 6 DR. MELIUS: If you could communicate with the 7 workgroup on that, it just would be helpful and 8 -- and when we --9 DR. GLOVER: Yeah, I apologize, I was trying --10 DR. MELIUS: -- clear that up, that's -- I'm 11 just -- we'd like to see (unintelligible) ask 12 it. Thank you. 13 DR. ZIEMER: Okay. Any other questions at this point? Yes, Mr. Presley. 14 15 This is Bob Presley. Sam, on the MR. PRESLEY: 16 uranium slide from -- looks like 1971 to 1984, 17 we had a low level of bioassay analysis per 18 year. Is that from low work level or is that 19 from no records? Can you tell? DR. GLOVER: I -- off the top -- I don't know 20 21 off the top of my head if that was associated 22 with reduced operations, which did happen 23 starting in the '70s. We had a switch from --24 separating reactor-based fuel from Hanford 25 irradiated fuel, so there were some changes.

1 MR. PRESLEY: Thank you. 2 DR. ZIEMER: Thank you. Other questions? 3 (No responses) 4 Okay. Let's then hear from the petitioners. 5 We have Ms. Hoyt on the line. MS. HOYT: I'd like to thank the Board for 6 7 their time and thank Dr. Glover for sending us 8 a copy of his presentation. One of the things 9 that I'd like to start with is the proposed 10 class definition as it is written in the SEC 11 evaluation report. It really is not clear, it 12 just is specifying buildings, and in conversation with Dr. Glover he assured us that 13 14 all employees are included in this class. 15 is not made clear in the body of the report. 16 The fact that all employees are included needs 17 to be made clear to all parties, especially to the Department of Labor. 18 19 Another item, NIOSH has specified in the 20 evaluation report buildings in the 300 area in 21 which thorium was located. At the Advisory 22 Board meeting in July we recall Dr. Ziemer 23 asked why they listed each building. As we 24 recall, the reply was that it covered all of

the 300 area. We contend that the whole area

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is contaminated, not just specific buildings or limited to inside the buildings.

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NIOSH has specified certain buildings in the 200 area in which americium was located. We contend that americium was across the site, not limited to specific buildings or limited to the inside of buildings.

There is an EPA radiation protection program, and I have an excerpt here. It says, quote, People may be directly exposed to gamma radiation from americium-241 by walking on contaminated land. They may also be exposed to both alpha and gamma radiation by breathing in americium-contaminated dust or drinking contaminated water. Living near a weaponstesting or production facility may increase your chance of exposure to americium-241. It is our understanding the findings in the SC&A report still have not been resolved by the Hanford working group. Without resolving all of the findings, there cannot be a defensible claimant-favorable evaluation of any petition. We dispute the fact that NIOSH claims that SEC-57-1 and 57-2 that external dose reconstruction is feasible.

1 The evaluation report states, quote, All 2 interviewees indicated that employees who 3 entered radiologically-controlled areas wore 4 external dose monitoring devices, end quote. 5 We question whether or not the interviews were conducted in accordance with the SC&A

quidelines.

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Also, at the worker outreach meetings former employees stated that not everybody wore monitoring devices. They told how they would pick up monitoring devices at the buildings, and at the end of the shift they would throw them all in a bucket. There is no monitoring during transportation through the areas. Buses drove through noxious vapors and yellow clouds. Former workers stated that they wore monitoring devices under various layers of clothing and protective gear, and the monitoring devices were not on the areas of the body that were exposed.

One of the rad techs, people that do radiation monitoring out there, stated that the -- at one of the worker outreach meetings he said that records of his personal exposure incidents were not accurate.

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The transcripts for the June and July meetings are still unavailable. This is burdening and -- this burden-- this is burdensome to all involved and hinders the process. Lack of funding and workload assignments are no excuse for not producing these in a timely manner. A publication entitled "Hanford Site Occupational Internal Dose, ORAUT (unintelligible) 00-5", there are serious flaws in this document. The review of the NIOSH/ORAU procedure and method used for dose reconstruction dated January 17th, 19-- or 2005 by SC&A report states that both the internal and external are deficient. SC&A's review of these procedures identifies a number of technical inaccuracies and errors. The evaluation report states in general information obtained through the interviews with former employees and facility experts was consistent with that found in NIOSH documents regarding the Hanford facility. We ask which experts are you referring to. There's a lot of credibility given to interviewed experts, and they are referenced repeatedly. Nowhere in this evaluation report does it deal with the

1 affidavits of falsification of records. 2 are affidavits stating monitoring records were 3 falsified, supervisors coerced employees to 4 change records or be sent home without pay. 5 There was a -- another meeting at Hanford and which NIOSH attended and the publication was 6 7 NIOSH dose reconstruction project meeting at 8 Hanford atomic metal trades council 9 (unintelligible) dated January 13th, 2004. On 10 page 4 of this document, quote, 11 (unintelligible), before good readings were 12 kept, a lot of people were exposed due to 13 fooling with exposure to get overtime. People needed exposure time to make the money they 14 15 wanted. In the '90s Navy came in and things 16 improved, but many people are gone, end quote. 17 That is another -- this also shows that it's 18 been common knowledge that the records have 19 been falsified and are not accurate. 20 The evaluation report states current and past 21 Hanford workers have access to their records at 22 any time upon request, end quote. We have no 23 confidence in this statement. This is a 24 prevailing concern. 25 An excerpt from the national advocate's call

dated August 16th, 2007. Caller, raise the question if we are sure that DOE is giving NIOSH and DOL all the documents in their position. Another caller said the program lacks the necessary transparency, and suggested that perhaps Sanford Cohen & Associates, through the Advocacy Board -- or through the Advisory Board, could be tasked with quality assurance of DOE records. I blanked out the name to protect those people's privacy. If the Board needs those names specifically, I can give them to them.

Jack J. Fix is mentioned in many of the documents regarding Hanford. He has a conflict of interest, having been the project manager/principal investigator for the Hanford external dosimetry problem -- program from 1979 to 1995. He is still active with a contractor to NIOSH/ORAU, which is Dade Moeller & Associates.

It appears NIOSH continues to churn out dose reconstructions to crunch the numbers -- so many were submitted, so many were reviewed, so many were approved. NIOSH states that they are under-funded, so they can't get the transcripts

of the meetings out. But they can continue to do dose reconstruction, even though there are serious deficiencies, inaccuracies and errors.

Neutron exposure continues -- neutron exposure dose reconstruction continues to be an area of unresolved findings with SC&A. This evaluation report is confusing, unorganized and does not address lost or destroyed records or affidavits supporting the SEC petition 57 in all of its forms.

Thank you.

DR. ZIEMER: Thank you very much, Ms. Hoyt. Is

Ms. Carrico also wishing to make a statement?

(No responses)

Apparently not. Okay, thank you very much.

I -- I do want to remind the Board that earlier this year we did take action on a petition from Hanford that covered the earlier years, '42 to '46, and you ma-- I just want you to have that in the back of your mind. This particular evaluation report most Board members just got within the last few weeks. It's a fairly extensive report, fairly complex. We earlier had tasked our contractor to review this report. That review is just barely underway.

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I'd like to call on the chair of the working group -- the Hanford working group, Jim Melius, if you could give us a status report and kind of outline the path forward from this point for the Hanford petition.

DR. MELIUS: Yeah. First can I ask -- I think it's a question for -- for Larry 'cause -- an issue that the petitioner brought up and I wanted to ask it also is -- is the issue that the process for attribution in the reports. As I recall, NIOSH was committed through their contractor to go through all the site profile reports and provide attribution of -- you know, sources for various information and so forth. I believe that was done for the Rocky Flats report and I'm -- I'm just curious what the status of that is for the Hanford report, at least on the -- I confess I'm just looking at it on the -- the web site now and it's in bits and pieces and different time frames, so I'm trying to get a sense of when that will be complete for that -- the Hanford report.

MR. ELLIOTT: I believe -- I believe it's complete.

DR. MELIUS: Okay.

1	MR. ELLIOTT: Hanford reports that we have
2	posted there have been fully annotated and
3	attributed.
4	DR. MELIUS: Some of it goes back to 2004,
5	Larry. That's all I it's in sections.
6	That's why I was
7	MR. ELLIOTT: Oh, the evaluation report is
8	fully attributed and annotated.
9	DR. MELIUS: You know I knew that. No, I was
10	asking about the site profile.
11	MR. ELLIOTT: Oh, we'll have to check on that.
12	I know
13	DR. MELIUS: Okay, if you could just
14	MR. ELLIOTT: that the site profile chapters
15	are all fully annotated.
16	DR. MELIUS: Okay.
17	MR. ELLIOTT: I believe that there or a
18	recent one that should have been.
19	DR. MELIUS: Yeah.
20	MR. ELLIOTT: But I don't know that the earlier
21	ones would have been. I'll check on that.
22	DR. MELIUS: And and given that I mean
23	one is the petitioner's obviously raised the
24	issue, but it's it's also that if we're
25	in the evaluation report you're referring back

to the site profile report a lot. I just think it's helpful and there was a commitment to do that. I understand it's time and effort and so forth, but for something like this that we'll be reviewing, I think it --

MR. ELLIOTT: Yes.

DR. MELIUS: -- it's helpful and makes the
process more transparent.

MR. ELLIOTT: We'll make sure we report to the working group on that.

DR. MELIUS: Okay, I appreciate that. Yes, the

-- had some -- some discussions this morning,
and actually prior to this meeting I actually
- once we -- I received this report, not only
was checking with -- with Sam about some sort
of logistical issues, but also with John

Morowitz (sic) and Arjun regarding the review
of this eval-- evaluation report and based on
- on those discussion -- we think this is going
to be a -- one, it's a large task. We were -had started to do the site profile review and
it -- and the major issue there was the -- a
major issue was the neutron/photon ratio issue
and that point, which is several months, if not
a year ago, NIOSH was then starting to -- to do

a revision of the -- the methodology for that, which I think, as Sam has indicated, is in some ways still -- still under-- underway in terms of gathering additional information and -- and so forth.

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What we thought would be the -- this -- a way forward for this task, in order to try to, one, expedite what we can 'cause petitioners and claimants are -- are waiting on this, at the same time, given the scope of this report -- it covers a lot of years over a very large facility where I think NIOSH's work is still some extent under-- underway and so we may have a -- so to speak, a moving target to -- to evaluate. What we thought we'd do, and I'd be interested in feedback on this, is task the -our contractor with first initially doing initial scoping effort on the evaluation report to identify key issues that they identified from an initial review of the report. We would then hold a meeting of the workgroup, probably by conference call, to then mutually determine a schedule for that review. And rather than trying to do a complete review of the evaluation report -- you know, deliver it at

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one point in time -- we would just deliver it in -- incrementally in sections by issues in a way that would hopefully expedite the -- the review process, make it easier to -- to handle and easier to do the -- the workgroup meetings and so forth to try to re-- resolve comments and -- and reach some conclusions on that. I believe we can start that process within the next month or so with a -- with a con-conference call meeting of the -- the workgroup. Initially a rep-- refers to initially a report coming from SC&A that would scope out the issues and give us some idea of where they thought -- according to review. same time as part of that scoping process, that would also give NIOSH an opportunity to give us feedback on where they were with any parts of the evaluation report or the site profile that they were updating. So rather than, you know, waste our efforts and time and money and so forth on reviewing something that was already being revised, we could wait till the revision is done if that's not going to delay things in-- inappropriately. But at the same time I think we have to recognize that this will not

be a -- a quick process to resolve the full evaluation report. Again, we may be able to break off parts of -- and so forth and look -- look at it that way, so I would appreciate any -- any feedback on that suggestion or -- or comments. Thank you.

DR. ZIEMER: Certainly seems like a logical approach. Board members, you have any comments on this for Jim and the working group?

And I might add, while you're thinking about your comments, that as we move forward in the proposed manner that the petitioners would be kept fully informed of all of the issues and invited to participate with the workgroup on -- on these various issues.

DR. MELIUS: Yeah, ab-- absolutely and -- and I think it would also make it better for them to be able to, you know, focus, you know, on a particular issue as we go forward so they would be able to provide whatever additional information and if we need to seek out additional information from people wor-- who've worked at the site and can provide information to us, that would facilitate that process and -- rather than putting out a broad call for all

1 the information and expect people to completely 2 understand this very, you know, lengthy and 3 complicated report. 4 DR. WADE: Dr. Poston. 5 DR. ZIEMER: Dr. Poston? 6 DR. POSTON: I have two general comments. 7 First, even though the report is dated the 14th 8 of September, my recollection is we just 9 received a review this week -- this past week 10 and so I don't know about the rest of the 11 Board, but I haven't had a chance to read that 12 and digest it, so we need time. 13 Secondly, if we're going to move ahead, and it 14 sounds like Jim has a good plan, it seems to me 15 that we need to do a better job with this 16 privacy clearance. It's been almost four 17 months since that meeting and the information 18 has not been released to the petitioners, and I 19 think that is absurd in terms of delaying 20 getting the information out. So I would urge 21 whoever has the stick there to get these things 22 done in a more expeditious manner. 23 DR. WADE: That's my task. Tomorrow I'll be 24 providing you with a matrix that updates all of 25 the transcripts and all of their status, and we

can look at that and we can focus our efforts and we can see what the reality is. But I do understand that as a -- is a problem.

DR. ZIEMER: And thank you for your comment, John. And indeed most Board members have not had a chance to digest this report, and for that reason we're not in a position to take action on the report. Obviously the workgroup has to get engaged with it, as does the -- the -- our contractor. So this basically, although we have a recommendation from the -- from the agency, from NIOSH, a recommendation on the petition, it certainly appears to the Chair that we're not in a position to act on that other than to agree that we will continue to study their report. We will garner the information from our contractor in the manner as described by the workgroup chair and proceed on that basis. And I'm going to take it by consent that that's what we'll do unless I hear strong objections to that.

If not, let me ask if the petitioners have any additional comments, having heard from the chair of the working group. Ms. Hoyt, if you're still there, do you have any additional

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comments for us at this point?

MS. HOYT: I think that it sounds like a good plan and we thank the Board for keeping us informed and asking for our input. We would appreciate that.

UNIDENTIFIED: We like working with the working
group. That would be a very effective thing.

DR. ZIEMER: Thank you. Then we will -- we will keep you informed and in the loop as we move forward on this.

(Pause)

BLIND REVIEW CASES WORKGROUP

Okay, before we take the break I'm going to use a few moments here to take care of a kind of a housekeeping matter, and that is I'm proposing to -- on advice from counsel, actually -- to appoint a new workgroup which shall be known as the workgroup on the selection of blind re-- dose reconstruction blind review cases. This workgroup will have the express task, which I'm expecting them to complete by tomorrow, of receiving from the Subcommittee on Dose Reconstructions suggestions on the list of proposed blind reviews that I believe has been distributed to the subcommittee. I'm -- I

would like Mark Griffon to chair the new 1 2 workgroup and Wanda Munn to be the other 3 member. We only ne-- it's a small workgroup. Your task would be to solicit from the members 4 5 of the subcommittee their recommendations on the cases for the blind review to come up with 6 7 a final recommendation for the Board on those 8 cases. And we'll hopeful that you will have 9 your report ready for this Board tomorrow, at 10 which point we will dissolve the working group. 11 It's my understanding that doing it this way 12 will meet our legal requirements as far as 13 confidentiality and other related matters, and 14 will allow the selection to move forward. So 15 that workgroup is hereby appointed. 16 Now let me ask for questions. 17 MR. GRIFFON: Just one -- one question. 18 said the subcommittee members -- are we 19 including alternates? I -- they were --20 everybody was here yesterday, kind of like --21 DR. ZIEMER: Sure. 22 MR. GRIFFON: And I don't know that Bob and --23 DR. ZIEMER: The alternates are members of the 24 subcommittee. 25 MR. GRIFFON: Right, and I don't know that --

1	Bob, did you get the list? I'm not sure
2	maybe
3	MR. PRESLEY: (Off microphone) (Unintelligible)
4	MR. GRIFFON: Oh, okay. Oh, yeah, you gave me
5	your suggestions.
6	MR. PRESLEY: (Off microphone) (Unintelligible)
7	MR. GRIFFON: Yeah, that's right.
8	DR. ZIEMER: In anticipation of the workgroup,
9	he has given you his suggestions. Okay.
10	And let me ask counsel I want to be assured
11	that this this will meet our legal
12	requirements as far as gathering the
13	information by this workgroup and making a
14	recommendation.
15	MS. HOMOKI-TITUS: Yes. I just want to be
16	clear that the suggestions that are being
17	provided are being provided by individuals, not
18	as a group recommendation.
19	DR. ZIEMER: The recommendations to Mark will
20	be provided individually by various Board
21	members, without collaboration with each other,
22	yes.
23	Any other questions on this matter?
24	(No responses)
25	Okay, thank you. It is so ordered.

While we are -- I'll take a further couple of minutes -- well, I -- oh, are we past our break time?

DR. WADE: No.

DR. ZIEMER: Well, one -- one -- this is -- be very rapid. Two of our newest members now have reached a point where they feel like their -- their time and abilities are not being fully utilized, and they have actually volunteered to participate in some additional workgroup activities. And with that in mind, I'd like to add Phil Schofield's name to the workgroup on the Nevada Test Site and Savannah River Site and the chair -- chairs of those groups can make a note and ask Dr. Lewis (sic) to add them --

DR. WADE: Nevada Test Site --

DR. ZIEMER: Nevada Test Site site profile workgroup and the Savannah River Site workgroup. And then we'll -- we'd like to add Josie Beach to the SEC issues group, Dr. Melius. So -- and those appointments the Chair's authorized to make and I so make them. Yeah, we'll add added manpower and womenpower to those workgroups. Thank you very much.

1 We'll now take a break, and when we return we 2 will proceed with the Sandia Livermore 3 petition. 4 UNIDENTIFIED: 11:15? 5 **DR. WADE:** 11:15. 6 DR. ZIEMER: 11:15. 7 (Whereupon, a recess was taken from 10:45 a.m. 8 to 11:15 a.m.) 9 DR. ZIEMER: Okay, I'd like to call the meeting 10 back into session. We're going to have a brief 11 comment from either Larry or Kate from ORAU. 12 Kate, are you -- this -- this is in answer to 13 Dr. Melius's question on attribution, so I 14 think we have the latest update on attribution of the Hanford material. Kate, ORAU, thank 15 16 you. 17 MS. KIMPAN: Thank you. I need the short 18 microphone. Hi, this is Kate Kimpan from ORAU. 19 It's a pleasure to see you all and actually a 20 pleasure to respond to this question which I 21 believe, Dr. Melius, if I heard it right via the phone part of the Board, you asked about 22 23 the Hanford TBD and whether A&A was complete on 24 that.

Right.

DR. MELIUS:

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MS. KIMPAN: The Hanford TBD, as with many of our large TBDs, has multiple sections. All sections but the medical, which is the smallest of them, has been fully completed. The medical section is still in the resolving of comments and questions and so it has not been signed by OCAS. When we complete our review, of course, we are only doing this on behalf of OCAS. They have final say. So when I say completed, I'm typically talking about what's been done, blessed, signed and posted onto the web site. The medical portion is not yet.

Regarding this issue, because it was a -- a very -- it is a very important issue and I spoke about it at every meeting, if -- if -- if you'd like, I can give you a one-minute fuller update on an-- annotation and attribution.

DR. ZIEMER: Go ahead.

MS. KIMPAN: I know that -- let me get this (off microphone) (unintelligible).

(On microphone) Conflict or bias, conflict of interest has been an absolutely important issue to this group and you've been concerned. I wanted you to know that I've spoken about what these types were before. When the new policy

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came into force, part of the ORAU team's compliance was to do a comprehensive review, as required by the policy, of our documents. We then did full annotation and attribution on any document where the original document owner would have been conflicted under the policy that didn't exist at the time. So we took the new policy, the current one; we looked through that lens, in an abundance of caution, and looked at the places where an owner would have been conflicted. We did those first for full annotation and attribution. I will tell you we included a couple of sites that we've much talked about at this table where there wasn't an actual conflict of interest, but where questions were raised -- the Paducah TBD, questions were raised and at the end of the day the lawyers and the -- the legal was that there wasn't really a conflict or bias problem there, but there were adequate questions raised, we included it in that first run of reviews. We are in full compliance, the ORAU team is, with the policy as it is written right now. And we can obviously continue to remain in compliance with that. Our new required

postings are all listed on our web.

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For those of you -- and I know this group will be among them -- in order to not have a lot of confusing things on the web, if someone does not work for the ORAU team now, their conflict or bias information is removed. As you're going through documents -- we obviously don't go back and take a document that was written four years ago and say it wasn't written by who's -- who was on the document. If you're ever looking through these documents and you find a name and you go to the web site and you don't see their information, we have retained all of that information. It is both available -- it's available to anyone, including the public, but we don't have it up on the web because it would be bad information. might have left my employ three years ago and the information wouldn't be current. For my employees, for everyone who works with and for ORAU team, our info is current. But if you see a name on a document from the first day of this program that you're not finding their information, let me know through OCAS and we'd be more than glad to provide you with those

1 historical conflict of interest declarations by 2 those folks. There's -- there's no intent to 3 remove that from you all seeing it. It's just 4 having in on our web site now, with inaccurate 5 information, just isn't the right thing to do. Any questions or other concerns? 6 7 DR. MELIUS: Yeah. Thank you. Okay, other question? 8 DR. ZIEMER: 9 Go ahead. 10 DR. MELIUS: Yeah, I -- I don't want to prolong 11 this neces-- but I would ask you to reconsider 12 that last policy because I think it's -- for 13 those of us on the Board who have those 14 questions, we've heard you and we know --15 MS. KIMPAN: Yes. 16 DR. MELIUS: -- how to pursue that. But for --17 somebody on the outside will see something with, you know, Joe Smith's name on it and have 18 19 no way of, you know, finding out about that 20 person's background at the time. And I suspect 21 that they'll end up having certain sections of 22 the report attributed to them as a source --23 MS. KIMPAN: Yes. 24 DR. MELIUS: -- and I think it's important that 25 people still be able to get that information.

1 And I would think that their conflicts of 2 interest or bias, whatever, statements at the 3 time, when they were in your employ, would 4 still be relevant 'cause that would be their 5 conflicts when they wrote the document. 6 I mean I think they should be properly 7 caveated, this is not up to date, so that if 8 somebody has a question about what Joe Smith's 9 done in the last four years, it's not on there 10 11 MS. KIMPAN: That's right. 12 DR. MELIUS: -- but -- but it might be just a 13 little bit more -- in terms of transparency to 14 have that available such -- you do have an area 15 on -- on the web site where old documents are, older versions of documents --16 17 MS. KIMPAN: Yes. DR. MELIUS: -- are -- are referenced, and I 18 19 think if there was some link there, some way of doing that -- so I'd ask you to consider doing 20 21 that. I don't need an answer now, I don't --22 MS. KIMPAN: Thank you for that suggestion --23 DR. MELIUS: -- but I think it'd be feasible... 24 MS. KIMPAN: -- Dr. Melius. I will tell you 25 what's happened several times, for what it's

worth. Now it's mostly people who have a web access where either Larry or myself will get those questions, so there are folks that have asked that are getting those answers. But you're right, we certainly want to make those - - that information is intended to be publicly available. We're just not trying to create any confusion as we proceed.

DR. MELIUS: No, I -- I appreciate that and I would just add that -- that -- I know we're having an update on the web site I believe tomorrow. But the web site is confusing to navigate, at least up until now. It's improving and it -- in some ways, but -- but I think not everyone will know where -- knows where to look and -- and --

MS. KIMPAN: Right.

DR. MELIUS: -- so forth. I think it's hel-the more we can do to link things and have a
complete information there, the better, 'cause
I think it's probably the best way of -- of
making this information available and it's the
least burdensome to NIOSH and ORAU in terms of,
you know, people having to request things and
so forth, so...

1 MS. KIMPAN: Very good suggestion, Dr. Melius. 2 DR. ZIEMER: Okay. Thank you, Kate. 3 DR. WADE: Always a pleasure. 4 SANDIA NATIONAL LABORATORY-LIVERMORE SEC PETITION DR. ZIEMER: Okay, let's proceed with the SEC 5 6 petition evaluation report for the Sandia 7 National Laboratory Livermore. Sam Glover is 8 going to present that. One -- the petitioner, 9 Gerald Giovacchini, if I've pronounced his name 10 correctly -- if not, forgive me, Gerald -- but 11 are you on the line? 12 (Pause) 13 Yes, he is on the line. After Dr. Glover gives 14 his report, Gerald, we'll give you an 15 opportunity to make your comments and then 16 proceed from there. Thank you. Dr. Glover. 17 DR. GLOVER: Okay, now it -- can we -- I've got this back on. Is it working okay? All right. 18 19 Now I've got no excuse not to properly use the 20 equipment, so my laser and my clicker. 21 All right. I'm going to pre-- this is an 22 update to a presentation that we did in May of 23 2007. It is SEC Petition 59 and it deals with

X-ray diffraction units at the Sandia Livermore

-- Sandia National Lab Livermore facility.

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1 A little bit -- this will be some repetitive 2 information -- some people weren't here, but 3 since it has been four or five months since the 4 presentation, I thought I'd go ahead and 5 reiterate those slides. 6 Site history, Sandia Lab Livermore was 7 established in 1956. It was to provide 8 assistance to the Lawrence Livermore National 9 Lab regarding nuclear weapons design. 10 primary mission from '56 to '89 was design and 11 testing of non-nuclear components for 12 Livermore. 13 A little bit about the petition. On May 5th, 14 2006 a petition was submitted to NIOSH on 15 behalf of a class of employees who included all 16 X-ray technologists and materials scientists 17 who worked in the X-ray Diffraction and 18 Fluorescence Laboratories in the buildings 913, 19 room 113, room 128, and in building 941, room 20 128, from the period December 1st, 1967 through 21 December 31st, 1990. 22 On October 4th, 2006 the petition was 23 qualified. On March -- March the 29th, 2007 24 evaluation report was issued. Immediately 25 before the meeting, on April 25th, NIOSH

1 received new addition -- new information from 2 the petitioner. On May 2007 the evaluation 3 report and new information was also provided by 4 the petitioner at the May 2007 Advisory Board 5 meeting. At that time the Advisory Board asked NIOSH to provide an update that addressed the 6 7 new information. And September 6, 2007 an 8 addendum to the evaluation report was approved 9 -- or was issued, is probably the most correct 10 term. 11 Briefly, the evaluated class included -- I was 12 -- it was modified by roo-- by removing 13 Building 940 room -- 941, room 128, because it 14 occurred after -- outside the period that was 15 covered, 1992. NIOSH evaluated the following 16 class: All X-ray technologists and material 17 scientists who worked at Sandia National Lab 18 Livermore in those buildings and rooms so 19 specified from December 1st, 1967 to December 20 31st, 1990. 21 I want to be clear. This is a very small 22 class, approximately three people. 23 The information was provided but I did want to 24 update it. At the time of this we had a draft 25 site profile. This was actually at a -- an

official Rev. 0 was actually issued after the initial prep-- preparation of the material. It was issued May 1st, 2007. A number of Technical Information Bulletins was -- were evaluated as part of this, including maximum internal dose estimates for DOE claims, dose reconstruction from occupations related to X-rays, and also internal dose reconstruction, OTIB-60.

As I said, this is a very small class. Cases which meet the class definition -- these are cases that have been submitted to NIOSH -- are one. Dose reconstructions which are completed are zero. Dose re-- cases which include internal dosimetry and external dosimetry, we had information for both. Of course the Computer-Assisted Telephone Interview was conducted as part of this.

So petition provi-- the petitioner provided a letter April 25th which was read at the Board meeting. On June 7th a follow-up call was conducted with the petitioner. On July 16th a petitioner letter and an affidavit were also received. September 11th an additional letter from the petitioner and an affidavit from a

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health physicist/industrial hygienist at Sandia National Lab Livermore was also received. This was received after the issuance of the evaluation report, but it was evaluated prior to the -- correct me -- as preparation for this presentation.

Several other sources of information include CD-- the Centers for Disease Control web site, cutaneous radiation injury, facts for physicians and radiation emergencies. basically to show -- provides information (unintelligible) effects caused by acute incidents of X-ray exposures to the skin. We -- and in preparation for this we also had a physician evaluate the -- the lymph nodes associated with this extremity dose. another report was the U.S. Department of Health, Education and Welfare radiation safety in X-ray diffraction and spectroscopy report. September 6th evaluation findings of the petition were submitted as an addendum to the SEC report.

Just to reiterate what the basis was is that -the basis was that unmonitored, unrecorded or
inadequately monitored exposure incidents

occurred. They associated with an incident that occurred in 1978 that was alleged by the petitioner and others that was documented in 1979 that was an accident with the beam turned on and at low power. This petition provided evidence that potential unmonitored exposure with no personal or area monitoring data for the first exposure incident. Further, SNL did not provide -- Livermore did not provide permanently mounted instrumentation for continuous recording of the ionaz-ionizing radiation that was being emitted. statement by the petitioner about the -- the type of instrumentation that was used. A little bit about the radiological operations. X-ray diffraction -- as we discussed, XRD is a very high-dose possibility, on the order of tens of thousands of rads per minute. very -- it's like a laser beam, essentially. It's a very small beam that's used to evaluate samples, so the -- very high, intense radiation source in the lo-- in a -- in a very localized area. And -- and fluorescence laboratories

1 preparation of samples and testing with this X-2 ray diffraction and fluorescence equipment. 3 Radioactive sources include depleted uranium, 4 small sealed sources, and obviously this X-ray-5 generating equipment. These were issues discussed with the 6 7 petitioner. There's approximately 14 points. 8 I'm just going to briefly discuss what they are 9 without providing you a great deal of detail. 10 These are addressed in the report. 11 discussed -- these are the points that were 12 provided, that personal monitoring records that are unavailable. However the class records are 13 14 available. That the directional nature of the 15 X-ray radiation emitted from the unit was --16 was outside the monitoring device or the badge 17 -- it wasn't in a badged area. That workers 18 devised makeshift shielding because the shields 19 could not be used for oversized samples. 20 The unrecorded exposure incidents associated 21 with the operation of the X-ray unit, 22 specifically a 1978 undocumented exposure and a 23 -- and a documented exposure incident in 1979. 24 The ability to bound exposures, that we were 25 unable to -- he felt we were unable to bound

1 the exposures as described using the 2 information that was available. He also 3 discussed differences in the workload among the 4 potential class, and impact of this difference 5 on the ability to reconstruct dose. Information was discussed regarding the use of 6 7 sealed sources and the preparation of samples, 8 essentially using a mortar and pestle 9 (unintelligible) homogenize or to have these 10 samples (unintelligible) specifically prepared 11 and the exposures that were part of that. 12 provided statements and discussed by --13 statements made by two doctors that ex--14 exposures resulted in cancer for the petitioner 15 and inappropriateness or inadequacies 16 associated with the risk models for radioactive 17 material exposures and the determination of 18 probability of causation. 19 So exposure data was forwarded to petitioner in 20 June 2007, and actually additional information 21 -- which was recently provided by the site --22 was forwarded, I believe within the last week, some later post-1990, I believe tritium and 23 24 uranium data. And so this was not pertinent to 25 the work that he performed in the X-ray

So

1 diffraction lab. Concerns about attempting to 2 reconstruct dose without ascertaining the 3 predominant energy of the X-ray beam -- one of 4 the issues with the beam is that they're 9 keV 5 copper X-ray, so it is an extremely low-energy 6 X-ray that's not very penetrating. 7 He expressed concern about the security badge 8 location in relationship to the dosimeter and 9 the shielding of the dosimeter by the security 10 badge. 11 Again, I believe we've already discussed the 12 use of copper X-ray target, and sometimes iron, 13 but not at the time of the major exposures. 14 That the dosimeter -- he also believed the 15 dosimeter used at the time would not provide a 16 valid account of the radiation dose, and the 17 lack of specific monitoring data, either 18 personal or area, prevents the adequate 19 reconstruction of dose. 20 So let's maybe talk about what we do have. 21 we do have bioassay data from all potential 22 members of the class have uranium bioassay. 23 External data for the class are available for 24 whole body badge dose. Sandia National Lab did 25 not do extremity monitoring till after 1990.

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Incident information includes a report -- let's see, shallow dose to the extremity was not recorded in the dose of record, and it was calculated from reports from the incident. So as part of this supplement I want to talk a little bit about what the bounding is. is an extremely large -- the dose -- dose rate. Essentially, deterministic effects bound the dose. If you don't see blistering -- this is actually an X-- an X-ray diffraction accident that occurred and these are the deterministic effects that oc-- occurred after -- as a function of time. You see 24, day 30, 64, day This kind of X-ray dose rate causes extreme deterministic effects, and so that the external dose is bounded by these deterministic effects and doses in the range of 15 to 40 Grey result in reddening of the skin, skin des-skin desquamation and blistering; that those effects were not observed and therefore deterministic effects would bound the dose. The report in some cases wasn't very clear on some of the calculations. I provided them, just to make it a little simpler on some of the math, where some of the numbers may have come

from but I won't belabor the issue here. The shallow dose came from a 1979 incident report, essentially 30 rads per 20 seconds. This was done at a lower power and a lower current, and so there are equations to scale that up to full power. The 1970 dose was for a full instrument done at 40 kilovolts and at 20 milliamps. So essentially the time that would be required is somewhere between the order of 32 seconds to 85 seconds to produce this 15 Grey to 40 Grey exposure that would have resulted in deterministic effects.

So we normalize the values. We took them from

So we normalize the values. We took them from the measured re-- the measured results at the accident. We brought those up to full operating values. Those were in -- in R or exposure, and then they were converted to dose rates for the shallow dose and also deep dose. So what can we do? We actually can do -- evaluate the direct beam exposure to the organs via the diffracted dose. The dose is added annually to XRD aper-- operators in addition to the missed dose. Based on the measured beam exposure of the 1979 Sandia incident report and scaled up to full operational power, this

1 results in a direct beam dose on the order of 2 1.96 times 10 to the 5th R per hour. 3 the unit of exposure. 4 Data show the diffracted dose is approximately 5 3.3 times 10 to the minus 6 of the direct beam. Okay? This is diffracted off the sample as it 6 7 comes to the shield. This is what you're going 8 to see in your body. 9 A health physics report written in the late --10 in the early '70s actually measures the 11 (unintelligible) instrument and what its 12 diffracted dose is. We came up with about .65 13 R per hour. That report determines something 14 on the order of .35 R per hour, so our number 15 is conservative, meaning it is higher than --16 yes, sir? 17 DR. POSTON: Sam, I don't like numbers without 18 What's the unit on the 3.3? 19 though you said diffracted dose, I still need 20 to know the units. 21 DR. GLOVER: That's just the -- it's a -- it's 22 a fraction. This is diffracted dose relative 23 to the direct beam dose, and they measured the 24 diffracted measure and this is its relative 25 value. It's just a ratio of -- of --

1 UNIDENTIFIED: (Unintelligible) 2 DR. POSTON: So it's not the diffracted dose. 3 DR. GLOVER: I'm sorry? 4 DR. ROESSLER: It's not a dose. 5 MR. GRIFFON: The parentheses are the 6 diffracted dose, the -- the (unintelligible). 7 DR. GLOVER: The direct beam dose is -- isn't -8 - is -- I expect the laser to be on the bottom 9 part; this should be the trigger. This 3.3 10 times 10 to the minus 6 times this results in a 11 .65 R per hour number, and I apologize for not 12 making that clear. 13 All right. So we're -- I hit too many buttons at the same time, apparently. Maybe if I just 14 do this. 15 16 All right. One of these is -- again, this is a 17 small beam so correction for time spent in the 18 beam based on a 10 centimeters squared beam, 19 upper front torso, approximately 25 percent of 20 the sk-- the skin exposed, the total skin area 21 is about 4,500 square centimeters. 22 Finally, exposure is multiplied by the organ 23 dose correction factor, so that R value needs 24 to be corrected to dose, so it's in Roentgens, 25 now we need to come up with rem or millirem.

1 And also then, using a 50 percent occupancy 2 factor, the instrument would have been used 3 roughly half the 1,000 hours per year, to 4 determine the organ dose. 5 If you do that, you come up with around -- a lymphatic dose around .08 rem per year and a 6 7 skin dose of about 1.25 rem per year, and the 8 re port discusses other organs and I -- but 9 they are very low and those are included in the 10 addendum. 11 Sorry about that, this... I don't know how to 12 make it stop. Did it go away? All right. 13 There's too many buttons -- too many options. 14 I need better training. I need to be... 15 All right, so it's -- uranium exposure can be 16 reconstructed using actual bioassay data from 17 missed dose and so that's a pretty well-18 established discussion. I'm not going to 19 belabor that. 20 External deep dose can be restructed (sic) from 21 the reported dosimetry results. We're going to 22 use that -- whatever reported dosimetry results 23 would be -- in addition to the missed dose that 24 would be occurred from that standard NIOSH 25 practice.

The shallow dose can also be conre-reconstructed based on the actual reported
dosimetry, supplemented with what we talked
about a little bit ago, and so this is just the
-- if you had a shallow dose, we would still
give you a missed dose for the -- for material
that you would have -- that would have worked
with. That's not going to include that 9 keV
scattered X-ray.

So in summary with that -- I want you -- that the internal source of exposure included the depleted uranium, the deep from mixed sources from the badge, the shallow dose that's badged, this assigned diffracted beam dose -- that missed dose that we talked about that -- if -- depending on the organ if it's the 1.25 rem per year or if it's a skin -- assigning extremity dose as appropriate. If you report that you were in an incident, then you would have an assigned extremity dose based on -- basically saying that you were part of this incident. There were no neutron sources that we're aware of.

I don't want to belabor each of these. They're available to the Board. Basically we used a

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consistent -- consistent with the previous set of dose reconstructions that were done, a male; date of birth, 1932; when -- the date of diagnosis and some -- varying the -- some of the different -- employment's determined, what is the effect of the addi -- additional dose. Essentially what we have is an XRD operator involved in the 1978 incident, cancer located in the beam during the incident -- located in the beam, and also the uranium bioassay. it's a basal cell carcinoma on the hand, this is an underestimate using a very small fraction of the dose, you're going to exceed the probability of causation of 50 percent. An XRD operator if the cancer's not located in the beam, uranium bioassay -- it's using a couple of different things -- of the lung, we received a 29.74 percent and a basal cell carcinoma on the chest around 30.51 percent. Obviously those change, depending on the exact circumstances that you would -- for of these dif-- these things, but just to try to give you a feel for what's going to go with the probability of causation.

(Pause)

1 Maybe that just times out. 2 (Pause) 3 NIOSH evaluated the petition using the 4 quidelines established in 42 CFR 83.13. 5 issued a report March 29, 2007. This addendum 6 report was issued on September 6, 2007. 7 We evaluated whether it's feasible to estimate 8 the level of radiation doses to individual 9 members of the class with sufficient accuracy. Is there a reasonable likelihood that such a 10 11 radiation dose may have endangered the class. 12 NIOSH found that the available monitoring data, 13 process descriptions and source term data are 14 adequate to complete dose reconstructions with 15 sufficient accuracy for the proposed class of 16 employees; and a health endangerment 17 determination is not required. 18 So at this time we feel it's feasible to 19 reconstruct dose for all sources of -- and I 20 apologize for this. It was pointed out to me 21 that I again -- this is -- this is not a --22 this is my second time that I've done this --23 I've left Fernald in the presentation and we have not caught it either time, but -- so that 24 25 was brought to my attention and we do use a

1	template and so I unfortunately, I left it
2	in there twice.
3	DR. MELIUS: (Off microphone) The third time
4	you get (unintelligible)
5	DR. GLOVER: The third time I get
6	(unintelligible).
7	UNIDENTIFIED: (Off microphone)
8	(Unintelligible)
9	DR. GLOVER: Additional documentation and
10	sample dose reconstruction scenarios are
11	available on the Advisory Board's review and
12	the share drive located as said, so I
13	appreciate your all's attention. I'll take any
14	comments or questions.
15	DR. ZIEMER: Okay, thank you, Dr. Glover.
16	Let's open the floor a moment here for
17	questions or other comments, additional
18	information needed.
19	(No responses)
20	If not, we'll proceed oh, I'm sorry. Jim.
21	DR. MELIUS: Well, go ahead with the
22	petitioner, then I'll I'll ask my question.
23	DR. ZIEMER: Okay. So Dr. Giovacchini, if I'm
24	pronouncing that correctly, are you on the line
25	still?

1 DR. WADE: Yes. Ask him to make his comment. 2 DR. ZIEMER: If you'll proceed with your 3 comments. 4 (Pause) 5 MR. GIOVACCHINI: Hello? Can the Board hear me 6 now? 7 DR. ZIEMER: Yes. Yes. 8 (Pause) 9 Upon review by the reporter of the 10 recorded telephone comments, it appears there 11 were breaks in the transmission which may have 12 led to an inaccurate transcription of portions of the petitioner's statement.) 13 14 MR. GIOVACCHINI: Okay. I'm assuming I'm 15 coming in loud and clear. I thank you very 16 much for the presentation and your devoted work 17 to acquire a dose reconstruction. My question 18 to you is, is your dose reconstruction 19 accurate? Is it precise? And is it exact in 20 every detail, 'cause that's exactly what the 21 SEC laws stipulate. 22 I have three documents that I wanted the 23 Advisory Board to be aware of -- two documents, 24 and -- and I might be repeating some of the 25 issues that you've already presented to the

Board, but two of the documents support the fact that a dose reconstruction cannot be reconstructed to any degree of accuracy.

Crucial exposure data is missing. And one of the documents from my oncologist supports a POC, a probability of causation, that my cancer stems from my ionizing radiation exposure.

Now if the Board would like to hear these, I would be thrilled to share them with them, but I would first like to make a statement on my own -- my own behalf.

Once again, my name is Gerald M. Giovacchini. I am the petitioner. This Special Exposure Cohort, SEC 00059, was filed for just three individuals that worked in the X-ray laboratory at Sandia, California. And bear with -- piece of the information that you already know. I just -- 18 years after first exposure, one of the individuals contacted (sic) one of the 22 cancers specified by the SEC guidelines at the age of 30-- just 39 years old. This person contacted (sic) a chronic cancer. It -- that cancer is called non-Hodgkin's lymphoma. He contacted (sic) that disease five times over a (unintelligible)-year period and has received

radiation, chemotherapy or a combination of the two five times. His oncologist has told him to expect additional tumors, followed by additional treatment, for the rest of his life. By the time he was 48 years old his job and his ability to support his family were taken away from him. He was considered 100 percent disabled by both Sandia Medical Department and Social Security.

To date he has provided three affidavits from highly qualified individuals stating that it would not be feasible to reconstruct the dose to any degree of accuracy. Without exposure data, any dose reconstruction would be a guess and certainly invalid. He has also provided two letters from doctors and -- that clearly demonstrate a health endangerment from radioactive occupational exposures. documentation supports a probability of causation linked to radioactive work exposures. Yet another research report submitted into the record spells out the dangers of biological effects of ionizing radiation. That's the fourth research report recently report-recently restudied confirms the link between

ionizing radiation and non-Hodgkin's lymphoma,
yet NIOSH contends that the dose can be bound,
not to any degree of accuracy, as the law
states. The Advisory Board contends that there

was no health endangerment.

At this point our understanding of our legal rights under the EEOICPA of 2000 allows us to make the two following requests: Appeal to the Advisory Board NIOSH's decision that they have accurately reconstructed dose; and two, we appeal to the Presidential Advisory Board and their technical consultant, Sandy Cohen & Associates, to audit the NIOSH (unintelligible).

This letter represents the written appeal of the class. The following paragraphs demonstrate in greater detail the underlying facts that substantiate these appeals. This letter and the following 18 exhibits form the basis of our appeal. Therewith -- with this S-- when this SEC was submitted according to criteria in 42 CFR Part 83 clearly states (a) it is not feasible to estimate with sufficient accuracy the radiation dose that the class received; and (b) there is a reasonable

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likelihood that such radiation dose endangered the health of members of the class. These two issues will be discussed separately, with emphasis on the words "accuracy" and "reasonable," as this is how the EEOICPA law was written.

Continuing on, part A says it is not feasible to estimate with sufficient accuracy the radiation dose that the class received. The class believes a dose reconstruction be reconstructed because intent of the law states that it must be performed accurately. Please keep in mind that accurate means exact or precise. SEC was filed because all exposures, daily and accidental, went -- monitored, unrecorded and/or inaccurately reported. Submitted criteria is as follows. Therewith I am repeating some of what you already know. The class consists of three members. exposures to ionizing radiation were incurred on a daily basis. Three, personal monitoring records for one class member is missing. Four, two members of the class incurred an actual elevated exposure. Five, the incident report for one member of the class -- including

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medical reports, after his incident. actual exposures were over and above the daily exposures that routinely occurred. And seven, the dosimeters were worn during the exposure periods. Eight, inadequate shielding was utilized. Nine, radioactive and toxic materials were analyzed. Ten, the location of the tiny dosimeter chip in relation to the Xray exposure was either totally blocked or filtered from the X-ray beam. Eleven, finger rings (break in transmission) by Sandia. Sandia also did not see a need saw no need. for area monitors. The type of radiation produced by these X-ray (unintelligible) was highly collimated. Please also keep in mind, and this is very important, when speaking of one member of the class, that member represents 33.3 percent of the class; therefore 33.3 percent of the exposure data for this class is missing and is not available to include in any type of calculation. Three notarized affidavits from qualified (break in transmission) report these circumstances, yet NIOSH insists that a dose reconstruction can be accurately calculated. And this is Exhibit 1

1 I'd like to present of the appeal, that 2 scientific facts were blatantly suppressed and 3 ignored in light of our supporting evidence. I would also like to inform the Advisory Board 5 that two individuals confirm my elevated 6 (unintelligible) accidents or exposure in their 7 affidavit. Please do not refer to my accident 8 as being alleged. It happened. It's been 9 That is Exhibit 2, the accidental verified. elevated exposure of Gerald M. Giovacchini --10 11 that is myself -- has not been alleged. 12 In addition I would also like to inform the Advisory Board that the affidavit of one 13 14 individual states a comment that he received 15 from health and safety department at the 16 Sandia, California site. This comment is: 17 work with X-rays; that's your job; you ought to 18 be willing to take your turn in the barrel. 19 Well, a comment of this nature clearly 20 testifies that daily ionizing radiation 21 exposures were incurred. 22 And this leads me up to Exhibit 3. The class 23 would like to submit the fact that daily 24 ionization exposures were inevitable and un--25 unknown. The exposures cannot be accurately

1 quantified because the element of exposure time 2 cannot be determined. We don't know how long 3 the exposure time is and because of the daily ac -- daily exposures. 5 In the addendum evaluation report dated September 6, 2007 (break in transmission) 6 7 contends that the dose can be bound by 8 researching characteristics and parameters, 9 without taking into account the amount of time 10 a class -- exposed. 11 This leads me up to Exhibit 4. I would like to 12 submit into the record that without knowing the 13 amount of time an individual was exposed, 14 either the daily or (break in transmission) 15 exposures, dose reconstruction calculation 16 lacks crucial data. When exact dose exposure 17 time cannot be accounted for, the 18 reconstruction would be an invalid calculation. 19 The law specifies an accurate dose 20 reconstruction, one that is precise in every 21 detail. Please keep that in mind. 22 I'm on a -- got a little bit more to read, so 23 bear with (break in transmission). 24 The class would also like to stress the fact 25 that when exposure data is missing for one

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member of the class, it represents 33.3 percent of the data. Only 66 percent of the exposure data is available for NIOSH to calculate a dose.

This leads me up to Exhibit 5 of the appeal. This further demonstrates that sufficient information is missing from which a dose reconstruction can be calculated to any degree of accuracy. The dose reconstruction would be baseless and unfounded, as stated in the affidavit.

The original evaluation report that the class received on March 30th, 2007 clearly stated that assumptions, estimations and correction factors were utilized.

This leads up to Exhibit 6. How accurate in every detail, how precise, how (break in transmission) can the dose be (break in transmission) postulated and unsubstantiated The class strongly objected to this methology (sic) and when it was challenged the NIOSH responded three months later with an interim evaluation report stating that they now use an alternative method to bound the dose. The class senses a lack of pride in NIOSH's

decision when utilizing their original approach.

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Exhibit 7 specifies the class to know why NIOSH (break in transmission) only to relent on it when it was challenged. The class believes NIOSH again to fail to follow the intent of the The class would like to see a scientific reasoning approach, as Congress intended. One issue that was not included in the evaluation report was brought to my attention by Laurie Breyer on September 24th when she informed me that tritium bioassay exposure records were retrieved by one of the health physicists doing the dose reconstruction. At first I was a little confused and not sure why tritium exposure pertained to the years I worked in the X-ray lab. But after careful consideration, I do remember working eight hours per week overtime for a two-year period in the tritium research laborat -- (break in transmission) [Name Redacted]*. I believe time frame was 1975 to 1977, but I do pay stubs to verify the (break in transmission) frame when the time comes.

I apologize for not remembering, but this leads

1 me up to (break in transmission) eight. I am 2 requesting that radioactive tritium exposures 3 (break in transmission) into the (break in 4 transmission) reconstruction record. 5 requested this bioassay data from Dave Sundin 6 on September (break in transmission) 2007. At 7 this point we (break in transmission) if it 8 relates to the 1975-1977 time frame or 9 exposures that count to even more 10 (unintelligible) lost data. 11 On June 7, 2007 I had an extensive conference 12 call with four individuals identifying themselves as Pat T., Joe G., Elsie T., Dan S., 13 14 and that is all I know of these individuals. 15 find this particularly disturbing for two 16 reasons. Reason number one, conflict of 17 interest. To avoid the potential for actual or 18 perceived conflict of interest, a class has the 19 right to written conflict of interest 20 statements. 21 Exhibit 9, the class is requesting these 22 conflict of interest statements. 23 And the second reason pertains to the 24 qualification of the individuals processing the 25 claims. During the interviews the discussion

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was focused upon my (unintelligible) elevated exposure to my right hand and fingers. One of the individuals asked if the Sandia health physicist investigating the incident held a Geiger counter up to my exposed hand and fingers to record a reading. (Break in transmission) and responded by asking one of the other interviewers to answer the question. The response from that individual was a Geiger counter would not show a reading. This comment by one of the interviewers is especially troublesome as -- and is an insult to me. comment makes a statement regarding the lack of qualifications of the individuals supposed to be performing a fair and uniform dose reconstruction attempt. When I enlist a doctor, a coworker, for an affidavit I am required to obtain notarized documents. (break in transmission) DOL knows the qualifications of the individuals out of respect for the class and especially the sick applicants and NIOSH show the same courtesy. And this leads me to Exhibit 10. The class would like to point out that this further reinforces the fact that not having qualified

1 individuals processing claims (break in 2 transmission) of the dose is strongly in 3 question and is more likely going to be inaccurate and flawed. 4 5 We go to part (b) where it states that there is 6 a reasonable likelihood that such (break in 7 transmission) in dose may have endangered the 8 health of members of the class. 9 believes the health of one member was 10 endangered because the intent of the law 11 states, and I quote, there is a reasonable 12 likelihood. 13 This SEC was also filed because there is an 14 obvious health endangerment. That'd be myself. 15 Be informed that I have contacted (sic) one of 16 the two specified cancers, non-Hodgkin's 17 lymphoma. 18 This leads me up to Exhibit 11 of the appeal. 19 The term "specified cancer" is defined in the 20 SEC criteria for eligibility. I quote directly 21 from the SEC law a -- these having been 22 acquired in the performance of duty while 23 exposed to ionizing radiation. The facts speak 24 for themselves. 25 The following three documents have previously

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been submitted into my medical records but are highlighted here to clearly demonstrate the link between my radioactive work exposure and my disease. Dr. (unintelligible) stated that Mr. (unintelligible) -- and I quote, I'm not making this up -- stated that Mr. Giovacchini's non-Hodgkin's lymphoma was more likely than not related to his radioactive work exposures at Sandia, California. And two, the second doctor, [Name Redacted], states that lymphoma has been linked to occupation exposures to ionizing radiation. [Name Redacted] further states that Mr. Giovacchini's most recent cancer was clearly distinct from his initial lymphoma, which was (unintelligible) and not nodular, rather than diffuse and clearly distinct in (break in transmission). further identifies the link between lymphomas and ionizing radiation. He supports the fact that distinctly different cancers suggests a second primary cancer. The third document that relates to health

endangerment to work exposures is biological effects of ionizing radiation. This is a reference that I've used. BEIR VII illustrates

1 that no dose is only -- is the only safe dose. 2 This is available on the Internet. I'm sure 3 we're all aware of that. This supporting documentation was recent 5 retrieved and has not been sent to NIOSH for 6 inclusion in the amended evaluation report. 7 This information is from the Collaborative on 8 Health and the Environment. They are called 9 the CHE. The CHE report is a toxicant and 10 disease database that pertains to non-Hodgkin's 11 lymphoma and how strong the link is to various 12 causes. Please be informed that ionizing radiation raised a (unintelligible). 13 14 Exhibit 12 so states the evidence presented 15 supports the fact that there is a reasonable 16 likelihood that such radiation endangered the 17 health of one member of the class. 18 supporting facts are from reliable and 19 trustworthy sources. I am requesting that this 20 supporting documentation be factored into my 21 medical record as proof that supports a 22 probability of causation link into the -- to 23 the radiological work exposures that I 24 incurred. 25 Moving on, I'd like to say a little bit more.

1 We already established that I have had non-2 Hodgkin's lymphoma five times. The NIOSH 3 report that previ-- previous (unintelligible) 4 submitted into the record. [Name Redacted] 5 confirms my lymphoma sites are extremely rare. 6 Keep this in mind. Lymphoma doesn't usually 7 occur at the sites (break in transmission) 8 occurred. All of my ionizing radiation 9 exposures from working in this lab have been to 10 the upper trunk of my body and most (break in 11 transmission) side as I am right (break in 12 transmission). Aligning my lymphoma sites to 13 my exposure area, there appears to be a 14 striking similarity. I do believe this (break 15 in transmission) more than a (break in 16 transmission) mere coincidence, especially 17 after five (break in transmission). 18 Exhibit 13 of our appeal, I would like to 19 submit this medical information into my file as 20 supporting a health endangerment and a probability about -- of causation that my 21 22 lymphoma is related to my work exposures. 23 I'd like to say a little (break in 24 transmission). Hopefully everyone (break in 25 transmission) still hear me.

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Now it has been established that the type of ionizing radiation that was utilized in this Xray lab (break in transmission) highly collimated. It would therefore strike smaller targets, mainly (break in transmission) of the body. It is not a broad X-ray beam like the medical X-ray. It stands to reason that it wouldn't be likely for this type of radiation to strike a tiny target like a dos-- tiny dosimeter chip head-on. However, I did wear a dosimeter for the unlikely incident. Unfortunately, most of the time the dosimeter was worn behind the security badge or at the waistline, in which case it was either filtered or totally blocked. The evaluation report states that the dosimeter and security badge was all one. That may have been true for the current employees, but that was not the case during the tenure of the class. Exhibit 14, I'd like to state that the class (break in transmission) to (break in transmission) the statement amended. The class would also like to point out that this argument substantiates an inaccuracy of monitoring of external exposures to the upper torso, head and neck when the dosimeter is filtered, blocked or not pointed (break in transmission) toward the approaching beam. This practice resulted in exposures that were unmonitored, inadequately

5 reported.

One additional point that I would like to (break in transmission) of the Advisory Board is the Sandia, California site profile, and this is very important. The evaluation report refers to the -- the amended evaluation report refers to the profile (break in transmission) circumstances. NIOSH has acknowledged that all of the exposure data for one member of the class is missing. It stands to reason that if this data is missing, then it wouldn't be included in the profile. Why access the site profile.

Exhibit 15 (break in transmission) of the appeal, it is not a (break in transmission) fair practice as the law so states to refer to the document that doesn't contain the exposure (break in transmission) question. Furthermore, it is (break in transmission) fair practice to utilize exposure data of another individual to determine exposure of another individual. My

1 job duties may have been similar, but not 2 performed to the same degree as my successor. 3 During my tenure in this X-ray laboratory I 4 utilized the X-ray machines (break in 5 transmission) often and for longer periods of 6 (break in transmission). 7 Exhibit 16, I mention this because if my 8 workload was greater it stands to reason that 9 my exposures would have been greater. Once 10 again, sufficient information (break in 11 transmission) lacking to find a precise dose. 12 (Break in transmission) I am currently working 13 (break in transmission) and many other Sandia 14 retirees to bring the Sandia, California site 15 profile up to date so that it will accurately reflect the working conditions (break in 16 17 transmission) that I can (break in 18 transmission). 19 Exhibit 17, the class is requesting that this 20 document be given adequate time to be reviewed, 21 updated and not be referenced until former 22 employees are given the right to update its 23 contents. After all, it was these former 24 employees like myself (break in transmission) 25 the environment in which they worked.

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Referencing data from an inaccurate, incomplete document would unfairly judge a sick applicant's dose exposure and medical condition and ultimately jeopardize his or her EEOICPA claim.

Now we are all aware that the Advisory Board's expert (break in transmission) -- and I didn't put these words in. I'm just quoting them from the information I received. We are all aware that the Advisory Board's expert contractor, San-- Sanford Cohen & Associates, has identified many concerns with NIOSH's approaches. Specifically, SC&A stated that it has concern over NIOSH's ability to implement the stated methods, approaches and coworker models to enable dose reconstruction with sufficient accuracy as provided in 42 CFR Part 83. Even Shelby Hallmark, the DOL (break in transmission) Assistant Secretary for the Office of Workers Compensation, publicly criticized the validity of dose reconstructions. Mr. Shelby (sic) is concerned that with the development of new coworker models, added adjustment factors, creation of new technical guidance documents, et cetera,

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the EEOICPA process has gotten far too convoluted to allow accurate dose reconstruction. The Associated Press cited strategies by adjusting a -- ajucating (sic) agencies to contain the grown and benefits under the EEOICPA program. More pointed critism (sic) comes from our elected officials. For example, Representative John Hostettler, Indiana, at the December hearing of the House Committee on Immigration and Border Security (unintelligible) cited memos and e-mails showing that DOL (break in transmission) pressuring NIOSH to limit claims. Congressman Tom Udall stated that the agencies appeared to have assembled small bureaucratic empires, spending millions to devise a maze of regulations that ensure that hundreds of people enjoy (unintelligible) and prosperous career administering the pro-- this program. there are more comments from high-ranking officials regarding the validity of the Rest assured I did not make these program. comments (break in transmission) and do not want to stoop to this level to prove my point, but it is this type of correspondent that

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reinforces the fact that dose reconstructions are being improperly computed. No wonder sick applicants are stating appeal after appeal and seeking legal (break in transmission) assistance. At what point will Congress recognize the fact that administrative costs vers-- versus the benefits are way out of proportion.

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Please hold on one second.

(Pause)

Okay, I'd like to continue. I'm almost done. Bear with me. Thank you for listening. Sick claimants represent a class of people who have put their lives on the line during their employment at nuclear facilities throughout the United States. These people jeopardized their health and safety while being exposed to radioactive and toxic substances so that the United States could research, fabricate and maintain their nuclear deterrent. I was part of that. I am proud of that. These sick applicants performed the job that was asked of them. They worked in these laboratories for their families, their coworkers, their friends, and all who enjoy the freedom of living in the

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United States. For all those listening, this includes you, too. Some are grateful; some take their freedom for granted. And then you have some, namely those individuals response for ajucating (sic) worker claims, instead of supporting the individual who become ill after working in nuclear (unintelligible) are choosing to make it extremely difficult for these sick applicants or their survivors to obtain the benefits they rightly deserve. Numerous people who are aware of my medical condition, including some former supervisors, have all asked me three simple questions. Do you think your illness was caused from your radioactive work exposure? Two, if you had to do it over again, would you still work in the nuclear industry? Three, knowing what you know now about your exposures, do you think that your radioactive exposures could have been prevented? I would like to answer these three because I get these questions all the time. Question number one, do you think your illness was

caused from your radioactive work exposures?

The EEOICPA program has placed the burden of

proof on the sick applicant. I personally have done my research and I obtained the supporting documentation from coworkers, doctors, et cetera, and presented the facts supporting the Congressional intent of an SEC. Yes, I am confident my disease stems from my radioactive work exposures.

Question number two, if you had to do it over again, would you still work in the nuclear industry? Let me answer this by saying, you know, I raised children and now I have grandchildren. I get all choked up here, sorry. Excuse me. My parents raised me. Their parents raised them. Our ancestors -- hold on one second, please.

(Pause)

Our ancestors were willing to take a stand.

They would stand up and fight. Whether they fall in the battlefield or in the laboratory, they backed the good old -- they backed the good old USA and what it stood for. I am thankful that our ancestors preserved and paved the way for all of us and those yet to come.

The answer to this question is definitely yes, I would in the nuclear industry again.

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Former Sandia employees won their battle in the laboratory. They are extremely proud of their accomplishments, and that is something no one can strip them of. Regrettably, many are currently sick, like myself, and many have died. Whether the ajucating (sic) agencies listen to scientific reason now or later, those sick applicants that are confident that their illnesses were attributed to their work exposures will continue to appeal their EEOICPA claims until the ajucating (sic) agencies process these claims as Congress and the President intended. Until that happens, we as a nation are stripping ourselves of our integrity. In the opinion of this working class, and I'm sure other SEC classes would agree, it is vitally important that ajucating (sic) agencies and sick applicants and/or their families all play this game by the same set of rules. I do not believe we are playing by the same set of rules. And the third question, knowing what you know now about your exposures, do you think that your radioactive exposures could have been

prevented? Without a doubt, I would answer

1 this question yes. If I had worked in another 2 (break in transmission) laboratory, the chances 3 of being exposed to the degree I was exposed 4 would have been significant reduced. To do it 5 over again today, I wouldn't have been exposed 6 to the degree that I was back in the Cold War 7 era, the reason being nuclear workers these 8 days are protected by much stricter and (break 9 in transmission) exposure guidelines. If these 10 guidelines of the Cold War years were adequate, 11 why were they changed? 12 One health physicist recently told me, by 13 today's standards, the exposures in those days 14 would have been sufficient to set the stage for 15 health endangerment. 16 This leads me to the last exhibit, number 18. 17 I'm requesting that this statement be submitted 18 to my dose re-- exposure record. If you want 19 his notarized affidavit and qualification, I'll 20 be happy and delighted to submit his statement 21 into the record also. 22 In summary, the facts supporting the 23 Congressional intent of SEC 00059 have been presented to NIOSH and now the Advisory Board 24 25 knows them as well.

1 Four things that I know for sure. Number one, 2 I have a chronic type of cancer, non-Hodgkin's 3 lymphoma. 4 I am confident -- number two, I am confident my 5 non-Hodgkin's lymphoma stems from my 6 radioactive work exposures. 7 Number three, this petition is valid. 8 evidence presented adequately supports SEC 9 00059. 10 And number four, I do not agree with the 11 conclusion of NIOSH that they can perform a 12 dose reconstruction to any degree of accuracy 13 based on all of the above that I have 14 mentioned. 15 Finally, please be informed I am sick and I am 16 dying, and right now I feel pretty stupid that 17 I even contacted (sic) cancer. But I would 18 feel even more stupid if I did not set the 19 record straight before I move on to meet my 20 Maker. At my graveside I want it said that 21 Gerry was a good husband, a good father, and 22 someone who took a stand for principle to right 23 a wrong. 24 I worked under government contracts for 20 25 years. To the best of my ability, I was a good

steward of taxpayers' money. I'm extremely proud of my tenure at Sandia. If the ajucating (sic) agencies had acquired precise exposure details and processed my claims, Part E, Part B and this SEC, with honesty, integrity and the respect for the individual, I would have accepted that. The Advisory Board now has in its power to take a first step and tell Congress (break in transmission) an injustice. We must amend this program and right this wrong. Hopefully the nine people who are on the distribution list of this letter will take action and contribute, to the best of their ability, to res-- resolve the inequities of this program.

I am personally loca-- looking forward to resolving this matter without too much more time and expense. Please note, though, that there are other inaccurate statements within the addendum evaluation report that I would like to correct for the record. These additional corrections could very well have an impact -- a positive impact on the SEC. These will be formulated and mailed as soon as I have a chance to update them.

Just

1 Thank you to everyone for allowing me the time 2 to, one, request an appeal to NIOSH decision 3 that they have accurately reconstructed the 4 dose. And two, request an audit of NIOSH 5 methology (sic). On behalf of the working 6 class and all sick applicants, this is Gerald 7 Giovacchini. Thank you all for listening. you have any questions, I will state them. 8 9 DR. ZIEMER: Thank you very much, Gerald. 10 know it was very difficult for you to relate 11 some of that information and we appreciate your 12 input. 13 Board members, it is now the lunch hour. 14 think we do need to go ahead and take our 15 break, and following the break we will have a 16 discussion of this petition and determine a 17 path forward on it. But let's take -- we have 18 an hour for lunch. Actually the time is 19 squeezing past us already, but get back as 20 quickly as you can and we'll try to reconvene 21 as close to 1:15 as we can. Thank you very 22 much. 23 (Whereupon, a recess was taken from 12:23 p.m. 24 to 1:40 p.m.)

DR. ZIEMER: We're ready to reconvene.

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1 prior to the lunch break we had heard from the 2 SEC petitioner for the Sandia petition which 3 involves the X-ray diffraction workers. Board 4 members, this SEC petition and the evaluation 5 report now are open for discussion and/or for 6 any appropriate motions. 7 Josie. 8 MS. BEACH: I have an ap-- I have a question 9 for Sam on the urinalysis data. Was that done 10 in-house or by an outside laboratory? 11 DR. GLOVER: It would depend on the time frame. 12 I believe those were in-house data, but I 13 couldn't give you a clear answer. I don't 14 think that was out -- out-sourced. 15 MS. BEACH: Okay. 16 DR. GLOVER: But I don't know off -- I can't 17 give you the date. 18 MS. BEACH: Can you -- can you let me know if 19 it was out-sourced, though, at some other time? 20 DR. GLOVER: Sure. 21 MS. BEACH: And the reason I'm asking is the 22 question for the S-- or the CEP labs. 23 UNIDENTIFIED: (Off microphone) 24 (Unintelligible) 25 DR. GLOVER: No, that's outside that scope.

1	MS. BEACH: It is outside? Thank you.
2	DR. GLOVER: And this is Sandia Livermore, not
3	
4	MR. GRIFFON: Right.
5	DR. GLOVER: versus Sandia Albuquerque.
6	MS. BEACH: It wasn't clear in the documents I
7	read so I was
8	DR. GLOVER: Okay. Sorry about that.
9	MS. BEACH: Thank you.
10	DR. ZIEMER: Further comments? Yes, Jim
11	Melius.
12	DR. MELIUS: I I have a question. Have we
13	received the information that the petitioner
14	referred to in his phone call? He referred to
15	a letter and a number of appendices, and
16	DR. ZIEMER: I I don't think that I've seen
17	the material that he's referred to. I'm not
18	sure, Sam, even whether you have or LaVon,
19	can anyone help us out?
20	MR. RUTHERFORD: No, I I don't believe we've
21	re we've received a package that he's
22	identified, no.
23	DR. ZIEMER: Is this somebody help me out on
24	sort of in terms of the rules of engagement.
25	Normally is the petitioner allowed to submit

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1 supplementary information after the ER, or --2 or... I'm sort of asking whether we're 3 obligated to wait for such information. 4 don't think under the rules that that's 5 permitted, but --6 DR. MELIUS: Well, I -- I -- I don't know what the rules are, but certainly our practice has 7 8 been --9 DR. ZIEMER: Well, I know --10 DR. MELIUS: -- and that -- since some of 11 that's always -- actually som-- some of the 12 information today was in response to, you know, 13 a new report -- an addendum report from NIOSH, 14 I mean I don't think there'd be an issue with 15 it. 16 MR. RUTHERFORD: Yeah, I'm not going to say 17 anything about what Dr. Melius said, but I 18 think what the rule says is -- and I agree with 19 what Dr. Melius said, we haven't exactly 20 followed that -- the rule -- and our legal team 21 can correct me if I'm wrong. The rule says once that a petition has been submitted to the 22 23 Board, if the person wants to provide new information, that would be provided in a new 24 25 petition.

1 DR. ZIEMER: I -- I guess one of the questions 2 is is the petitioner allowed to provide 3 information in response to the ER itself. I 4 think probably we have allowed that, just as we 5 have allowed additional input from -- from NIOSH and from the contractor. So I'm not 6 7 sure, in terms of the strict interpretation of 8 the rule; but in general we've been fairly 9 flexible on that. I think we leave it to the 10 Board whether you want to see those exhibitions 11 (sic). And if so, that would require delaying 12 action on this. Or do you feel you have 13 sufficient information now? 14 NIOSH has indicated that -- that they can bound 15 this dose with sufficient accuracy. 16 claimant doesn't agree with that, apparently. 17 So what is -- what is your pleasure? 18 DR. MELIUS: Well, I mean I'll -- I mean -- if 19 I understand your question, Dr. Ziemer -- I 20 mean I would certainly like to see the 21 information and be ab-- be able to review it if 22 it's -- you know. And I believe it --23 petitioner said he would make it available to 24 us. I thought he had, that's why I was asking. 25 I thought maybe it -- something had come in and 1 (unintelligible) --

DR. ZIEMER: I'm not aware that we have it, so -- I've certainly not seen it. NIOSH apparently has not seen it. Certainly as a courtesy to the petitioner, why, we can delay action if you so wish.

DR. GLOVER: Just to be clear, in -- in the report I -- you know, there was a series of -- he provided additional information back in April 25th, and then immediately before the Board meeting he provided information in the conference call and additional affidavit last week that -- they were all taken into consideration as part of this. I believe this is all new information -- or not new, necessarily, but a separate packet that he's prepared in response to this supplement ER report.

DR. MELIUS: Yeah -- yeah, I was trying to figure this out as he was speaking and -- and I believe some of that -- most -- mu-- much of what he's referring to was in -- though some referred -- he did refer back to some earlier stuff that I think you had considered. And one reason I wanted to see it was I was confused by

1 some of the -- what was new, what was old and -2 - and make sure we understood the points 3 (unintelligible). 4 DR. WADE: I do think, from my perspective, 5 this Board has always operated on the premise that it wants to have all of the information in 6 7 its possession and that it wants to make the 8 appropriate decision. So I don't know that 9 waiting for information is necessarily at all 10 out of character for this Board. Whether or 11 not you think that information would sway you 12 or not, that's a judgment you can each make. 13 DR. ZIEMER: So appropriate actions, one motion 14 would be to defer action until the material has been received and we have a chance to review 15 16 Another possibility would be to make a 17 motion either to accept the NIOSH 18 recommendation or a motion to reject it, and --19 there's three possibilities there. So the 20 Chair's open to some sort of motion to get 21 things moving. 22 Phil, do you wish to make a motion? 23 MR. SCHOFIELD: Yes, I do. I'd like to make a 24 motion that we postpone a decision on this 25 until we do see -- receive the documents from

1 the claimants. 2 DR. ZIEMER: Okay, is there a second to that 3 motion? 4 DR. POSTON: (Off microphone) (Unintelligible) 5 DR. ZIEMER: The motion's open for discussion. 6 Okay, Dr. Melius. 7 DR. MELIUS: Well, this -- I'm not -- this may 8 be out of order, but the question I also have 9 that's related to that is do we want to ask 10 SC&A to review some of this information. 11 would -- we do have a category called a 12 targeted review, and I think there are a few 13 issues that could be addressed that would not 14 require a lot of time, though frankly, some of 15 that would -- may depend on some -- what those 16 issues are may depend on us seeing this submission from him. 17 18 DR. ZIEMER: Let me partially respond to that. 19 I -- I think we also have to be cognizant of 20 the use of resources in terms of -- this, in 21 effect, is a one-person petition, and I'm a little concerned about how much resources we 22 23 spend on a one-person petition. Not that it's 24 not an important petition, but as opposed to a

-- you know, a group of several hundred

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1 petitioners where the level of effort from the 2 contractor may be relatively substantial. 3 Let's hear from others. Wanda? 4 MS. MUNN: We've had a significant amount of 5 information provided to us, both by the petitioner and by NIOSH. It does not seem 6 7 reasonable that this Board would need to 8 involve our contractor further in investigating 9 what we have already seen and what is certainly 10 going to be well-covered by our postponement in 11 order to review the additional information the 12 petitioner has asked. 13 DR. ZIEMER: Okay. Other comments? 14 DR. MELIUS: Yeah. 15 DR. ZIEMER: Jim? 16 DR. MELIUS: I guess I'm -- I'm very leery, Dr. 17 Ziemer, of using a cost-benefit analysis to 18 apply this. I mean I -- and I'll actually --19 think that NIOSH did an excellent job of, you 20 know, putting an appropriate amount of 21 resources into this. They -- they've done two 22 -- two reports now and -- I mean I -- it's a 23 fact there's three -- we have another petition 24 coming up that we -- covers one person or 25 something and -- difficult. But at the same

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time, if there are, you know, valid technical issues that -- that need to be addressed, then I -- I think they should be addressed regardless of the numbers of people involved. Does that mean that we spend millions and millions of dollars? No. But -- which is why I suggested they partially -- suggest a focused review as -- as a possibility. I'm -- again, I'm not sure we're ready to make that decision now nor to know what to focus on now 'cause I -- I would -- personally would like to see the -- while -- what's been -- will be submitted from the petitioner, but I think we just need -- a little bit careful of using the cost of something as being the -- the driving force

And I would also add that I don't think that Congress intends us to, you know, have some ceiling as to whether -- where we would -- in terms of the use of our contractor to do -- do technical reviews. And if we --

DR. MELIUS: -- if there's not enough money, then we should be asking for more. We should not be trying to ration this amount out.

does not mean we don't use that resource appropriately and wisely, but...

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DR. ZIEMER: Well, one of the -- let me indicate that I don't disagree with that. However, I was cautioning us that we do nonetheless have to be cognizant of the judicial use of our resources. They are not unlimited, and I was simply cautioning the Board to take that into consideration. When I said the fact that it was one person does not make it unimportant, I was hoping to convey that part of the message. But nonetheless to be cognizant that we at the same time have a -and it's not necessarily a money resource. Wе have some -- also time constraints for our contractor which, in many respects, are equally as important. So I simply caution the Board to be cognizant of those constraints 'cause neither the time nor the money is unlimited. If we -- if we decide that this is worth pursuing, I don't object to that at all. I just want us to be -- on any of these, whether it's one person or 500 -- to make sure that we actually need the assistance of the contractor before we make -- or task them to do that.

you may be right, we may want to see these documents first and then make the decision at that point. I'm certainly personally a little reluctant to make such an assignment without having some idea of what -- well, we have an idea of what's in these documents --

DR. MELIUS: Yeah.

DR. ZIEMER: -- but it's a little vague without having them. Okay. So I don't think we're in basic disagreement there, I --

DR. MELIUS: No.

DR. ZIEMER: -- just want us to -- not to approach these things as if things are unlimited, either in time or resources. Yes, we'd like to have more resources and more time, and probably more people.

Okay, Wanda.

MS. MUNN: Because this particular petition addresses, in many ways, an entirely different set of circumstances than what we work with usually, it's clear to me that the agency has spent a great deal of focused energy on identifying what the salient points are in the petition and has tried to outline those to us, I think very well. I continue to feel that we

1	have a good explanation of what's in the
2	documents and where NIOSH can go with those.
3	Certainly reviewing the petitioner's documents
4	is well within our purview and will be
5	appreciated. The information that we've
6	received seems to be clearly adequate should
7	be for most
8	DR. ZIEMER: Well, are you speaking against the
9	motion to postpone to get the documents from
10	the petitioner?
11	MS. MUNN: No
12	DR. ZIEMER: No?
13	MS. MUNN: I'm not.
14	DR. ZIEMER: Just cautioning us about what
15	happens after that, perhaps.
16	Any others, pro or con? The motion before us
17	is to postpone action until we have a chance to
18	review the petitioner's additional documents,
19	as described in his presentation.
20	Yes?
21	MR. GRIFFON: Yeah, I'll I'll speak in in
22	support of the motion, but I I just wanted a
23	clarification I see one one case affected
24	by this petition, but I thought I heard three
25	and I'm I'm maybe I'm confused as

1 to -- somebody could help me out there. 2 DR. GLOVER: There are -- I have -- one thing I 3 -- this -- as we get to the oneness and 4 singularity, we begin to be careful about 5 Privacy Act stuff --6 MR. GRIFFON: Yeah. 7 DR. GLOVER: -- and so I have to be careful on 8 what we -- and how I couch presentation 9 materials, and I did want to caution the Board 10 as we review data, many of the things that he 11 spoke to were discussions about his particular 12 experiences and the singularity of his 13 experience. And so --14 MR. GRIFFON: But (unintelligible) --15 DR. GLOVER: -- it does -- well, the class is 16 three people who operated the X-ray diffraction 17 equipment and X-ray fluorescence in those 18 areas. They were in a later time frame, some 19 of the other people, and so there are numerous 20 21 DR. ZIEMER: Potentially there are three 22 individuals in the class. Is that what I'm 23 hearing? 24 DR. GLOVER: But not who were part of the 25 incidents involved as stated. I try to be

1 generic --2 MR. GRIFFON: Yeah. 3 DR. GLOVER: -- in this. 4 MR. GRIFFON: And then -- and then as I'm 5 looking at the uranium urinalysis, then -- then that -- obviously the -- the raw results that 6 7 I'm pulling up here in the reference that you 8 had in your evaluation report includes uranium 9 data from various areas, obviously, 'cause 10 there's more than three people. There's --11 there's ten -- at least tens of people that are 12 covered in there, and I think it looks more 13 like 40 or 50, you know, people in the uranium 14 urinalysis raw data. 15 DR. GLOVER: Well, for this partic-- you're --16 all the data from the class -- we have uranium 17 data for all members of the class. Those were 18 not specific to -- in the individual person. 19 All data was less than the MDA. 20 MR. GRIFFON: Right. 21 DR. GLOVER: I think I can -- so -- but that --22 not (unintelligible) one person's data. 23 MR. ELLIOTT: Again, we're trying to be very 24 careful here and protect the privacy of an 25 individual, but I believe that this -- this

1 petitioner had time outside of this class 2 definition. Is that correct, Sam? 3 DR. GLOVER: Yes. 4 MR. ELLIOTT: And so that urinalysis applies to 5 other employment and exposure experience that 6 he had. 7 MR. GRIFFON: Okay. But -- I -- I guess what 8 I'm getting at is -- and -- and understanding 9 the class, there's a -- a reference in the evaluation report that -- I think it's --10 11 anyway, it's one of your last on your reference 12 list, uranium bioassay results, '65 through '90 or something like that. Clearly, when I look 13 14 through all that raw data -- I don't know that 15 you've put this in any kind of spreadsheet 16 format, but when I glance through it, there's a lot of -- a lot of -- more than three and 17 18 probably, like I said, 40 or 50 individuals. 19 But they wouldn't have been in this particular 20 facility. Right? 21 MR. ELLIOTT: Were not in that X-ray 22 diffraction unit. 23 MR. GRIFFON: Okay. Okay. 24 MR. ELLIOTT: That's -- that's the distinction 25

1 MR. GRIFFON: So you -- you didn't parse that 2 document out to support this evaluation re--3 right? All right. I -- I think I understand 4 now. It's broader than just that facility. 5 MR. ELLIOTT: Yes. 6 MR. GRIFFON: Okay. MR. ELLIOTT: These are --7 8 MR. GRIFFON: When I glanced at it, I --9 MR. ELLIOTT: We do individual dose 10 reconstructions that are based upon the 11 circumstances of experience and exposure that 12 an individual had. And so when we start 13 talking about a --14 MR. GRIFFON: Yeah. 15 MR. ELLIOTT: -- singular class member, that's 16 where we get -- that's where it leads us to 17 talking about those circumstances, and that 18 becomes very difficult in a public forum 19 because of the Privacy Act. So I think you've 20 got it. I think you understand that the 21 uranium urinalysis is representative of 22 exposure outside of the class definition. 23 MR. GRIFFON: Okay. 24 DR. ZIEMER: Thank you. 25 DR. MELIUS: I --

1 DR. ZIEMER: Another comment, Jim? 2 DR. MELIUS: Yea-- yeah, I mean one of the 3 other reasons that I suggested having a -- a 4 focused review is -- is because of this very 5 difficulty. There's some questions I have that 6 I can't ask here, or at least they -- NIOSH 7 can't answer here -- here in public because of 8 the -- the small number of -- of individuals 9 involved and -- and so forth, and I think some 10 of the pursuit of some of these technical 11 issues involve privately -- information 12 involving individuals and -- and so I'm trying to think of a way of -- that we can pursue this 13 14 that doesn't -- al-- allows us to answer some 15 of these technical issues, but -- you know, can 16 -- can do it, not -- not in a -- a public 17 forum. Now there's a possibility of a 18 workgroup. There's a possibility of just doing 19 that -- that in-- that individually. I -- I 20 would suggest that the -- I would certainly 21 speak in favor of the motion to postpone. I 22 would suggest that we maybe talk about this at 23 our -- I believe it's December --24 DR. WADE: December 6th.

DR. MELIUS: -- phone call and just try to

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resolve a -- what the way forward will be. By that time hopefully we will have seen the additional information and -- and we can reach a resolution.

DR. ZIEMER: Okay. Wanda?

MS. MUNN: It would be wise, I think, for every member of the Board to make sure they've had an opportunity to read all the current existing documents on this prior to that phone call. It is the belief of some that those documents will contain the answers to the questions that people want to pose, if they are read carefully. Most of us I think have an opportunity only to scan these things when they come to us. Since this has become an issue, it would behoove us all to take careful care, read it, and be prepared to see if we still have the same questions when we make our phone call.

DR. ZIEMER: Thank you. Other comments or recommendations? Okay, Larry, you have an additional comment?

MR. ELLIOTT: I just want to make sure that it's stated here for the record that we have followed the law in our regulations, and those -- the law and the regulations require us to

answer that two-part -- two-pronged question. You heard the petitioner talk about the perception he holds that we have not accurately reconstructed his dose. We feel that we have accurately reconstructed his dose on a dose reconstruction, as well as accurately can reconstruct dose for this class -- not only by bounding, but by more precise estimation of dose. So that's one comment I feel is important for the record.

Another comment I would offer is that there's risk here. Our policy has been to advance petitions that qualify so that we can give full explanation and rational, clear understanding - if we can impart that -- to petitioners on how we go about doing our work, whether it's dose reconstruction or an evaluation of the petition. So in that -- in that effort, we have not held back and -- and denied these one-party petitions, if you will, where the class is so small, it's so narrow, that it represents an individual who's not happy with perhaps the outcome of a dose reconstruction. So I think that needs to be considered as -- as well.

It's been our policy to advance these forward,

1 but you could find yourself dealing with a lot 2 of individual-represented petitions. This is 3 not an appeal board, as you know, but it 4 certainly -- there's risk there toward that. 5 DR. ZIEMER: Thank you. Are you ready to vote on the motion --6 7 DR. MELIUS: Can --8 DR. ZIEMER: -- to postpone? 9 DR. MELIUS: Can I just respond? 10 DR. ZIEMER: Yeah. 11 DR. MELIUS: I think Larry did -- is making a -12 - a good point there and concur, and some of the problem with this, the -- dealing with this 13 14 petition is that we get into -- may not be 15 dealing with SEC issues, per se, but into the reconstruction issues related to that 16 17 individual and it -- it's awkward and -- in 18 some ways and -- and -- and does have some 19 peril. So I mean we are, I think -- at least 20 I'm -- and I think others are cognizant of --21 of that. At the same time I certainly think in 22 this case, as I read the petition and you--23 your res-- your response to it, that -- I mean 24 I think it was justified to handle this as an 25 SEC petition. I think a legi-- a very -- a

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legitimate issue came up about capability of doing accurate dose reconstruction, and so I --I find nothing, you know, wrong with what you're doing. I don't think you're erring (unintelligible) or not. That may not always be, you know -- I mean these are -- these cases are difficult, I think, simply by the nature of how you have to deal with them and -- and this is one avenue for so-called appeal. But -- but I -- I mean I do think it was justified to follow the steps. It -- it is harder to -- for us to deal with it because of the privacy situation, as well as this issue of getting into -- sort of commenting on an individual dose reconstruction that's in process, which is something we try to avoid.

DR. ZIEMER: As a caution, we need to duly note that we don't want this to be a back-door appeal process for every failed dose reconstruction.

Now are we ready to vote on the motion to postpone? The motion to postpone carries with it an implied -- well, not implied; an agreed-upon discussion at least of where we are at the next meeting in determination of next steps at

1	that point, assuming we have the documents all
2	in place by that time.
3	Those who favor the motion, raise your right
4	hand.
5	(Affirmative responses)
6	Any opposed to the motion?
7	(No responses)
8	Abstentions?
9	(Indicating)
10	One abstention.
11	The motion carries. Thank you very much.
12	DR. WADE: Do you wish to have your vote
13	recorded?
14	DR. ZIEMER: I'll vote in favor.
15	DR. WADE: Okay, so that's 11 in favor, one
16	abstention. And I put it on our tentative
17	agenda for a December 6th call. I guess I
18	would ask someone on the NIOSH staff to
19	interface with the petitioner to be sure that
20	we get that information.
21	DR. ZIEMER: Okay. Next
22	MR. ELLIOTT: Laurie will contact him and make
23	sure we get the information, distribute it to
24	the Board and post it on the O drive.
25	Y-12 SEC PETITION

1	DR. ZIEMER: Thank you. Next we're ready to
2	proceed on a Y-12 SEC petition. LaVon again is
3	going to present the evaluation report for
4	NIOSH.
5	MR. RUTHERFORD: (Off microphone) Can you check
6	and see if (unintelligible)?
7	DR. ZIEMER: Yes, [Name Redacted]* is to
8	represent the petitioners. Is she on the line?
9	(Pause)
10	UNIDENTIFIED: No.
11	DR. ZIEMER: At the at the moment she's not
12	on the line. Do we need to contact her first,
13	or shall we proceed?
14	DR. WADE: Well, we should try to contact her.
15	Do you have the contact information
16	DR. ZIEMER: Yeah, before we do that, we have a
17	conflict on Y-12?
18	DR. WADE: Yes. They've stepped away from the
19	table.
20	DR. ZIEMER: Okay, Mr. Presley and Dr. Poston -
21	- is that it, just the two, have reclused (sic)
22	themselves. They are conflicted on this one.
23	MR. RUTHERFORD: We'll have to check with
24	Laurie Breyer to see if she's got [Name
25	Redacted] phone number.

1 DR. WADE: Okay. (Unintelligible) do that? 2 Larry's (unintelligible). I think we should 3 proceed. 4 DR. ZIEMER: Yeah, 'cause she has the report. 5 Let's go ahead then, LaVon, if you'll proc--6 proceed with the presentation. Perhaps she'll 7 come on shortly. 8 MR. RUTHERFORD: All right. I'm LaVon 9 Rutherford and I'm going to present the Y-12 10 SEC petition, that would be SEC Petition number 11 00039. 12 We received this petition on July 28, 2005. Our initial review of that petition was that it 13 14 did not qualify and we issued a proposed 15 finding that the petition did not qualify on 16 September 28. In January 26th of the following 17 year the petitioner requested an administrative 18 review of that petition. 19 From the time that the petitioner requested the 20 administrative review, there's a -- as I 21 mentioned yesterday, a number of things that 22 went on. We did an internal assessment of our 23 own process to ensure that we were communing 24 well -- communicating well with the 25 petitioners. We also got Laurie Breyer on

1 board as our SEC petition counselor. 2 Lockey's group looked at the SEC petition 3 process to -- at petitions that did not qualify 4 and provided recommendations as well -- ways 5 that we could communicate better with the petitioners, so -- Dr. Lockey's group did not 6 7 get a chance to look at that actual petition 8 because it was in the administrative review --9 review process at the time. 10 The administrative review panel came back with 11 a recommendation that we should qualify the 12 petition because they felt we did not provide 13 clear justification to the petitioner for not qualifying that petition. 14 15 So on January 11th, 2007 we qualified the 16 petition and moved forward with our evaluation. 17 On June 29th we issued our evaluation report 18 and -- to the Board and the petitioners. 19 The petitioner's proposed class was all 20 statisticians who performed statistical 21 analysis of biological experiments related to radiation who worked in all locations at Y-12 22 23 from the period of January 31st in 1951 through 24 June 30th, 1959. 25 There were a number of reasons we modified that

1 class definition. One, we had a -- a previous 2 SEC petition that had completed evaluation that 3 actually evaluated class up through the end of 4 1957, so this person was included in that 5 portion of the class. And then we also --6 because the petition basis was an acute 7 incident occurring in the first quarter of 8 1958, we modified the class definition to --9 for the years -- or for the period January 1, 10 1958 through June 30th, 1958. 11 All right, a little (unintelligible) on Y-12. 12 Y-12 National Security Complex is located in 13 eastern Tennessee. It was part of the 14 Manhattan Project. Its function was to produ-process uranium for the first atomic bomb. 15 16 Construction of Y-12 started in February of 17 1943. Enriched uranium production started in 18 November of that year. 19 The first site mission was to separate uranium-20 235 from natural uranium by the electromagnetic 21 separation process. 22 Since World War II Y-12 missions have included 23 uranium enrichment, lithium enrichment, isotope 24 separation and component fabrication. After World War II the -- there was a 25

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moratorium on construction of facilities at the Y-12 facility -- or down in the Oak Ridge complex. Oak Ridge National Lab needed facilities to do work with their Biological Science division. There was the -- there was facilities at Y-12 that were no longer in use because of the Calutron operations had ceased and left some buildings available, so the Oak Ridge National Lab used some of the Y-12 -- was given some of the Y-12 facilities to con-conduct their animal research concerning carcinogen used at -- carcinogens. Biological Science division used sealed radioactive sources of cesium, cobalt, californium to do their experiments. In our evaluation we looked at a number of sources for information. We looked at the existing Y-12 site profile. We looked at Technical Information Bulletins that we currently had. We had interviews with former employees and case files in the NIOSH database, looked at the site research database specifically for incidents and things that may have led -- that would have been indi-- been indicative of this event. We looked at the Y-

1 12 Delta View Imaging system. We -- in that 2 system we -- again we looked for incident 3 reports of things from the ear-- the '58 time 4 period to see if we could, you know, find an 5 incident where this occurred. We also reviewed 6 documentation and affidavits provided by the 7 petitioner. 8 Other technical documents, we looked at 9 dosimetry documents on fogging or light leaks 10 that had -- was identified in this petition. 11 We also looked at medical reports on acute 12 radiation syndrome. 13 Radiation exposures occurred through -- to the 14 class. The principal internal exposures would have been from the residual uranium from the 15 16 (unintelligible) production operations that 17 occurred in the facilities. We did look at the 18 possibility of leaking sealed sources --19 leaking of the sealed sources. We had no 20 indications of any source leaks during the time 21 period, or in the few years after, of this 22 petition. 23 We looked at principal external radiation 24 exposures were beta -- beta exposures from 25 residual uranium contamination, gamma exposures

from the cesium and cobalt sources, and neutron exposures from a californium-252 source.

As of November 20th, 1951, all Oak Ridge

National Laboratories, regardless of work area, were required to wear a combination security badge and film dosimeter. NIOSH has external monitoring data for -- for members of the class.

NIOSH's evaluation on this class -- we -- we focused on external monitoring because of the exposure scenario identified. The exposure scenario was identified was an acute exposure occurring in the early 1958 period and that the -- a film badge reading was falsified which I - I actually identified this acute exposure. Therefore, our focus was -- like I mentioned, we did look at the -- looked for indications of failure, leakage sealed sources, and had no indication. So our focus on internal monitoring was on uranium then, and we had internal monitoring data for some members of the class, and we also have a Y-12 coworker model.

Issues identified by the petitioner and -- and our findings with -- with respect to those

issues. Petitioners submitted medical evidence of a depressed white blood count for a member of the class, and actual -- the -- actually submitted a number of reports on the -- this white blood count that actually started from -- records indicate that they started in '58 up through -- all the way up until the person acquired a form of leukemia later in -- around 1990.

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We also looked at -- and -- and based on that information, there was the expo-- as I mentioned, there was a film badge reading that indicated roughly 4.3 rem exposure. That film badge reading was lined out and identified with 100 percent light leak, and it also included in the interview with the employee that -- that employee indicated that they were not aware of being involved in any radiological exposures. The technicians and the -- the individuals that looked at the film badge made the determination that it was caused from a light leak in the film badge and therefore it was lined out and identified 100 percent light transmission and they were given a zero on the badge reading to -- to be consistent with their other badge

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readings they had previously received.

I want to point out that the -- the actual -the levels that -- of a film badge are not consistent with what you would see for a reduced white blood count. 4.3 rem is roughly on an order of magnitude below what you would see blood changes and could be much higher than that to see the reduced white blood count indicated. Also, for this acute incident to have occurred in this area, you would had to had a failure of interlocks and administrative controls. The highest source at that time -- I know if you look at the report you'll see a source that indicates that it releases -- or the exposure rate of over 400 rems per hour. However, that source was not in place in -- in the 1958 period. It went into place in 1962. That should have been reflected in the report but it's not.

The sources that were available, the highest exposure source was 26.5 rem per hour. That source would have had to been exposed for a considerable period of time, for hours, in order to -- an individual at one meter to receive the exposure indicated.

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As I indicated earlier, we reviewed the Oak
Ridge National Lab and Y-12 incident -incident reports. We had no indication of an
incident that would have delivered an exposure
of this level.

We also looked at the monitoring data for other -- the biological science division employees to see if we had indications of higher readings during a given time, and we had no indications of any higher readings for those individuals. And we looked at the -- the medical evince -evidence in itself does not support an acute Typically seen from acute exposure exposure. to high level of radiation causing a reduced white blood count you will have a drop in the white blood count to a -- a level, and subsequently that white blood count, over the weeks following, will return back to a normal What we've seen from the medical evidence provided was that a chronic expo-- a chronic low white blood count for a number of years. And if you actually look at one of the medical reports, it indicates that the individual in question is -- white blood count was at the low level in -- if you would look at

1 a range of white blood counts being 5,000 to 2 10,000 being the normal range, this individual's white blood count was around 5,000 3 4 in the years prior to this supposed incident 5 occurring. So based on what we've seen -- and we also 6 7 noticed that if you looked at the levels of the 8 white blood counts in the subsequent years, 9 they ranged from roughly 1,950 up to 5,150 in 10 the following years, and then they were fairly 11 constant from that point on. 12 As we mentioned, the petitioner identified a 13 film badge reading that they felt had been 14 modified that was indicative of this exposure 15 incident occurring. This film badge reading 16 again was investigated by Oak Ridge National 17 Lab and determined to be caused by a light 18 leak, and I did review that report. 19 We also looked at other reports and we did 20 actually find another report at Oak Ridge 21 National Lab and at Y-12 that had similar 22 findings for 100 percent light leak on four 23 individuals, so this is not a -- necessarily an 24 isolated incident. 25 We -- again the employee was consulted and

1 indicated no past history of radiation 2 exposure. We did look at their individ -- that 3 individual's readings, and they did have some 4 external monitoring exposure. I think 17 5 millirem was identified in a quarter, and 6 possibly 100 millirem in another quarter. So 7 although that was indicated at that time, that 8 they had no -- no experience of exposure 9 history, I did want to point out that we did 10 review their records to look at that. 11 And we did look at the indication of tears and 12 pinholes and stuff in the film badge. 13 -- a phenomenon that is ki-- that was seen in the industry at that time, so that is not 14 15 something that was just identified at Y-12 or 16 Oak Ridge National Lab. 17 Our evaluation process -- I also wanted to go 18 back to -- among the concerns -- I apologize. 19 One concern was -- is a criticality incident 20 that occurred at Y-12 in 1958. And if you look 21 at the report, in the report it clearly lays 22 out that the inci-- that the 1958 criticality 23 occurred I think in -- it was August or 24 September --25 DR. ZIEMER: June.

MR. RUTHERFORD: June, thank you, I couldn't remember -- which was later in the year from -- from the supposed first quarter 1958 occurrence.

So two-pronged test, is it feasible to estimate the level of radiation dose for individual members of the class. If we answer that yes, we don't ask the second question, is there a reasonable likelihood of health endangerment. We found that the available monitoring data, process, source term description -- source term data are sufficient to complete dose reconstruction for the proposed class. NIOSH determined it is feasible to complete dose re-dose reconstruction with sufficient accuracy; therefore a health endangerment determination is not required.

In summary, the internal exposures from uranium and all external exposures can be calculated for the individual -- or for the proposed class. And our recommendation is that we can reconstruct dose.

Questions?

DR. ZIEMER: Thank you, LaVon. This now is open to questions. Let's find out if the

1 petitioner is on the line. Is [Name Redacted]* 2 on the line now? 3 DR. WADE: (Off microphone) (Unintelligible) 4 someone calling her immediately outside the 5 door. We've been trying to contact her all day and have not been successful. 6 7 DR. ZIEMER: Apparently not at this moment. 8 DR. WADE: Laurie -- Laurie is trying. 9 her to try at the end of LaVon's comments. 10 DR. ZIEMER: LaVon, I'd like to ask a couple of 11 questions. I think you implied that the -- the 12 large cesium source was not in -- I assumed it 13 was the cesium source -- was not in use at that 14 time. 15 MR. RUTHERFORD: Yeah, in fact, you know, in 16 preparations for the -- my presentation, when I 17 drew in all the information, I -- I was 18 concerned because our report did not identify 19 that and I actually went back and verified that 20 the large cesium source did not come into 21 operation till -- it was like 1961 or '62. 22 DR. ZIEMER: Right. Now I also want to ask 23 about the californium source during this period because my recollection of -- is that 24 25 californium wasn't -- sources weren't really

1 being used till the '60s. Am I right? 2 that too early for californium as well? We can 3 call on Dr. Poston as a site expert. DR. POSTON: Well, I -- you're correct. 4 5 There's a couple of things -- and I am conflicted, but I have to tell you that I 6 7 participated in the installation of that 8 californium source in the biology division, and 9 it didn't occur before about 1967. 10 MR. RUTHERFORD: Okay. 11 DR. POSTON: I'm estimating probably 1971. 12 The health physics division at Oak Ridge 13 National Lab installed that and calibrated that 14 source. There was no source. Also, it could 15 not have been 3.9 curies -- maybe 3.9 16 microcuries -- 'cause in those days we were 17 making small quantity sources, not -- I don't 18 think we made four curies total since we 19 started making californium. 20 Yeah. Well, californium sources DR. ZIEMER: 21 weren't available anywhere in the country, or 22 perhaps the world, till later. So the only 23 source I think that would be under considered -24 - consideration would be the cobalt source --25 MR. RUTHERFORD: Okay.

1	DR. ZIEMER: possibly. Is that correct? I
2	I think you were saying that the 80-curie
3	cesium probably wasn't in play until
4	MR. RUTHERFORD: The way I understood was the
5	cesium source was the cesium the 65-curie
6	cesium source was in play. The
7	(unintelligible)
8	DR. ZIEMER: (Unintelligible)?
9	MR. RUTHERFORD: Yeah, the actual cobalt one
10	was, but that the 65-curie ce or cesium
11	source actually had the higher dose rate at
12	well, actually I'm sorry
13	DR. ZIEMER: Couldn't the
14	MR. RUTHERFORD: curie.
15	DR. ZIEMER: cobalt runs roughly four times
16	
17	MR. RUTHERFORD: Uh-huh.
18	DR. ZIEMER: on dose rate
19	MR. RUTHERFORD: Yeah.
20	DR. ZIEMER: at at a distance than
21	cesium, so
22	MR. RUTHERFORD: But it we actually have a
23	dose rate in there on the cobalt source if you
24	look take a look at it.
25	UNIDENTIFIED: (Off microphone)

1	(Unintelligible)
2	MR. RUTHERFORD: Yeah. Yeah, and it's only a
3	3.75 curie source. There was five
4	DR. ZIEMER: I'm so it seemed like
5	MR. RUTHERFORD: R per hour.
6	DR. ZIEMER: the cesium source would give
7	more output is what
8	MR. RUTHERFORD: Yeah, the cesium source we've
9	identified is rated at 26.5.
10	DR. ZIEMER: Oh, that's the rate.
11	MR. RUTHERFORD: Yeah, that's the one I called
12	out.
13	DR. ZIEMER: Oh, I see. Oh, okay. I gotcha.
14	MR. GRIFFON: And that was there at the time?
15	MR. RUTHERFORD: Yes.
16	MR. GRIFFON: Okay. I I (unintelligible) on
17	that.
18	MR. RUTHERFORD: Yeah.
19	DR. ZIEMER: Okay.
20	MR. RUTHERFORD: And that's the one that's
21	called out in the report, later on in the
22	report, if you look at the feasibility section.
23	DR. ZIEMER: And can you also tell us whether
24	that was a fixed source such as in a fixed
25	irradiator, or was it portable?

1 MR. RUTHERFORD: I believe it was a fixed 2 source in a fixed irradiator is the way I 3 understood it. Now I've -- I would probably 4 have to go back and -- and double-check on 5 that, but the way I understand, it was a fixed 6 source. 7 DR. ZIEMER: Thank you. Other questions? 8 DR. WADE: We've -- we've called just this 9 moment and the woman is not available. 10 not there. 11 MR. GRIFFON: Was that -- just trying to get my 12 bearings remembering the buildings in Y-12, the 13 criticality accident that you identified that 14 did happen later in June of '58, was that in 15 the same area? 16 MR. RUTHERFORD: No, I think if you look in 17 your report there's actually a map on that, and 18 they are considerably -- a considerable 19 distance between them -- buildings, no--20 DR. ZIEMER: What you mean by same area, it's a 21 couple buildings over. 22 MR. GRIFFON: Yeah. 23 DR. ZIEMER: Probably several hundred meters. 24 MR. RUTHERFORD: Yeah, if you -- and it's 25 actually laid out in the -- in the report and -

1 2 DR. ZIEMER: The criticality accident, though, 3 was quite well-characterized and --MR. GRIFFON: Yeah -- oh, yeah. 4 5 DR. ZIEMER: -- the dose rates at various buildings are quite well-known. 6 7 MR. GRIFFON: Right, right, but my -- I 8 guess my question was were -- were they doing 9 different types of research activities or --10 MR. RUTHERFORD: Well, the --11 MR. GRIFFON: -- you know. 12 MR. RUTHERFORD: -- criticality incident -- and 13 -- and I think --14 MR. GRIFFON: Yeah. 15 MR. RUTHERFORD: -- Dr. Ziemer could give you 16 better information on that, but was more of a -17 - a liquid tank coming together and coming to a 18 criticality. 19 DR. ZIEMER: It was a cleaning operation. 20 MR. RUTHERFORD: Right. 21 DR. ZIEMER: Sometimes referred to as the 22 impromptu barrel reactor, because we were 23 draining stuff into a barrel. 24 MR. GRIFFON: Yeah. No, I -- I -- I've seen 25 all the reports on that. I guess what I'm

1 asking is, the research activities in the area 2 where this person -- these statisticians were. 3 MR. RUTHERFORD: No. There was actually no 4 other work like that --5 MR. GRIFFON: Right, right, right --6 MR. RUTHERFORD: -- going on. 7 MR. GRIFFON: -- okay. 8 DR. ZIEMER: And this was X-10 work --9 MR. RUTHERFORD: Yes. 10 DR. ZIEMER: -- and the other was Y-12 work. 11 MR. GRIFFON: Right, right. 12 MS. BREYER: I did want to give the Board an 13 update on the petitioner. I e-mailed her on 14 the 14th with the information, in Sep-- on 15 September the 14th, and I also called her on 16 September 19th and verbally gave her that 17 information as well (unintelligible) interested 18 in listening. I called and was unable to get 19 anybody to answer, but they might -- you might 20 want to check just one more time, make sure 21 nobody on the phone -- that she's not 22 listening. 23 Thank you. Wanda Munn? DR. ZIEMER: 24 MS. MUNN: I'm prepared to move that we accept

NIOSH's evaluation of this petition as it is

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1 described in the documentation. 2 DR. ZIEMER: Okay. There's a motion to accept 3 the NIOSH recommendation, which would then be a 4 recommendation to the Secretary that the 5 petition be denied. Is there a second to that motion? 6 7 DR. ROESSLER: Second. 8 DR. ZIEMER: Seconded. Okay, discussion? 9 MR. GRIFFON: Can I -- can I ask a -- a --10 DR. ZIEMER: Yeah --11 MR. GRIFFON: -- follow-up on --12 DR. ZIEMER: -- sure. 13 MR. GRIFFON: -- on the -- you -- you mentioned 14 in one of your la-- later slides that you do 15 have the data to reconstruct --16 MR. RUTHERFORD: Yes. 17 MR. GRIFFON: -- you know, I -- I'm curious, 18 this one individual who had the -- a -- at 19 least apparently, you know, erroneous badge 20 reading --21 MR. RUTHERFORD: Right. 22 MR. GRIFFON: -- which was corrected to zero, 23 you -- so how would you reconstruct for the 24 statisticians? Would you use a coworker model 25 or do you have sufficient data individually or

-- or --

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MR. RUTHERFORD: We would use coworker mo-- I mean we could use the coworker model from the statisticians, but I think we need to -- I mean we could also give them, you know, a -depending on what we determine, we could use the LOD-type detection because of the fact that their exposures up to that point were zero. mean -- so I think that the data -- we have enough data that we could use -- if we determine that it wa more appropriate to give them a -- a coworker model external exposure, we have that data to do it. All right? Otherwise -- and really, to be -- you know -well, I shouldn't even say it, but you know, we -- we've done dose reconstruction on it already.

DR. ZIEMER: LaVon, do any of the members of the -- or any of the petitioners allege that they actually were aware of an incident -- for example, there are cases where -- in these kind of facilities where the source gets stuck in the out position and somebody goes in -- you know, they're irradiating mice or something, and then they think the source is back in the

1 shield and they wander in and -- and get 2 substantial exposure -- by substantial, perhaps 3 50, 60, 70 -- which happened in the -- oh, one 4 of the animal facilities at Oak Ridge that's 5 operated by the U. of Tennessee and --MR. RUTHERFORD: Yeah, I did that dose 6 7 reconstruction, by the way. 8 DR. ZIEMER: Okay. Well, so -- so then -- but 9 normally when that occurs, the -- the 10 individuals involved know that that has 11 occurred. 12 MR. RUTHERFORD: Right. We have --13 DR. ZIEMER: Do any of these individuals allege 14 that they inadvertently walked in when the 15 source was out or anything like that? MR. RUTHERFORD: No, we have -- and -- and 16 17 that's the thing -- you know, just like you 18 said, if -- if it occurs, it's going to occur -19 - I wouldn't expect it to occur necessarily to 20 a statistician as much as if it was going to occur it would be to the actual technician 21 performing the activities. And in the typical 22 23 (unintelligible) that occurs, you know, it is 24 known because interlocks, the administrative 25 controls, things have been violated for it to

1	occur, you know, so you know, and and
2	when as I mentioned, when I did the dose
3	reconstruction for the individual that the
4	one you were talking about, there were a number
5	of interlocks that were violated in that
6	situation, so
7	DR. ZIEMER: We have a motion before us.
8	Anyone wish to speak for or against the motion?
9	Or are you ready to vote?
10	(No responses)
11	I take it we're ready to vote. All who favor
12	the motion, which would be to concur with the
13	NIOSH assessment and recommend that the
14	petition be denied, raise your right hand.
15	(Affirmative responses)
16	One, two, three, four, five, six, seven, eight,
17	nine, ten I see ten.
18	Are there apparently no noes, and no
19	abstentions. The motion carries.
20	DR. WADE: Ten zero with two members away from
21	the table.
22	DR. ZIEMER: Right. Okay, thank you very much.
23	The two members are may now rejoin us.
24	Welcome back.
25	DR. WADE: We feel whole again.

Florence Black, are you -- can you come up and join us?

PLANS TO PROCURE BOARD CONTRACTOR FOR FY09 AND BEYOND

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DR. ZIEMER: Okay, we're going to talk about our Board contractor for FY '09 and beyond, and Flo is -- is going to give us the scoop on how we proceed here. Lew, take a moment and introduce Flo more formally to the group.

DR. WADE: Florence Black, who might be known to some of you, is a contracting officer who works in Pittsburgh. David Staudt has done the work on the SC&A contract, but David has been detailed to Atlanta for some four months and during his absence Flo is taking on this part of his portfolio. We fully expect David to be back, possibly in time to proceed with most of these actions. But as I mentioned to you before, this is such an important contract and we all understand the vagaries of government progress on these things that we wanted to start very, very early in the process. rather than wait for David to be back, we thought we would begin now the process of moving towards the recompetition of the contract that provides technical support to the

Board. That contract runs through the fiscal year we're in now, which means that SC&A will be on the job through September of 2008. We have time, but best that we begin now.

What we've done to sort of prime the pump is we've provided you with the statement of work - - tasks -- that were used the last time, with one modification. Flo and I added in the -- the new task for SEC work that was added to the original contract. So the statement of work you have is basically the statement of work that was used to compete the last time, with the task added in for SEC work, as the contract was modified.

We've also included the evaluation plan that was in essence used last time. There were some slight modifications to put it into the format that CDC uses now. So you have the statement of work and the evaluation plan.

I've also shown to you -- and Flo has posted this on the public web site so that the world can see everything that we do. I would rather have all of the discussions regarding our pursuit of a new contractor public. And I think the best way to do that is to put all the

documents on the public web site so that the world can see and comment at the same time we discuss them.

What -- what will happen, I think, is that we can have the discussion now related to those documents. Individual Board members can make comment to Flo or I between now and the December 6th meeting. At the December 6th meeting we can have another discussion of the statement of work and the evaluation plan. Hopefully after that we would be in a position to move forward with an announcement of our intent to solicit that would hit the street in January, and then the full solicitation would be out when, Flo?

MS. BLACK: Hopefully by the end of March.

DR. WADE: The only other thing that we want to lay before you is -- the process as it -- the selection process and evaluation process, as it will take place, will have the formation of a technical advisory panel. In the past that panel was chaired by the technical project officer -- that would be my position. The last time we had, I think, three Board members who participated on the technical evaluation panel,

1 and we ne-- we'd like to --2 MR. GRIFFON: Two, I think it was two. 3 DR. MELIUS: Paul and Mark, I think. 4 DR. WADE: Paul and Mark? I don't know the --5 I don't know the third name, so Paul and Mark at least -- do you think --6 DR. MELIUS: (Off microphone) (Unintelligible) 7 8 remember being on it? 9 DR. WADE: Do you remember? 10 MR. GRIFFON: (Off microphone) (Unintelligible) 11 two. 12 MS. BLACK: I -- I thought there were -- a -- a 13 third one when I looked in the file, but maybe 14 I read it wrong, perhaps. 15 DR. ZIEMER: Tony might have been on it. 16 you think it was... 17 MS. BLACK: There were two people that had the 18 training that's required, and one didn't have 19 to have it because the Board composition was 20 okay, but I -- that's what I was thinking was 21 from the Board. 22 DR. WADE: Maybe it was Tony. 23 DR. ZIEMER: Well, I --24 DR. MELIUS: Could have been Tony, yeah. 25 DR. WADE: So -- so if you would like to

1 recommend two or three -- when a technical 2 panel is put together of any members, it's 3 necessary that at least half of them have gone 4 through training. So depending upon the size 5 of the panel and the Board members we select, it's possible that a Board member selected 6 might have to take training, which is a five-7 8 day class that can be taken on line. 9 I'll say this off-record -- we'll try and 10 shield you from that, but it might be necessary 11 for you to have to take that. 12 DR. MELIUS: Five days? 13 DR. WADE: Five days. Take it on line. 14 Yeah -- well, we hope you can take MS. BLACK: 15 it on line; I can't guarantee that. And if you 16 take it for five days, you take a test. 17 tested class. 18 DR. WADE: Now my records show that Dr. Ziemer 19 is duly tested --MS. BLACK: 20 Yes. 21 DR. WADE: -- trained and tested. Mark is not. 22 So we would hope that, based upon the 23 arithmetic, maybe the Board members wouldn't 24 have to take the training.

So that's everything. We can talk about it.

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You can question Flo.

MS. BLACK: Okay, I -- I did want to make a couple of comments. First of all, a few of you know that actually because of things that occurred in the contracting office back in 2003, I actually was the specialist that made the original award. It had been transferred to me during a transition of -- of staff, and then I trans-- it was then assigned to David when he was hired, so I'm a little familiar with -- with the -- the basics of the support for the Board. And I think some of you have heard my name in other contexts, too.

There are a couple -- if you read these announcements, these were posted. The www.fbo.gov is a public web site, anybody can go to it. I've had -- already had one inquiry and the -- it's of a very general nature, just asking me was this a recompete. But anybody can go to it. This will be up there for quite a while.

I do want to caution you that if you go to the statement of work and print it, you're going to find out that, because I cut and pasted it from a Word document, that particular web site takes

1 all things like apostrophes and -- and 2 quotation marks and turns them into question 3 marks, so you're going to see a lot of question 4 marks if you -- if you print that out. Or if 5 anybody you know prints it out please tell them we didn't fill it with question marks. 6 7 -- it's a fluke in the system and feel free to 8 share this document with anyone now that it's a 9 public document, it's not restricted in any 10 way. 11 If you'd like I'll -- I'll do a -- a quick 12 review of this, just --13 DR. WADE: Please. 14 The statement of work starts out MS. BLACK: 15 with --16 DR. WADE: Everybody should have it in their 17 book --18 MS. BLACK: And there are copies in the back 19 for -- out there. This is the standard format, 20 it starts out C.1, because the statement of 21 work in a contract or in a request for proposal is section C, so it starts out C.1. And to the 22 23 extent that you have any comments that -- that 24 you want to make on them, if you can reference 25 the section, that's really helpful 'cause then

we'll know you're -- you're -- that's what
you're talking about.

Yes?

MS. BEACH: I have a question. Are the inquiries that are made on the web site, are those posted, and the answers?

MS. BLACK: No, this is not a formal synopsis.

MS. BEACH: Okay.

MS. BLACK: Okay? The formal synopsis we hope to post the first week of January, depending on -- on how that falls out with ev-- everybody's schedule. This is -- this was published under what's called a special notice, but the number that's given to it is the request for proposal number that will follow it through the whole process, that 2008-N-09682. Good question, because when it's -- when the proposal goes on it and inquiries are made, all of those answers This was just to ask for public are posted. comment, and we can -- as -- as a Board, you can include the comments. We, as a -- you know, a contracting office, as Lew and his staff, can read the comments and say they don't apply, and we don't have to respond to them when it's a special notice. It's just to get

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feedback. And -- and because Lew and the rest of us are very concerned about making the process as transparent as possible because of the nature of the work, so we put it out as a special notice.

This is not usually done in -- this -- it's not a -- it's not a standard contract procedure. It's completely acceptable, but it's not a standard contract procedure. Usually the first thing you see out there is the synopsis. But the statement of work is the one with all the Cs, and it's again in the standard format. The purpose of the contract, the -- the background and need, which talks about all the regulations and the -- and the establishment of Then the specific tasks, which are the Board. divided into sub-parts under C.3, and that's the lengthiest part -- as it always is since that's where we describe what's necessary. And that's the formal statement of work as -- as it will go -- this could be the final one, if -if no one has any comments that need to be incorporated, this may be what ends up as section C of the actual request for proposal.

And then, unless we would make changes as a

1 result of the proposals we would get, this 2 could be the final contract. 3 DR. WADE: Now under the explanation we have A, 4 B and C. A is individual dose reconstruction 5 reviews, and there we talk about the blind reviews, the advanced reviews, the basic 6 7 reviews -- as we did before. B is NIOSH/OCAS 8 site profiles and procedures reviews. 9 C is review of SEC petitions, so it's -- it 10 covers all the work we're currently doing, 11 consistent with the terminology you currently 12 use. 13 DR. ZIEMER: Does this automatically cover --14 there is another task, which is ma-- mainly a 15 tracking task. I think it may be Task IV; I'm 16 looking for John Mauro --17 DR. WADE: Task II. 18 DR. ZIEMER: -- or Task II? 19 DR. WADE: We don't do that. 20 DR. ZIEMER: No, but they do a --21 MS. MUNN: We have other tracking that we're --22 DR. WADE: They developed a tracking system. 23 DR. MELIUS: Yeah, they developed a tracking 24 (unintelligible). 25 DR. MAURO: In the previous -- the first

1	contract there was a Task II for the tracking
2	system. That has been completed and delivered.
3	DR. ZIEMER: Oh, here we go oh, we've got
4	procedures and site profile in one here, so
5	that's that
6	DR. WADE: The procedures.
7	DR. ZIEMER: I was looking for a fourth task.
8	DR. WADE: Correct, the procedures task is
9	there.
10	DR. ZIEMER: Okay.
11	DR. WADE: We don't call out a project
12	management task. I think that's up to the
13	offerors to to propose back to us.
14	DR. ZIEMER: Oh, that was the other one,
15	project management, but that's kind of built in
16	here. Right?
17	MS. BLACK: Right.
18	DR. ZIEMER: Yeah.
19	MS. BLACK: That's actually when you look at
20	the criteria, you'll see that's the second
21	evaluation criteria. Any other
22	DR. ZIEMER: Well, what do we do you need
23	specific comments today or are you just
24	soliciting
25	MS. BLACK: That's your that's your choice.

1 DR. WADE: If you want to guide us in a 2 different direction immediately --3 MS. BLACK: Right. DR. WADE: -- that's fine. If Board members 4 5 want to give us individual comments, that's fine. On the 6th we'll have you together again 6 7 as a duly constituted group. You can comment 8 then. 9 DR. ZIEMER: I believe this -- now that I see 10 the hard copy, I got this recently by 11 electronic copy, I don't know if the Board 12 members have had a chance to go through this 13 and digest it. It looks a lot like what we had 14 before --15 MS. BLACK: Uh-huh. 16 DR. WADE: It's the same. 17 DR. ZIEMER: -- almost verbatim. 18 MS. BLACK: Uh-huh. 19 DR. ZIEMER: But do you wish to -- to submit 20 your comments individually, which we can do, or 21 you can do it -- do them here? DR. MELIUS: Well, I -- I'd like to get this on 22 23 the agenda for the December meeting conference 24 call. 25 DR. ZIEMER: Well, it will be.

DR. MELIUS: Yeah, well, but then let me follow up. I think there's some issues that we need to think about. I'm not sure we'd make changes, but number one is the -- for the dose reconstruction reviews, I think that subcommittee members should certainly think are there changes that we want to have. Are we -- are we satisfied with the -- the current mix and -- and -- of types of reviews. Is there something we -- we need to, you know, think about in -- in terms of --

DR. ZIEMER: So you would be suggesting that we ask the subcommittee, for example, to give us input on that particular issue at the December meeting.

DR. MELIUS: Correct. The second thing I -I'd like to have us think about would be the -something we -- we talked about this morning.

Are we -- and I think it applies to site
profile reviews, also. I think -- when we
talked about the Special Exposure Cohort
evaluation reviews for large sites, we talked
about getting away from, you know, a single
large review to -- to -- I don't know if it's
focused review, but a series of -- of -- of --

sort of a step-wise process to this and -- and I don't know if that needs to be reflected in the -- the contract document or not. It may -- I'm not even sure it's something everybody agrees with. We haven't really tried it yet, though.

DR. ZIEMER: I'm wondering if it would be worthwhile -- and you could do this off-line-- is describe that a little more for Florence so that she might be able to tell you how much specificity we need in here. It -- it really is a description of how the contractor currently on the large sites would be carrying out the site profile review. It's sort of the direction that Hanford's going and sort of what was done at -- at Nevada Test Site, and I think that's what you're talking about, should that be reflected in the document.

DR. MELIUS: Correct. I -- correct, and I think for site profile -- I think we're -- I don't believe NIOSH is going to be preparing too many more new complete site profiles, but they're -- they're, quote/unquote, living documents and they would, you know, continue to evolve and so, again, may-- that may be a more

1 focused review and in order to -- I think to 2 have a fair, you know, competition, that --3 that we need to specify exactly what -- what 4 we're looking for. 5 Another area I think that I would suggest NIOSH 6 think about -- and again, there -- there may no 7 -- not be any changes planned, but if NIOSH is 8 going -- thinks it's going in a new direction 9 in terms of how it's going to be, you know, 10 doing its work or -- or -- or something, that -11 - then, you know, maybe there's some changes 12 there. I -- I'm not sure I can even think of 13 any, but it's -- it's a possibility. 14 DR. WADE: The PER possibly. 15 DR. MELIUS: Yeah, exactly. That -- that's --16 that's one. I think, again, we also have a 17 procedures workgroup which --18 MS. MUNN: (Off microphone) (Unintelligible) 19 DR. MELIUS: Huh? 20 MS. MUNN: Stop -- stop right there. 21 DR. MELIUS: Well, I'm just going to suggest 22 that -- I'm not going to tread -- I'm not, I'm 23 not trespassing. 24 DR. ZIEMER: Well, she's -- you're going to

take away what she was going to say.

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1 DR. MELIUS: Well --2 MS. MUNN: Go ahead. I'm being facetious and I 3 shouldn't. Go ahead. 4 DR. MELIUS: What I was going to suggest is the 5 procedures workgroup should also think about, 6 you know, are there -- you know, something 7 about the way that task is described, should it 8 be a separate task. Again, that's going to be 9 a question of updated procedures, new 10 procedures, if it's a -- that, and so forth. 11 And -- and I don't know how all this affects 12 contract, but -- contracting process, but it 13 seems to me that a lot of this'll be, you know, 14 smaller reviews in terms of scope rather than -15 - than somebody like a complete site profile 16 review, so the task would be, you know, for, so 17 18 DR. WADE: That's exactly what we need to have 19 a discussion about. 20 DR. MELIUS: Yeah, I --21 DR. ZIEMER: And --22 DR. WADE: We just used the same numbers, 40 23 procedures, six --24 DR. ZIEMER: Right. 25 DR. WADE: -- site profiles, as a starting

point. Now based upon experience to date, we need to modify (unintelligible).

DR. ZIEMER: Right. The other thing we didn't know at the front end when we entered into all of this was what the resolution process would look like. And it might be of value to the bidders or the potential contractors to have some idea of what that -- that's become a substantial part of what we do. It is part of dose reconstruction reviews and site profile reviews, but the resolution process itself is a substantial effort and we need to make sure that it is covered in the description here in some way.

Wanda, you have a comment.

MS. MUNN: Oh, a couple of quick things. When we have our -- our reports later in the day, the procedures workgroup has a couple of things to say about changes that might go on that probably wouldn't affect this SOW.

It would be very helpful for me, and I think perhaps some others, if -- if I could get a feel for the changes that have occurred from the original SOW that we processed in years past and this one. I can see that there are

1 several specific changes, but I -- I guess I --2 I understand you're asking us to say and what 3 else besides this --4 DR. ZIEMER: This is almost unchanged. 5 DR. WADE: That's the only -- that's the only thing I did -- 'cause I cut and pasted -- was 6 7 take the original and then I took the SEC task, 8 which was not part of the original contract, 9 and I took the words out of it and pasted it in 10 here as Task V. 11 MS. MUNN: Okay. So I'm essentially looking at 12 the same thing, with the addition of --13 DR. WADE: Right. 14 MS. MUNN: -- the SEC. 15 DR. WADE: Yeah, I could have tried to rewrite 16 it, but I thought that would have been 17 presumptuous of me. We need to -- I need to 18 hear your comments and then we'll take those 19 comments. But just as Dr. Melius is 20 enumerating, those are the kinds of things I 21 think need to be changed. We'd like to do it 22 based upon your wisdom. 23 MS. MUNN: Yes, and -- and what you said, Jim, 24 triggered something in -- a question from me, 25 as well. We -- we should be pretty far along

1 in terms of the large site profiles now, are we 2 not? We --3 DR. MELIUS: Yeah. 4 MS. MUNN: -- we're -- we're pretty much --5 DR. ZIEMER: Yeah, so it's those revisions that he's referring to. 6 7 MS. MUNN: Yeah. 8 DR. MELIUS: (Off microphone) (Unintelligible) 9 revisions and those revisions tend to be done 10 by --11 **UNIDENTIFIED:** In cycles. 12 **DR. MELIUS:** -- chapter. 13 MS. MUNN: Yeah, I --14 UNIDENTIFIED: In cycles. 15 DR. MELIUS: (Off microphone) (Unintelligible) 16 done piecemeal (unintelligible) -- being fair 17 to -- it's -- they -- done by chapters. 18 some are quite large and -- and involved and so 19 forth, some are minor. But I -- but I think 20 it changes how these'd be assigned and -- and 21 so forth --22 MS. MUNN: Yeah, that's probably true. 23 DR. MELIUS: -- Yeah. 24 DR. WADE: We're also getting further and 25 further behind the target of two and a half

1 percent dose reconstructions --2 DR. MELIUS: Right. 3 MS. MUNN: Uh-huh. 4 DR. WADE: -- which is something else you could 5 put in. 6 DR. MELIUS: Yeah. MS. MUNN: Yeah. 7 8 DR. ZIEMER: Okay, so we need some input --9 we'll -- we'll need input from the dose 10 reconstruction subcommittee, probably some 11 input from the -- the procedures review 12 workgroup --13 MS. BLACK: Dr. Ziemer --14 DR. ZIEMER: -- general input from everybody on the issue of resolution of matrices and so on. 15 16 MS. BLACK: Yeah, I -- I do want to say 17 something, and maybe I'm not understanding 18 'cause this is my first Board meeting, but we 19 don't want to tell the peop-- the potential 20 offerors exactly how we want them to do 21 something. That's what they tell us. That's 22 what we evaluate on. 23 DR. ZIEMER: Right, right, okay. 24 MS. BLACK: 'Cause when you start talking about 25 processes, what we want to tell them is what we

1 want them to -- the products we want from them, 2 what we want them to do. Then they come back 3 and tell us this is how we would do it. And 4 those who tell us -- you know, and that's 5 That's what the -- the technical scored. evaluation committee scores. 6 7 DR. ZIEMER: Well, for example, let -- let me 8 take the conflict -- rather the issue 9 resolution process. I'm not proposing that we 10 tell them exactly how we're not -- how we're 11 doing that now. 12 MS. BLACK: Okay. 13 DR. ZIEMER: But it seems to me that we -- we 14 should at least say that they -- they need to 15 tell us how they would suggest we do issue 16 resolution if they raise issues in their review 17 process. 18 MS. BLACK: Right. Yeah, I -- you just -- when 19 you started about process, I --20 DR. ZIEMER: Because we -- we know factually 21 that that's become a substantial part of the 22 work of our contractor. 23 MS. BLACK: And maybe we need to add something 24 else in section C --25 DR. ZIEMER: Right.

MS. BLACK: -- and even in the evaluation criteria 'cause although we haven't talked about those yet, the evaluation criteria are supposed to feed off of the statement of work. And so to the extent that we add something substantive to section C, we might add another -- we don't even have to change the points. You just add another component to the evaluation criteria.

MS. MUNN: Yes, that resolution process bears heavily on the work we do in procedures workgroup and creates a --

DR. ZIEMER: Well, all the groups -- workgroups, yes.

MS. MUNN: Yes.

DR. WADE: Another reason I'd like you to -- to focus on now for discussion later is we have an evaluation plan of 100 points. We're proposing an additional plus or minus 20 points for past performance, think about that. That's a weighty subj-- a weighty amount to give to past performance. Of the 100 points, we're breaking it out -- ten points for understanding purpose and objectives, ten points for management approach, 25 points for the technical approach,

1 and then 25 points for corporate experience 2 broken down into conflict of interest plan and 3 work history -- and I neglected to say 30 4 points for personnel. So those are numbers 5 we're proposing --MR. GRIFFON: How does --6 7 DR. WADE: -- talk to us about. 8 MR. GRIFFON: How does this compare with the 9 numbers -- the grading system we used last 10 time? I don't recall. MS. BLACK: The format is different. CDC has a 11 12 -- a different format now where we have these 13 five umbrella subjects, and then we fit 14 everything else under that. 15 DR. WADE: I tried to be consistent. 16 MS. BLACK: Right. 17 DR. WADE: No, the -- the past performance I 18 think is a little higher than last time, or is 19 it the same? 20 No, I think it's the same. MS. BLACK: 21 DR. WADE: Same as last... MS. BLACK: Sometimes we do -- and that would, 22 23 again, be something you could tell us what you want. We -- we never do lower than ten -- plus 24 25 or minus ten on past performance, but we've

1 even done as high as 25, plus or minus. would depend on -- on how much weight you want 2 3 to give to past performance, and that's over 4 and above the 100 technical points. 5 UNIDENTIFIED: (Off microphone) (Unintelligible) sounds --6 7 DR. MELIUS: Well --8 **UNIDENTIFIED:** -- (unintelligible) weighty. DR. ZIEMER: Well, we don't need to decide that 9 10 today. 11 MS. BLACK: Right. 12 DR. ZIEMER: She's saying --13 MS. BLACK: Yeah. 14 DR. ZIEMER: -- heads up on that, another issue we need to resolve at --15 16 MS. BLACK: Right. 17 DR. ZIEMER: -- perhaps at the next meeting. 18 DR. WADE: And again, just to -- to be clear on 19 the -- people's expectations, the Board can say 20 what it wishes and the contracting officer will 21 listen and then do what the contracting officer 22 thinks is appropriate. The Board doesn't hold 23 the final decision here. I can't imagine the 24 Board won't hold great sway over this process,

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though.

1 MS. BLACK: Right. 2 DR. MELIUS: Can I raise an issue related to 3 that? 4 DR. WADE: Sure. 5 DR. MELIUS: And again, pardon my memory and also I hope I don't offend anybody, but I -- I 6 7 -- I think we also have to remember that there 8 is a -- you know, the perception of how this 9 process is done is -- is -- is important, and 10 that to be fair to I think NIOSH and to the OCAS staff and so forth, I think we -- just be 11 12 -- be careful in terms of how we make up the 13 evaluation committee. I don't recall -- I -- I 14 believe Jim Neton was on it last time. I don't 15 remember --DR. WADE: I think Jim chaired it last time. 16 17 DR. MELIUS: Yeah, and -- and I would --18 I don't have any objection to Jim personally, 19 but -- but I think there'd be -- I'd have a 20 concern this time about someone from NI--21 NIOSH/OCAS chairing the evaluation committee. 22 I'd also have a concern about there being a 23 significant representation from NIOSH/OCAS on that committee, I -- 'cause not -- last time we 24

had the -- the perception of the OCAS reviewing

1 who's going to be -- who is going to be 2 evaluating them. This time we would be -- have 3 the perception of -- of OCAS reviewing who has 4 been ev-- who has been evaluating them. And 5 although I don't expect there would be any 6 problems, I'm not making that assertion, I 7 think for reasons of the perception of -- of 8 how this process is and -- and to be fair to 9 people applying and so forth that we should 10 avo-- avoid that potential -- that perception 11 of bias. 12 DR. WADE: Understood. 13 DR. MELIUS: And so having -- and I get -- I 14 don't know the numbers and I can't remember --15 DR. ZIEMER: Well, it is an appropriate point. 16 DR. MELIUS: Yeah. 17 DR. ZIEMER: The contractor, in a sense, is critiquing NIOSH. So to the extent to which 18 19 NIOSH chooses their critiquer (sic) --20 DR. MELIUS: Yeah. 21 DR. ZIEMER: -- that would raise questions. 22 DR. MELIUS: Yeah. 23 DR. ZIEMER: We -- I don't -- I don't know if 24 we can even think about -- is it possible to

have non-voting people on the board, like if we

1 said okay, we don't want NIOSH to even have a 2 vote but we may want some input or something. 3 But the contracting officer can advise us on 4 that --5 DR. WADE: I can tell you right now --6 DR. ZIEMER: -- but that's a big -- the 7 independence is an important issue. 8 DR. WADE: Right now the plan would be that I 9 would chair --10 MS. BLACK: Right. 11 DR. WADE: -- with several Board members and 12 several other technical experts, to be 13 determined. But we'll be very clear to you 14 about that before we do it. 15 DR. MELIUS: Yeah, well, we -- we need to -- I 16 think we need to discuss that specifically as -17 - as we get -- get further along. 18 DR. ZIEMER: Other comments today, Board 19 members? Yes, Wanda. 20 But that issue would be a matter of MS. MUNN: 21 degree, certainly. It would appear logical 22 that you would want the agency who was the 23 primary agency to have a significant voice in 24 the individual groups that they are going to be 25 expected to interface with over a long period

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of time. That's a -- you know, it's -- it would -- it would be -- it would be unwise to go too far the other direction, as well as going too far in the -- extremes are -- never serve us well.

DR. ZIEMER: Thank you.

DR. MELIUS: Can I just com-- I think it's something we can work out. I would personally find that having someone from OCAS chairing that evaluation committee would be, you know, at least -- again, appear-- appearance, and it may be how -- may be a totally valid review and -- and, you know, Jim or whoever would do an excellent job, I -- that, but then, you know, people on the outside are going to interpret that, you know, based on -- on appearance or something --

DR. WADE: Sure.

DR. MELIUS: -- and I think that would be, you know, troublesome for -- for -- for this program and -- and I think we have to also recognize there's a significant handicap in terms of trying to find other people that -- to do this. There are a limited number of health physicists within the federal government and --

1 can do this and within the agency and -- and we 2 -- we've struggled with that issue for a whi-while, so it's -- how to balance sort of the 3 4 technical input, the -- the bias and so forth 5 and I think we -- I agree with Wanda, we have to reach a -- a balance and I think the best 6 7 way to do that is to have some discussion of 8 that and be open about it. 9 DR. WADE: I -- I'd be compelled to say also I 10 think the technical evaluation panel last time 11 did an excellent job. 12 DR. ZIEMER: When do we need to select that 13 panel? 14 MS. BLACK: When Dr. Wade gives me what we call 15 a formal request for contract. 16 DR. ZIEMER: Okay. 17 MS. BLACK: He has to include a list. to tell me those that do have training and 18 19 those that don't. 20 DR. WADE: Probably December --21 MS. BLACK: Yeah, prob--22 DR. WADE: -- after our -- after our meeting. 23 MS. BLACK: After the meeting he'll submit 24 that. 25 DR. WADE: So we'll talk about that at --

1 robust (unintelligible) --2 DR. ZIEMER: So we'll need to know -- you know, 3 if all 12 Board members are interested in being 4 on the panel, then we'll have to make some 5 decisions on that. If none are interested, then we'll have to do some arm-twisting. 6 7 Somewhere between that, we may have a 8 combination of volunteers or others who can be 9 10 DR. WADE: Some. 11 DR. ZIEMER: -- compelled to -- to participate. 12 DR. WADE: It seems like people with experience would be wise to include. 13 14 DR. MELIUS: I -- I -- I concur. I -- also we 15 can do like a bidding process, like, you know, 16 when airlines overbooked, you know -- you know, 17 will you take the five-day course, will you take the ten-day course -- we'll see how that -18 19 20 DR. ZIEMER: I was going to --21 DR. MELIUS: -- if we narrow it down. 22 DR. ZIEMER: I was going to claim ignorance of 23 having remembered taking the course nor its 24 content, but... 25 DR. MELIUS: But my recollection was a one-day

1 course when I was in the government. Maybe 2 I'll just (unintelligible). 3 DR. ZIEMER: I think we're all a little slower 4 nowadays. Okay, do we need anything further 5 today then on that? I think we -- it's a good 6 opening. Thank you very much, Florence, for 7 getting us underway and our thoughts, and we'll 8 proceed at our next meeting to take the -- the 9 next steps to bring this to fruition. 10 DR. WADE: And we're ahead of schedule, but we 11 want to stay there. 12 DR. ZIEMER: Very good. 13 MS. BLACK: Right, yes. 14 DR. ZIEMER: Okay, it's time for a break. 15 We'll take a roughly 30-minute break, or a 16 rough 30-minute break, or something to that 17 effect. Thank you. 18 (Whereupon, a recess was taken from 3:05 p.m. 19 to 3:35 p.m.) 20 DR. ZIEMER: Translation of that coded message 21 is that we're ready to resume our 22 deliberations. Before we return to the agenda 23 let me introduce several folks who've joined us 24 this afternoon. John Nowack* who's with 25 Senator Biggert's office -- John, where are you

1	welcome. Also Robert Stephan is here from
2	Senator Obama's office and
3	DR. WADE: Robert's over there
4	DR. ZIEMER: Robert
5	DR. WADE: against the wall.
6	DR. ZIEMER: There he is, okay. Couldn't see
7	you in the glare there. And then Deb Deb
8	Detmers from Representative Shimkus's office is
9	here.
10	DR. WADE: Deb still might be working
11	(unintelligible).
12	DR. ZIEMER: Okay, we'll catch her later and
13	we'll hear from Robert later. John, did you
14	have a comment you wanted to make as we get
15	underway? Okay, we're pleased to have you
16	here, nonetheless.
17	(Pause)
18	SEC PETITION STATUS UPDATES
19	We have have a series of SEC petitions that
20	we want to get updates on that are in various
21	stages of review and consideration. They're
22	listed on your agenda and we'll just go down
23	through the list.
24	BLOCKSON CHEMICAL

The first is Blockson Chemical, and we'll get a

report from the chair of the working group.

And that's Wanda Munn, and she'll give us an update on the deliberations of the workgroup and path forward on Blockson.

MS. MUNN: The Blockson review from SC&A has been under consideration for our last couple of meetings. Most of the issues that they raised have been resolved. We met last on August the 28th in Cincinnati. The largest outstanding issue was the issue of the path of the thorium through that chemical process. We've had several communications with respect to that in the interim, and NIOSH has issued a white paper with respect to an additional review of the literature on the finer points and with respect to the thorium itself.

Just earlier this week, on the 4th, we met to discuss -- to get a verbal response from SC&A from their very cursory review of that -- that NIOSH white paper. They're going to take a look at it, give us a written report on their reactions to it so that NIOSH can have an opportunity to again respond to their reactions. That information we expect to discuss on a conference call which we will have

1	on November the 2nd.
2	It is our goal on that call to reach a
3	resolution of those last outstanding items and
4	have a recommendation for the workgroup to
5	bring to the Board at our next full meeting in
6	January.
7	DR. ZIEMER: Thank you, Wanda. And Robert, did
8	you have some comments that you wanted to make
9	relating to Blockson at this point? Thank you.
10	MR. STEPHAN: Thank you, Dr. Ziemer. Robert
11	Stephan, S-t-e-p-h-a-n.
12	UNIDENTIFIED: (Off microphone)
13	(Unintelligible) hear you.
14	DR. ZIEMER: I'm make sure the is the
15	mike up?
16	MS. MUNN: We're having a problem with our
17	system, Robert.
18	MR. STEPHAN: Okay. How about now?
19	DR. ZIEMER: Good.
20	MS. MUNN: Better.
21	MR. STEPHAN: That works? Okay. Robert
22	Stephan, S-t-e-p-h-a-n. Dr. Ziemer, a couple
23	of things is one you know, we started
24	from our office on this Blockson issue in terms
25	of a public way last year this time, in

1 November. So I want to thank the working group 2 and the Board for being so deliberative on the 3 Blockson issue because this all could have been 4 decided a year ago and -- and maybe in a way 5 that we didn't have that much confidence in. So I feel like the -- you know, the effort that 6 7 needs to be made is being made and -- and we 8 are thankful for that. We look forward to the resolution that comes from this issue about 9 10 thorium. And so our understanding is, if I 11 heard correctly, that no vote today on the 12 Blockson SEC, but you're going to hopefully 13 vote -- if this thorium issue is resolved to everyone's satisfaction -- in January. Is that 14 15 right? 16 MS. MUNN: That's our goal. 17 DR. ZIEMER: That would be the --18 That's the goal. MR. STEPHAN: 19 Right. That's, I think, the final DR. ZIEMER: 20 issue to be addressed on Blockson. As far as 21 we know, that will close all the issues and 22 we'll be ready for action. 23 MR. STEPHAN: Okay. As we have said in the 24 past and we will say in January, you know, from 25 our perspective the issue before the Board -- a

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major issue before the Board is do you accept the data being used from the Florida site and -- and applying that to Blockson. And our position is that -- that you should not. I mean I know NIOSH disagrees with that. respect that wholeheartedly. If -- if you are a -- a NIOSH staffer, if you are a health physicist, if you're a Board member even, maybe, you know, it may pass muster that scientifically you believe it's okay to use data from one site and apply it to another. From our perspective, and I think the perspective of the claimants and the workers, that doesn't pass the smell test for them. And so that's the one thing we would continue to ask you to keep in mind is whether or not we can apply data from this Florida site to Blockson with integrity and have integrity in So we -- we believe that you the process. can't, and so that's the one thing I want you to keep in mind as you continue to go forward with this and -- and you head towards a vote in January. Okay? Thank you.

DR. ZIEMER: Thank you, Robert, and certainly we'll remain cognizant of that issue as we

1 proceed. 2 Board members, any other questions for the 3 working group on Blockson, or comments? 4 DR. MELIUS: Yeah, I --5 DR. ZIEMER: Dr. Melius, you have a --6 DR. MELIUS: I just have one thing that's come 7 up since the workgroup meeting this morning. 8 want to sure that well before our January 9 meeting that we've made available workgroup 10 minutes and the -- the record of those meetings 11 that are available to the petitioners and so 12 forth. I don't believe that our last workgroup 13 minutes have been -- are publicly available yet 14 and I -- I think that's something we can talk 15 tomorrow when --16 DR. ZIEMER: In fact that --17 DR. MELIUS: -- on that issue --18 DR. ZIEMER: -- it is on the agenda tomorrow. 19 DR. MELIUS: -- the agenda, but I just want to 20 make sure that everyone's alerted to that as a 21 -- as a issue in terms of scheduling for 22 Blockson. 23 DR. ZIEMER: Thank you. Any other comments on 24 Blockson? 25 (No responses)

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Okay, let's proceed with Fernald and the chair of that workgroup is Brad Clawson. And Brad, give us a status report and a little bit of your idea of path forward there on Fernald. MR. CLAWSON: Okay. And August 8th was the first time that we met as a working group. as usual in the start of this, we had a lot of new information that working group nor -- nor SC&A have been able to review or go forth. originally made it clear through the matrix, but with all this new information we wanted to be able to have time to be able to get it on the O drive, plus be able to review it. So at this time NIOSH has put -- it appears to be most of the information that was requested onto the O drive. We've got a working group scheduled for October 25th -- 24th, and we're going to continue on from there.

DR. ZIEMER: Okay, thank you. Any questions for this workgroup?

DR. WADE: We do have on the phone I believe
Ray Beatty, who would like to make a comment --

DR. ZIEMER: Okay.

DR. WADE: -- on Fernald.

1	DR. ZIEMER: We'll hear from Ray Beatty then.
2	DR. WADE: He's a very nice gentleman who's
3	spoken to us.
4	DR. ZIEMER: (Off microphone) (Unintelligible)
5	petitioner?
6	DR. WADE: Yes.
7	DR. ZIEMER: Ray Beatty I believe is one of the
8	petitioners.
9	UNIDENTIFIED: Or a representative.
10	DR. ZIEMER: Or is he a representative?
11	(Whereupon, several Board members spoke
12	simultaneously.)
13	DR. ZIEMER: Representing the petitioners, yes.
14	UNIDENTIFIED: (Off microphone)
15	(Unintelligible) Sandra Baldridge.
16	DR. ZIEMER: Okay.
17	(Whereupon, several Board members spoke
18	simultaneously.)
19	DR. ZIEMER: No
20	DR. WADE: Okay, Sandra Baldridge, she's the
21	petitioner.
22	DR. ZIEMER: Yeah, she can speak, sure.
23	DR. WADE: Okay.
24	MS. BALDRIDGE: I have a question. At the
25	working group NIOSH said that they would be

1 sending revisions of the site profile as they 2 were developed. I was wondering if the Board 3 has received any of those revisions. 4 DR. ZIEMER: Okay. Thank you, we'll -- Brad, 5 can you or someone from NIOSH tell us the status of those revisions that they're -- are 6 7 referred to? 8 MR. CLAWSON: Not -- not at this time that --9 we haven't received anything on that. 10 DR. ZIEMER: Are there some Fernald revisions 11 that are in the pipeline that -- okay, Jim 12 Neton is going to --13 DR. NETON: I'm -- I'm not aware, although I'm 14 not 100 percent certain, that there are any 15 revisions that have been released recently. 16 They're being worked on, but I don't think that 17 we formally released any --18 **UNIDENTIFIED:** No, they haven't. 19 DR. NETON: -- updates at this point. 20 DR. ZIEMER: Okay. Well, the main thing will 21 be to assure the petitioners that any revisions 22 that are forthcoming are provided. So as far 23 as we know, the petitioners should have 24 everything that the Board has in terms of 25 documents at this point.

1	DR. NETON: Yeah, we had just finished on and
2	it's it's under the internal it's
3	under review
4	DR. ZIEMER: Okay, there is
5	DR. NETON: but it's not released. And I
6	would say that any revisions, as they come out,
7	are automatically posted on our web site, so
8	they will be available and then
9	DR. ZIEMER: As soon as they're released,
10	they'll be available.
11	DR. NETON: As soon as they're released, within
12	a day or so, they're published on our web site.
13	DR. ZIEMER: When do you anticipate that? Is
14	that is that imminent? Are we talking a few
15	weeks or several months or just a I
16	(unintelligible)
17	DR. NETON: It's under internal review. It
18	should be a matter of several weeks.
19	DR. ZIEMER: Okay, so there should be something
20	coming out we're expecting then by the end
21	of October, perhaps.
22	And here's Larry Elliott to add to that.
23	MR. ELLIOTT: I haven't seen this document
24	myself, but it it may not be released until
25	we're through with the deliberation of the

1	working group, either. There may be some
2	issues that are being discussed in that working
3	group that could hold this document up, so
4	DR. ZIEMER: Yeah, but the point is, there's
5	not another
6	MR. ELLIOTT: There's not
7	DR. ZIEMER: document out there
8	MR. ELLIOTT: there's not any revisions that
9	have been produced
10	DR. ZIEMER: Right.
11	MR. ELLIOTT: to date.
12	DR. ZIEMER: Okay.
13	MR. ELLIOTT: And and I'm hesitant to say
14	when we're going to produce this one that was
15	just raised.
16	DR. ZIEMER: Right, and also we are making sure
17	that the that the petitioners are notified
18	of the workgroup meetings, I believe, and are
19	given an opportunity to participate.
20	Now another comment from the petitioner.
21	MR. CLAWSON: (Off microphone) (Unintelligible)
22	MS. BALDRIDGE: (Unintelligible) to the
23	revision information before it is posted on
24	line.
25	DR. ZIEMER: Could you ask her to repeat

1 that, please. 2 MS. BALDRIDGE: (Unintelligible) I have access 3 to the revision information before it is posted 4 on line to the public. 5 DR. WADE: She would like access before it's 6 posted. 7 DR. ZIEMER: None of us have access to the 8 drafts before they're posted for the public. 9 The Board doesn't -- these are agency 10 documents. I believe that's correct. They are 11 not available to the public or to the Board 12 until they are posted. 13 DR. WADE: Could we call the petitioner when 14 they're posted? 15 UNIDENTIFIED: (Off microphone) 16 (Unintelligible) 17 DR. WADE: Let's commit to that. 18 DR. MELIUS: But -- can I just comment? 19 DR. ZIEMER: Yeah. 20 DR. MELIUS: I believe that if there were 21 issues in the revi-- revised site profile that 22 are relevant to the SEC evaluation, those'll 23 come up as part of the discussion to the 24 evaluation --25 DR. ZIEMER: Or the workgroup --

1	DR. MELIUS: the SEC workgroup's evalua so
2	through that process, that information should
3	be make that
4	DR. ZIEMER: That would be available
5	DR. MELIUS: (unintelligible) available
6	available (unintelligible)
7	DR. ZIEMER: in the workgroup meetings.
8	DR. MELIUS: available to the petitioners
9	and and to the public, so it it may not
10	be the entire site profile revision, but it may
11	be
12	DR. ZIEMER: The sort of issues, yes.
13	DR. MELIUS: Yeah, that's been our past
14	practice and
15	DR. ZIEMER: Right, thank you. Okay, any
16	further comment from the petitioner on this?
17	(Pause)
18	No. Thank you.
19	DR. WADE: What about Ray, is does Ray still
20	want to make a comment?
21	(No responses)
22	Ray Beatty?
23	(Pause)
24	MR. BEATTY: Yes, Dr. Ziemer?
25	DR. ZIEMER: Yes okay.

1 MR. B

MR. BEATTY: Hello?

DR. ZIEMER: Tell him to go ahead, yes.

(Pause)

MR. BEATTY: Yeah, Dr. Ziemer, this is Ray Beatty calling on behalf of Fernald petition, and my comments are the fact that when we were at the last workgroup meeting there were several mentions of revisions that have occurred. And I'm not real sure that we have been afforded the privilege to see these revisions prior to the -- the next meeting, or when these revisions are being made, we -- we learn of them at the workgroup meetings. I just think that there ought to be an opportunity that the petitioner should get to review these prior to the workgroup meeting, if that is possible. Not so much the date of -of putting them on line.

DR. ZIEMER: The day that they're put on line is basically the day they're released and the day that the workgroup gets them so that they're available to everybody at the same time. That would not necessarily coincide with a workgroup meeting, so -- in other words, if they're -- if they're available a month before

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1	the workgroup meets, that's when you'll get
2	them. Basically they will be available to you
3	the very same time they're available to the
4	workgroup.
5	MR. BEATTY: (Unintelligible) satisfactory,
6	sir. Thank you.
7	DR. ZIEMER: Yeah, thank you. Okay, let's move
8	on then to oh.
9	DR. WADE: Larry needs to clarify.
10	DR. ZIEMER: Oh, Larry
11	MR. ELLIOTT: I just want us to be clear here.
12	DR. ZIEMER: (Unintelligible) if I misstated
13	that.
14	MR. ELLIOTT: No, no, I I don't think you
15	did. I I want us to be clear. We want the
16	petitioners to have all the available
17	information that the working group's working
18	with. But I think what Ray was speaking of may
19	have been the matrices that the workgroup deals
20	with rather than revisions to the site profile.
21	DR. ZIEMER: Well, the matrices will be
22	available certainly.
23	MR. ELLIOTT: But the matrices, in some
24	instances, have to be privacy
25	DR. ZIEMER: (Unintelligible)

1 MR. ELLIOTT: -- reviewed and redacted --2 DR. ZIEMER: (Unintelligible) 3 MR. ELLIOTT: -- I just want it known that 4 that's not my responsibility to make happen 5 quickly. I will put it on the web site and 6 share it with the petitioners as soon as it's given to me in a redacted form. 7 8 DR. ZIEMER: Yeah, if you're talking about the 9 revisions of the resolution matrices, insofar 10 as that has Privacy Act information, that has 11 to go through the redaction process. And Board 12 members may have that sooner than the 13 petitioners since they're permitted to have 14 those documents. MR. CLAWSON: Also too there was -- there was 15 16 many things that we were going to have put on 17 the O drive and we -- we need to make sure that 18 they realize that that's not public --19 DR. ZIEMER: The O drive materials are 20 materials that are not public materials, for a 21 variety of reasons. 22 MR. CLAWSON: Right, and we had that --23 DR. ZIEMER: But insofar as those materials are 24 -- can be made public, I guess they will also 25 appear in a public forum, yeah.

1 MR. CLAWSON: Right, we have --2 DR. ZIEMER: Insofar as we can legally make 3 them available. 4 MR. CLAWSON: We had a lot of different 5 information that was coming up -- our raw data, the OTIB-25, an Excel spreadsheet, Dr. Petty's* 6 7 report, lab procedures and so forth, and 8 several white papers that are going to be on 9 the O drive. But -- but those are not --10 DR. ZIEMER: Those aren't available --11 MR. CLAWSON: Right. 12 DR. ZIEMER: -- at this point, yeah. Okay, 13 thank you for clarifying that. 14 Yes, Jim. 15 DR. MELIUS: Again this is for discussion 16 tomorrow, but I'd repeat that I -- I think we 17 need to make sure that we make those available 18 to the petitioners in a timely fashion. 19 not be contemporaneous with the -- the 20 meetings, but in a way that they have access to 21 them and a way -- can provide input and -- and 22 comment on them, and certainly well before any 23 decisions are -- are reached on a -- on a 24 petition. And we've not been doing a good job

of that up till now and we need to be doing

better, so...

CHAPMAN VALVE

DR. ZIEMER: Thank you. Okay, then let's move on to Chapman Valve. We do want to hear from Sharon Block, who's on Senator -- is she on the line? We'll have to check and see. She's on Senator Kennedy's staff and wanted to make a brief statement prior to the Chapman Valve discussion.

MS. BLOCK: Thank you. I appreciate the opportunity to make a -- a brief statement. I just wanted to express the Senator's concern about, you know, the ongoing delay in making a decision on this petition. Obviously these petitioners have been waiting years, and I'm anxious to see if any progress has been made since the July meeting. We were somewhat concerned that -- that we didn't hear or that the Board didn't hear from the Department of Energy till September 25th, when I believe [Name Redacted]* letters went out at the beginning of August.

And also I -- I have some questions about some documents that the constituents have provided to us suggesting some activity related to

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Chapman Valve as far back as 1945, but maybe it makes more sense to hear the update on -- on what the Department of Energy and Department of Labor have come up with during this intervening time, and maybe then after that we can talk about whether these documents are -- are things that -- that the Board is aware of or not. DR. ZIEMER: Okay. Thank you, Sharon. We'll -- if you'll stay on the line you'll hear some updates, and then if you have additional comments, we'll be pleased to hear those. Board members, you should have the packet that Dr. Wade distributed -- actually two packets. One -- the first part of that packet is a letter from Dr. Wade to Dr. Worthington at DOE. Then there's a copy of Dr. Wade's letter to Pete Turcic of Labor. And then you'll see a copy of Pete Turcic's response relative to Chapman Valve, and a copy of Pat Worthington's response. And then there are some supporting documents as well in the other packet. And I think Pat Worthington is prepared to tell us ba-- basically I think what Labor has said is that if -- if Department of Energy comes up with additional information that would cause

1 them to change the -- the definition of the 2 covered periods and so on, that they would then 3 have to consider that. But let's hear from Pat 4 Worthington on the status of what Energy is 5 doing relative to Chapman Valve. 6 DR. WORTHINGTON: Good afternoon, and thank you 7 again --8 Tip the mike down, if you would. DR. ZIEMER: 9 DR. WORTHINGTON: Just a little bit? Can you 10 hear me better now? 11 DR. ZIEMER: That's good. 12 DR. WORTHINGTON: Is that much better? 13 afternoon, and we wanted to give you an update. If you have the package, it'll be a little bit 14 15 more thorough than -- than my discussion. 16 In terms of the time that it's taking us to 17 answer the question, we certainly don't -- we want to be thorough. We want to be as complete 18 19 as possible and to follow all the leads that we 20 have so we can -- so that was one of the 21 reasons that it took so long to do that. 22 NIOSH asked us to -- to research whether or not 23 contaminated manifolds which -- were 24 transferred from Y-12 back to Chapman -- to the Deen* Street location, and we've be-- we're 25

1 working on that. We queried a number of 2 sources and could not find documentation to 3 support this activity. We have a lot of 4 evidence and documents that it was clear that 5 they were -- purchased valves and manifolds 6 from Chapman Valve. 7 We were also able to substantiate the War 8 Department -- Navy had contracts with Chapman 9 Valve. We also found documentation from the 10 Navy that Deen Street did actually exist. 11 Rad work was for Brookhaven, we did find that 12 in the main Chapman location. We found no 13 evidence of AEC work that took place at the 14 Deen Street location. 15 You have in your package the document that will 16 list all the various sources that we used, and 17 I had mentioned yesterday in an earlier 18 discussion that we also engaged the Office of 19 Legacy Management at the Department of Energy. 20 They have experience and knowledge and depth in 21 doing these kinds of things. They helped us to 22 further research the various documents that you 23 will see listed in the package here. 24 I don't know if you have any other questions or 25 if there are any specific things about this

1 particular activity, but I will -- will go now 2 to sort of -- again, to re-emphasize the 3 conclusion section of the report in terms of 4 what we did find, that each known record 5 collection in the custody of the agency with 6 information about Chapman Valve Manufacturing 7 Company has been thoroughly (unintelligible), 8 and any documents identified were retrieved and 9 reviewed. It's clear that they -- they did 10 have numerous contracts, but we didn't find the 11 evidence that we were specifically asked to 12 look for by NIOSH. 13 DR. ZIEMER: Let me ask, Dr. Worthington, from 14 DOE's point of view, do you then consider this 15 issue closed or is it -- is there something 16 ongoing yet on Chapman Valve? 17 DR. WORTHINGTON: We have nothing ongoing on 18 Chapman Valve. This is the conclusion of our 19 query. Unless there's some other specific 20 question or document that we have not looked at 21 or looked for, we would have to be given an 22 additional source to request. We think we've 23 looked at --24 DR. ZIEMER: Right. 25 DR. WORTHINGTON: -- all the sources, based on

1	the information that we had.
2	DR. ZIEMER: And lacking that, then Labor's
3	position would be as stated by
4	DR. WORTHINGTON: I would have to
5	DR. ZIEMER: Pete Turcic
6	DR. WORTHINGTON: defer to Labor
7	DR. ZIEMER: I believe.
8	DR. WORTHINGTON: on that.
9	DR. WADE: Jeff Kotsch.
10	DR. ZIEMER: Jeff is nodding yes. And so I
11	think with that that was the only open issue
12	on Chapman, was it not?
13	Now as far as I know, the workgroup has has
14	not met further because there was not any
15	additional information for them to review. Is
16	that correct, Dr. Poston?
17	DR. POSTON: Yes, that's correct.
18	DR. ZIEMER: So basically on Chapman we are
19	back to where we were originally, as far as the
20	workgroup's recommendation was concerned.
21	Let me ask Sharon if she has any additional
22	questions at this point.
23	MS. BLOCK: I do actually have a question
24	because I believe it was a constituent had made
25	available to us some documents related to

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Chapman Valve, as I said before, dating back to 1945. So I guess I would have a question for -- for Dr. Worthington as to whether the Department has seen these and -- and whether that influences their decision that there's nothing else to look at with regard to Chapman Valve. These are Stone and Webster Corporation correspondence to the district engineer at Oak Ridge, and they refer to contracts with dates 1944 to 1945, and the contract numbers are the same as the contract numbers listed -- let's see, it's in the last page of the attachment on Dr. Worth-- Dr. Worthington's September 25th letter, the DOE environmental management file room list that lists some contracts for Chapman Valve with the dates 1948, but these Stone and Webster documents that -- that were made available to us have that same contract number but dates 1944-'45.

DR. ZIEMER: Okay, stand by.

DR. WORTHINGTON: We would actually need to see that document. If we could -- could get it we could cross-reference against the list that we have and we could get back to you on that. So if someone can make that available to us, we

1 would -- we would certainly look at it and make 2 sure it's something that was included in our 3 search. It didn't seem obvious from the quick 4 list that I was looking at here. 5 DR. ZIEMER: So it's not clear that you've seen the document that they refer to --6 7 DR. WORTHINGTON: That she's talking about, yes, that's correct, so we would need to look 8 9 at that. 10 DR. ZIEMER: Okay. Thank you. 11 DR. WORTHINGTON: Thank you. 12 DR. ZIEMER: Comment? Yeah. 13 MR. GRIFFON: Yeah, I guess there's other 14 information, too. The interview -- I mean the 15 main source of this, you know, concern about 16 the Deen Street facility came from an interview 17 conducted by SC&A, and in that interview the 18 individual interviewed identified a few people 19 from the Manhattan Project or Stone and Webster 20 -- at least that's my understanding. And I 21 think -- you know, I don't know if that thread has been pulled at all, you know, to see if 22 23 there's any documents that exist for these --24 from these individuals or...

DR. WORTHINGTON: Is your question did we look

1 at the information from the affidavits or 2 interviews to help us target our searches? 3 MR. GRIFFON: Yeah. 4 DR. WORTHINGTON: We certainly looked at that 5 information, used it to help focus our review and to come up with the list of documents that 6 7 we felt we needed to look at. 8 MR. GRIFFON: Okay, and -- and I think she also 9 indicated in the interview that they received 10 parts from, and I -- I don't -- I -- I know you 11 haven't -- just haven't been able to find or 12 confirm that, is that what you -- what your 13 investigation's showing? Received parts from 14 Y-12, not -- not only made parts for -- you 15 know, and they received parts from... DR. WORTHINGTON: (Off microphone) 16 17 (Unintelligible) comment on that. 18 MS. CANO: Right, right. We actually had our 19 Legacy Management staff research, and what they 20 were able to find was that there were Navy 21 contracts for Deen Street facility. However, 22 in querying they looked for Y-12 connection, 23 ORNL connection in regards to anything going 24 back to Chapman. They could not find any 25 documentation to substantiate that.

1 There were many contracts where Oak Ridge did 2 purchase valves and manifolds from Chapman. 3 But actually going back to Chapman, we couldn't find that information. 4 5 DR. WORTHINGTON: We believe that we have pulled the string on all of them, but if 6 7 there's some specific information or specific 8 reference that you're questioning or you 9 believe was not included, please make us aware 10 of that. 11 MR. GRIFFON: Okay. 12 DR. ZIEMER: Okay. Well, it appears that the 13 only -- the only possible one is this one that 14 Sharon has just raised that might be 15 interpreted as an open item yet. This Board 16 was deadlocked, as it were, on -- on the 17 Blockson (sic) issue. 18 DR. WADE: Chapman Valve. 19 DR. ZIEMER: Or Chapman Valve issue, and I --20 it's not obvious to the Chair that we would be 21 well-served to take another vote today, particularly if there's some question on this 22 23 particular document. But let me hear from the 24 Board. What is your pleasure? 25 DR. WORTHINGTON: Well, we do have individuals

1 back there in -- in Germantown, so if they make 2 this available -- if you tell us the number, 3 we'll certainly double-check that. 4 DR. ZIEMER: Yeah, Dr. Poston -- oh, excuse me. 5 MS. CANO: Can I say something else? 6 regards to the contract, it is possible on the 7 '45 that it might have been for purchasing of 8 valves or manifolds. But in regard to actual 9 rad work, that's something we'd have to take a 10 look at. We do have -- we have established the 11 '48 time period for the Brookhaven work, but in 12 regards to the '45 contracts, that's something 13 we'd have to look at. It could be just for the purchase of, you know, manifolds or valves. 14 15 But again, we'd -- we'd have to look at that. DR. ZIEMER: Thank you. Dr. Poston? 16 17 DR. POSTON: I just have a procedural question. 18 Since the period we were considering, the 19 working group was considering, started in 1948, 20 do we have to extend the time period to --21 DR. ZIEMER: Well, not unless there's evidence 22 that it should be extended. I think that was 23 the question at this point. 24 DR. POSTON: But it is clear -- I participated 25 in those interviews and it is clear that at

1 least one, and perhaps more folks, did testify 2 that there were manifolds coming back from Oak 3 Ridge into the facility by rail, and they were 4 transferred to truck and then taken to another 5 facility. They were not -- as far as we could 6 tell from the interviews, not taken into the 7 Chapman Valve's facility under -- that's in 8 question, but there's -- the testimony in the 9 interviews made that clear that they did come. 10 DR. ZIEMER: Uh-huh. 11 MR. GRIFFON: Right. 12 DR. ZIEMER: Thank you. 13 MS. CANO: Right, and that's something we'd 14 have to do is we would have to change the covered time period. We -- if we could find 15 16 that information, we'd provide that to 17 Department of Labor. But the question that 18 came in from NIOSH was to substantiate whether 19 or not there were contaminated manifolds from 20 Y-12 back to Chapman, and so that was what we 21 were querying. 22 DR. ZIEMER: Thank you. 23 THE COURT REPORTER: Could I get your name, 24 please?

MS. CANO: It's Regina Cano.

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1 THE COURT REPORTER: Spell the last name, 2 please. 3 MS. CANO: C-a-n-o. 4 DR. ZIEMER: Okay. Other comments, Board 5 members? What is your pleasure here, do you 6 wish to postpone further action till we trace 7 this last document down, or do you wish to --8 DR. POSTON: Of course as the working group 9 chair I'd like to resolve this issue, but right 10 now we're deadlocked on the recommendations 11 from the working group to the Board. 12 DR. ZIEMER: The deadlock has the effect of turning down the petition, in -- in effect. 13 14 DR. POSTON: Yeah. 15 DR. ZIEMER: Huh? 16 DR. WADE: In essence, yes. 17 DR. ZIEMER: In essence. So unless there is 18 evidence that would cause Board members to 19 change their votes, then that's where we are 20 and the Chair's suggesting perhaps this 21 additional piece of evidence might have some 22 impact. Otherwise, I -- unless someone wishes 23 to call for a new vote, why --24 DR. POSTON: Yeah. Well, this was just passed 25 around. I haven't had a chance to read it,

1 obviously. 2 MR. GRIFFON: Right. 3 DR. POSTON: I got it a couple of minutes ago, 4 so perhaps we can talk about this tomorrow if 5 we get a chance, or... Okay. 6 **DR. ZIEMER:** Or at the next meeting. 7 DR. POSTON: Or at the next meeting. 8 DR. ZIEMER: Jim Lockey. 9 MR. GRIFFON: And I don't know that the 10 petitioners have seen this document, either, so 11 we -- we have that same question of --12 DR. ZIEMER: Okay, Jim Lockey? 13 DR. LOCKEY: And can -- can the additional 14 information that the petitioner has be 15 researched by Department of Labor by tomorrow, 16 or is that not possible. 17 DR. ZIEMER: Department of Energy? 18 DR. LOCKEY: Department of Energy, excuse me. 19 DR. ZIEMER: That -- I think that's putting a 20 fair burden on them to try to research that in 21 one day. Plus if the petitioners have not seen 22 any of this, that may be an issue as well. 23 MR. PRESLEY: Well --24 DR. ZIEMER: Okay. Robert? 25 MR. PRESLEY: Did you all check any of the

1 shipping documents, or were you able to find 2 any shipping documents at Y-12 where these 3 things did go back or where they did drop --4 where they were received down there, what they 5 were? MS. CANO: My unders-- my understanding is that 6 7 we found purchase orders for the manifolds and 8 valves, so they were going from Chapman to Y-9 12, but nothing going from Y-12 back to 10 Chapman. 11 MR. PRESLEY: Okay. 12 DR. ZIEMER: Okay. Jim? DR. MELIUS: Well, you've sat down, I'll --13 14 have another question. I mean does your 15 investigation include going to the field at all 16 and asking people questions or something? I --17 is this all just a paper exercise in 18 Germantown? I'm just trying to understand what 19 you -- what you do, and I'm sorry, I missed the 20 session yesterday and I may have -- may have 21 talked about this. I'm -- does... 22 DR. WORTHINGTON: Our responsibility was to 23 retrieve the documents, research and look for 24 the documents. We certainly take the 25 information that we get, if there've been

1	witnesses or affidavits or whatever that would
2	lead us and help us to focus our our actual
3	document reviews in in a better way. But
4	we have not, Gina, been engaged in any
5	interviews. We've been looking we were
6	asked to look for documentation and so we used
7	the in the information we received about
8	witnesses, workers, to help us to focus those
9	reviews, to help target where to look for these
10	actual documents.
11	DR. ZIEMER: Thank you.
12	DR. MELIUS: Thank you.
13	DR. ZIEMER: Okay.
14	DR. WADE: I would like us to spend just a
15	moment and be very specific about what's going
	3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3
16	to happen next so we don't lose this again. As
16 17	
	to happen next so we don't lose this again. As
17	to happen next so we don't lose this again. As I understand it, Senator Kennedy's
17 18	to happen next so we don't lose this again. As I understand it, Senator Kennedy's representative has documents that have been
17 18 19	to happen next so we don't lose this again. As I understand it, Senator Kennedy's representative has documents that have been provided to them that raise issues about the
17 18 19 20	to happen next so we don't lose this again. As I understand it, Senator Kennedy's representative has documents that have been provided to them that raise issues about the period 1942?
17 18 19 20 21	to happen next so we don't lose this again. As I understand it, Senator Kennedy's representative has documents that have been provided to them that raise issues about the period 1942? MS. MUNN: '44 and '45, I thought.
17 18 19 20 21 22	to happen next so we don't lose this again. As I understand it, Senator Kennedy's representative has documents that have been provided to them that raise issues about the period 1942? MS. MUNN: '44 and '45, I thought. DR. WADE: '45, so we need to make sure that

1 spent on that would be -- would be wise. 2 DR. ZIEMER: My understanding is that DOE would 3 examine those to determine if they impacted on 4 the covered period. Is this the issue we're 5 looking at? Or is it both the covered period and whether or not some manifolds actually went 6 7 to the site or left contamination at the site 8 during the transfer, I suppose is part of the 9 issue. Perhaps both of those, and the -- the 10 transfer part might be easy to come -- more 11 easily identified than this contamination 12 issue, which would -- was there an indication 13 that these were contaminated, John? 14 DR. POSTON: There was no indication that they 15 were contaminated, as far as I remember. 16 Arjun's here and John's here, maybe they 17 remember. But during the site survey, when the 18 site was being decommissioned, there was at 19 least one sample that was identified as 20 slightly enriched uranium. And all the 21 documents show that what they did at the 22 facility was machine uranium metal that was not 23 enriched, so --24 DR. ZIEMER: So the question of where did that

come from.

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DR. POSTON: Yes. And when we discussed this with folks at the facility, they said well, we did have these manifolds come in. And then I guess the -- we started putting two and two together and said well, these manifolds probably came from Y-12, probably came from the Calutrons, and that could be the source of the enriched -- slightly enriched uranium. There's also a question of whether it really was slightly enriched uranium or not, or was a false positive.

DR. ZIEMER: Okay.

MS. CANO: I have a question for clarification purposes on our part. It was -- it's our understanding what -- DOE's role is to designate the AWE, which we have with Chapman Valve. We were under the understanding that we were trying to figure out whether or not Deen Street performed radiological work, and that's the basis for AWE designation. And what we've come up with basically is that we were not able to substantiate that that actually took place, that the Navy actually did have contracts with the Deen Street facility. And based on our search, we could not find any documentation to

1 lead us back to Deen Street pertaining to 2 radiation -- rad work. Chapman is designated 3 as an AWE from '48 to -- I just blanked on it -4 - beginning in '48 based on work done at 5 Brookhaven, and it was -- it was uranium. so I'm -- I'm -- I guess what I'm thinking is 6 7 that some of the contracts that they might have 8 was for the purchase of maybe manifolds or 9 valves, which indeed were just steel-based. 10 They were not radioactive whatsoever. So I'm 11 just trying to figure out what -- what we -- I 12 mean we can look at the document. We will --13 we will do that. But it's just trying to 14 understand what else you want us to do. 15 DR. ZIEMER: Yes, thank you for raising that 16 point. I don't think we were asking about Deen 17 Street per se because that would not affect 18 this petition. 19 DR. POSTON: Right. 20 This petition is unique to the DR. ZIEMER: 21 main -- other facility. And I think --22 MR. GRIFFON: Unless that was considered part 23 of the Chapman facility, that's -- that's a 24 question, too, (unintelligible) --25 DR. ZIEMER: Well, but it was -- physically it

1 would be a separate facility, as I understand 2 the way these get defined. So even if it's 3 part of Chapman, it would require a separate 4 petition, I believe -- as I understand it -- if 5 that were the case. MR. GRIFFON: Well, somebody explain that to 6 7 I'd like to understand that. 8 MR. ELLIOTT: It's my understanding the Deen 9 Street facility is a Chapman Valve facility. 10 DR. POSTON: Yeah. 11 MR. GRIFFON: Right. 12 Okay? But the Chapman Valve MR. ELLIOTT: 13 facility that's been designated an AWE, I don't 14 know which street it's on -- I don't recall 15 that -- but that is the contiguous AWE 16 facility. Deen Street is not part of that. 17 DR. ZIEMER: And even if they had received some material, it doesn't automatically become part 18 19 of this because of physical location, or would 20 it? 21 MR. ELLIOTT: What -- what we asked DOE to do 22 was to look at the Deen Street facility and 23 determine whether or not it should become an 24 AWE. And the evaluation report you have before 25 you for this SEC petition only deals with the

1	Chapman Valve AWE facility. So we were trying
2	to seek out from from DOE whether or not
3	Deen Street should be designated an AWE in of
4	itself.
5	DR. ZIEMER: If it were, does that affect this
6	petition? That's sort of what I'm asking.
7	MR. ELLIOTT: It would not. It would have to
8	be another petition for that facility.
9	DR. ZIEMER: Right. So the outcome of this
10	question doesn't, in a sense, affect this
11	petition. Mark, do you think it
12	MR. GRIFFON: Yeah, I guess I'm going a little
13	uncertain, but
14	DR. ZIEMER: Unless the unless the
15	contamination issue
16	MR. GRIFFON: My quest
17	DR. ZIEMER: enters into it.
18	MR. GRIFFON: Yeah, my question was if and
19	this gets a little murky for me between DOL and
20	NIOSH's function in you know, defining the -
21	- but if in a normal situation, if NIOSH
22	finds that the time period for a petition
23	should be extended, then they'll extend it.
24	Now here the time period's set by DOL already.
25	Right?

1	MR. ELLIOTT: The time period of a covered
2	facility
3	MR. GRIFFON: Of a cov I'm not talking about
4	a covered facility, though. I DOL
5	establishes that, I understand that.
6	MR. ELLIOTT: Yes.
7	MR. GRIFFON: But in an SEC, if you find reason
8	to believe that, you know, you should extend it
9	longer than the than that identified in the
10	petitioner (sic), you'll you'll self-
11	identify that sometimes.
12	MR. ELLIOTT: If it's within the bounds of a
13	covered facility designation, yes
14	MR. GRIFFON: So here we have
15	MR. ELLIOTT: we can do that.
16	MR. GRIFFON: a circumstance where we're
17	saying, you know, there may be other work prior
18	to the defined time period
19	MR. ELLIOTT: But the AWE does not cover 1945.
20	It starts at '48. And I'm not sure when this -
21	_
22	MR. GRIFFON: But it's the same facility, and
23	if DOL changed their designation, then the same
24	I don't know, I I
25	DR. WADE: Right. Well, I let's we let's

1 clear them up one at a time. 2 MR. GRIFFON: Yeah. There seems to be information that -3 DR. WADE: 4 - that goes to the issue that the period should 5 be extended from '48 back to '45. information needs to be looked at by DOE and a 6 7 judgment made, so that's one issue we've heard 8 today. 9 The second issue we've got is that there's this 10 elevated reading that appears that might have 11 resulted from something on its way from 12 somewhere to Deen Street being off-loaded at 13 the Chapman Valve covered facility, and that's 14 the other part of the issue. 15 What we need to hear from DOE, if this 16 information comes from the good Senator's 17 staff, then you can look at it and determine 18 whether or not it goes to the fact that the 19 covered period should be extended back to '45. 20 That's something you can do? 21 MS. CANO: We would be happy to take a look at 22 In regards to the DOE designation, DOE has 23 already designated Chapman Valve. In regards 24 to extending the covered time period, that's

something that Department of Labor would have

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1 to -- to make that decision. 2 DR. ZIEMER: But they would do that based on a 3 recommendation from you, or from whom? 4 MS. CANO: They -- Jeff? 5 DR. WADE: Well, let's --6 MS. CANO: Sorry, I can't speak for Department 7 of Labor. 8 DR. WADE: So if someone presents a bit of 9 evidence that -- that argues strongly that 10 there -- that the period should be extended to 11 1945 --12 MR. KOTSCH: Right, that's --DR. WADE: -- do you act unilaterally on that? 13 14 MR. KOTSCH: Yeah, right, Department of Labor's 15 responsible under the program for the --16 additional covered period, if you want to say. 17 Usually it's not on an individual basis based 18 on evidence provided by, you know, a particular 19 claimant. But yes, we extend -- or determine 20 the covered periods. 21 In -- in the case of like Deen Street, if that were an issue, DOE has the responsibility to 22 23 determine that that's an AWE, and then we would 24 determine the covered period for that 25 particular facility.

1 DR. WADE: But that hasn't happened to this 2 point. 3 MR. KOTSCH: Yeah. 4 DR. WADE: So what we have is the information 5 from Senator Kennedy's staffer that potentially 6 goes to 1945, so that needs to come to you to 7 look at. 8 MR. KOTSCH: Yeah, I mean I would recommend it 9 gets sent -- gets sent to Pete then. 10 DR. WADE: Send it to... 11 MR. KOTSCH: Send it to Pete Turcic. 12 DR. WADE: Send it to Pete Turcic. So if that 13 information gets to you, then that's an action 14 item you would look at and could report back 15 on? 16 MR. KOTSCH: Yes --17 DR. WADE: Okay. 18 MR. KOTSCH: -- I guess (unintelligible). 19 DR. WADE: So that's one resolved. Jim? 20 DR. MELIUS: I also believe that there are 21 people that -- former workers from the facility 22 that have information from that, and I would 23 hope -- about these early contracts, and I 24 would hope that somehow DOE can manage to 25 interview them or obtain information from them.

1 Our contractor, SC&A, has talked to them, I 2 believe. I'm not sure if the -- the workgroup 3 talked to all of them or whether NIOSH staff 4 has, but -- but certainly that information can 5 be made available and I think can provide some helpful information for this follow-back. 6 7 DR. WADE: And this is --DR. ZIEMER: Arjun, did you have a comment on 8 9 that, or John? 10 DR. WADE: Arjun was trying to get out. 11 DR. ZIEMER: Oh, no -- no comment? 12 DR. POSTON: Well, the -- the workgroup -- the 13 workgroup in its entirety did not participate, 14 but I did participate with Arjun and John. And 15 Arjun wrote a report which was the results of 16 the interviews that was distributed to the 17 workgroup, so... 18 DR. ZIEMER: Arjun? 19 DR. MAKHIJANI: Dr. Ziemer, the only thing I 20 was going to say is that the published 21 interview is redacted for privacy, and there 22 might be a couple of things in the redactions 23 that might be helpful to the Department of 24 Energy, and we'd be happy to make that 25 available. I don't know what the process is,

1 but --2 DR. ZIEMER: They can have an unredacted --3 DR. MAKHIJANI: Do you -- do you have the --4 I'm not sure, you have the unre-- okay. 5 MR. GRIFFON: Okay. 6 DR. WADE: So to get our agencies straight, 7 though, the information -- and this information 8 is going to the Department of Labor to look at 9 the issue of extending the covered period. 10 Okay. 11 Now we still have the contaminated manifold 12 issue to deal with if we want to. Or is that 13 done now in everyone's mind? 14 MR. GRIFFON: Well, it's not done. 15 DR. ZIEMER: Well, I'm not sure what we'd do 16 with that, other than -- that -- that issue was 17 the one that triggered this whole question, 18 where did the U-235 come from. Would that in 19 any way affect the Chapman Valve findings, the 20 fact that there was this contamination at the 21 transfer point? 22 MR. GRIFFON: It -- it only affects it in -- in 23 my -- in my mind, anyway, it only affects it in that it's something that we -- we can't 24 25 explain. And I agree with John that -- that we

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-- you know, we -- we've also heard that it could be due to laboratory error. I mean it could be within the error of the lab. But it's something that we haven't been able to explain. And with this additional -- very detailed interview, I might add, of -- of the accounts of materials coming from Y-12, it -- it raises the question of was there something else going on at Chapman. I mean we -- we didn't even know about Deen Street -- I still think of Deen Street as part of the Chapman facility, quite frankly, just wasn't known when they put -when DOL put their site list together. But now this raises a question of were there other activities. I think if we knew what they were and they were -- they were dealing with some slightly enriched uranium and that was it, I don't think it would change the conclusion. But it just raises that question of what else went on there and can we make sure we have covered the breadth of the operations that were going on at Chapman.

DR. ZIEMER: Well, it appears to me that we've gone as far as we can today on this, that we need to hear the outcomes --

1 MR. GRIFFON: Yeah. 2 DR. ZIEMER: -- and then go from there. you have an additional --3 4 MR. ELLIOTT: I just want to be --5 DR. ZIEMER: -- comment or --6 MR. ELLIOTT: I want to be --7 DR. ZIEMER: -- advice for us? 8 I want to be clear for the record MR. ELLIOTT: 9 that we have provided Department of Energy and 10 Department of Labor the non-redacted 11 information that has been assembled both from 12 our levels of effort and SC&A's levels of 13 effort, and we've provided that to them in an 14 unredacted form. 15 The Deen Street facility goes to whether or not 16 it should be designated as an AWE, and that 17 goes to DOE to decide. 18 The extension of time for the current AWE at 19 Chapman Valve goes to DOL, and I think if --20 you're right, if the -- if Senator's staffer 21 can provide that information to DOL, they can 22 look at that and determine whether or not the 23 covered period for the current AWE is 24 appropriate and accurate, or needs to be 25 adjusted.

I don't know what you do about the Deen Street facility from this point. I think DOE is still looking -- as you've heard them, they're still looking for information that would tell them whether or not it should be an AWE or not. That's -- I just wanted to (unintelligible) to say to you that we have given up everything that we have, in an unredacted form, to the right Departments for the right decision-making.

DR. ZIEMER: Okay, thank you very much. Yeah,
Brad?

MR. CLAWSON: Well, and I just wanted to go back to some of the comments that have been made by some of the petitioners and so forth, my understanding of it is they -- they actually felt that the Deen Street was just part of Chapman Valve, it was doing another process of Chapman Valve. And this is -- this is what has -- I -- I guess convoluted some of the things that have gone on there, but in the petitioners' eyes also, too, Deen Street was just part of Chapman Valve and they had processes that were going on and they were showing -- they were telling us, you know, and

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they -- they've been pretty good to be able to show us so forth and that -- that's where I think some of the confusion comes in.

DR. ZIEMER: Okay, thank you. Jim Lockey, do you have an additional comment? No, okay.

MR. GRIFFON: I only had one -- one last thing.

DR. ZIEMER: Yeah.

MR. GRIFFON: One -- one last thing, a little -- it was the other aside that we asked about follow-up on and that was the documentation from -- from later activities, the -- when this facility was cleaned up and when they disposed of it did anyone find any more records related to the cleanup process or the waste that was shipped or any of that information. I -- it might be in your report. I'm glancing at it while I'm trying -- while we're trying to talk here, but did DOE or NIOSH find anything related to the cleanup -- I think it was '90 or '92 when they -- they did the cleanup process. We have the -- the cover report, but is there any detail. And my curiosity on this is if we see a certain number of grams of U-235 manifested, that at least supports this question of was there, you know, really U-235

1 there -- maybe it supports it, anyway. 2 DR. NETON: We don't have any information to 3 add on that time period. That's still listed 4 as reserved, I think, in the evaluation report. 5 It was not specifically evaluated as part of this SEC petition. 6 7 MR. GRIFFON: Oh, but I had asked for the 8 information so that it could maybe answer some 9 questions about quantity and type of materials 10 11 DR. NETON: Well, we don't have any more 12 information to offer at this time. Okay. All right. 13 MR. GRIFFON: 14 This Board is not in a position of DR. ZIEMER: 15 tasking either the Department of Energy or the 16 Department of Labor to do things, but I --17 MR. GRIFFON: Yeah. 18 DR. ZIEMER: -- I do want to ask if both Labor 19 and -- and Energy would be willing, if you get 20 these doc -- I think you're going to get the 21 documents from the Senator's office, and if 22 you'd be willing to report to us at our next 23 meeting what you find from those documents, and 24 if it has any impact on Blockson, either in 25 terms --

1 DR. WADE: Chapman. 2 DR. ZIEMER: -- or Chapman, I get -- get off of 3 Blockson here -- Chapman in terms of either 4 time period or the -- or the Deen Street 5 location, that would be helpful to us. 6 Basically this is going to leave things hanging 7 for another meeting. But in the absence of 8 that, I think we're going to be at the same 9 place anyway. We're not going to resolve that 10 today, and perhaps this additional information 11 will help us come to some kind of closure if 12 you'd be willing to at least tell us what 13 you've learned 'cause clearly the Senator has 14 indicated that they're going to provide this --15 these documents to you. So I assume you'll 16 need to follow up, in any event, since the Senator has asked that that be done. 17 18 The Board has a call on December 6th 19 as the next time the Board will be together. 20 DR. ZIEMER: If -- if that information is 21 available by then, that would be good. 22 Otherwise we'll have to wait to our full 23 meeting. 24 DR. WADE: Which is January 8th, 9th, and 10th 25 of next year.

1 DR. ZIEMER: Okay. Jim, you have a comment 2 here? No. 3 DR. MELIUS: My only comment is that it's not 4 clear from the DOE report that -- that the 5 interviews and so forth that NIOSH made available to them were reviewed. They may 6 7 have, but just in glancing through, I don't --8 I don't see that and so... 9 DR. ZIEMER: But that could be clarified. 10 DR. MELIUS: May be confusion, but again, all 11 the reason to let's have some follow-up and 12 give that some time. 13 DR. ZIEMER: And Jim Lockey? 14 DR. LOCKEY: Just for my own clarification, by 15 the next meeting are we going to get additional 16 information about Deen Street? 17 DR. ZIEMER: What I've asked is that both -- if 18 both Labor and DOE are willing to tell us what 19 the outcome is when they see the documents from Senator Kennedy's office, if that -- in their 20 21 judgment -- has any impact on either the 22 location designation, Deen Street, or the time 23 of the covered period. 24 DR. LOCKEY: Okay, so if there's -- if there's 25 no additional information on Deen Street, then

1 2 DR. ZIEMER: Deen Street is -- as I understand 3 it, is not a covered facility. Is that -- is -4 - it's -- it's differentiated, even though some 5 of the workers may regard it as part and 6 parcel, Brad, of the same thing, apparently 7 it's not covered as a physically separate 8 entity -- as I understand it, and I think 9 that's how NIOSH understands it and -- and 10 Energy, as well. And Labor, as well. Okay. 11 So --12 DR. WADE: But Dr. Lockey's -- looks puzzled. 13 I think the issue is if there were things going 14 on at Deen Street that caused radioactive 15 material to be shipped through the covered 16 facility, that's important for us to know. 17 DR. LOCKEY: I think that -- you said it much 18 more articulately than I could. I think that's 19 the question and I think --20 DR. ZIEMER: That's what we're hoping to learn, 21 if that in fact, in Energy's opinion, changes 22 the status of Deen Street -- if they're able to 23 make that judgment from the documents that they 24 get. 25 DR. LOCKEY: I think the real question is

1 what's -- what's really happening at Deen 2 Street. That's really the most important 3 question. Then a second question, do we extend 4 it back to 1945, and that's a --5 DR. ZIEMER: Yeah, those are the --6 DR. LOCKEY: -- DOE/DOL decision. 7 DR. ZIEMER: -- two issues, right. 8 DR. WADE: The two issues. 9 DR. ZIEMER: We're saying if these documents 10 shed light on either of those two issues, then 11 that would be helpful to us, perhaps in coming 12 to some kind of closure on this. 13 With that I'm -- I think we're going to move on 14 unless someone has some additional words of 15 wisdom. 16 DOW CHEMICAL 17 DR. WADE: Dow. 18 DR. ZIEMER: Dow Chemical -- this is Dow 19 Madison, more specifically. We have some 20 documents -- Lew, can you step us through the 21 documentation that -- the response to your 22 inquiries? 23 DR. WADE: No, it was your inquiry, but --24 DR. ZIEMER: Well, my -- okay, I --

DR. WADE: -- that the Board wrote to the

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1 Secretary of HHS. 2 DR. ZIEMER: Tell me what I did, will you? 3 DR. WADE: Well, you did well, as always, and 4 you led the way. The Board -- the Board, under 5 Dr. Ziemer, wrote to the Secretary of HHS 6 asking the Secretary to -- to interact with the 7 other agencies relative to the questions that 8 had been raised on Dow Madison. The Secretary 9 asked Dr. Gerberding, the Director of CDC, to 10 reply to the Board's letter, which she has now 11 done and that's in your materials. 12 I also am aware of the fact that our colleagues 13 from DOE are here to make some additional 14 comments about Dow Madison. Dr. Branche is 15 passing out the -- the handouts that you 16 brought to us. And I also believe that our 17 friends from the Hill wish to make some 18 comments concerning Dow Madison. 19 DR. ZIEMER: Okay, we can start with Robert 20 Stephan again. Robert, do you want to --21 DR. WADE: And Deb Detmers. 22 DR. ZIEMER: -- make your remarks now -- or 23 Deb, or both? Do you want to comment at this 24 point? 25 MR. STEPHAN: We're just going to come up

together, but I think it may be more useful to just go through the discussion about the site and then we'll comment --

DR. WADE: Okay.

DR. ZIEMER: Sure.

MR. STEPHAN: -- after. If our -- if our comment is relevant as your discussion goes, we'll jump up, but I think we want to hear the discussion first, if you don't mind.

DR. WADE: Okay.

DR. ZIEMER: Well, one of the -- the issues on -- the Dow issue really had to do with the extension of the covered period, and there was -- the petitioners raised some issues which we're all aware of that suggest that perhaps it needed to be looked into. And so the request went to the Secretary's office to ask both Labor and Energy to look at certain documents that might be considered in -- in changing the covered period. So we -- we did basically get replies from Pete Turcic and Pat Worthington, and the letter from -- from Dr. Gerberding on behalf of the Secretary indicating that the request had been made to Labor and DOE.

currently are not in a position to change anything unless Department of Energy so designates, as I understand it -- or so suggests. And Department of Energy has been looking into some of the documents. I think that's still in process. And Pat, I don't know if you have any comments on -- tell us where you are -- it's my understanding this is still ongoing.

DR. WORTHINGTON: It is still ongoing. We want to make sure that we do the best job we can in terms of answering the questions. We've engaged other organizations to help us out. We've actually -- we went to the FBI to ask them to look at those documents. They have some unique techniques, I understand, to be able to -- to see and -- and describe to us what's there. We have not yet heard back from them, but we're pleased that they accepted the assignment and they would do this for us on behalf of the workers.

Also we've engaged NNSA within the Department of Energy for them to explain their process to make sure that we understand what was going on at the various sites to see if that could in

1 fact provide some additional insights. 2 We've also contacted the -- the lawyers from 3 the contractor's side of the house, from Dow 4 Chemical, and we -- we've had some exchange and 5 we're -- we're hoping to be able to close with them to get more information from that side. 6 7 And also there were a number of FOIA requests 8 related to these activities. We -- we 9 expedited those things and also looked at -- at 10 those documents, as well. And so we are at 11 this point still trying to get closure and we 12 hope to hear back from the working group, with 13 NNSA, back with the lawyers, and also to see if 14 there's something on those documents that will 15 give us some -- some real insights. And so as 16 soon as we have that, we will report back to 17 everybody. 18 DR. ZIEMER: Okay. 19 DR. WORTHINGTON: Thank you. 20 DR. ZIEMER: And the timetable on that is 21 probably uncertain. DR. WORTHINGTON: It's uncertain because --22 23 DR. ZIEMER: We will hear from you --24 DR. WORTHINGTON: -- the FBI didn't --25 DR. ZIEMER: -- when you have something.

1 **DR. WORTHINGTON:** -- give us a schedule. 2 Thank you. 3 DR. ZIEMER: Board members, do you have any 4 questions regarding this issue? We don't have 5 an outstanding action to take on Dow unless the time period is extended --6 7 DR. WADE: Correct. 8 DR. ZIEMER: -- in which case then that would 9 have to be considered. So this is, in a sense, 10 kind of a pending issue till we see what 11 Department of Energy learns from this 12 investigation. And again, we appreciate the 13 input that -- that your office has had, Pat, to 14 follow up on this, to pull -- pull the strings 15 and give us some level of -- of confidence as 16 to what was or wasn't done, so -- and now we'll hear from both Robert and Deb. 17 18 MS. DETMERS: Yeah, hi, welcome to Illinois. 19 I'm Deb Detmers, district director for 20 Congressman Shimkus -- and I'll spell that for 21 you, it's Debra, D-e-b-r-a, Detmers, D-e-t-m-e-22 r-s. And I'd like to introduce two people that 23 rode up with me this morning, Homer Simmons --24 Homer, you want to stand up? And Homer is a

former employee of Dow who's had a pending

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claim since 2001. And also Mr. Bill Hoppe -you want to stand up, Bill? -- who's also had a pending claim with Dow Chemical since 2001, so I do want to state that they do have pending claims for six years. I do want to point that out. Yes, that is six years that they've had pending claims, and we've been working on it. Neither of them will qualify under the current SEC as both of them started working about six months after the time frame, so neither of them are qualified under the pending claim. Just one comment that Robert and I do want to bring up, and I think both of us do want to bring this up, is -- we do have a question about worker testimony. We have provided a great number of documents to this Board and to Department of Energy. And these are documents that we have produced ourselves, including the document that's now at the FBI. documents that, through Dr. McKeel's efforts and through our office, working with Senator Obama's office -- and it's really taken a village and the law firm that we work with -that works with us on a pro bono basis -- we have provided boxes and boxes of documents, and

all of these documents have not gotten us to the point that we need to be on for this extended period.

What we do have -- 11 workers, at a minimum, that have provided worker testimony that all matches. There is no worker testimony that says anything different than what those 11 workers say. We have worker testimony that we've taken. We have worker testimony that the working group has seen. At what point -- I guess the question comes -- do we believe worker testimony? At what point does worker testimony get taken at face value, and who makes that final decision? That is, I guess, my stan-- my question.

I do want to state one more time, these guys rode out with me to Ohio. They did not ride out with me to Denver, but they did ride out with me to Ohio. They're here again with me in Naperville. They come and see me all the time, and there are lots of guys like them, with cancer, that I see every day. And my question is, at what point do we take worker testimony - that's not been contradicted -- at face value? And that's my question.

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MR. STEPHAN: That is -- that is exactly the issue from Senator Obama's perspective. He's in total agreement with Congressman Shimkus. But to take that just a step further, at what point do we take worker testimony when there is no document to disprove what they say? it's an important distinction between no document which proves what they say and no document which disproves what they say. And so, you know, our understanding -- correct us if we're wrong -- is that the whole purpose of the SEC is that when there are no documents, there's -- there's no, you know, exposure data, there's no monitoring data to do a DR, that then we go the -- do go the SEC route. So here we are where we have a document being examined by the FBI. Who knows what the conclusion will be. Either it will -- either it will prove our point or it will simply not prove what the workers have said. It is not a document that's going to be a smoking gun to show that what the workers have said is not accurate. So, Dr. Ziemer, we would, you know, inquire to the Board how -- how can we resolve this question? Because in the Senator's mind and

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the Congressman's mind, this is not a question that simply can continue to be resolved on a case by case basis. What -- what is the guiding, you know, theory that the Board uses to evaluate worker testimony? Is it -- is it a decision that simply is up to each Board member and we come to a vote and we see how it's evaluated? Or is it something that the Board can -- can reach consensus and say when we have worker testimony and it cannot be disproved by NIOSH, DOE or DOL, then -- then the Board will take the worker testimony at face value. And then further, could the -- could the Board, realizing that you guys are -- are just dealing with NIOSH and not DOE or DOL, could the Board send a letter to the three agencies -- NIOSH, DOE and DOL -- and ask them what their practice is or how they evaluate worker testimony when there are no documents of any kind that So two questions, one about the Board and one about can we inquire to these agencies. DR. ZIEMER: Well, first of all, on the particular issue that we're talking about which is the designation of the facility, this Board

in fact does not get involved in that directly. We have inserted ourselves into it in this letter to the -- to the Secretary. But in reality, it is outside of our purview. Both Energy and Labor have described in their letters how they weigh worker testimony and affidavits -- I believe it was in both; I know it was in Pete's and I think, Pat, you may have addressed it as well. But in any event, my understanding is -- and Energy and Labor will have to speak to this -- but in the case of this designation, the ball is in their court in terms of how they weigh the worker testimony vis-a-vis the other documents that are examined and so on.

MR. STEPHAN: Okay.

DR. ZIEMER: As far as worker testimony for this Board in terms of say dose reconstruction, we don't have a cut and dried rule that says worker testimony counts a certain percentage.

I think Board members weigh -- weigh this in a sense individually. We -- I think we -- we try to take worker testimony seriously and if -- if it gets ignored, we raise the issue with the agency and say basically -- for example, it's

1 one of the questions that has come up in some 2 of our dose reconstruction audits if -- if we 3 raise the question was the worker's testimony 4 taken into consideration in the dose 5 reconstruction. That question has been asked a number of times in the audits. So I think -- I 6 7 think we try to do that. But this particular 8 issue, I don't think the Board is directly 9 involved. 10 Others may add to that, but --11 MR. STEPHAN: But... 12 DR. ZIEMER: We -- we -- as I say, we have sort 13 of inserted ourselves outside of our --14 MR. STEPHAN: Right. 15 DR. ZIEMER: -- charter, as it were, and it --16 to -- to get involved in this and both DOE and 17 Labor have been shall we say courteous enough 18 to say even though it's not your business, we 19 will respond to it. 20 MR. STEPHAN: I'm not sure that we have a 21 response from DOL -- maybe Jeff can -- from 22 DOL. Maybe Jeff can help us as to -- and maybe 23 that's in this letter; it's a pretty long 24 response --25 DR. ZIEMER: There was a letter from --

1 MR. STEPHAN: -- as to exactly how DOE (sic) 2 weighs worker testimony. 3 DR. ZIEMER: It may have been a letter to Dan 4 McKeel. 5 MR. STEPHAN: Right. DR. ZIEMER: Because I know that Dan has asked 6 7 that ques-- Dr. McKeel, are you here? Yeah, I 8 -- I know that you have asked that question of 9 both Labor and Energy, did you not, and --10 DR. MCKEEL: Yes, sir. 11 DR. ZIEMER: And you may have not been 12 satisfied with their answer, but there was -- I 13 know that Pete gave a response. I couldn't 14 remember if Labor did or not -- or if Energy 15 did or not. 16 DR. MCKEEL: Well, my -- my view of it is that 17 that may be considered a partial answer, but 18 what I'd proposed to Pat Worthington yesterday, 19 and still propose, is what's needed is a very 20 direct question such as Robert just posed, and 21 a very direct answer. And that really has not 22 been forthcoming. I would say the answer I got 23 from Labor was convoluted, and it's a very 24 simple question and I think Robert posed it 25 quite well. And you know, Pat said yesterday

that maybe the DOE lawyers were going to have to address that and I said that would be great, let's put the question to them in -- a couple of sentences is all it would take, and let's get a similar kind of answer back in a couple of sentence from their lawyers, and then we can present it to the Board and show them. So I -- I think that's an important issue.

DR. ZIEMER: Thank you.

MR. STEPHAN: And Dr. Ziemer, I just want to draw -- draw a distinction. I mean we -- we respect the -- the role and the responsibilities of the Board and NIOSH and Labor and DOE. But there is a -- a much -- there's a big difference between asking these agencies if they have weighed worker testimony and asking them what weight they give worker testimony when there are no documents that -- that disprove what the workers have to say. So it's an important distinction.

The position of the Senator is that it -- it would be preferred if the Board would adopt a guiding principle as to how you address this issue as a Board if you can reach consensus about how to do that. And then it goes to the

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issue of what does -- you know, knowing that the Board does not have purview over DOL or DOE, regardless, how do -- how do you evaluate when deciding to vote on an SEC what their decision-making process was relative to worker testimony. So I guess that's a question. - maybe you can answer it, maybe you can't. We're not trying to put you on the spot; we just don't know. Do we -- do you accept DOL --Department of Labor and DOE at their -- at face value when they say that they have accepted worker testimony and you go with that, or do you not? Is it a case by case basis, and is there room to discuss that the Board would -would take up this notion of trying to reach consensus about how we deal with worker testimony -- is -- ra-- rather than always going down the route of it's up to each individual Board member. And if we can't reach consensus, that's obviously how you would have to proceed, but -- but there may be consensus here about how you deal with -- with worker testimony when there's nothing to disprove what they say.

DR. ZIEMER: Well, should we -- let's -- let's

say we could reach such a consensus -- we may or may not be able to do that, but should we be able to, I'm not sure that would have any bearing on what either DOE or Labor actually does because we cannot impose our view of that on them. And -- and once -- once they make a determination of let's say an AWE, we may or may not agree with it, but I think -- and I'll ask others -- it's like many other things in this law.

We -- we are mandated, in a sense, to proceed as designated in the law. For example, we're not in a position to say well, I don't care what Congress said about the 22 cancers; we're going to use a different number or a different list. So -- so we may disagree on a number of things, either technically or philosophically, but in a sense are bound by sort of the boundaries that are put around us. Sometimes -- sometimes to our dismay and sometimes to -- maybe we are in agreement with, but -- so I'm being a little evasive here 'cause I don't know fully how to answer your question. Other Board members may have some views on that, and Dr. Lockey can -- can articulate something here.

DR. LOCKEY: In relationship to SEC petitions that the Board reviews, there are -- rightfully so, there are DOE worker representatives -- I mean workers on this Board to -- to help I think the Board as a whole understand the job tasks involved when working in the Department of Energy facilities.

Second of all, I -- when we heard NIOSH review how they go through an SEC petition, they have outreach programs where they -- as I understand, workers come in and they interview workers and they take notes and they have affidavits to review and NIOSH reviews those with the Board as part of the review process. And so when we look at an SEC petition, there's opportunity to look at those -- the things that workers are saying. And when an issue is raised such as with Chapman Valve about the Deen Street facility, our -- we go back and say we need further clarification about that.

I don't know if that helps or not.

MR. STEPHAN: No, it -- no, it's good to know.

Thank you. It -- it sounds like what we need to do then is -- Jeff, maybe if you guys are willing, and Pat is -- to inquire to both DOE

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and DOL as to -- with this specific question. And if you would -- would not mind, is providing a specific answer as to how worker testimony is weighed when there are no documents to disprove what they say. that is the specific question. There's no other question. That's the specific question. And to have both agencies respond in turn, because you both have different roles and responsibilities, we respect that, but the -the document -- the letter you reference from Mr. Turcic is not related to this question, and that question is not answered. This is a very important question for us related to Dow -- a very important question.

DR. ZIEMER: Right. Thank you.

DR. WADE: Just to always be receptive as -- as a board, the other avenue is for you to ask the Secretary of Health and Human Services to expand the charter of this Board to allow it to function in ways that you would like to see it function. That's not the char-- that's not within the charter of the Board now, but you always have that prerogative.

MR. STEPHAN: Appreciate that. There -- there

1 is no dispute then that the Board must follow 2 the decisions about site designations, et 3 cetera, as they are made by DOL and DOE. You 4 are not -- you do not have the authority to 5 choose to ignore those designations. 6 correct? 7 DR. WADE: That is my interpretation of the 8 Board's charter. 9 MR. STEPHAN: Okay. Okay. And I believe that 10 Dr. McKeel has a couple more points on this 11 point, or maybe another point, if --12 DR. ZIEMER: Sure, you bet. 13 DR. MCKEEL: Oh, I'll -- I'll make it brief, but I do have a couple more things to say about 14 15 this petition, and they do have to do with the 16 Board. And this gets back to something you all 17 are going to take up tomorrow, which is about 18 the timeliness of transcripts. 19 As I remember the discussion on May the 4th, 20 and as I remember the discussion on July the 21 19th, we all talked about whether the Board had 22 the authority, without any further input from 23 DOE or DOL, to cover the residual period under 24 an SEC. And I made the statement that --25 bolstered by some input some time ago by [Name

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Redacted] -- that the law itself which we're talking about does not preclude doing that. And the discussion that I remember is when you were going to write to the Secretary of HHS, that letter was going to result in a legal opinion from HHS on whether in fact my opinion was -- could be upheld or not. In other words, what -- what -- if the law prevents covering a residual period in an SEC, what specific provision of it does that. And I think -- and I -- if that transcript would ever come forward, I think we could all read it. I think you said that there was a question in your mind about whether that was a valid -- that might be a valid point. You didn't say it was; you didn't say it wasn't. But you -- you certainly didn't say that you could rule that out. make it as an assertion that I -- it's a testable hypothesis, we would call it. And -and the solution to the hypothesis really needs to be a -- a -- a definitive legal opinion from So I still think that's on the table. But I would also like to mention that for resolving this whole issue of extending the SEC, there are many other things that I'm still

waiting for and that we need. We all need to really evaluate this. When the -- SC&A, for instance, in between the July meeting and this meeting, has furnished an evaluation report of NIOSH's evaluation of the SEC. NIOSH has made an addition to their SEC evaluation. The Board has not considered ei-- any of -- either of those documents.

I'm still waiting -- four days after the NIOSH

I'm still waiting -- four days after the NIOSH evaluation came out for the SEC, I wrote 14 questions to Larry Elliott. He answered six of them and he made eight of them into FOIA requests. I'm still waiting for the final answers to those FOIA requests. I -- I got an interim response May 17th that had many documents that are up on the FUSRAP web site and were not useful to me. But the specific things I asked about about that evaluation report that are down at CDC now in their -- in their FOIA office, I haven't gotten that report.

Now I wrote to Dr. Wade about that and he said he had no jurisdiction over the CDC FOIA office. But as a petitioner for Dow, you know, I can't do my part for these men. I feel badly

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about Homer Simmons. I feel badly about Bill Hoppe. I don't know what to tell them anymore. You know, I -- we've done a lot of things. We've expended ourselves. Now we need some answers and results. And so I would just mention there are a lot of outstanding issues and to the extent that the Board has jurisdiction -- and I think they do have jurisdiction over my central question -- you know, we will follow up with DOE and DOL, and I -- I believe, to put Robert's question even more in perspective, if DOE -- if DOL would accept worker affidavits in the absence of contravening documents, then they would have to conclude that DOW did send thorium-magnesium alloys in large quantities to at least Rocky Flats. And you know, the workers have also testified they went to two other AEC installations.

And another issue that I've got to take up with them, they've imposed an additional burden on us. They say well, if you sent thorium alloys to Rocky Flats, even if that was acknowledged, you would have to prove -- I would have to prove, Robert -- our group would have to prove

that that thorium alloy was used in nuclear weapons production. And the answer that I've given back is well, I think we can do that. I think we can make a common sense argument that uses that. Even -- we haven't found any documents from Rocky Flats that would support what that was used for, or even that it was received.

But my point is -- and I use the analogy of the research I did at Washington University. I had NIH grants that had direct costs that were -that paid for the research, they paid for the test tubes, they paid for the microscopes and all that kind of stuff. We also had indirect costs, and those indirect costs, which were 70 percent of the direct costs, went to maintain the building, to have secretaries, to have heat, all that kind of stuff. Well, did that -- did that money contribute to my research? was essential for my research, and that's why I say that if thorium alloy was sent to Rocky Flats, it had to be used in some way to support -- if it didn't go directly into a nuclear weapon per se, it certainly went to support nuclear weapons and went into nuclear weapons

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production because that's all Rocky Flats did.

That's all the Mallinckrodt uranium division

did. There weren't any other functions of

those institutions.

And so I say logically using, you know, normal, intellectual, intelligent reasoning, that argument should prevail. And I -- I think we're due at this point a similar reasoned, intelligent argument back from DOE and DOL why I'm wrong, and -- and we haven't gotten it yet. So our job, as I see it, is to pursue that vigorously in a straightforward way and expect a prompt and equally rigorous answer back and then we can come to that.

But I -- the -- the thing I would ask the Board is to please consider asking HHS for a similar direct answer, can the Board approve an SEC to cover the residual period of contamination, yes or no? And I think that's a straightforward question and I -- I -- I honestly don't think it should take more than a couple of weeks to get the answer. So I'm actually begging you to please think about doing that before we get together in January. And please at that point, one way or the other, let me give Homer and

Bill a final answer.

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DR. ZIEMER: Okay. Thank you, Dan. Larry's got some comments on this and it has to do with the rule.

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MR. ELLIOTT: Well, yeah, I want to make it clear, there's some con-- there's a lot of confusion around what's happened here with Dow. The NIOSH SEC evaluation report that you took a vote on and -- and the Secretary has designated a class on dealt with our inability to reconstruct thorium exposures during the covered period. Thorium was not an AEC activity, according to the designation of the facility by the Department of Energy, during their operations for the AEC. So in the residual period we did not come forward with a recommendation to add that time frame into the class for -- for this facility because thorium was not covered under the residual period. was not a covered activity under the residual period. It was covered under the covered period and so we included that in our evaluation and concluded that we could not reconstruct it. So I just want to be clear on that.

If we had come forward with an SEC evaluation report that said during the residual period there's a component of dose that we cannot reconstruct, we would have done so and I'm sure that the Board, in its wisdom, would have accepted that and moved for the designation of such a class.

Yes, Dr. McKeel, the Board can pass a SEC class in a residual period. However, the constraining point here is that thorium activities at this facility were not considered AEC-related, so the residual period is not covered in that regard.

I just want to be clear about that. Within its purview, NIOSH has done everything it possibly can do, unless the facility designation for DOW is changed in some regard.

DR. ZIEMER: Thank you. Robert.

MR. STEPHAN: And not to belabor this point beyond what we already have, but La-- Larry is exactly right. But the -- our quibble here is not with the decision that NIOSH has made. We respect the roles and responsibilities here. But when we're talking about what -- what you just said, that this thorium beyond the covered

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period is not related to AEC work, that -- that is what we are disputing because no one has given us any information to tell us that that is actually true. What -- we -- we don't have any. I have another -- I don't know, guys, how many pages of documents did you send from DOE, a couple of hundred maybe on our latest request? Yeah, about 500. We have 600-someodd from Dow, but none of them actually -- from -- from DOE or from DOL can show us that they have information which says that what the workers say is not true and what -- you know, what Larry just alluded to is in fact -- fact true. It's being accepted as fact, but no one can give us any information to show us that it is fact. So unless DOE or DOL can come to us with some other principle, we have no choice but to think -- to use an analogy here -- if this were a courtroom, that they're not actually going by -- that an eyewitness account is a pretty valuable account, as it would be in a courtroom. The -- the burden of proof in DOL or DOE related to worker testimony is above and beyond a preponderance of the evidence. I don't -- I don't know what -- principle agree -

that is totally counter to what the SEC is supposed to do, which is supposed to help when there are no documents. And what DOE and DOL are saying is if you don't have documents to prove what the workers say is true, then it's not true. So this is -- this is -- this is a big -- a big issue. So I'm just trying to make the point that you have no reason to believe about this thorium beyond the covered period, that it's not related to AEC work, until someone gives us information that shows that it's not. Okay? Thank you.

DR. ZIEMER: And we understand that and we've heard the argument before, and I think Larry is pointing out that it's not an -- NIOSH doesn't have the purview to make that designation, that the thorium was or was not part of the -- the weapons program. They have been given the AWE designation as it was. And unless that gets changed through DOE and -- and Labor, that's the parameters he's working under.

Dan, you have an additional comment on that? Yeah.

DR. MCKEEL: I promise this is two quick -- Dr.

1 -- I'm sorry, but --

DR. ZIEMER: That's all right.

DR. MCKEEL: Yeah, but just the final conclusion is -- but NIOSH does have the wors--worker testimony to consider about -- that -- that we contend that Dow Madison -- that some of the thorium activities were AEC-related. That's one point.

And the second point is that the other agency involved in all this, the one who originally said that none of the thorium activities were AEC-related, was the U.S. Army Corps of Engineers who remediated the site. And as I've told this Board repeatedly, you know, we went and met with the Army Corps of Engineers in -in June of 2005 and directly asked Mark Bunche*, who's the assistant counsel, for exactly what Robert was talking about: what is your proof? You made this statement in your FUSRAP report of 2000. What -- what was the basis for it? What document do you have that can show that? And they were unable to do that and -- and I -- I have invited the Board, DOE, DOL -- I -- I took your suggestion. You said Dan, you've got to do some work after May 4th,

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and I've done that. And I've invited them -- I gave both agencies Mark Bunche's address, his telephone number. They can do what we did, call him up and -- and see if he can produce those documents. So I -- I agree, we've talked about it enough. I think we'll do our part and I hope we'll come back with some good news for you.

DR. ZIEMER: Thank you. Any other comments?

That sort of brings us up to date, as it were,
on Dow. But here, another comment, okay.

MS. CANO: Again, it's --

DR. ZIEMER: DOE.

MS. CANO: It's -- it's Regina Cano from DOE.

Mr. McKeel, I believe with the Army Corps of
Engineers, we did contact the program manager
for that evaluation, and I believe on Monday or
Tuesday she did send a letter to Dr. McKeel and
also cc'd us that she went back and looked
through her records and could not figure out
why she stated that in the public meeting, that
she misspoke. That was a mis-- a misstatement
on her behalf at that meeting, but -- so...

DR. MCKEEL: Gina's getting something else

confused. Sharon Cotner*, who was -- is the

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program manager for FUSRAP activities in the St. Louis district, made a comment in a February 2000 meeting that uranium processing at Dow Madison took place from 1957 to '62, so we followed up with her and said we've never heard that before. Nobody's ever heard that uranium processing took place in '61 and '62. But if that's true, of course, it would make a difference in the covered period. And she promptly went back and looked in her records, said they looked and looked and she can find no other evidence other than they processed uranium up through '60, just like the SEC class has it, and that -- in her response back she said I was mistaken. That had nothing to do with the thorium activity at all.

DR. ZIEMER: Okay, thank you. Who did -- Jim, did you have a comment?

DR. MELIUS: Yeah, I would just like to thank

DOE for coming here to the meeting today and

presenting -- I know we've been maybe giving

you a little bit of a hard time, but appreciate

their efforts and I think it's -- it's useful

and would hope that -- could continue to

interact and -- in a positive way to -- to

1 settle some of these issues. DR. ZIEMER: Thank you. We do appreciate it, 2 3 Pat, and I've mentioned to Pat to convey also 4 to Glenn Podonsky our thanks as well. 5 Okay. 6 BETHLEHEM STEEL 7 DR. WADE: One last -- Bethlehem --8 DR. ZIEMER: Bethlehem Steel, we have a letter 9 from Senator Schermer -- Schumer's office, I 10 think. Is that going to be read into the 11 record? 12 DR. WADE: Yes. Introduce yourself, please, 13 Richard. 14 MR. WESTON: I will. DR. ZIEMER: Okay. 15 16 MR. WESTON: My name is Richard Weston, W-e-s-17 I'm employed by the Centers for Disease t-o-n. 18 Control and Prevention. I work in the 19 Washington, D.C. office of the Director as a 20 public health advisor. My colleague, Jason 21 Broehm, usually attends these meetings. He 22 couldn't. I'm here in his -- his place. 23 Senators -- Senator Charles Schumer of New York 24 has a one-page statement that his office has

asked to be presented to the -- to the meeting,

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and I would like to read that. It's a onepage, double-spaced statement so it might take me three or four minutes to read that. (Reading) Thank you for the opportunity to address this Board on the petition of the Bethlehem Steel Plant in Lackawanna, New York. I appreciate this chance to share my views with you, and I'm going to take this moment to again urge you to add a class to the Special Exposure Cohort for these former nuclear workers. At its last meeting the Board decided to delay any decision on Bethlehem's petition until the working group makes its recommendation regarding the appropriate limitations on the use of surrogate data and site profiles. Though I'm disappointed by the delay, I remain optimistic that the working group's efforts will bring clarity to a process that until now has felt arbitrary and at times capricious. I firmly believe that a policy that establishes limitations on surrogate data, rather than the current ad hoc decision-making process, will lead to a favorable decision on Bethlehem's petition. Any reasonable limits on the use of surrogate data would fall well below the

excessive level at which it is employed in Bethlehem Steel's profile.

While I do not dispute the usefulness of surrogate data in limited circumstances, over-reliance on it, as in the case of Bethlehem

Steel, is unacceptable. As I and many others have expressed repeatedly, the inordinately heavy reliance on surrogate data in Bethlehem's site profile renders the profile unusable. It is not a reliable representation of the plant's real conditions. Under the circumstances, the Board should void the site profile, release the CDC from its futile attempts at dose reconstruction, and declare the employees of Lackawanna Bethlehem Steel a new class of the Special Exposure Cohort.

The Energy Employee Occupational Illness

Compensation Program was established to repay in some small measure the America's -
America's debt to these former Energy workers.

Their work was critical to building the nuclear arsenal that brought the Soviet Union to its knees, keeping the Cold War from erupting into a hot war which could have killed thousands upon thousands of people.

But tragically, though their hard work saved our nation from violence, America still has wounded veterans of the Cold War. These sickened Energy workers have borne the weight of injury that they spared the rest of the nation. They are fallen heroes of the Cold War and deserve to be treated with the dignity and veneration that a great nation always affords its wounded warriors. It is the least that we can do to fully compensate them for their terrible illnesses, which they have contracted through service to their country.

Finally, many of the men and women who are awaiting compensation from this program are

awaiting compensation from this program are aging and unwell. Time is, unfortunately, of the essence now. For these workers, justice delayed will be justice denied. Please move with all due haste to establish compensa-- to establish and compen-- excuse me. Plea-- please move with all due haste to establish compensation for our fallen heroes.

And that's the statement of Senator Charles Schumer.

DR. ZIEMER: Thank you very much. And the Senator of course in his letter did focus on

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the issue of surrogate data, and as you know, we do have a workgroup that is addressing the surrogate data issue. And Jim Melius, perhaps you can give us a brief report because that impacts essentially on -- ultimately on the Bethlehem Steel issue.

DR. MELIUS: I was just helping my colleague find one of our documents. We actually -- it -- it's a brief update. We've had a -- SC&A do an inventory on the use of surrogate data among our site profile procedures, I believe it's SEC evaluations and so forth, and they had provided that to the workgroup about three weeks ago. And the workgroup will -- hopefully is having a brief meeting -- it's getting briefer as time goes by -- this afternoon immediately following this meeting to establish our -- our work plan for -- for going forward. But I'd like to certainly thank SC&A for doing the inventory 'cause I think it provides a basis for us to be able to evaluate and -- in a fair fashion rather than selecting out arbitrarily -- we'd (unintelligible) better way of -- of approaching this.

DR. ZIEMER: Thank you very much. So let me

1 ask, Board members, do you have any additional 2 questions on Bethlehem Steel? We're obviously 3 -- be awaiting the workgroup's recommendation 4 relative to this particular case, as well as 5 some others perhaps as well. Questions or 6 comments? Requires no action today, it's 7 basically an update on the status of that 8 particular petition. 9 (No responses) 10 If not, I think we are ready to recess until 11 the public comment period. Do we have any 12 housekeeping issues to take care --13 DR. WADE: No, we do -- we do have our science 14 presentation that we will hear tomorrow. 15 DR. ZIEMER: Jim, you can't go home tonight yet 16 then. Okay. 17 DR. WADE: You can't go home again, Jim. 18 DR. ZIEMER: Okay, let's recess then until 19 7:30, at which time we will have our public 20 comment period. 21 (Whereupon, a recess was taken from 5:15 p.m. 22 to 7:30 p.m.) 23 PUBLIC COMMENT 24 DR. ZIEMER: Good evening -- good evening, 25 everyone. We're going to go ahead and begin

1 our public comment session this evening. 2 just going to go through the list in the order 3 that we have it. 4 I do want to remind the speakers that the Board has imposed a ten-minute time limit on the 5 speakers. I like to remind people that that 6 7 should be seen as an upper limit, not a time to 8 be achieved. So -- but that's mainly so that -9 - as a courtesy to those that are later on the 10 list have ample time to give their remarks as 11 well. 12 So Marilyn Schneider, we have you first, if you 13 want to begin, and then we'll take up from 14 there. 15 MS. SCHNEIDER: Are you ready for me? 16 DR. ZIEMER: Go ahead. 17 MS. SCHNEIDER: My name is Marilyn Schneider. 18 I worked as -- as a secretary at Mallinckrodt's 19 Destrehan and Weldon Spring sites in '57 and 20 '58 while they were refining radionuclides for 21 the Cold War and was unknowingly exposed to 22 radioactive material. I was not monitored for 23 exposure and had no idea what was being 24 produced. 25 As an office worker my skin was exposed to

airborne radionuclides and I inhaled contaminated air through the ventilation system and from the open window next to my desk, and drank the contaminated water. My desk and any paperwork I handled were exposed to radioactive chemicals in the air. I was also a mouth breather because I had a deviated septum at that time.

Plant workers with and without uniforms ate in the same cafeteria with the office workers and visited the offices. Tables in the cafeteria were often coated with yellow dust.

Deformed frogs from ponds on the Weldon Spring site, some with two or three heads and extra or missing limbs due to probable carcinogens in the pond, were brought into the office by plant workers. Other office workers at Weldon Spring also developed cancers. When the radioactive waste was buried, the cleanup crew developed skin cancers.

I was in a carpool with plant workers and drove the family car every third day. My car and other cars in the carpool were parked five days a week in the plant parking lot. These vehicles were contaminated in and out with dust

from the smokestacks. The plant workers did not wear uniforms and rode to and from work in their street clothes. I rode in these contaminated cars, and was also exposed to dust from the workers clothing. In addition, my family was also exposed from use of the family car.

In 1975, 17 years after exposure to radiation, I developed colon cancer with penetration of the cirrhosa* and metasis (sic) to two of eight nodes. The first surgery removed eight inches of colon. I wore a colostomy bag until my second surgery resected the bowel. I was given a 30 percent chance of surviving one year.

Despite severe nausea, vomiting, mouth sores and hair loss from two years of high-dose chemo in the veins, followed by two years oral chemo, I did survive. Because my veins blow up due to the two years of chemo in my veins, I now have a port inserted to take the chemo.

In 1977 I developed episodic loss of consciousness of undetermined origin, which still continues. After many tests, the causes cannot be determined and I'm taking medication for seizures.

1998 I was diagnosed with hypothyroidism and take thyroid medication. Hypothyroidism can be caused by radiation exposure.

In July of 2000 I had a pulmonary test which indicates my inspiratory loop is slowed, suggestive of variable intrathoracic obstruction. On the tests I could inhale but could not completely exhale.

I was cancer free until diagnosed with breast cancer in 2000, 42 years after exposure. I was treated with a lumpectomy, sentinel node biopsy and radiation.

Then in 2001, after pain in my right calf, I was diagnosed with a very rare cancer of the smooth muscle cells called leiomyosarcoma.

Soft tissue sarcomas are wildly growing cells from the soft tissue part of the body and include fat, blood vessels, nerves, muscles, skin and cartilage. Lab results didn't show a clear margin after the first two surgeries.

The third surgeon said he would take off my leg if he didn't get a clear margin. My surgical chart said it was my left leg. I was very concerned and wanted this error corrected, so the anathesiologist (sic) marked my left leg

"no" and my right leg "yes," and the third surgery removed five inches of fibula, which controls foot movement. The bone was removed from my knee to mid-calf. The bone was not replaced. I was told I would be able to walk, but may have foot drop. Twice a day for one week after surgery I received internal radiation through plastic tubes inserted into the surgical site -- they called it brachytherapy -- then external radiation for another 35 days.

Two months after surgery I had excruciating pain in the surgical area and wanted to die. Even morphine was not effective. I could not walk. I could not be carried. I could not be touched. Every test possible was run at the Barnes-Jewish Siteman Cancer Center and there was no diagnosis other than probable nerve damage.

Upon research I found that -- I'm calling this LMS for short, the leiomyosarcoma -- is a very rare cancer in the U.S., but a major cancer in Japan because of exposure to radiation from the atomic bomb. LMS is very unpredictable. It can be quiet for a long time, then erupt after

20 years. It's a resistant cancer, not responsive to chemo or radiation. This disease progresses from stage one to stage four. I had stage three. I will be monitored by specialists for the remainder of my life. My specialist, [Name Redacted], now tells me if I break this leg they will have to amputate because the only bone in my leg may not heal. A schoolmate who lived within five miles of the Weldon Spring site also developed LMS in a kidney. She did not work at the Weldon Spring plant.

The three cancers I've had at this point are totally unrelated. In December 2001 [Name Redacted], a genetic counselor in St. Louis, Missouri, stated that none of my cancers were family related. Her report states most carcinomas arise from somatic mutations that are acquired after birth, such as exposure to carcinogenic agents.

In 2004 I developed a fist-sized benign tumor on my uterus. The doctor was going to biopsy until he was told about my leiomyosarcoma. He immediately reacted and said it would require a laparotomy -- I guess I'm saying all these

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words right -- adheliosis, total abdominal hysterectomy and bilateral salpingoophorectomy. I fully expected this to be another cancer. In March 2006 [Name Redacted], a cancer specialist at Barnes Jewish in St. Pete-- St. Louis, sent a letter to the Department of Labor stating that my cancers were likely environmentally caused, and there is certainly a known association between exposure to environmental carcinogens and radioactive material and the development of cancer. He also stated soft tissue sarcomas are relatively uncommon cancers and there are only 8,000 known cases in the United States. According to [Name Redacted], director of statistics at the Radiation Effects Research Foundation headquartered in Hiroshima, cancer risk from radiation exposure continues throughout life. In May 2007 when my skin and eyes turned yellow as a banana, I was immediately admitted to the hospital and a stint was placed in my bile duct. I was diagnosed with another rare cancer, adenocarcinoma of the bile duct. days later when my bilirubin had decreased from 20 to eight -- this is the yellowness -- I

underwent Whipple surgery. The surgeon removed my gallbladder, half of my pancreas, part of my stomach and the entire duodenum.

Recovery from this cancer was pretty rough.

Food couldn't enter the stomach because of the surgery. My oncologist and radiologist tell me there are no statistics on how to treat this cancer. As previously stated, I had the port inserted for the chemo because my veins have collapsed from previous chemo in 1975. This port now enters the carotid artery. It takes a highly skilled phlebotomist to even draw blood. IVs now have to be put in my neck.

I had just started chemo and was recovering from surgery when I noticed a small black mole with tendrils on the calf of my right leg. My doctor remarked, looks like trouble, and sent me to a dermatologist for a scraping and lab work. The diagnosis was junctional melanocytic proliferation. He consulted with my oncologist, and my chemo was put on hold until this precancerous melanoma could be excised. I was taking chemo, dealing with a confirmed cancer of the bile duct and another possible cancer, melanoma, at the same time.

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Then I receive another denial from HHS stating I wasn't exposed to enough carcinogens to warrant four, possibly five, unrelated cancers. I've been denied compensation seven times, more times than Judas denied Christ. In the official report proceedings before the Final Adjudication Branch of the U.S. of Labor (sic) dated 12/14/06, Tom Daugherty, the hearing representative, states that NIOSH and Department of Labor found a 34.63 percent combined probability that my colon and breast and leiomy-- leiomyo was casually related to impairment under the Act. This was prior to the adenocarcinoma of the bile duct. scheduled for a closing interview on September 13th, 2007 for denial number six. When the interviewer called I told him I had submitted a claim for adenocarcinoma of the bile duct, which he didn't know about until he checked. Then in a letter dated the very next day, September the 14th, HHS denied my claim. This was denial number seven. How could dose res-be -- reconstruction be done so quickly? addition, this rare bile duct cancer was dosed as gall bladder.

1 My last two cancers are extremely rare. 2 leiomyosarcoma, about one in a million chance 3 of getting this, but leiomyosarcoma is now 4 being seen in several other sites. 5 The last dose reconstruction states there is no existing model to calculate dose for soft 6 7 tissue of the calf. How is the rarity of this, 8 or my current cancer, being addressed. Or is 9 it? 10 NIOSH states external doses from stack releases 11 or other radiation sources may have been 12 unmonitored at this site, and there was no 13 workplace data done on exposure to radioactive 14 material. Department of letter -- Labor letter 15 dated 8/16/07 reporting my bile duct cancer to 16 NIOSH states due to my job description I was 17 not exposed to radiation, that I did not handle 18 radioactive materials, and that my job 19 description and probably work -- probable work 20 location would not involve exposure to airborne 21 radionuclides higher than that reported 22 environmental lever -- levels. 23 Let me tell you about my possible exposure. 24 did not handle radioactive material, but had 25 chronic exposure from breathing the air at the

1 plant and in my car, handling paperwork on my 2 desk, eating in the cafeteria with the plant 3 workers, exposure to the paperwork brought into 4 the office by plant workers. The denial did 5 not take into account my breathing airborne 6 radionuclides from the open window by my desk 7 and breathing air from the ventilation system. 8 I never wore a dosimeter badge; therefore there 9 is no record of my internal or external 10 exposure to airborne radionuclide 11 concentrations. 12 I have never smoked and I do not drink. 13 all my cancers are unrelated to heredity. 14 common tie is the carcinogens I was exposed to. 15 Lastly, a coworker at the same sites with two 16 of the same cancers, colon and breast, was 17 approved. I had chronic exposure to 18 carcinogens without informed consent. I was a 19 human radiation experiment. My medical bills 20 and emotional trauma have been astronomical. 21 If I survive this cancer, I will probably get 22 another. 23 During one closeout with HHS I was told to be 24 sure to let us know if you get another cancer. 25 If this cancer doesn't kill me, you can be sure

1 I'll call when I get another one, even though I 2 was told I wasn't exposed. 3 I physically and emotionally cannot keep up the 4 fight for compensation. I'm about ready to 5 throw in the towel. How many cancers must I get in order to meet the 50 percent probability 6 7 of causation? 8 And I thank you for your attention. 9 DR. ZIEMER: And thank you, Marilyn, for 10 sharing that with the Board. 11 Next we'll hear from Susan Pru* -- did I 12 pronounce that correctly? Yeah. 13 MS. PRU: Thank you. That's tough after 14 hearing that. My name is Susan Pru. I'm the [Identifying 15 16 information Redacted] of [Name Redacted]*. 17 [Identifying information Redacted] is now 18 deceased -- I'm so upset by hers, I'm sorry. 19 DR. ZIEMER: Could you get a little closer to 20 the mike? Thank you. 21 MS. PRU: My [Identifying information Redacted] worked at the Y-12 plant. The Department of 22 23 Energy and Labor confirmed her employment 24 during the covered time of [Identifying 25 information Redacted]. Her diagnosis of breast

cancer, one of the 22 covered cancers, was submitted with her claim. We haven't yet received her medical records for the surgical remover -- removal, excuse me -- of the cancerous growth on her tongue, and we're still awaiting her de-- detailed earnings report.

We know that [Name Redacted] was a [Identifying information Redacted]. My husband and sisterin-law know that she was in the plant. And according to DOE and the CDC, it wasn't until January of 1951 that all employees were required to wear a badge, regardless of where they worked. [Name Redacted] left employment [Identifying information Redacted] years prior to this.

My question for the Board is, why was the claim recently sent to NIOSH for dosage reconstruction when she fit the very criteria for the SEC? She had one of the 22 cancers and worked over the 250 days, and we know that she was in the plant. So I would just love for someone to find out why. Thank you.

DR. ZIEMER: Thank you very much, and perhaps one of the NIOSH staffers can help you track that down.

1 DR. WADE: Department of Labor is here. 2 DR. ZIEMER: Or -- or Labor, okay. Thank you. 3 And we'll get you connected with somebody here. Dan McKeel -- Dr. McKeel? 4 5 DR. MCKEEL: (Off microphone) (Unintelligible) 6 DR. ZIEMER: I'll give it to Ray, yeah. 7 Thanks. 8 DR. MCKEEL: Good evening. These comments 9 tonight are about GSI. No more Dow, I promise. 10 I -- I'd like to complement in these remarks 11 what John Ramspott had to say about GSI and the 12 appendix BB for that site last night. 13 Our group, SINuW, believes this document, 14 appendix BB, is technically highly flawed. And 15 we are happy that SC&A has been tasked to review it and thank the Board for that. 16 17 John did not mention two reasons we believe 18 appendix BB needs to be changed. These facts 19 were both mentioned in my critique to that 20 document now posted in redacted form on the 21 OCAS web site. Neither of these facts is 22 mentioned in appendix BB, nor was the second 23 Betatron, both cobalt-60 and iridium-192 gamma 24 sources, nor the 250 kVp portable X-ray unit

that all contributed to worker dose at GSI.

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First is the fact known since at least 1950 that 20 to 25 MeV Betatrons used for both industrial non-destructive testing and medical treatment emitted neutrons as well as photons. Attached to my remarks is a neutron curve from one of the three medical Allis Chalmers

Betatrons in the St. Louis area, and this data is from 1973, a very similar unit to that used -- to the two used at GSI. This is important data because GSI worker badges did not record neutron data and the relative biologic effectiveness of neutrons is ten-fold that of gamma photons.

Second is the fact that individual film badge dosimetry data is available at Landauer for 30 GSI Betatron and isotope workers. The years covered are 1963 to mid-1973 when the plant ceased Betatron operations. I informed OCAS about this dosimetry data months ago. One worker had a cumulative dose for one year of 38,000 millirems, indicating significant over-overexposure occurred at the site.

Many GSI workers who have recently undergone dose reconstruction by NIOSH are aware of and are concerned about these serious technical

deficiencies in appendix BB. One of those workers has asked me to read into the record tonight the letter he wrote to Larry Elliott as a result of his exit interview. This letter bears out the report that SC&A just released on the same topic, the exit interview before signing OCAS, that NIOSH doesn't pay enough attention to worker input at this end of the process.

[Name Redacted], gave me explicit permission to state his name and disclose his NIOSH tracking number, and [Name Redacted] writes as follows to Larry J. Elliott. I want to read into the record this statement. (Reading) I believe that it has been preordained that my claim is to be denied, based on these facts. One, my Social Security record of employment for the years 1951 through 1961 was changed. I worked at General Steel Industries, which my Social Security record showed at the time of retirement in 1996, and was changed to roll capital CO prior to or after my claim was filed.

Number two, the name of the site was changed from General Steel Industries to Granite City

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Steel. My claim was denied because I did not work at Granite City Steel during the years that uranium was being X-rayed at General Steel. Congressman Jerry Costello aided in getting this corrected.

I have -- three, I have not been given any factors being considered for the dose reconstruction, such as (a) my work station and task performed, (b) the amount of dust in the air that could be radiated (sic) by the sunshine effect, or (c) the amount of residual radiation in the area. And in parentheses, a cleanup of the site of the old Betatron was completed in 1993 because of radiation contamination and the local city authorities were not notified, end of parentheses. Number four, General Steel Industries' own railroad cars that were used throughout to transport the uranium castings, and they were also used throughout the plant. No one measured to see if they were contaminated. don't know if that was considered. Number five, NIOSH letter dated June the 27th, 2007, signed by April Jenkins, said that Joe

Dickey was to do my dose reconstruction and I

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had to let her know if I had any reason to want someone else. I called on July the 6th at 12:48 p.m. central time and said that I wanted a hold on the dose reconstruction to have an expert review the qualifications of Mr. Dickey. She said, quote, okay, end quote, and I was to call her when I wanted it to proceed. On July the 9th, 2007 I called and talked to Nancy Vander Ahe, A-h-e, and she said my case was put on pending status, in quotes, but would not sent me a letter to confirm. On August the 2nd, 2007 I received a NIOSH letter stating that my case was in dose reconstruction. letter was simply dated July 2007. I called at 1:30 p.m CT and asked to talk to Nancy or April, and was told they were not available. then asked who made the decision and was told a, quote, decider, end quote. I asked who, and they said a leader named Richard McCarthy. asked to talk to him and was told he would call me back. He didn't call. On August the 3rd I called and was told Mr. McCarthy was not in on Fridays. On August the 6th Mr. McCarthy called and told me my dose reconstruction would go forward and I couldn't stop it.

Six, I am now told I'm to sign a form saying I don't have any other information to submit. I don't know what information they have, or how they perceive that information. If I don't sign the form and submit it in the allotted time frame, they may, quote, administratively close my dose reconstruction, end quote. This sounds like a, quote, done deal, end quote, and there's nothing I can do. I can tell you I don't like it, and I will share this information with Senators Durbin and Obama, as well as Congressman Costello.

And his final paragraph says (reading) My work station was out in 10 building among all the castings and burners, the welders, the chippers, the grinders, the sandblast operators, the inspectors, the four foremen and laborers. I was there every day and was exposed to everything they were. I went to check on castings at the Betatron site on many occasions. During lunch breaks in good weather, we would go out and sit on the company rail cars to eat our lunch. No one told any of us about radiation. Inside the building if someone was working inside of a casting -- and

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1	in parentheses, tank (unintelligible) and
2	turrets, end end parentheses it was
3	necessary, due to the noise, to lean against
4	the casting to get information from them. It
5	was normal practice to lean on the castings
6	while writing information on the cards.
7	Respectively (sic), [Name Redacted], NIOSH
8	tracking number [Identifying Information
9	Redacted].
10	Thank you very much.
11	DR. ZIEMER: Okay. Thank you very much, Dr.
12	McKeel.
13	Next Deb Detmers, and Deb we heard from earlier
14	and is she back this evening? She's with
15	the Congressman's office.
16	DR. MCKEEL: (Off microphone) I think they may
17	have (unintelligible).
18	DR. ZIEMER: Yeah. We did hear from her
19	earlier today.
20	Bev is it Marcoski*?
21	MS. MARCOSKI: Yes.
22	DR. ZIEMER: Yes.
23	MS. MARCOSKI: I'm I'm Bev Marcoski and I
24	just have a few comets comments on the SEC
25	petition with Olin Chemicals. My comments

revolve around four areas. One is water testing at the plant, which was not done. Two is part of the chlorination process for the calcination process. Three, you talk about specific assumptions versus the general. And then four, I'd like to touch base on the thorium.

I was doing some general reading in Aviation

Medical and it referenced ww.epa.gov/radon

(sic), and what is stated was drinking water

deaths are primarily due to lung cancer due to

the radon. And I started thinking about

Blockson and all the information I've read over

the past six years, and nothing was mentioned

about water testing on the site. I'm assuming

the men worked -- the workers in the plants and

the ladies in the administrative offices drank

the water. Also there's a possibility that

some of the men also showered in this water,

and there's no objective evidence of any water

testing.

When I was in Joliet a couple of weeks ago one of the gentlemen sitting next to me said that Olin did have six of their own wells.

Further to think through the process and in

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your SEC petition, the most recent one from July, it talks about washing down the uranium area, and then what happened to that waste water? And also what happened to the waste that they swept up? That was something that was unanswered, how that left the plant or where it went. And again, I guess there's open questions on how polluted was the water. Secondly, chlorination is known to have a carcinogenic effect as well, and I'm sure they used high levels of chlorin -- chlorine to increase this calcination process. I know you've only looked at radionucleides (sic), but I know chlorine does have a carcinogenic effect as well, especially when you're inhaling the fumes, and there's nothing said about that, and that's page 31 of the SEC. Specific versus general, and I guess I did my own little analysis versus my father's job, and -- and they also did one on page 41 in the SEC for only one person, a filter operator. Most of the assumptions made for this job analysis

were general, assuming that a person stood 30

case I assumed that my dad took some of the

centimeters away from the contamination. In my

waste away from this building and that he handled it for approximately 20 minutes two times a week. That millirem exposure is 160 at that, and when I did the math it came down to -- over a 10-year period -- 51,150 millirems of exposure versus the assumed 24,000. So even in his small job of taking the waste away from this area where they processed the uranium, specifically looking at it, his exposure was twice what was assumed. And they only did this for one other person, and I guess if you're going to analys -- analyze the jobs, you might have to look at the specific jobs each of these people did, which may be very encumbersome (sic) to do to get exact exposures versus general.

And then fourthly, to talk about the thorium, and I guess in the third Technical Basis

Document, I believe page 13, there is an unknown value of what the matrix is exactly on it. And I don't know how you can go back to the '50s. It says in thorium-230 the matrix may not dissolve, assuming that it didn't go to the phosphoric acid stream, but maybe a larger portion could have gone to the sulfuric acid

1 gypsum pond; and if so, how would it change the 2 technical assumptions. 3 Those are the four things that I had. Thank 4 you. 5 DR. ZIEMER: Okay. Thank you, Bev, and the --6 the Blockson working group has heard your 7 remarks and can consider them further as they 8 continue -- at least many of them. Certain 9 ones, such as chlorine, actually are outside 10 the purview of this group. We recognize there 11 are many carcinogens in the workplace, and 12 we're somewhat restricted in what we can 13 address in terms of the legal framework that we 14 work in, but that's -- that's one of the issues 15 that is always a concern. But be aware of that 16 at least also. 17 Cyril -- looks like G-u-r-e, Gure? Close 18 enough for government work, as they say --19 right? 20 MR. GURA: Well, my name is Cyril Gura*. 21] Identifying Information Redacted] was [Name 22 Redacted] and he is tracking number 23 [Identifying Information Redacted], and [Name 24 Redacted] is my [Identifying Information 25 Redacted]. And had an opportunity to review

SEC 00058, and I believe it's more favorable and more in line with the intent of federal litigation, but reviewing it I did see some things that I didn't see answered.

In one particular case, personal protective equipment, looked -- personal protective equipment was issued to employees as comparison to what personal protective equipment employees would have to wear now when handling -- working around these noted hazardous materials identified at Blockson's.

And then secondly, on page 27 of 50, talk about urinalysis. It was sampled from April 1954 to February 1958. The petition class definition was from January 1st, 1951 to December 31st, 1962. The sampling occurred over three years and ten months. However, if there was any records, there was no sampling indicated for seven years and one month. What would DOL do if their inspectors went out on site to review compliance records and a seven year one month period were missing now? Is this one of many latent conditions that is indicative of poor safety oversight, or is this something more blatant like hiding something or keeping unsafe

1 working conditions unknown? 2 And lastly, I understand in order to calculate 3 exposure certain assumptions need to be made, 4 and some of these newer assumptions like 5 increasing the radon level from 50 percent to 95 percent does help. And including other 6 7 cancers -- lung, liver, kidney -- also helps. 8 And some of the other exposures from thorillium 9 (sic) and uranium are important. But it should 10 be remembered that employees' health 11 disabilities, like this woman over here, just -12 - dif-- different types of cancers, fatalities and also -- should be a heavily-weighed factor, 13 14 even more so than calculations based on general 15 assumptions, even though that these assumptions 16 are the best that could be ascertained at the 17 time. 18 That's all I have. 19 DR. ZIEMER: Okay. Thank you very much. 20 I think we have on the phone Terrie Barrie from 21 Colorado. Is Terrie Barrie on the line? 22 Terrie Barrie is on the line. Terrie, you may 23 give us your comments, please. 24 MS. BARRIE: Well, good evening, Dr. Ziemer and

members of the Board. My name is Terrie Barrie

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and I'm with the Alliance of Nuclear Worker
Advocacy Groups. I'd like to thank you and Dr.
Wade for allowing me to call in my comments
tonight.

I want to address the draft report that SC&A submitted on the closeout interviews. summary is very similar to what I hear from individual claimants. Claimants are asked during the initial interview to supply names, for example, of coworkers that could help verify an exposure or workplace condition. the claims I have tried to help I have yet to hear of NIOSH contacting those workers -- those coworkers. It appears to the claimants that the initial interviews, as well as the closeout interviews, are for show only. The claimants do not feel that NIOSH ever intended to investigate and find the whole truth of how much radiation the workers were exposed to. [Name Redacted], an advocate for some Los Alamos claimants, contacted me this morning. She requested and I agreed that the Board should instruct SC&A to review a much larger sampling of randomly-chosen claims (unintelligible) exit interviews from each DOE

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facility from across the country. This'll give the Board a more accurate assessment of the problem with the closeout interviews. If the sampling results in a systemic problem, then each and every denied claim needs to be reopened to ensure that all evidence was considered in reconstructing doses. I shudder to think of the cost, but reopening claims will preserve the due process rights of the claimants. However, if this dose re-- if this does come to pass, I strongly recommend that the ORAU team not be permitted to perform the new dose reconstructions. They should not receive financial compensation when it was their failure to produce adequate procedures for dose reconstructions. There is one other issue I'd like to raise tonight. SC&A had concerns on NIOSH/ORAU's ability to apply their new procedures correctly. I had one Rocky Flats claimant contact me last week -- last week with a disturbing story. Her husband had died of lymphoma years ago. When the new procedure, target organs for lymphoma, was issued she

petitioned and was granted a reopening of her

1 claim by DOL. DOL sent her claim back to NIOSH 2 for another dose reconstruction. However, the 3 new dose reconstruction applied the super S 4 model instead of the lymphoma target organ 5 procedure. Some of -- Board members may not realize that 6 7 she cannot appeal to DOL that NIOSH used the 8 wrong procedure to reconstruct dose. 9 the appeals process DOL will not consider the 10 claimant's objections to NIOSH's procedures or 11 calculations. 12 The claimants have heard nothing but how 13 claimant friendly this program is. Ignoring 14 evidence is not claimant friendly. The 15 inability to question NIOSH's procedures during 16 a DOL hearing is not claimant friendly. 17 strongly recommend that the Board do everything 18 in its power to rectify these injustices. 19 Thank you again for allowing me to speak 20 tonight. 21 DR. ZIEMER: Okay. Thank you very much, 22 Terrie. 23 Let me ask if there are any other individuals 24 on the line that did wish to address the 25 assembly?

1 Yes, we'll take -- please tell us who you are 2 and then you can proceed. 3 MS. BALDRIDGE: Okay. This is Sandra 4 Baldridge. DR. ZIEMER: Okay, very good. Proceed. 5 UNIDENTIFIED: (Off microphone) 6 7 (Unintelligible) 8 MS. BALDRIDGE: I was listening to the 9

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discussions earlier today, and the issue came up in discussion about worker statements being taken and used as evidence, and concerns about whether -- you know, what were -- what statements would be listened to, how they would be considered and so forth. And I recalled reading in the rules and regulations on 42 CFR 82, and this is the statement as it appears in the rules and regulations. It says it is well recognized from health, behavioral and social research that there are substantial limitations and variations in the ability of people to accurately call -- recall past events and that these limitations generally increase with the time elapsed since the past event. However, all other sources of information available to NIOSH in conducting dose reconstructions

1 potentially involve substantial limitations. 2 To conduct dose reconstructions NIOSH will 3 apply procedures available to it to mitigate 4 these limitations to the extent possible to 5 improve the recall of employees. 6 Now the statement, as I read it, shows that it 7 is written into the rules and regulations a 8 prejudice against statements and testimony 9 taken from workers. I don't know how anyone 10 else sees it. 11 Another issue I would like to address is the 12 timeliness. In filing the petition for Fernald, much of the basis of that was the 13 14 discovery that there had been thorium processing in plant six where my father worked 15 16 for three and a half years. That had been 17 omitted from the site profile and actually was 18 without -- had been conducted without NIOSH's 19 knowledge. They claimed that records had been 20 destroyed. 21 And again in the rules and regulations, after 22 having presented this information in 23 [Identifying Information Redacted] case and claim, I got no -- nowhere with it on his 24 25 personal case. It says HHS has added

provisions under 82.27 of this rule to authorize NIOSH to review com-- to review completed dose reconstructions on its own initiative upon obtaining new information or changing scientific elements underlying dose reconstructions. HHS has targeted the added provision to circumstances in which new -- the use of new information or scientific element could increase the levels of radiation doses previously estimated since the purpose of these provisions is to provide new information to DOL on claims that were denied based on outdated information.

When I have contacted NIOSH about this they have chosen to make their prerequisite the revision of the site profile. Now they said today that they didn't know when their site profile for Fernald would be revised. So they have chosen to increase the time for the consideration of [Identifying Information Redacted] claim based on the new information that they have -- that they received in the spring of 2006. And to this point [Identifying Information Redacted] case is closed and none of the information that has been presented has

ever been considered.

My third point, at the Fernald working group meeting, Dr. Ziemer, you indicated that OTIB-2 was being looked at for its appropriate use, but you didn't elaborate on that. My con-- my continuing concern is that its application to dose reconstructions for workers who don't meet the criteria for its application, specifically an initial hire date after 1969 as stated in the document, and a start date prior to 1970. I was wondering if you could give me a little more information about what is being done to look into the use of inappropriate technical basis information bulletins.

DR. ZIEMER: Okay. Thank you, Sandra. I -- I think what I will need to do is get back to you separately on that particular thing. I need to talk with -- with Mr. Clawson, the head of the workgroup, and then get some clarity on how that relates in this particular case.

But let's see, do -- do we -- well, we'll do this off line. I think we -- we may have your phone number in the records already, but if not, we'll -- we'll track that down and try to get you a more specific answer. I -- I don't

1 know the answer to that as I sit here right at 2 the moment. 3 MS. BALDRIDGE: You know, in [Identifying 4 Information Redacted] case, he was an employee 5 at National of Ohio in 1971 -- actually in 1951, not 1971. So you know, I would like to 6 7 know how it applies and --8 DR. ZIEMER: Yeah, and as I said --9 MS. BALDRIDGE: -- so forth. 10 DR. ZIEMER: -- I don't know on that specific 11 case, but --12 MS. BALDRIDGE: I'll be expecting a response. 13 DR. ZIEMER: -- in general -- right, and in 14 general if -- if the Board identi--15 MS. BALDRIDGE: Do you have any questions for 16 me? 17 Right. Thank you. In general, if DR. ZIEMER: 18 -- if the Board identifies what we think is 19 inappropriate use of -- of any of the 20 documents, that's raised and we -- we try to 21 learn what NIOSH's perspective is on it and --22 and we also have our contractor look at these 23 and then we try to reach some kind of 24 resolution. But on specific cases, I think our 25 preference is not to try to resolve those in

1 the public forum since there are often privacy 2 issues involved. But we will get back to you 3 and try to be more specific in answering this 4 question for you. 5 MS. BALDRIDGE: I was just -- you know, hadn't 6 really understood what was going on and, you 7 know, thought I would take this opportunity to 8 ask. 9 Fine. We'll -- we'll follow up DR. ZIEMER: 10 with you, Sandra, on this. Thank you. 11 Is there anyone else on the line that wishes to 12 address the group tonight? 13 (Pause) 14 We have another person on line? Yes, we do. 15 Please have them proceed, identify themself 16 (sic). 17 MR. DUTKO: (Unintelligible) G. Dutko. 18 from Granite City, Illinois. I was a Betatron 19 operator at GSI between 1963 and 1966. Sir, my 20 question is -- and it's not malicious or 21 aggravating -- any intent whatsoever. But sir, 22 50 years ago we weren't exactly told the truth 23 of the hazards of the Betatrons. I was one of 24 the fellas that turned the key on and simply I 25 don't know the damage or sickness caused by

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holding back the -- the truth about these machines, but I don't understand -- going back to Robert Stephan's and Dr. McKeel's statements -- why aren't the statements of the working -working people -- carry any weight with the Board or with NIOSH or DOE? Sir, we fired thousands and thousands of Roentgens back in that time. We fired thousands of Roentgens, sir. And I don't know of any firing sheet possessed by NIOSH of these logs of radiation or Roentgens. I -- I -- they might have film badges. We never did trust a film badge. wore dosimeters quite a bit. They burned the dosimeter logs. We never saw them. We never saw reports on -- on any kind of blood tests or urine tests. How, sir, can accurate -- how can accurate dose recon-- dose reconstructions be done in this case? And my -- my remarks are not intended to be malicious.

DR. ZIEMER: Okay. Okay, thank you for that question. I might tell you that in fact both NIOSH and our contractors are looking at the Betatron issue to determine the extent to which the doses can be re-- reconstructed. I don't think we know the answer to that yet. It's

I don't

1 possible that they may say they can't be. 2 possible that they may determine that they can 3 -- can reconstruct. So certainly we've taken 4 seriously the -- the issues as raised by John 5 Ramspott and -- and by Dan McKeel. We are 6 looking seriously at the Betatrons and we hope 7 to come up with a -- an answer to the very 8 question that you have asked. 9 MR. DUTKO: One thing I -- I do know and the 10 only thing I got out of that 50-year-old 11 experience is a lot of us people that turned 12 the keys on the machines don't know really what 13 kind of life expectancy we're going to have, 14 sir, after finding out 50 years later what we 15 didn't want to hear. 16 DR. ZIEMER: Yeah, thank you, and I have to 17 take that as a rhetorical question. 18 think any of us know that. 19 Our court reporter does need to get your name, 20 we missed that, if you could repeat it, please. 21 MR. DUTKO: My name is John G. Dutko, D- as in 22 dog u-t-k-o. I was a Betatron operator, 24 and 23 25-million-volt Betatron operator at GSI 24 between November '63 and November '66. Thank 25 you, sir.

1 DR. ZIEMER: Yes, thank you very much. Anyone 2 else on the line tonight that needs to speak to 3 us? 4 (No responses) 5 Apparently not. Let me go back here to those 6 assembled. Is there anyone who wishes to 7 address the assembly that didn't have a chance 8 to sign up and -- and put your name on the 9 list? We'd be pleased to hear anyone else who 10 wishes to address the group tonight. 11 (No responses) 12 If not, I thank you all for being in attendance 13 and sharing with us. I would remind you that 14 the Board will be meeting again tomorrow. 15 You're welcome to join us again at that time. 16 We'll recess now until tomorrow morning at 17 8:30. 18 (Whereupon, an adjournment was taken to Friday, 19 October 5, 2007 at 8:30 a.m.)

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CERTIFICATE OF COURT REPORTER

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 Oct. 4, 2007; 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 7th day of November, 2007.

STEVEN RAY GREEN, CCR
CERTIFIED MERIT COURT REPORTER
CERTIFICATE NUMBER: A-2102