UNITED STATES OF AMERICA

CENTERS FOR DISEASE CONTROL

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NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

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ADVISORY BOARD ON RADIATION AND WORKER HEALTH

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60th MEETING

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WEDNESDAY, DECEMBER 17, 2008

The meeting came to order at 9:00 a.m. in the Oglethorpe Room of the Augusta Marriott Hotel and Suites, 2 Tenth Street, Augusta, Georgia, Dr. Paul L. Ziemer, Chair, presiding.

PRESENT:

PAUL L. ZIEMER, Chair

JOSIE M. BEACH, Member

BRADLEY P. CLAWSON, Member

MICHAEL H. GIBSON, Member

MARK A. GRIFFON, Member

JAMES M. MELIUS, Member

WANDA I. MUNN, Member

ROBERT W. PRESLEY, Member

JOHN W. POSTON, Member

GENEVIEVE S. ROESSLER, Member

(via telephone)

PHILLIP M. SCHOFIELD, Member

TED KATZ, Acting Designated Federal Official

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C-O-N-T-E-N-T-S

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Adjourn

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PROCEEDINGS

(9:11 a.m.)

WELCOME

CHAIR ZIEMER: Well, good morning, everyone. We are ready to reconvene the meeting of the Advisory Board on Radiation and Worker Health, meeting here in Augusta, Georgia.

This is day two of the meeting.

The agendas, again, are on the back table as well as a variety of documents in support of the activities of the meeting.

Also another reminder to register your attendance in the registration book in the foyer. Even if you did that yesterday, we'd like you to do that again today. We track the attendance every day.

And then finally, members of the public, if you wish to make public comment during our public comment session this evening, please sign up in the book that is

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1 there in the foyer. 2 I also want to confirm the presence of our remote Board member, Dr. Gen Roessler. 3 Dr. Roessler, are you on the line? 4 MEMBER ROESSLER: I am on the line. 5 CHAIR ZIEMER: Okay, well you don't 6 7 seem so remote. MEMBER ROESSLER: Well, you know the 8 connection is much better this morning. 9 Ιt 10 almost impossible to hear last night during the public comments. 11 ZIEMER: Oh, really. 12 CHAIR 13 sorry to hear that. ROESSLER: Well, I think 14 MEMBER 15 Ted's reminders always help. 16 CHAIR ZIEMER: Very good. And the record will also show that 17 Phil Schofield has joined the Board this 18 19 morning. That is, well, he's joined the Board much earlier in this career, but he is with us 20 this morning. We're glad to have you here as 21

well.

1 The only person really missing from 2 our deliberations today is Dr. Lockey who is not able to be with us. But we do have a 3 4 quorum. Mr. Katz, do you have any remarks 5 as we get underway? 6 MR. KATZ: Just a reminder for the 7 folks on the phone, just a general reminder, 8 please remember to mute your phones, and use 9 10 *6 if you don't have a mute button, and please don't put us on hold but hang up and dial back 11 in if you need to break from the call for a 12 while. Thanks. 13 CHAIR ZIEMER: Thank you very much. 14 15 We are going to follow the agenda pretty much as it's given, but let me indicate 16 to you a couple of changes. 17 Number one, you may recall that 18 19 yesterday we deferred the report from the Department of Energy because Dr. Worthington 20 was not able to arrive because of difficulties 21

in her flight, but she is with us now, and we

are going to schedule the DOE update at the 11:15 slot, which, on your agenda, is the location of the science update.

We will move the science update to the slot in the meeting which is labeled review close-out process, Dr. John Mauro. Because that report you will recall was given yesterday because we had the open time slot yesterday.

So with that slight juggling in the agenda, we will proceed.

The first item then, this morning is 83.14 SEC Petition for Vitro an Manufacturing. The petition evaluation report will be presented by Stu Hinnefeld. Also, we may hear from the petitioner if the petitioner It was not absolutely sure when we so wishes. last contacted the petitioner that that person wished to speak or comment, but we will have opportunity for that as well.

So let us proceed then with the evaluation report on Vitro. I don't know

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| 1 | actually if it is pronounced Vitro or Vitro. |
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| 2 | Maybe it's Vitro, Stu. What is the official |
| 3 | NIOSH pronunciation of this company? |
| 4 | MR. HINNEFELD: Vitro. |
| 5 | CHAIR ZIEMER: You call it Vitro. |
| 6 | Okay. |
| 7 | MR. HINNEFELD: I don't know that |
| 8 | that is the correct pronunciation. |
| 9 | CHAIR ZIEMER: I thought perhaps the |
| 10 | people who worked there were in vitro, but |
| 11 | Vitro sounds a little better, I think. So we |
| 12 | might have to get corrected on that. In any |
| 13 | event let's proceed. |
| 14 | VITRO MANUFACTURING 83.14 SEC PETITION |
| 15 | MR. HINNEFELD: Thank you, Dr. |
| 16 | Ziemer. I suggested to LaVon last night that |
| 17 | he might want to come and give a presentation |
| 18 | with my name on it, but he didn't take me up |
| 19 | on that, so here I am again. |
| 20 | This is, as introduced this is |
| 21 | our presentation for the Vitro Manufacturing |
| 22 | site, AWE site in Canonsberg, Pennsylvania. |

It's just a little ways south of Pittsburgh.

Here is a little bit of the history of the petition. The date, November 10th, when we notified one of the petitioners, or one of the claimants from Vitro Manufacturing that we were not able to do a dose reconstruction for his or her case, and we sent them a letter to that effect, and sent them a blank Form A petition, SEC petition.

They then returned the petition promptly and we qualified on November 26th, our finding being that we are unable to complete dose reconstruction with sufficient accuracy for the employees there.

The Vitro site is an atomic weapons employer. Its operational period is from 1942 to 1957. That is the period we have evaluated for this petition, as the active period, the covered period.

They conducted chemical processing to extract uranium from ores and scrap AEC materials, and they were, actually before

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World War II, they were already engaged in the extraction of uranium and radium for commercial purposes. And then when the war effort was looking for uranium, they went to places like Vitro that already knew how to do it, and had them specifically for the Manhattan Engineer District then extract uranium for the government's purposes.

Physical forms were uranium ores, concentrates, U-308 is typically the product of the purification process, and then the byproduct materials containing uranium progeny.

So in other words, of course when you purify the uranium out of the ore you got all the stuff that is not uranium from the ore that is left over, many of those are radioactive elements, the progeny of uranium.

The data capture efforts are actually summarized in the petition evaluation report. There is a table in the back that describes documentation that was obtained, and

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the records that -- where they were obtained from, you will see that. We have used our existing database which we have populated over the project course of from the DOE Germantown offices and National Archives and records centers, and a number of data capture activities. We have -- we looked at existing project technical documents to see if they would help to help inform us about this. looked at the legacy management, considered site's database, couple а of open databases, NRC ADAMS database, and a variety of other systems including the Hanford declassified document retrieval system, Science the Office of Technical and Information, OSTI.

And we evaluated the interviews that had been conducted with the claimants, the computer-assisted telephone interview that is conducted on all the claimants. Didn't see any information in that, in those interviews that would cause us to think that we could

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feasibly reconstruct all the doses at Vitro, and we also concluded from that that we were not likely to learn anymore from interviews, or learn enough from interviews that would allow these reconstructions to be feasible given the limitations that we've encountered.

The radiological operations Vitro Manufacturing, like I said, they were actually uranium purifier and radium а purifier prior to World War II. In 1942, the Manhattan Engineer District contacted them to purified produce uranium from ores and concentrates, and so the start date is actually date 1942, which an August in coincides with the establishment of the Manhattan Engineer District. So that is the start of the covered period.

In 1947 they received, essentially, some additional work from the AEC to process scrap and uranium-bearing residues. Now scrap materials as I understand it were often just off-specification manufactured products. It's

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not like there was scrap metal that contaminated; it was mainly uranium product that just didn't meet specification, so went back for the uranium to be reclaimed out of, and Uranium-bearing residues, et cetera, and a number of other products. I think there was something about Canadian slimes, which I guess is some sort of byproduct of milling. And these products then, or these were all processed for uranium extraction for AEC at this point.

In 1955 most uranium processing ceased. There were still some small-scale activities. The waste residue removal began in 1956, and the last AEC contract terminated in 1957, so that is the end of the covered period.

Not all radioactive waste was removed, and there is in fact a residual radioactivity period for this site beginning at the end of the covered period. Our evaluation doesn't extend into that at this

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point, and as I recall -- I don't know for sure -- as I recall, we don't have any claimants who are strictly in the residual period. I believe all claimants had some employment at least in the covered period, and then some of them do extend into the residual period as well.

The available monitoring data, we actually do have some monitoring data from employees there. We have some uranium urinalysis results for the years `50 to `54. It is not clear this is comprehensive. It's not clear we have all the samples, and there are some legibility issues with some of them. Some of those you can't necessarily read. But there are a number of legible ones as well.

And then there are a limited number of radon samples from 1944 and from 1950, not from the entire period, just from those two years. But even at the time, the people taking the samples didn't express a lot of

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confidence in those results. So we don't think that they are going to be of much value to us; that is a radium body burden bioassay technique that given at the time that they were taken, they weren't considered very reliable. We feel we would be hard-pressed to put much confidence in those.

On the external monitoring side we do have external monitoring periods from February of 1944 through March of 1954, and those appear to be complete. It appears we have essentially a comprehensive list of the external results of the people who were monitored for those years.

Workplace monitoring, we have some area air sampling, a very limited number of breathing zone air samples for certain years, '49 to '53, those are in total alpha activity. And the -- but the breathing zone samples are too limited to really draw conclusions about the entirety of the exposure situation, and the general area air samples are oftentimes

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difficult to correlate to what was actually in the breathing zone of workers from general area air sampling.

And there are some surface contamination surveys as well, but it is hardly a comprehensive survey, that we feel like we have a comprehensive knowledge of the contamination situation at the plant.

In talking about feasibility of dose reconstructions, the available bioassay that we have, the internal monitoring results, are for uranium only, and since this did purify uranium, it extracted uranium from ores and residues and other materials, the degree of disequilibrium between the uranium and the uranium progeny is variable, depending on the site location and the particular process. So we don't have a constant relationship between the progeny and the uranium, so we can't use the uranium bioassay results to deduce what the non-uranium intakes would be.

While I'm on the subject, and I

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don't think this comes up in a slide, I did want to mention also, we don't have, really, information that the material was well controlled or confined to some portion of the facility, and our indications are that anybody that worked at the facility could very well have been exposed to the material. It wasn't limited to certain job titles, for instance.

The limited breathing zone samples we feel are not sufficient to give us adequate view of what the an exposure situation was over the whole period of time, and the general area air samples, we do have more of those, but it is very difficult to deduce breathing zone concentrations from a general area air sampling program.

The breath radon samples, as I said, were not considered reliable at the time, and so we don't feel like it would be appropriate for us to use them in any manner, and I don't know if we would have -- I'm not sure we have the technique to even interpret

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those, and so we don't know very much about them.

So in terms of our determination for internal dose reconstruction, we have determined that internal dose reconstruction is not feasible for uranium progeny.

For uranium exposure, we intend to use the bioassay if necessary, when it is necessary to due dose reconstructions, members, for people who aren't compensated via the class, we intend to use their uranium bioassay core claimant. If a claimant has a uranium bioassay, we intend to use that to interpret their uranium internal exposure. But we don't have what we feel is a sufficient data set to allow us to build something like a coworker model to reconstruct internal doses for unmonitored people. If we have claimants who don't have bioassay data, we don't think we can do an internal uranium assessment for them.

For external, like I said we have

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external monitoring data for the years 1944 to 1954, so for those years, certainly we can -- we will use those if we need to do dose reconstructions, we will use those data for claimants who have it.

We have not really determined whether we feel it is a sufficient dataset to do coworker models. We might very well. We might be able to do a coworker study, satisfy ourselves that we feel like we have gotten a decent representation of the monitored people. It appears -- we don't know for sure we have all of them, but it appears to be a complete set that we have.

And it may very well allow us to do some sort of coworker approach, certainly for those monitored years. And I don't know about extending it. We haven't really decided if we would extend it. These decisions will have to be made before we complete the partial dose reconstructions for people who aren't compensated through the class.

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There's where I am -- we have not made a determination as to whether we can reconstruct the external dose for people without exposure records. We do believe we can reconstruct occupational medical dose using some existing complex-wide documents that we use for medical dose reconstruction.

In terms of feasibility, I kind of decided I'd throw a curve ball in here on the internal uranium and say it's infeasible and not feasible both - or it's feasible and not It's feasible for uranium feasible both. bioassay that we have. We intend to use the uranium bioassay we have if it pertains to a claimant and that person's internal dose, if we need to do a dose reconstruction. But for people who don't have their own uranium bioassay samples, we don't believe we can do internal uranium doses.

We don't find that it's feasible to reconstruct the internal dose from the uranium progeny for anyone, and from an external

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standpoint we believe we can reconstruct the medical X-rays and for beta gamma we believe we can reconstruct it for monitored workers for sure, or for the ones who have it, we will include that. For the unmonitored workers, we have not yet determined yet whether we would try to include them.

health endangerment And our determination, evidence we have no discrete incident that could result extremely high doses like you would receive from a criticality accident. But the evidence does indicate that workers in the class may have accumulated chronic radiation exposures sufficient to cause harm.

So we conclude that health may have been in danger for those workers covered by this evaluation who were employed for a number of workdays aggregating at least 250 in the class, or aggregated with other classes.

Our proposed class definition is all atomic weapons employer employees who

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worked at Vitro Manufacturing in Canonsburg, Pennsylvania from August 13th, 1942 through December 31st, 1957, for a number of workdays aggregating at least 250 workdays, occurring either solely under this employment or in combination with workdays within the parameters established for other classes.

And this summarizes our recommendation for the period. This is the entire covered period for the site. We don't believe it's feasible to do a complete dose reconstruction, and we are recommending that the class be added for this facility for this period.

I didn't include it in the slide, but I did look awhile ago, we have 21 cases from this facility, some six of those have been compensated through dose reconstruction already.

So I believe that's the end. If there are any questions?

CHAIR ZIEMER: Thank you, Stu.

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| 1 | Let's open the floor at this time for |
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| 2 | questions on Vitro. |
| 3 | MEMBER MELIUS: Yes, any idea of the |
| 4 | overall size of the facility? |
| 5 | MR. HINNEFELD: It was on the order |
| 6 | of 35 acres. |
| 7 | MEMBER MELIUS: Yes, but I mean, |
| 8 | number of people working there. |
| 9 | MR. HINNEFELD: I do not. I don't |
| 10 | know that we have a way to reconstruct how |
| 11 | many people worked there, but I do not know |
| 12 | right now. |
| 13 | MEMBER MELIUS: And you have not |
| 14 | tried to reach out or do any community |
| 15 | outreach there, talk to - I know you are not |
| 16 | directly involved, so it may be hard to answer |
| 17 | this. |
| 18 | MR. HINNEFELD: I don't remember any |
| 19 | specifically for Vitro, no. |
| 20 | MEMBER MELIUS: Okay. |
| 21 | MR. HINNEFELD: We've been in the |
| 22 | Pittsburgh area, but I don't believe that was |

associated with Vitro. 1 Okay, I'm just 2 MEMBER MELIUS: trying to sense of the number get а 3 potential claims, and some of the, like we had 4 with the canal site, there was a huge worker 5 population, but there hadn't been enough 6 7 outreach for people I think to know they could apply. And I'm just curious on this one also. 8 MR. HINNEFELD: I don't really know. 9 10 Like I said, it's a 35-acre site. that could be -- I guess you could put a lot 11 of people in 35 acres. 12 13 MEMBER MELIUS: You can't say. MR. HINNEFELD: You can't say from 14 15 that. 16 MEMBER MELIUS: What they did after, other than --17 MR. HINNEFELD: Yes, the site became 18 19 - well, it was a storage site for a while, and then it became essentially an industrial site. 20 There was a -- or actually it wasn't a FUSRAP 21 mediation. The uranium mill tailings remedial 22

1 action program, remediation was done I think 2 in the 1980s, and there were some vicinity properties involved in that as well. 3 ZIEMER: Stu, I'd like to 4 CHAIR follow up on that. Was it a multiple-building 5 6 site? I think you said we really don't know -7 MR. HINNEFELD: We don't know --8 CHAIR ZIEMER: -- the extent to 9 which it was even controlled, right? 10 MR. HINNEFELD: Right, we don't know 11 a whole heck of a lot about it. The tailings 12 13 that were -- I believe we know that the tailings piles were just stored in piles at 14 the site. I mean it wasn't like confined in 15 16 any particular way, but there were tailings piles on the site, and some erosion from 17 those, as well. There was some migration. 18 CHAIR ZIEMER: And I think we should 19 recognize, this was at a time period when a 20 lot of uranium was being used in consumer 21

I've wondered on this particular

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products.

1 site, both from its name and location if this 2 might have been the source of the uranium which was sent to the Homer Laughlin Company, 3 near Pittsburgh, and which made 4 which is Fiestaware during this -- just prior to the 5 Manhattan Project. The folklore says that 6 7 Homer Laughlin Company made the red or what looks like orange Fiestaware right up until 8 the Manhattan project, at which point their 9 10 source of uranium was diverted to the project, and they ceased to make that particular 11 Fiestaware. 12 13 But my point is that uranium was widely used in consumer products, and the idea 14 15 that it could be hazardous probably wasn't in the picture very much at that time. 16 MR. PROCTOR: Dr. Ziemer? 17 CHAIR ZIEMER: Yes. 18 19 PROCTOR: This is John Proctor I grew up in Morgantown, West 20 in Las Vegas. Virginia. That's home of the glass country. 21

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CHAIR ZIEMER: Yes.

1 MR. PROCTOR: We used a lot of that 2 uranium for making yellow glass. CHAIR ZIEMER: Yes, yellow glass is 3 so-called Vaseline glass 4 the which contains uranium, and of course a lot of that 5 material still exists, mainly in antique 6 7 stores. MR. PROCTOR: There's a lot of it in 8 the old warehouses, a lot of glass company. 9 10 I've got a friend who's got cobalt, gold, and uranium still to this day they -- the AEC 11 comes by and checks it every year to see if he 12 still has it. 13 CHAIR ZIEMER: Well, it's a good 14 15 collector's item. Actually, the chair of this 16 committee has a large collection of Fiestaware which is almost at critical mass in his house. 17 MEMBER ROESSLER: Paul, this is Gen. 18 CHAIR ZIEMER: Yes, Gen Roessler. 19 20 MEMBER ROESSLER: Yes, to answer some of the questions that Jim had, whoever 21 to follow through on them I'd would want 22

| recommend health physicist Joel Lubenau. He |
|---|
| has done a lot of work and a lot of writing |
| about this particular site. |
| CHAIR ZIEMER: Well, it certainly |
| will be important to try to identify actually |
| what the total workforce was there who might |
| be eligible for this particular cohort. |
| Let's see if there are other |
| questions? Dr. Melius, do you have a follow |
| on? |
| MEMBER MELIUS: No, not really. |
| CHAIR ZIEMER: Other board members? |
| I want to ask if the petitioner is |
| on the line, and if so does she wish to |
| comment? |
| PETITIONER: Well, I don't think I |
| really have any comments. It's all been very |
| interesting. |
| CHAIR ZIEMER: Okay, thank you very |
| much. |
| MEMBER PRESLEY: Paul? |
| CHAIR ZIEMER: Yes, Bob Presley. |
| |

| 1 | MEMBER PRESLEY: I have a question |
|----|---|
| 2 | for the petitioner. Can you hear me, ma'am? |
| 3 | CHAIR ZIEMER: Use your mike there, |
| 4 | Bob. It might help. |
| 5 | MEMBER PRESLEY: Sorry. Ma'am, can |
| 6 | you hear me? |
| 7 | PETITIONER: Barely. |
| 8 | MEMBER PRESLEY: Can you tell us how |
| 9 | large the company was? |
| 10 | PETITIONER: Well, I would say they |
| 11 | must have had over 200 employees, I would |
| 12 | guess. I was in the lab all the time, and I |
| 13 | wasn't really so many of the workers came |
| 14 | into the lab, but usually they were the |
| 15 | foreman who would bring samples in. So I |
| 16 | really don't know how many laborers were |
| 17 | there. But I would imagine it had to be |
| 18 | around at least a couple hundred. |
| 19 | MEMBER PRESLEY: Thank you, ma'am. |
| 20 | CHAIR ZIEMER: Mike Gibson. |
| 21 | MEMBER GIBSON: Stu |
| 22 | CHAIR ZIEMER: Use the mike, Mike. |

| 1 | MEMBER GIBSON: In Section 10 of the |
|----|--|
| 2 | evaluation report, you talk about the |
| 3 | evaluation of a second similar class that may |
| 4 | be needed. Could you just give us a few more |
| 5 | details about that? |
| 6 | MR. HINNEFELD: Well, that would |
| 7 | have to be for the residual period, because |
| 8 | this class covers everybody during the |
| 9 | operational period. Any additional |
| 10 | consideration would have to be applied to the |
| 11 | residual period. |
| 12 | And the residual reconstruction |
| 13 | approach is sort of a topic of discussion and |
| 14 | debate elsewhere, and we kind of rely on much |
| 15 | the same technique, and it is being reviewed |
| 16 | elsewhere. |
| 17 | So we are kind of holding a |
| 18 | determination or a judgment on whether it's |
| 19 | feasible for that period. |
| 20 | MEMBER MELIUS: And that is also |
| 21 | sort of boilerplate language in these reports? |
| 22 | MR. HINNEFELD: Yes, the whole |

1 thing, we don't want to shut out -- if later 2 on we learn something else we don't want to shut anybody out by writing this without that 3 in there. 4 MEMBER MELIUS: One of the other 5 reasons was, I kept looking trying to find out 6 what the other class was. 7 I finally called LaVon and he explained to me that there wasn't 8 really one. 9 MR. HINNEFELD: Yes, I think you are 10 right, it's just to make sure that what we are 11 writing here doesn't close anybody out, that 12 13 we might learn something later on. PETITIONER: Could I add something? 14 CHAIR ZIEMER: Yes, certainly. 15 16 PETITIONER: When I said about 200, you might consider that a wild guess, because 17 I am really not sure about that. 18 19 CHAIR ZIEMER: No, we appreciate I think we wanted to get a feel for 20 whether it's a handful of people or thousands 21

At least it gives us kind of a

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of people.

| 1 | ballpark idea. |
|----|--|
| 2 | PETITIONER: It was more than a |
| 3 | handful of people; I know that. |
| 4 | CHAIR ZIEMER: And were there - |
| 5 | PETITIONER: There was a little |
| 6 | community right in that area called Strabane, |
| 7 | and so many of the workers were from that |
| 8 | particular area, because it was right adjacent |
| 9 | to the plant. |
| 10 | CHAIR ZIEMER: Okay, thank you. |
| 11 | Other questions? If there are no |
| 12 | other questions, it would be in order to have |
| 13 | a motion concerning this particular site. |
| 14 | Mr. Presley? |
| 15 | MEMBER PRESLEY: I move that we |
| 16 | grant the class petition. |
| 17 | CHAIR ZIEMER: Okay, the chair will |
| 18 | interpret that motion as a motion to recommend |
| 19 | to the secretary that this become a class of |
| 20 | the SEC. And seconded by Dr. Poston. |
| 21 | Is there any discussion on the |
| 22 | motion? If there is no discussion we will |

| 1 | have a roll call vote. |
|----|---|
| 2 | We will, after the vote, also seek |
| 3 | the vote of Dr. Lockey, and his vote will be |
| 4 | recorded as soon as we are able to get it. So |
| 5 | let us proceed. |
| 6 | Mr. Katz. |
| 7 | MR. KATZ: Ms. Beach. |
| 8 | MEMBER BEACH: Yes. |
| 9 | MR. KATZ: Mr. Clawson. |
| 10 | MEMBER CLAWSON: Yes. |
| 11 | MR. KATZ: Mr. Gibson. |
| 12 | MEMBER GIBSON: Yes. |
| 13 | MR. KATZ: Mr. Griffon. |
| 14 | MEMBER GRIFFON: Yes. |
| 15 | MR. KATZ: Dr. Melius. |
| 16 | MEMBER MELIUS: Yes. |
| 17 | MR. KATZ: Ms. Munn? |
| 18 | MEMBER MUNN: Yes. |
| 19 | MR. KATZ: Dr. Poston. |
| 20 | MEMBER POSTON: Yes. |
| 21 | MR. KATZ: Mr. Presley. |
| 22 | MR. PRESLEY: Yes. |

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1 MR. KATZ: Dr. Roessler. 2 MEMBER ROESSLER: Yes. MR. KATZ: Mr. Schofield. 3 MEMBER SCHOFIELD: Yes. 4 MR. KATZ: Dr. Ziemer. 5 CHAIR ZIEMER: Yes. 6 The motion carries, and we will 7 again develop the precise wording that will go 8 to the secretary and have that ready for the 9 10 Board's perusal tomorrow together with other action that we took yesterday. 11 Now we are just slightly ahead of 12 13 schedule. I am looking to see whether or not we will have Mallinckrodt petitioners on the 14 15 According to my notes, we will not have 16 any Mallinckrodt petitioners on the line, so I think we can probably proceed then with the 17 next item on the agenda, and that is the 83.14 18 19 SEC Petition, and Dr. Neton from NIOSH will present the evaluation report for this one. 20 MALLINCKRODT (1958) 83.14 SEC PETITION 21 DR. NETON: Thank you, Dr. Ziemer.

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Good morning, everyone.

I'm going to continue on the theme of discussion of uranium processing facilities and SECs by bringing forth the evaluation report for the Mallinckrodt Chemical Company, Destrehan Street plant for a very specific time period, that is calendar year 1958.

Okay, a little bit of a petition overview here. This was a petition that NIOSH evaluated in accordance with the requirements of 83.14, 42 CFR 83.14, that is, which is a petition submitted by a claimant whose dose reconstruction could not be completed by NIOSH because we didn't have sufficient information available to do the reconstruction.

This particular claimant was employed as a clerk-typist at Mallinckrodt from 1957 through 1960.

A little bit of the background here. Mallinckrodt should be very familiar to everyone; it was one of the first SEC classes granted under this program, and there were

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actually two classes added for Mallinckrodt. There was a period of time between 1946 and 1948 where it was determined that some data were technically unreliable. There were questions regarding the integrity of the data, and a determination was made that we couldn't reconstruct dose except occupational medical dose during that timeframe.

The latter period at Mallinckrodt was covered, 1949 to 1957, and it -- after some lengthy discussion with the Board and others, it was determined that the class would be added because there was insufficient information to reconstruct dose from the radium progeny, the long-lived progeny of radium or uranium, in particular, thorium-230, protactinium-231, and actinium-227 that's present in the raffinate material, which is a byproduct of uranium processing.

We did conclude, though, that we could construct external dose and some internal dose, depending on the availability

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I just want to this qo over raffinate term a little bit, because it is a term of art that is not, maybe, understood by It's used specifically to find the all. residues created from the refinement of ore in facility. And in fact the chemical extraction process that creates this material as Stu pointed out in his presentation creates a disequilibrium in these streams.

It's most important for these isotopes, radionuclides listed here, that is radium, actinium, thorium and protactinium.

To refresh your memory, I've just presented a slide that we went over in some detail during our deliberations for the SEC on Mallinckrodt originally, and you will see that the pitchblende ore is cleaned up through a chemical extraction process outlined here -- there's a pointer -- and as the pitchblende ore goes through it is resolved in the sulfuric acid material, and you create these

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cakes on the right-hand side that are lead sulfate and barium sulfate. The radium comes out of the process there, so you have an extraction of radium there, and then as you go further down the process, you end up on the right-hand side with the Sperry Cake and Airport Cake, and that is where the disequilibrium products occur for the thorium, actinium and protactinium in these cakes here. for And that in fact is the basis inability to reconstruct dose at Mallinckrodt, very much like the Vitro facility. don't know how much of this material was generated and became airborne, because there was no monitoring program, bioassay, or area monitoring program that could be used establish internal exposure the to those workers.

We reviewed the documentation and it indicated that operations similar to those that granted the class between 1949 and 1957 existed at Mallinckrodt into the 1958 time

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period. And in fact, no substantial difference could be found in those operations, and in looking at it knew that the we operations terminated some time in 1958, but it wasn't very clear as to when in 1958 that stopped. So we decided to include the entire calendar year to bound this particular class.

In addition to that, there was insufficient information to limit the class based on work location. These people often did not in changed jobs; work the same location of the plant at any given time. very much consistent with what happened in the Mallinckrodt; early class at couldn't we for instance, of raffinate create a class, workers, because we just couldn't tell who was a raffinate worker at any given point in time during the operation of the plant. And this is the same limitation that existed in 1958 in our opinion.

After `58 however the plant operations went into decontamination and

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decommissioning activities, very much different type of monitoring occurred. The large raffinate materials were no longer being generated, so we believe that we can do dose reconstructions in the D&D period at Mallinckrodt.

Much of this I have gone through for the feasibility of dose reconstruction. They continued to handle raffinate through `58. No individual bioassay does exist. And as I talk about it, records indicate workers rotated jobs.

The last bullet is important as because well, believe that workplace we materials and controls were insufficient to provide us any confidence that exposures were limited what. would consider to we traditionally occupational radiation exposed categories, that is rad workers, chemical operators and those types of people. really don't know whether administrative personnel were located very close, adjacent to

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the production activities, how often they had to traverse the work area, that sort of thing. So this class has no limitation based on job category at all.

We do believe however that we have sufficient data for reconstructing external and medical X-ray doses. We have external film badge data, and at a minimum we can apply the technical information bulletin we have for reconstructing medical X-ray doses, that we have used at many other facilities.

And internal exposure to radionuclides, for instance, the uranium, will be reconstructed using the data that we have available. So we are going to do whatever we can do with the remaining data that exist outside of t.he raffinate materials t.o reconstruct doses in this class.

So to get to the health endangerment issue, we do have evidence that these workers accumulated chronic radiation exposures. There is no evidence that there

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was an acute incident such as a nuclear criticality that created -- the class should be based on presence.

So therefore we are recommending the class be based on a chronic exposure scenario for workers who have aggregated at least 250 days within the parameters established for this class.

And the proposed class definition here would be, all employees of DOE, its predecessor agencies, and their contractors and subcontractors who worked in the uranium division at the Mallinckrodt Chemical Company Destrehan Street plant in St. Louis, Missouri, from January 1, 1958, to December 31st, 1958, for number of workdays а aggregating at least 250 workdays occurring either solely under this employment or in combination of workdays with the parameters established for one or more other classes of employees included in the SEC.

And the final slide is our

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recommendation that dose reconstruction is infeasible for members of this class, and health was endangered, and the covered period would be calendar year of 1958.

That completes my presentation.

I'll be happy to answer any questions if there are any.

CHAIR ZIEMER: Thank you, Jim.

Is it safe to assume that actually a good portion of the people in this time slot would already have been covered by the previous SEC? I'm just wondering --

DR. NETON: I believe that is a very good assumption. There are very few people in this class. In fact part of the delay in getting this class out was, we had identified one person who was a plant worker, a regular traditional-type rad worker, and they ended up being compensated in the first class.

So we had to go back to the table and identify another candidate for the petition.

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| 1 | CHAIR ZIEMER: So this would be |
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| 2 | mainly people who, perhaps, didn't have enough |
| 3 | time in the earlier class? |
| 4 | DR. NETON: Correct. |
| 5 | CHAIR ZIEMER: Or maybe started at |
| 6 | this time period. |
| 7 | DR. NETON: If they started in 1958 |
| 8 | early, or maybe late 1957. |
| 9 | CHAIR ZIEMER: So the numbers |
| 10 | involved here, we don't know exactly. |
| 11 | DR. NETON: Yes, I'm sorry I don't |
| 12 | have an exact number here, but it's a very |
| 13 | small number of people that are affected by |
| 14 | this designation. |
| 15 | CHAIR ZIEMER: Okay, thank you. |
| 16 | Other questions? Dr. Melius? |
| 17 | MEMBER MELIUS: That was actually |
| 18 | I was trying to understand why because I |
| 19 | thought we'd discussed this a little bit three |
| 20 | or four years ago. I thought we were sort of |
| 21 | aware of this potential issue. |
| | |

DR. NETON: I think it picks up a

few more people.

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MEMBER MELIUS: I was just trying to figure out why it took so long.

DR. NETON: Well, part of it was that we lost our what we called our litmus candidate who ended up being compensated. We had to go back to square one.

had to other part was, we substantially rewrite the site profile after the Mallinckrodt class was added. I think it final in November of 2005. So we became entire site profile pulled back that rewrote it so we could do non-presumptive And when we started to apply it I cancers. think the site profile was issued in 2007. was only then that we realized that this 2008 period was substantially the same as the class between `49 and `57, because originally if you recall it was our contention that we could do all of Mallinckrodt until the SEC finalized, determination then and we recommended that we add the class.

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| 1 | MEMBER MELIUS: My memory is too |
|----|--|
| 2 | foggy to remember. There was some issue about |
| 3 | we weren't quite sure about where to draw the |
| 4 | line. I can't remember why we ended up |
| 5 | choosing `57. |
| 6 | DR. NETON: Actually I think the |
| 7 | petition only requested the class through `57, |
| 8 | so we only evaluated through `57. Then it |
| 9 | dawned on us later on, in trying to |
| 10 | reconstruct `58, that it substantially had the |
| 11 | same characteristics as the class through `57. |
| 12 | CHAIR ZIEMER: Denise can add to |
| 13 | this, please. |
| 14 | MS. BROCK: That is correct. The |
| 15 | original petition ran from 1942 to 1957. I |
| 16 | think the slide said `46. But the petition |
| 17 | was from 1942 on. |
| 18 | CHAIR ZIEMER: Was the first class - |
| 19 | MS. BROCK: Yes. I think I had `42 |
| 20 | through `57, and I think the first estimate |
| 21 | was `42 |
| 22 | DR. NETON: `42 through `46 based on |

| 1 | the data reliability issue. |
|----|---|
| 2 | MS. BROCK: Then we went for the |
| 3 | rest of it. |
| 4 | DR. NETON: Then the raffinate. I |
| 5 | apologize for that error. But for some reason |
| 6 | `46 to `48 did seem like kind of a short |
| 7 | period. Thank you, Denise. |
| 8 | CHAIR ZIEMER: Josie. |
| 9 | MEMBER BEACH: When did operations |
| LO | end there; what year? |
| 11 | DR. NETON: That is a good question. |
| 12 | The cleanup persisted for some time, and I |
| 13 | believe that the residual contamination goes |
| L4 | through the 1990s or somewhere thereabouts. |
| L5 | MS. BROCK: That is correct. I |
| L6 | believe that that actually stopped in 1962, |
| L7 | and Weldon Spring started in 1955, at least |
| 18 | that was prior to production, but when they |
| L9 | were getting everything ready. And I think |
| 20 | the production actually started at Weldon |
| 21 | Spring in 1957/58 timeframe, and went into |
| 22 | like 1967 or `68. |

DR. NETON: That's part of our problem is identifying who actually worked at Mallinckrodt, Destrehan Street versus Weldon Springs, because same company.

MS. BROCK: Same company, and then there was also United Nuclear which ran about the same time, that was in the hematite area, and that ran at the same time as Weldon Spring did, so it's very confusing.

CHAIR ZIEMER: Thank you. Other comments or questions? Jim.

MEMBER MELIUS: Yes, I have one just general question. It seems that we get in these situations where we are sort of stuck by what the petitioner has put in for numbers of years, and they are -- depending on who is involved in the petition they may not know the exact time frames that appropriate. are Sometimes NIOSH modifies it it as comes forward, but then we end up like this in situation where we closed off at `57, and then you have to wait until you find an

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person.

DR. NETON: Well, this is a little unique in this respect. If you remember, NIOSH's official position was that we could do dose reconstruction through `57. It was only until the Board deliberated, and the decision was made through the secretary that we couldn't do it. And that's why -- we would normally expand the class beyond the `57 and look to see, is that an arbitrary date or is there reason to continue forward. In this particular case it was NIOSH's position we could do it, so there was no reason to look about that at that juncture.

CHAIR ZIEMER: Denise may have an additional comment.

MS. BROCK: If I remember correctly, and it's been quite some time ago, I think that the designated timeframe was 1942 through `57, then sometimes it happens that additional years are added. Because I would have actually petitioned for that entire timeframe.

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| 1 | So it was my understanding that, during that |
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| 2 | initial period, it was just showing `42 |
| 3 | through `57, and I think after that we started |
| 4 | seeing additional time in the Federal |
| 5 | Register, that DOE had actually extended that |
| 6 | if I remember correctly. |
| 7 | CHAIR ZIEMER: But I think the |
| 8 | comment that Jim made probably is has been |
| 9 | the pattern where NIOSH says we can't |
| 10 | reconstruct dose, and they have expanded in |
| 11 | some cases based on what they know about it. |
| 12 | DR. NETON: We would normally do |
| 13 | that, to look about that the proposed end |
| 14 | date to make sure that some other situation |
| 15 | doesn't exist. |
| 16 | MS. BROCK: I thought that was the |
| 17 | situation in this one. |
| 18 | DR. NETON: But this one I think we |
| 19 | just flat out thought we could do it from the |
| 20 | very beginning, and there was no reason to |
| 21 | look past `57 because if we could do `57 we |

thought we could surely do `58; that was our

position at that time.

MS. BROCK: But at the beginning didn't they believe there really wasn't production going on after that, that it had actually went out to Weldon Spring? That was my thought.

DR. NETON: That was also part of it, the production we thought ceased in `57, and then when we looked closer at the records, there is a piece going into `58. We are not sure when it stopped in `58, but it clearly did continue into 1958.

MS. BROCK: Thank you.

CHAIR ZIEMER: Well, in any event, we are sort of correcting past oversights here.

MEMBER MELIUS: No, I just worried sort of about the potential petitioners, that they are told `57, so anybody that started working in `58 -- and there has been all this outreach and publicity; Denise was part of that. Now we have this new sort of very small

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class, and might have -- in retrospect it would have been better. So I was just thinking procedurally -- and again I don't think we all remember -- at least, I don't remember the details of this, but whether we are better off leaving open the possibility to go and review just rather than start doing a total close-out and trying to keep some of these open for review. I'm not sure it was possible in this one.

CHAIR ZIEMER: Well, in fact we've had a couple of fairly recent SECs where we were told at the time we approved them that the end date was somewhat uncertain and that, if necessary, it would be extended even after we had approved it.

DR. NETON: Los Alamos falls in that category right now.

CHAIR ZIEMER: So I think to the extent that NIOSH is able, at least they are trying to accomplish that, perhaps not always successfully. We obviously don't want to

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| 1 | piecemeal it year by year and month by month. |
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| 2 | MEMBER MELIUS: And if we rely on |
| 3 | the 83.14 then NIOSH has to wait until it |
| 4 | finds the right candidate case, track them |
| 5 | down, and as Jim said, this case wasn't the |
| 6 | right case. So they are compensable, and it's |
| 7 | difficult. |
| 8 | CHAIR ZIEMER: But your point is |
| 9 | well taken, thanks. |
| 10 | Other comments on this? If not, it |
| 11 | would be appropriate to have a suitable motion |
| 12 | for action. |
| 13 | Brad Clawson. |
| 14 | MEMBER CLAWSON: I move to accept |
| 15 | it. |
| 16 | CHAIR ZIEMER: A motion to accept, |
| 17 | which is a motion to recommend to the |
| 18 | secretary that this class be added to the SEC. |
| 19 | MEMBER MUNN: Second. |
| 20 | CHAIR ZIEMER: And seconded by, I |
| 21 | think, Wanda Munn. |
| 22 | Further discussion? Then let's |

| 1 | proceed. Again, a roll call vote, and we will |
|----|---|
| 2 | seek Dr. Lockey's vote as soon as possible. |
| 3 | MR. KATZ: Ms. Beach? |
| 4 | MEMBER BEACH: Yes. |
| 5 | MR. KATZ: Mr. Clawson. |
| 6 | MEMBER CLAWSON: Yes. |
| 7 | MR. KATZ: Mr. Gibson? |
| 8 | MEMBER GIBSON: Yes. |
| 9 | MR. KATZ: Mr. Griffon? |
| 10 | MEMBER GRIFFON: Yes. |
| 11 | MR. KATZ: Dr. Melius. |
| 12 | MEMBER MELIUS: Yes. |
| 13 | MR. KATZ: Ms. Munn? |
| 14 | MEMBER MUNN: Yes. |
| 15 | MR. KATZ: Dr. Poston. |
| 16 | MEMBER POSTON: Yes. |
| 17 | MR. KATZ: Mr. Presley? |
| 18 | MEMBER PRESLEY: Yes. |
| 19 | MR. KATZ: Dr. Roessler. |
| 20 | MEMBER ROESSLER: Yes. |
| 21 | MR. KATZ: Mr. Schofield. |
| 22 | MEMBER SCHOFIELD: Yes. |

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1 MR. KATZ: Dr. Ziemer. 2 CHAIR ZIEMER: Yes. There are no abstentions, at least 3 not so far, unless Dr. Lockey abstains. 4 But in any event, the motion carries, and we will 5 6 proceed with the recommendation to the 7 secretary, and the exact wording of the letter to the secretary will be provided later in 8 this meeting to the Board members. 9 10 Now I'm wondering if we can go ahead with the Blockson report. According to 11 my notes there may be a Blockson petitioner on 12 13 the line to comment, also. And since we are ahead of schedule, I'm wondering whether we 14 15 need to delay. 16 Ms. Munn, as I understand it, you may not have a specific recommendation for us 17 but rather a report this morning, is that 18 19 correct? MEMBER MUNN: That is correct. 20 CHAIR ZIEMER: I wonder if we could 21 go ahead and proceed with the report. 22

petitioner is not on the line now, we would 1 2 allow that petitioner make additional to 3 comments. if 4 MR. KATZ: Let me see the Are either of the petitioner is on the line. 5 6 two petitioners for Blockson on the line? Ιf 7 you would let us know if you are already listening. 8 (No verbal response.) 9 MR. KATZ: Apparently not. 10 CHAIR ZIEMER: Okay, let us proceed, 11 I think, with the Blockson report, and then if 12 13 petitioners are on the line later we will give them the opportunity to add any comments. 14 BLOCKSON CHEMICAL SEC PETITION STATUS UPDATE 15 16 MEMBER MUNN: Thank you, Dr. Ziemer. Let me refresh your memories for 17 those of you who are not a part of 18 19 workgroup involved in this particular site. have begun our deliberations 20 based on seven specific findings which our 21 contractor brought to us when they reviewed

the site profile. And the bulk of those centered around the extraction process and concerns over what radionuclides were contained in the raffinates, and specifically whether thorium followed one stream or the other during the process.

We fairly quickly resolved all of those outstanding issues. After consultation with workers, two worker group meetings, and extensive deliberation among the members of the workgroup, the contractor and our NIOSH representatives, at a meeting earlier this year, we brought this to the Board with an unformed consensus from the workgroup.

At that time it was the direction of the Board for us to go back, address very specifically the concerns that had been raised with respect to radon, and focus on that issue and see if we could resolve it yet further down the line.

We did that. We had a workgroup meeting in Cincinnati on October $15^{\rm th}$, and left

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that meeting with a number of questions relative to a few technical issues that had evolved from those discussions.

Throughout the month of November, there was a significant amount of information exchange: emails and technical conversations between NIOSH and the contractor. And on the 3rd of December we had a technical call, not the Work Group itself, but the technical folks who were specifically concerned with what's possible and what is not possible in the real world.

Then on the 12th we had scheduled
- it was the only available date that we had
following that technical call, that we could
arrange to have the Board members who were
involved in the Work Group present, and gave
both NIOSH and SC&A an opportunity to produce
a couple of additional background papers, one
involving an outside expert on air movement.

At this juncture, we are down to a primary concern with respect to radon and air

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exchange rates inside Building 40, because of power outages and problems that some of our work group members had with the weather last week were unable to conclude we our 12th deliberations meeting our on the on primarily because some of us had not had an opportunity to fully digest the material which had just been produced in the technical papers that I mentioned.

At that time it was suggested that, rather than try to move further until the Work Group members themselves had all satisfied themselves with respect to the content of that material, we would instead continue to delay this process. The Work Group Chair was really very concerned about that, because we have been attempting to bring this to closure now for quite some time, and I'm concerned about any further delays on behalf of the claimants.

But we didn't see any way around it in this particular instance. So what we expect to do at this juncture is, during our

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Board administrative time tomorrow, we want to set up work group call time for all of us to have had an opportunity to review all of the material from the beginning to satisfy ourselves that there is no further item of interest to anyone to be addressed, and present this matter for а vote the Albuquerque meeting.

That is the current plan. My apologies to all of you, and especially to the Blockson claimants for the length of time that is being involved here. We are trying to follow every thread that has been presented to us, and every potential concern that has been raised to its ultimate end.

It remains my personal opinion as the chair that we have the body of evidence our supporting ability to address reconstruction here, but that is not Work opinion of the Group. I'm expressing a personal opinion; that is not the position of the workgroup.

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We will certainly, of course, address that issue at our work group meeting when we determine that date.

We hope to be able to do that by telephone, hopefully toward the end of January, in adequate time to have a good presentation available for Albuquerque.

CHAIR ZIEMER: Thank you very much.

Let me see if there is any question that Board members have concerning the Blockson report, or any comments from other members of the workgroup, or others.

Mark.

Wanda, this MEMBER GRIFFON: at point I was following some of the documents and I was considering dialing in Friday but I had no phone or electric at that time. the radon, I know there was some discussion on the radon model -- has NIOSH put forward a new Or are we at this point are we still model? discussing SC&A's model for the radon? initially other words, know, in the Ι

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evaluation report, there is a certain approach described, and I don't know where -- whether NIOSH has revised that approach, or whether that is still the approach we are considering.

CHAIR ZIEMER: Jim Neton.

DR. NETON: During deliberations of the working group NIOSH put forth that they would adopt the model prescribed by SC&A with one exception, and that would be the lower bound air exchange rate for Building 40, we would adopt one air exchange per hour versus their I think recommended .25 or something like that, and then we would end up with a Monte Carlo generated distribution of possible potential air concentrations within the building. So that is where we're at.

MEMBER MUNN: This is why I said, we are down to the question, the sole question of air exchange.

CHAIR ZIEMER: Thank you. Further questions? Ted Katz.

MR. KATZ: Let me just add something

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to this report. I think there was a very illuminating dialogue as part of this process. So we will have a verbatim transcript, and I think one of the reasons for postponing also was so that the full Board could have the benefit of that transcript. And I have asked for it to be expedited. It will be available to us early in January, and we will distribute it to the full board. But I think it helps, because it was an extensive dialogue and very clear and raised a lot of important issues, thought that that would really be helpful to the full Board when it begins its deliberations as well. So I just want to add that point.

MEMBER MUNN: That's doubly important in light of the fact that several Board members made the comment in previous presentations that they had not investigated themselves the material that was available, and I was asking for a packet of information and I had directed them to the record that

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exists.

So thank you, Ted, for reminding me that one of the items that we hope you will take an opportunity to look at even before it comes to you is the transcript when it comes out.

CHAIR ZIEMER: Thank you.

Other comments or questions?

Perhaps let me check again to see if either of the Blockson petitioners are on the line at the moment.

Either Blockson petitioner on the line or wish to comment?

Okay, we will check again after the break to see if they are here.

I think what we will do at this time is go ahead with our morning break. We are a little bit ahead of schedule, but let's plan to reconvene at 10:40. That will give you a decent size break and still keep us a little ahead of schedule.

(Whereupon, the above-entitled matter went off

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| 1 | the record at 10:13 a.m. and |
|----|---|
| 2 | resumed at 10:42 a.m.) |
| 3 | CHAIR ZIEMER: If you would take |
| 4 | your seats, we will reconvene the meeting. |
| 5 | I'd like to double check the phone |
| 6 | line. Dr. Roessler, are you on the line? |
| 7 | MEMBER ROESSLER: Paul, you said |
| 8 | something about the phone line, and then I |
| 9 | unmuted. This is Gen. |
| 10 | CHAIR ZIEMER: Oh, Gen, I was just |
| 11 | checking to see if the phone lines were on, |
| 12 | and if you were on. |
| 13 | MEMBER ROESSLER: Yes, I am. When |
| 14 | you unmute, you can't hear anything for a |
| 15 | minute. |
| 16 | CHAIR ZIEMER: Okay, thank you, Gen, |
| 17 | we are getting ready to reconvene here. |
| 18 | Before we have our next |
| 19 | presentation, I do want to check at this time |
| 20 | to see if either of the Blockson participants |
| 21 | or petitioners are on the line. Either of the |
| 22 | Blockson petitioners. |

1 MS. PINCHETTI: Kathy Pinchetti is 2 here. CHAIR ZIEMER: Okay, Kathy, thank 3 you for being on the line. This is Dr. 4 Ziemer, and I want to tell you that just prior 5 6 to our break, since we got a little ahead of 7 schedule, Ms. Munn, the chairman Blockson workgroup reported to the 8 basically that the workgroup 9 had no recommendation at this time because they are 10 still dealing with the radon issue. 11 I'm assuming that the workgroup has 12 13 kept you apprised of the issues that they are working on; is that correct, Ms. Munn? 14 Have 15 the petitioners been involved in those 16 deliberations? MUNN: 17 MEMBER We have not. had specific communication with them. It has been 18 19 our desire and our hope that they have been joining us on our open -20 CHAIR ZIEMER: In any event, Kathy, 21

be

aware

wanted you to

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just

the

that

workgroup has indicated to us that they are not prepared to make a recommendation yet - are you still there? I'm hearing a lot of noise.

MS. PINCHETTI: Yes, I'm still here.

CHAIR ZIEMER: Still dealing with the radon issue and they are hopeful that they will be ready at the next Board meeting to make a more definite recommendation.

But we did want to give you an opportunity if you had some comments at this time to make those. You may or may not, but you are certainly welcome to do so if you had any comments for the Board.

MS. PINCHETTI: Well, I did get a copy of the SC&A report from Laurie Breyer, as well as the Harley report. And it still seems like the focus is on the 25-year-old spot of radiation that was found in `83, and that was 20 years after the contract ended. So in other reports it seemed like there was discussion about how the radiation dissipates,

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you know with the exchange of air over time. 1 2 And it just seems like a 1983 spot would have lost a lot of the radiation by the time it was 3 And I'm not clear if that spot was 4 found. found in Building 40 or Building 55. 5 CHAIR ZIEMER: Okay, I'll let Ms. 6 7 Munn address that, and also the issue of the air turnover rates in one of the buildings was 8 under discussion. But maybe Ms. Munn can add 9 10 to that. Did you understand the question that was being asked? 11 MEMBER MUNN: Yes, I did understand 12 13 the question. CHAIR ZIEMER: Can you hear Ms. Munn 14 okay? 15 16 MS. PINCHETTI: I can. MEMBER MUNN: The issue with respect 17 to the air sample is not as pertinent actually 18 19 as the issue of the air exchange. And that's what we've been focusing on. 20 The reason I made that is because 21 statement you are

correct, some of the measurements that have

been provided to us were made after the covered period. But they have been used as verification that the levels could not have been higher than that. The process is now quite well understood, and the amount of radon that could be possible is now - could be available from the process is now quite well understood.

Our issue is primarily how much of that stays in the building and how it is being transferred by the normal airflow through the building.

So that is our focus at this moment. So far as I understand that is the final aspect of the full production picture that has not yet been tied down.

CHAIR ZIEMER: I am going to suggest - the workgroup plans to schedule another meeting I believe by phone, and I'm going to suggest that you make sure that the petitioners are aware of the time of they meeting listen in so that can and

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1 participate with that. Kathy, is that 2 agreeable with you? PINCHETTI: Yes, that's fine. 3 MS. 4 It's just that there is a big time change. It's approximately 7:45 right now. 5 And I work. 6 7 CHAIR ZIEMER: So we need to make sure that we can find, perhaps can find a time 8 that is also convenient for the petitioner in 9 10 some way. MS. PINCHETTI: That's okay. 11 I'm not asking for any accommodations. 12 It's just 13 that I need to make arrangements at work so I can listen in. 14 15 CHAIR ZIEMER: Okay. MS. PINCHETTI: One other question I 16 had was about Florida. I don't know if we are 17 basing the Florida comparison based on that, 18 19 because that is where the rock came from. don't know if there are purchase orders that 20 show that data was actually from Florida or 21

somewhere else, or if the Florida comparison

is because - they also had a phosphoric process going on there.

But in the reports that I just got they are making reference to the 25-foot doors being open, and in Illinois I don't think that is possible for probably nine months out of every year, because it is either snowing, or there are sub-zero wind chills, or there's a tornado brewing. So I don't know.

And also that the vents were frozen shut in some of the buildings because it was so cold, the vents in the ceilings.

MEMBER MUNN: Let me address your first issue with respect to time. You certainly have this voice's sympathies. also live on the West Coast, and am not happy with being on the phone at six o'clock in the morning either. And so I feel fairly sure that we will be scheduling - we will be attempting to schedule a time that - certainly during the day, during the normal workday, and it will be at such a time that hopefully it

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will be convenient for you also.

We will have that time established after our work period tomorrow afternoon, and it will be posted on our information website.

I'm not certain, but it will appear on the website as scheduled workgroup call.

Yes, the material that was used at Blockson did in fact come from Florida. That has been fairly well documented now. And we are aware of the fact that the doors certainly would not be open throughout the year. That is a part of the discussion that - several of the workers did mention however that they were greatly relieved during the summer months to be able to have them open because of the chemical fumes that accumulated during the process.

So it was our understanding that when they could be open they made every effort to do that.

MS. PINCHETTI: Okay.

CHAIR ZIEMER: Well, thank you. Did

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you have any additional comments then, Kathy?

MS. PINCHETTI: No, no, I'm fine.

Happy holidays to the Work Group. I know you
guys have been putting in a lot of work on
this.

CHAIR ZIEMER: Thank you for being with us this morning. Thank you.

We will proceed then to the next item on the agenda, which is the Department of Energy update. And we are pleased that Dr. Patricia Worthington is with us this morning. She had a difficult time getting here with fog and other issues yesterday. But Pat, welcome, and we are pleased to have your update at this time.

DEPARTMENT OF ENERGY UPDATE

DR. WORTHINGTON: Good morning. I'm pleased to be here. I certainly was challenged quite a bit yesterday to try to fly here. A lot of fog in the Augusta area. But I am pleased to be here today, and thank you for readjusting the schedule to fit me in.

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I wanted to give you an update on where we are in supporting this program. No major changes in the roles and responsibilities of DOE, but I want to give you a few statistics, and talk about some initiatives, and to address any questions you may have. And I have two members of my staff that are in the audience as well, Greg Lewis is here and Isaf, she's here as well. So we are happy to give this update.

A little bit about the activities We have sort of three main areas, and we continue to focus on those areas. One is to provide information for the individual claims. Those primarily employment are verifications and exposure records. We provide support to the Department of Labor, to NIOSH, and to the Board and its contractors through research and retrieval of documents from various DOE sites, and we update covered facilities information.

I mentioned that I would give a few

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statistics, and here are just some, to give you an idea of sort of the magnitude of the support that we provide. In terms of employment verification, nearly 7,000 a year dose records for NIOSH, about 4,000, and then employee work history and exposure rates, about 7,500 per year.

Again just a few more statistics.

A little bit about what we did in 2007 and 2008. Basically we had almost 22,000 completed requests in 2007 and about 18,000 in 2008. Again those were the numbers for those years. It's not intended to reflect a trend that things are going down, but just to give you sort of the stats.

For example in 2006 I think we had about 16,000 completed.

The SEC support is certainly a huge effort for us. You see on this slide that we are supporting a number of them. Probably the biggest ones for us right now are Hanford and Savannah River, and I will talk a little bit

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more in detail about what we are doing in those areas.

The Savannah River one, we have certainly been active, and since June up to this point we've hosted seven visits for NIOSH, major visits there. And we've been doing a number of things to improve access to documents, and to make those things more available. And you see a term here called, electronic document work flow system. It's something that we made available to NIOSH so that they can on their own search it for key words and key phrases and be able to get a better feel for what kind of things they might need.

We have completed document reviews for over 2,000 documents, nearly 300,000 pages. That certainly was a significant effort on the part of getting things ready at Savannah River.

And the last bullet is intended to talk about document reviews for what we call

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classified documents. There were 157 of those documents.

Let me back up a little bit to this almost 300 pages of documents. When I say that we have conducted those reviews, we have actually conducted reviews, we've gathered information, we've scanned the materials and they have actually been transmitted to NIOSH for their use.

I will switch now and talk a little bit about the Hanford SEC, and the kinds of things that we are doing there, and continue to do. The first bullet is intended to talk about 400 unique boxes. There are many boxes at the site. There are certain things that characterize those boxes. We have pulled 380 boxes, and there was an opportunity to go through those boxes and determine exactly what is needed, because typically you may not need everything that's in a box.

And as a result of nearly a million pages, 1,000 specific documents were

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The same with Savannah River.

We've hosted six major visits at the site for

NIOSH, and a number of smaller visits.

I want to go to the next slide and talk a little bit about the tours in terms of hosting a visit and what does it Certainly talking we are always about documents and retrieving documents. But it's nothing like being on the ground and kind of getting a sense for the operations and what went on there and the size of the facilities and the types of operations. So at Hanford there were multiple facility tours in various kinds of buildings. Some of the buildings were actually contaminated. Some of the buildings were undergoing demolition or D&D, so a wide variety of buildings and activities.

As you know Hanford is a very complex site. There are many different contractors with different missions going on at that time. We have great cooperation from

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all of those contractors, and all of them are involved in participating in making the information available and looking for ways to better do that.

Again, about improving access, we are looking at record systems access so that NIOSH can search the systems themselves for key words and key activities, and look for ways to determine what kind of information might be needed.

In terms of additional staff, certainly when you have a big project like an SEC, the existing staff that were available at that site to do records retrieval might not be sufficient. So a number of people were added, I think six additional individuals added to help with that process, including individuals that are cleared and that would understand classification.

Office space and computing equipment, that is very critical that we make that available so that people can have space

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to look at documents to do staging of documents and so forth, and they have computers that they can use right there and work on the information.

So all of these are part of our initiatives to make the information more available, and make the individuals that need to do the review more knowledgeable about the kinds of things that went on at those sites and those operations.

The next slide is just intended to provide a quick overview of the things we do in terms of supporting the Department of Labor. Ultimately we want to make sure that they have all the information available to them to be able to make decisions on claims. So this was part of working our interface.

Again, I've talked before about tours. We've made sure that the Department of Labor had an opportunity to tour facilities as well.

We met with Department of Labor

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staff in Seattle to discuss issues related to the tank farms. There are a lot of complex issues associated with that, again, it's a way of making people more familiar with these unique activities that have gone on within the Department of Energy.

The last bullet is just making sure that where we have SMEs that are available and are knowledgeable about hazards, where it's appropriate we can make them available to the Department of Labor to provide some additional insights.

Again this is about the covered facilities. We continue to do that as more information becomes available, if there is a need for additional research to work on expanding that, we certainly take an opportunity to do that.

A little bit about initiatives: I talked about making people familiar with the operations on the sites and the facilities.

Another thing we want to continue to work is

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our interfaces, to make sure that communicating with the various groups we have to work with. We have designated a specific point of contact within our organization to regularly with work the Board and the contractors and NIOSH to make sure we understand all their issues. We hold weekly conference calls with the members of NIOSH and its contractors to make sure that they receiving the information, there aren't problems or concerns, or if there are things we need to do different or to work on those things. We don't want to wait until things build up. So we're looking to a designated individual, weekly interactions, and in some cases daily if things are - if we need to do that.

We talked at the last few meetings here about our Office of Legacy Management. They have great experts there who are very familiar with record retrieval that continue to support us in looking for covered facility

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information, any of the things that we need additional research, we certainly reach to that organization to help us do that.

little bit more about the This initiatives. process of retrieving information and developing reports and making sure that documents are reviewed certainly is one that we've worked on quite a bit this year, and we continue to collaborate with the various organizations, and we believe that we've streamlined that process and that we are able to do things much faster and much more efficiently than we were in the past.

We attended the NIOSH Advisory Board Meeting and the Department of Labor town hall meeting, so whenever DOE can be available to support those activities, we want to do that.

The next initiative I want to spend just maybe a moment talking about that, in terms of how do we do things better? How do we improve the record retrieval and retention

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process in the department? We wanted to start at the head of the Department of Energy, at the Department of Energy. The records issues are owned by the CIO's office. So we've been working with that organization, and working with procurement and contract our organizations to ensure that we have contract mechanism in place that will make sure that we can access and maintain ownership of records. We look to our contractors when we employ them at the sites to do certain things to be able to help us do that. want to make sure the contract language This certainly is very important in clear. the area of subcontractors, because they are workers too, and want to be able we to retrieve information regarding subcontractors when it's needed, and that's been a challenge, much more so than some of the other things that we've been doing. So we are trying to make sure that we have a process in place that we can do a better job in that area.

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The next bullet deals with a site specific one that we have been working on. And that is to work with the Los Alamos medical center to get records there of DOE Before 1964 the hospital was a part workers. of the DOE and its predecessor organization. After that time it was a separate private organization, records, and some records, were left at the hospital. been working with the hospital. We have a mechanism now. We have a process in place where we can go in and clean up those records, sort them, package them, retrieve them and have them available so that if people request them that information can be provided.

The next slide is intended to kind of summarize some DOE activities. This is primarily about the DOE oversight POCs. We have them at every one of our major sites, and they play a major role as part of the site and part of the federal organization to help in this process they attend public meetings.

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They work with DOE and NIOSH to facilitate the interviews with current and former workers.

In one of our sites in particular this particular POC actually goes out and meets with former workers in their homes there they are comfortable in having discussions, and talk with them about EEOICPA and what they can do and how they might apply.

So again, looking for other ways to reach out to these organizations, to these individuals, and make sure that they are aware of information and mechanisms and processes that might be available.

The next slide is intended to be one that will spark some questions and I'm here certainly to answer any questions that you might have. I do want to reflect back on one of the slides that I put up earlier, and that was about streamlining the process. And I want to mention sort of the role of our organization in terms of streamlining that organization in terms of HSS. If there are

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| 1 | documents that are subject or need to be |
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| 2 | reviewed, it is a high priority of our own |
| 3 | organization, the people that you see here |
| 4 | that you are working with to make sure that |
| 5 | they happen, but it's also a high priority |
| 6 | across the HSS organization. So at any time |
| 7 | if we need to involve the security |
| 8 | organization, this becomes the highest |
| 9 | priority for them, and there are individuals |
| 10 | that have been designated to work with us and |
| 11 | to quickly act upon these things that need to |
| 12 | be reviewed. |
| 13 | Having said that, I am available |
| 14 | now for questions or for further elaboration |
| 15 | on any of the quick topics that I've mentioned |
| 16 | today. |
| 17 | CHAIR ZIEMER: Thank you very much, |
| 18 | Dr. Worthington. |
| 19 | Let's open the floor for questions |
| 20 | or comments. Brad Clawson. |
| 21 | MEMBER CLAWSON: First of all, I'd |

like to thank you. I just returned from

Hanford, and the tours we had up there were tremendous. It was very beneficial to us, especially the 300 area, the 100N area, because these are very complex sites, and they are hard to be able to keep up.

But one of the problems that I do see is that up high they're understanding what we need as far as being able to get documents processed through and so forth, but whenever we review documents, as a subcontractor or as a Board member, before we can take notes or anything else like that, they have to be cleared by you, which we understand, but it's taking a tremendous amount of time.

Some of our records are six months old. I still haven't got any of my notes from Mound. And I know that we are just getting this started, but I feel like this is an area where we can make some improvements.

DR. WORTHINGTON: We believe that we have made some improvements. There is certainly a ways to go, and some of the things

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1 are legacy things that happened before we set 2 these particular processes in place. looking hear back from all the 3 are to organizations in terms of real time if they 4 believe that documents are held up. And Greg, 5 you may have a comment on that. I'm not aware 6 7 that there is a holdup on the Mound documents. You may want to comment on what. 8 Actually, 9 MEMBER CLAWSON: I'm 10 speaking more to the notes. DR. WORTHINGTON: To the notes? 11 MEMBER CLAWSON: The notes that we 12 13 took when reviewed some of the we documentation and stuff. And this was done in 14 15 the Federal Building in Cincinnati, and then 16 we turned over all of our notes, and we are still waiting for them. 17 DR. WORTHINGTON: What timeframe was 18 19 that? Was it several months? MEMBER CLAWSON: What, four or five 20 It's when we went to Cincinnati. months? 21

was SC&A and myself. What it came down to was

worker interviews.

DR. WORTHINGTON: Worker interviews?

MEMBER CLAWSON: Yes.

CHAIR ZIEMER: Greg, did you have a comment on that, or can you enlighten us?

MR. LEWIS: Yes, this is Greg Lewis. That may be an issue of what the notes ended up being marked as, and who we can release them to, which speaks to the security plan we're working on how. But to my knowledge all of the notes were reviewed and released back to the person who submitted them.

DR. WORTHINGTON: And there's no reason that they would be held up for six months. So we need to follow up on that and make sure that they weren't directed to the wrong individual, or that we have made a mistake and they are still there and we need to forward them on, because six months is extremely long. Our process isn't designed for it to take that much time.

MEMBER CLAWSON: Well, and part of

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it too I think that we'll come to find out is that sometimes when we - especially with a worker's interview that has to go through a clearance process, and sometimes they get redacted or whatever else like that, at some point we are falling into a glitch because they get classified as somewhat classified or whatever, and we are losing track of where they are going, because they are going to have to go through a redaction process, and we never figure out - it's kind of hard to track where they are at.

Now we have made some substantial changes with that because through our subcontractor they are starting a process to be able to track where stuff is at. It'll make it a little bit easier. I know this is kind of the grass roots. But I hope that we can make sure that we can process through this as easy as possible. But there have been some issues.

MR. LEWIS: This is Greg again. I

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understand tracking is an issue difficulty. But for the most part on our end we have had notes back to you within - we're working on a 7 to 14-day turnaround period. I'm not sure if we have always been hitting that, but certainly our recent experience is And in addition as far we have been. reviewing original copies of notes, that is also available for someone with the clearances and in the right setting cleared location. They can always review the if originals it happens that the specific piece of information that was removed something that is important - typically that's not the case - but they can be reviewed in their original form as well.

DR. WORTHINGTON: But this action we will take back and look for notes from Cincinnati about six months ago that were submitted for review never should last this long. So we will get back to you. We will look into that and see what happens.

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CHAIR ZIEMER: Maybe you can get the
exact date of that before you leave and that
would simplify this.

MEMBER CLAWSON: I can probably get
the exact date because what this covered was
on Mound. But and this is - you probably hit

probably a legacy of before we kind of got

into some of this process. And I think it's

the nail on the head when you said this is

trying to catch up with a lot of the notes and

interviews and so forth that we need to do,

and maybe we can sit down with our contractor

and make sure where all this stuff is at.

DR. WORTHINGTON: And make sure of it, because we aren't aware that anything has fallen through the cracks in that period of time. But if that is the case we need to fix it.

Joe has a comment, I think.

MR. FITZGERALD: Yes, this is Joe Fitzgerald. This is one I think I just talked to you a little earlier on. I've been to DOE

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Germantown twice on those interview notes. So they are being reviewed, but the issue is, when things are identified there is a dialogue that is needed just to kind of move things forward. And I think that part of that is something that we're working on. That process is not probably where it needs to be right now.

DR. WORTHINGTON: And that's the one that I said I would follow up with you, Joe, there shouldn't too, because on be any documents there from six months ago that was just for an interview and that they've been and an action should cleared by one group, have been taken, and a document that suitable for release should have been made available or whatever. We need to follow up on that and make sure.

MR. FITZGERALD: Yes, and we've discussed this process issue. There is a process issue though, and I think it was probably unduly delayed, but that I think

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needs to be addressed. 1 2 MEMBER CLAWSON: But we are addressing those issues though? 3 4 MR. FITZGERALD: Oh, yes. These interview notes are pretty critical time wise 5 6 for the SEC review. So I've basically been 7 following up and working with the reviewers in Germantown to make sure they don't get hung 8 But the one thing I particularly think we 9 up. 10 are looking at, and I mentioned this to Pat, is that if there is an issue, feedback needs 11 to come back. 12 13 DR. WORTHINGTON: We need it immediately. 14 15 FITZGERALD: quickly MR. As 16 possible, so we can go ahead and resolve the issue not when it hits the stand. 17 CHAIR ZIEMER: Let. me insert 18 19 something here, and we have several other But there has been some concern in 20 the last month or so, general concern on the 21 part of the Board that the process seems to be 22

developing in terms of how this is Board input; is, without that that contractors seem to be in the loop and NIOSH and others, but the Board in general has felt like it's not been in the loop on exactly how this process is going to proceed. And we can discuss that some more.

But let's get the comments here.

Josie?

MEMBER PRESLEY: My comment was just back on the notes. I actually got a clearance and review Kathy's, from SC&A's in notes, from Mound, from Germantown, and they weren't available. She had gotten Brant's notes, but not her own notes, which were the that had actually reason we met to specifically look at them.

CHAIR ZIEMER: Okay, thank you.

DR. WORTHINGTON: So this seems to be an issue on the Mound notes, and we need to get to the bottom of that and expedite getting that done and getting it back out to people in

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the form that they can use, or if there is a need for classified discussion of key individuals to arrange that as well. But yes.

CHAIR ZIEMER: Okay, Jim.

issue MEMBER MELIUS: МУ is the major issue that we have, this so-called security plan or agreement we have waiting on for many, many months now. appears to be in a draft form that goes back and forth. The Board has not been made privy to that documentation at all, and I think it's by itself causing major problems.

I have major concerns that somehow processes or procedures are being institutionalized that will cause major delays in our ability, the Board's ability, to get its work done. And we have sort of no involvement in this.

And secondly, I think to have to deal with these issues on a case-by-case basis, or individual basis like these notes or that note, or this situation and so forth, is

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difficult. It's difficult for you coming here to do that, because we don't know what the expectation should be. What's a fair amount of time that certain things should take? Which notes, which documents and so forth are subject to review, and what are ways that that can be appropriately expedited and so forth. And we are just being left in the dark on this.

I think it's a major problem, and continues to be a major problem. And no one from either the NIOSH or the DOE end seems to be willing to discuss it with the Board. Why we have not been able to see a draft document of this security plan is beyond me, since we are going to have to live with it and deal with it. We are not - and so forth, and we are going to have to explain and try to figure out why major reviews and so forth are going to be taking months and months longer because of these new procedures.

And I'm very disturbed by it, and

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mostly disturbed by the secrecy that seems to be surrounding this particular document.

I don't think anybody has any problem with security reviews as necessary. But the fact that we've been - it seems to me like it's over a year we've been aware of this issue, and the security plans being discussed, and we have yet to see a document.

DR. WORTHINGTON: Maybe this is a good opportunity for me to give a little bit more information, a little bit more insight into the process and what we are doing. And I'll start with the endpoint first.

And the endpoint is that various groups, NIOSH and Labor and DOE, we've been working together to come up with a security plan. And what we wanted to bring to the Board was a consolidated draft plan for Board comment and feedback. And we are still not at that point.

I believe that we are very close to doing that. We have some comments that we are

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working on with NIOSH, and so at some point I believe in the near future we would be in a position to be able to do that.

Your question about what kinds of things ought to be reviewed, and how much time should they take, because some things did occur over the last year or so, we have been working on that to make sure that we had a security plan in place that would clearly lay out a number of things. It would lay out access to the sites, how you get badges, how you get cleared, the kinds of documents that generated and at what point in the generation of those documents would they be subject to reviews.

been working on that. we've It's been lessons learned from some of the events that have occurred. And we think we are fairly close to having a plan that would fact national in meet the security also would expedite requirements and process, and would be one that would minimize

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the possibility of reports that were not appropriate being generated, and then therefore cause a delay.

And I believe that what we put together typically would mean that when documents are subject to review that they would be 10 to 14-day reviews, and that we've been able to get them down now to days and not even weeks in terms of those things.

So I think that we are close, and certainly we were looking for the right time to come with a plan that would in fact address all of those things.

CHAIR ZIEMER: Thank you, Pat, and I think the intent there is certainly good.

One of the sort of concerns as I sense it or have sensed it over the several weeks is that the agencies have sought input the process from the Board's on appeared contractor, and it that it assumed that the contractor was speaking on behalf of the Board. And although

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endpoint is that a document would come to the Board for a final review, a concern that perhaps there could have been input earlier from participating Board members, so that not that we don't think the contractor's input is important. But the sort of assumption that the Board's contractor was speaking on behalf of the Board when we really had no input on that, I think, was a sort of concern. I think I've expressed, based on some email exchanges that I've seen on this issue, between Board members, and I've told the Board members, we can't do the business by email.

if have those kind So you of concerns, let's get them out in the open, and I'm trying to express, I think, what I've seen exchanges between Board members, concern that it. assumed that our contractor was was speaking on our behalf.

MEMBER MELIUS: Can I just clarify?

I mean, Paul, we were informed after the fact
on a meeting. John Morrow went to a meeting.

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So it's hard to express our concerns ahead of time when nobody informs us about what's going on.

CHAIR ZIEMER: That's what I'm saying.

MEMBER MELIUS: Number two, it was presented to us in that email as I recall that we would essentially get to see the final security plan. There was no talk about the Board, any involvement of the Board in reviewing that or having any input into that final security plan.

I would also point out that this process has been going on at least since the St. Louis meeting which I believe was in June, and it started I think even a little bit before that. It's over six months, and if the three agencies can't get it together to produce a draft security plan, I'm sort of dumfounded that it should take so long. And I think it is - and the fact that we don't have one continues to cause delays and problems.

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And every time we raise these issues about delays or problems, they say, well, it's being taken care of in this plan, and the plan, which we've never seen, the process seems to go on and on forever.

CHAIR ZIEMER: Larry.

MR. ELLIOTT: Well, I think there is an opportunity here to apologize for any perception that was acquired by asking SC&A to be involved in examining the draft plan. And let me make it clear: there are two plans. There is an agency plan at DOL that serves right now to speak to an audience at DOL as well as an audience on our side, as well as DOL's side.

There is a plan that complements the DOL plan that is a NIOSH plan that is overarching that includes the activities of NIOSH as well as all of the contractors involved, SC&A, ORAU, and whichever contractor is named to be successors to those two.

We ask - I ask SC&A to review and

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NIOSH

1 on the draft plan that 2 prepared, and was in negotiation with, DOL and

DOE on, so that SC&A could comment on their 3

ability to comply with such a plan. 4

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We were in negotiations with the other two agencies, and primarily DOE here, to come up with a plan, and it didn't seem fair to share the draft with our contractor and not share the draft with your contractor to get their input on the ability to comply; also the ability to provide comment about difficulties that the plan might present.

There is а these are decisional documents, and until the agencies which have a management prerogative here come forward with their final negotiated and agreed upon plans, that is when I think the Board has an opportunity to provide comment on these plans.

Until then as they are being negotiated, they are considered pre-decisional and the agencies are not going to allow us to

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put them out into the public forum because they are in the process of negotiation and change.

DR. WORTHINGTON: And we wanted to have a consolidated view of the agencies that we would present rather than drafts that may have diverging views.

MR. ELLIOTT: And the fact that it has taken as long as it has I think is notable in the context that we are negotiating; we are going back and forth on some things that are fairly critical.

We recognize at NIOSH that DOE has the responsibility and the authority to protect national security information. what they are asking of us is to review documents that are prepared based upon source documents that they have given to us or our contractors to make sure there is not sensitive information. That's all they are reviewing, for sensitive information. that's about all I can say at this juncture.

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I wish the plans were out there, too. We both have been pushing to get these things finalized, and it's taken us this amount of time to come to a series of agreements as to what these plans need to look like.

CHAIR ZIEMER: Thank you, Larry.

Brad, any additional comment?

MEMBER CLAWSON: Well, one thing I wanted to bring up, Pat, I do want to tell you how much I do appreciate Greg and Regina because they have been a great help.

DR. WORTHINGTON: Thank you.

MEMBER CLAWSON: But one of the things that I have seen that really comes down, and it's difficult, and I hope that we think about this, is like at Hanford, the for be able to see certain Sigmas us to things, it's not being portrayed down to the site or wherever we are going at. They didn't understand what we could really see and what There is a breakdown at the very we couldn't. bottom end of it that when this is set forth

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| of what we can actually review. Because at |
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| Hanford I had the opportunity to be able to |
| look at something from Pantex, and it'd |
| already be prearranged. But the people at |
| Hanford did not - they were scared to give me |
| it, and it was through Greg and Regina that |
| then I was able to get to it. So I just |
| wanted to bring forth that there is a |
| breakdown at the site of what can and can't - |
| and nothing was sent to them saying, when my Q |
| clearance comes up what Sigmas I actually |
| have. And this may be something that I hope |
| that we can look at, because I saw a great |
| nervousness until Regina called and took care |
| of it with them of them of being able to allow |
| us to see anything. Because they felt fine |
| with their Hanford stuff, but how could they |
| speak for somebody else's site. And this is |
| an issue that is going to be coming up again |
| and again, and I hope that maybe we can work |
| through it and so forth. Because they didn't |
| understand what I had and what I could see. |

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There was a breakdown there. And they were very reluctant, let's put it that way.

DR. WORTHINGTON: Greg, do you have a comment?

MR. LEWIS: Yes, I was just going to say, that speaks to, this Board is very unique in terms of what they do and how they interact with DOE sites, especially at a site like Hanford where we facilitated a workspace where you can review both Hanford documents and documents related to other sites. That's a unique situation, we've done that at Hanford and at Livermore, and they are not That's very different than how used to that. And like you said, they are they operate. very used to making decisions on what and how release make Hanford documents t.o and It's a little bit different with accessible. other facilities. We are working that. was a bit of a delay in the instance you're speaking of. But we do understand that is a unique situation at those two facilities and

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any others where we facilitate similar types of arrangements. We will make sure they are getting used to how to handle that, and what preapprovals and things like that we need.

DR. WORTHINGTON: And we think that is the role of our office that we need to make this clear. We are the champions, we are the people working with the sites to make it happen, to make sure they understand the clearances and what is required.

So we will certainly be more aggressive in making sure that people understand the clearance level, and that those things are made available to you.

MEMBER CLAWSON: Yes, because - like I say, this was a new situation and so forth like that. And I don't ever want to go into a site and have them scared that they are going to mess up, and this is kind of what I saw. Because I asked her about it, and they took me down, and they said, this is what your clearance shows. And it shows no Sigmas.

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| 1 | We've got this and so forth. But I wanted you |
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| 2 | to be aware of those issues, because we are |
| 3 | probably going to get into it. And like you |
| 4 | said, this is still grassroots, and I |
| 5 | appreciate everything that you have done, but |
| 6 | I don't ever want to - I don't like the |
| 7 | situation where a site is very, very scared to |
| 8 | even talk with us because they are afraid of |
| 9 | some of these issues. |
| 10 | And I think if we address this up |
| 11 | front and stuff we will be able - it will work |
| 12 | out. It did work out at Hanford, and that's |
| 13 | where you and Regina came in, and I appreciate |
| 14 | it very much. |
| 15 | CHAIR ZIEMER: Jim, do you have a |
| 16 | comment? |
| 17 | MEMBER MELIUS: Yes, I would like to |
| 18 | try to get a timeframe when this security plan |
| 19 | is going to be made available and be |
| 20 | completed. |
| 21 | CHAIR ZIEMER: Any reliable |
| 22 | predictors, Pat or Larry? |

| DR. WORTHINGTON: I know that Larry |
|--|
| and I, we are working on it, and we will |
| continue to expedite it. I don't want to make |
| a commitment that we can't deliver. But it |
| certainly is of highest priority and it's |
| being worked on as we speak, as we are here at |
| this meeting right now, so we hope to get back |
| to the Board soon. |
| CHAIR ZIEMER: Can you give us an |

CHAIR ZIEMER: Can you give us an indication, at what level in the agency - for example at DOE, is this something that goes all the way up to the secretarial level, or is it an assistant secretary? Who approves, and also at NIOSH?

DR. WORTHINGTON: Within the Department of Energy it is our intent that it will be approved in Glenn's organization, the HHS organization. Yes, that is the intent unless something changes.

MR. ELLIOTT: And at NIOSH it will be approved at the agency level, NIOSH's level, not CDC; not the department. But we

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| 1 | will have to have - there are elements in the |
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| 2 | plan that speak to how certain types of |
| 3 | information are handled with regard to Privacy |
| 4 | Act information which in DOE's parlance falls |
| 5 | under official use only. We have to have our |
| 6 | FOIA office and our Privacy Act office review |
| 7 | what we are inserting into the plan in that |
| 8 | regard, and so while they have helped us with |
| 9 | language they haven't seen this yet in its |
| 10 | full entire form. So they will have to look |
| 11 | at that. |
| 12 | We also have - OGC has to put eyes |
| 13 | on this plan as well, and they have not done |
| 14 | that. |
| 15 | CHAIR ZIEMER: Larry, while you are |
| 16 | at the mike, you may not be able to answer |
| 17 | this, but to your knowledge is there anything |
| 18 | in the Department of Labor's plan that would |
| 19 | have a significant impact on what this Board |

MR. ELLIOTT: I don't know that Labor is putting a plan together. Labor is

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does?

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mentioned in DOL's plan - or DOE's plan.

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DR. WORTHINGTON: I think earlier you may have said Department of Labor one time.

MR. ELLIOTT: I'm sorry. DOE has a plan they are working up that speaks to the activities and an audience at DOE, but it also speaks to us and to DOL.

NIOSH has a complementary plan that speaks to the NIOSH audience and its an overarching plan, which contractors, as would include the Board's contractor and the Board's activities. And there would also be that will procedures be OCAS two based procedures that have to be followed. So you will have essentially two plans to look at, one DOE plan, one NIOSH plan. Both will speak - cross-walk each other. And then there are two procedures behind the NIOSH plan.

CHAIR ZIEMER: So the intent would be that we would have the opportunity to see both of those plans? I thought one was sort

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| 1 | of independent as a Labor plan. But you are |
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| 2 | saying it's not a Labor plan; it goes up |
| 3 | through their system. |
| 4 | MR. ELLIOTT: It goes - DOE is |
| 5 | producing a plan, a security plan, that speaks |
| 6 | to how NIOSH and Labor will interact with DOE. |
| 7 | DR. WORTHINGTON: Will interact and |
| 8 | protect the information, yes. |
| 9 | MR. ELLIOTT: But I don't know that |
| 10 | DOL is planning to put forward any kind of a |
| 11 | plan. |
| 12 | CHAIR ZIEMER: Does that require |
| 13 | approval by Labor, is what I am sort of asking |
| 14 | too. |
| 15 | MR. ELLIOTT: Well, I'll let Pat |
| 16 | answer that question. |
| 17 | DR. WORTHINGTON: The plan as |
| 18 | written, Greg, I don't believe there is a |
| 19 | line. We certainly will be sharing, and have |
| 20 | been sharing with the Department of Labor. |
| 21 | But try to remember that the plan as it exists |
| 22 | now, is there a line for the Department of |

| 1 | Labor, I don't recall. I know we have been |
|----|--|
| 2 | interfacing with them. |
| 3 | MR. LEWIS: I agree with you. I am |
| 4 | not sure whether they are going to be |
| 5 | officially signing off, but they have been |
| 6 | providing input, and they are going to be |
| 7 | working with it. |
| 8 | And just to clarify, I believe DOL |
| 9 | is drafting a plan, but it is much more |
| LO | limited, and it deals only with their |
| 11 | interaction with DOE. I don't believe it |
| L2 | would involve NIOSH and/or the Board's |
| L3 | operations. But again that - |
| L4 | CHAIR ZIEMER: But I guess if that |
| L5 | is occurring, my original question still |
| L6 | applies. Do we know whether there is anything |
| L7 | in their plan which directly impacts - well, I |
| L8 | guess impacts on NIOSH or this board? |
| L9 | MR. LEWIS: Yes, as I said, I don't |
| 20 | believe there is again - |
| 21 | DR. WORTHINGTON: I don't think we |
| 22 | can speak for Department of Labor. But we |

certainly have been working with them on elements of the plan to make sure, where there might be inconsistencies that we can work through those.

CHAIR ZIEMER: Thank you.

Josie, another comment?

MEMBER BEACH: This may be a separate situation, but I want to know if the plan will cover a situation that I encountered last week. I was trying to get clearance to view Mound documents at Hanford, and through second and third people was told I needed approval from NIOSH to be able to view those documents and have approval - will this plan help that?

CHAIR ZIEMER: No. I don't know who is on the line, but you don't need approval from NIOSH to view the documents. But you do need to go through a NIOSH point of contact to facilitate your getting assistance from DOE to review those documents. If you approach DOE on your own they are going to of

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course say, what authority do you have? And NIOSH has got a point of contact identified for these sites that you are aware of that needs to facilitate getting you into to see what you want to see.

It's not that we are sitting as a gatekeeper and giving approval. We are trying to facilitate, trying to answer the questions for DOE that have been raised, like Brad brought up about what gives you the right to see certain types of information. That's what we want to be out in front of, and be pushing to DOE and to the sites. This person has that authority to see that information based upon these needs.

So I know Jess is coming up here, and I can't speak for DOL either. But I would say - maybe he is going to say this - DOL's part in the DOE plan is to subtitle E, not anything on B. It's for their ability to get toxic chemical exposure information from DOE.

DR. WORTHINGTON: At Department of

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| 1 | Energy, as a way of getting a handle on these |
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| 2 | document requests. We were receiving at one |
| 3 | point document requests from many individuals, |
| 4 | and we look for a way to have some single |
| 5 | point of accountability, and to be able to set |
| 6 | some priorities, and to kind of manage and |
| 7 | schedule and budget for these activities. |
| 8 | So we've asked NIOSH to designate |
| 9 | points of contact or specific leads for those. |
| 10 | Because in the past we were receiving in some |
| 11 | cases multiple requests, the same thing from |
| 12 | different individuals. We needed a little bit |
| 13 | more control over it, so that was the idea |
| 14 | that NIOSH would have a designated individual |
| 15 | for those sites, and those things would be |
| 16 | better coordinated, and hopefully we could |
| 17 | deliver better services that way. |
| 18 | CHAIR ZIEMER: Thank you. |
| 19 | Jeff, did you have a comment from |
| 20 | labor? |
| 21 | MEMBER KATZ: Only to say that I |
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employment verification with DOE. I don't 1 2 think there is anything that impacts NIOSH. As Larry said, there is the Part E component 3 for the chemicals. 4 Thank 5 CHAIR ZIEMER: you. Any further questions or comments for Pat, or in 6 7 general on this issue? WORTHINGTON: If not, I would 8 want to make just a few closing remarks. 9 10 The intent of what we call process improvements, they are intended, in fact, to 11 12 improve the process. There was never 13 intent to have secrecy or to eliminate Board others from participating 14 or or 15 providing comments. 16 We were looking collectively, the agencies involved, for the right juncture, so 17 it would be meaningful, rather than coming 18 19 back to you many times with many different drafts, to say, this is DOE's position, but 20

labor and NIOSH, they have a third or a fourth

So we were trying to consolidate,

position.

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and to come back to you.

So we hope to be able to get back into the Board soon with some additional information and insights.

But your questions are always welcome, and we want to hear what you are thinking, and hopefully answer those questions or look for ways to figure out how to do that.

CHAIR ZIEMER: And we appreciate the efforts to streamline this process.

Jim.

MEMBER MELIUS: This is not - well, maybe it is a question for you - but is there any reason members of the Board can't be involved in these activities, in these meetings, and to be informed about what is going on? I'm talking about specific individual members of the Board.

DR. WORTHINGTON: I believe that some specific things that we discussed about the process, there was a Board member present and participated in at least one of the

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1 meetings. So certainly some of them 2 meetings that require cleared individuals. But again, the agencies are trying 3 4 to come up with a process and then present that draft. I don't know if that answers your 5 question. 6 7 Greg, you had a comment? MR. LEWIS: Yes, I was going to say 8 I believe Mr. Presley and Mr. Clawson 9 10 were involved in some initial discussions, at least as far as the role of the Board and how 11 you operate and how we could help facilitate 12 13 that. DR. There 14 WORTHINGTON: were 15 couple of meetings at least. MEMBER CLAWSON: That mainly 16 was though, that was how we were going to handle 17 some Mound issues that we were doing. 18 19 far what Dr. Melius as talking about - about the procedure and stuff 20 - I haven't been involved with it. But I was 21 involved in D.C. with the issues we 22

discussing with Mound and how we would handle those.

DR. WORTHINGTON: We want to look for the - the agencies to look for the best opportunity, the most timely manner, to get back to the Board with something for their review.

CHAIR ZIEMER: I suppose part of the issue - and maybe we can think of how to deal with this - I know the agencies are reluctant to have something out in public before they have developed their policy, and our process in the Board is the things we do have to be made public. So I'm not sure where the balance is between that. If a Board member participates, does that force us to go into the public arena or not?

MR. ELLIOTT: Well, again, these are pre-decisional documents until the agencies decide that they have come to an agreement on what the document contents are going to be. And again, that is the point in time when the

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Board should become involved. It's not going 1 2 to be helpful at this point in the process of negotiating these agreements. 3 We are very close. I was hopeful 4 that we would have something for the Board to 5 chew on at this meeting, and that just didn't 6 7 It didn't happen because, as I said earlier, the Privacy Act component of this, of 8 our document and DOE's document just weren't 9 10 ready for prime time, and we didn't have the authority to speak about that from the FOIA 11 office or the Privacy Act office. 12 13 CHAIR ZIEMER: Well, certainly -DR. WORTHINGTON: And this is a high 14 priority. It really is for us, and we are 15 trying to get it done. 16 CHAIR ZIEMER: Well, certainly, the 17 intent of both agencies I believe is 18 19 streamline the process and minimize its impact on your activities and our activities. 20 in that sense we have a common goal. 21

I guess one of the concerns is, if

| 1 | there is something in the agreement that |
|----|---|
| 2 | somehow is not palatable to the Board is it |
| 3 | too late? Are these - that is more of a - |
| 4 | it's almost a rhetorical question. |
| 5 | MR. ELLIOTT: I don't presume to |
| 6 | know what's going to be unpalatable to the |
| 7 | Board in this. |
| 8 | CHAIR ZIEMER: I know that. |
| 9 | MR. ELLIOTT: I can only - |
| 10 | CHAIR ZIEMER: I think our goals are |
| 11 | the same. |
| 12 | MR. ELLIOTT: Our goals are the |
| 13 | same. |
| 14 | CHAIR ZIEMER: So we will be |
| 15 | optimistic about the outcome. |
| 16 | MR. ELLIOTT: We don't want to see |
| 17 | any obstruction. We don't want to see any |
| 18 | delays. But at the same time we have to |
| 19 | recognize that DOE has an authority and a |
| 20 | responsibility to protect information here. |
| 21 | And so we are trying to work across the |

agencies to make sure that we are not in

violation of any national security issues in this whole process, and all I can say is, certainly we want to know what the Board's thoughts are on these two plans once the agencies have come to finalization.

DR. WORTHINGTON: And we want to listen to the Board and receive their comments. And certainly that is a strong statement from us that we want to do that; we want to bring it to you for you to look at and speak freely on what your thoughts are on that document.

MR. ELLIOTT: Depending on what your issues and concerns are, there may be room to move; there may be ways to change. But I will be frank and honest about it: in some instances there may not be. Our hands may be tied. And I'll give you an example.

We worked with one thing that Mr.

Presley did work with us on in the early
goings of the negotiations was to come up with
a list of sites that would represent

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1 facilities or sites where DOE review for sensitive information was mandated. 2 That didn't survive the day in the end. 3 So we are away from that list now, 4 and we are at where right now all sites -5 DR. WORTHINGTON: All sites. 6 7 MR. ELLIOTT: are considered to subject to this review for sensitive 8 information. 9 10 We have also struck an agreement, and I think also an accord in this agreement, 11 that as we move forward and DOE reviews what 12 13 they review, if there are examples instances where they find that they don't need 14 15 to see that kind of a document, or that site 16 has already been taken care of and we don't need to see that kind of information again, 17 they will be quick to tell us and we can then 18 19 draw the boundaries in on this. And we have some experience, some 20 examples to show to that effect. 21

We have also come to a place where

in the process, as we are working with DOE, we identify a critical need on a certain review, and they have demonstrated an ability to turn those around. The three 83.14s that you have before you at this meeting are an example of that where in our planning to deliver an 83.14 SEC or an 83.13 SEC; we had not anticipated DOE's need to have seven to 14 days to examine the document. So that had to be plugged in.

And for these three that you talked about at this meeting, Mallinckrodt, Vitro and the Met Lab, those weren't reviewed by DOE until late last week, early last week in fact, and they turned them around within a day.

DR. WORTHINGTON: But it is a commitment of our organization that these things get the highest priority. And so we will assign people to get on them and review them when they are needed.

And there has been, I think, sort of a feeling that this process is one for delay. Our lessons learned from the people

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that have been involved in it, is that it is a process that expedites things, and that it prevents a report that is generated that is inappropriate that will be investigated and delayed for long periods of time.

ensure that that wouldn't happen, so things are moving out in a timely manner. It is a process to help; it's not a process for secrecy or delays or whatever it is. And I think we've been able to demonstrate as we work through the plan that that can happen, and that we want to continue to move down that path.

CHAIR ZIEMER: Thank you, and we appreciate the commitment to that, and I think as I said, I think our ultimate goal for all of this is the same in that regard. So we look forward to receiving the documents soon.

Another comment?

MEMBER MELIUS: I did have another comment, but I will hold it.

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CHAIR ZIEMER: On this topic? 1 MELIUS: Ι just fail 2 MEMBER understand why expediting becomes as we had in 3 the case of Hanford SEC now over 4 waiting for access to documents. 5 And an SEC 6 evaluation that is going to go on for another 7 couple of years because of the way NIOSH laid it out. 8 So I understand you are trying to 9 10 I don't have any problems with your intent. But unfortunately, these things just 11 at least in the implementation phase and 12 13 the uncertainty over what is going to be in the plan and how these procedures would work 14 are leading to long delays. 15 And the credibility of this program 16 little to begin with, with 17 is very the claimants, and this only makes it worse. 18 19 MEMBER ZIEMER: Thank you. 20 Greq? MR. LEWIS: Yes, we have not to my 21 knowledge limited access documents 22 to any

relevant to Hanford to those with the appropriate clearances, and they can review them in the appropriate setting.

We have in some instances limited our release of certain documents until there are assurances in place to make sure that people have appropriate plans to handle those documents. But on site in the proper location they can be reviewed by anyone.

DR. WORTHINGTON: I think we have done a number of things to expedite, I know that concerns you, but really to expedite the process by getting clearances for past workers to come forth so they can talk openly and freely about any of the processes they need, making sure that space is available to do those kinds of things.

And I guess at this point only time will show that we have been working on this, and it in fact is to improve the process, and it's not to hurt it and to make things available for these workers.

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| 1 | CHAIR ZIEMER: Thank you. |
|----|--|
| 2 | Other comments? On this or |
| 3 | anything related to the DOE report? |
| 4 | Again, thank you Pat. |
| 5 | DR. WORTHINGTON: Thank you for your |
| 6 | attention. |
| 7 | CHAIR ZIEMER: And your other staff |
| 8 | members who are with you today, Greg and Isaf, |
| 9 | we appreciate your efforts as well. |
| 10 | I think it would be appropriate for |
| 11 | us to go ahead with our lunch break. We are |
| 12 | going to extend it. It is now quarter of |
| 13 | 12:00, and if we go until 1:15, that will give |
| 14 | us an hour and a half for lunch rather than |
| 15 | the hour that we had yesterday. |
| 16 | So let's recess for lunch, and plan |
| 17 | to be back here at 1:15. |
| 18 | (Whereupon, the above-entitled |
| 19 | matter went off the record at 11:45 a.m. and |
| 20 | resumed at 1:18 p.m.) |
| 21 | CHAIR ZIEMER: We are ready to |
| 22 | reconvene the afternoon session. I will check |

1 to make sure that everyone that needs to be on 2 the phone line is on there, in particular Board member Dr. Gen Roessler. Gen, are you 3 with us? 4 MEMBER ROESSLER: I'm with you. 5 CHAIR ZIEMER: Thank you very much. 6 7 We are going to begin our afternoon session today with the science update, and 8 that will be Dr. Neton from NIOSH. So Jim, we 9 10 are pleased to have you back for the science update. 11 SCIENCE UPDATE 12 13 DR. NETON: Thank you, Dr. Ziemer. It's my pleasure to provide the 14 15 Board what has sort of become а standard agenda item on the Board's schedule, and that 16 is an update of the science issues, where 17 NIOSH stands on science issues; past issues 18 19 that were identified; our progress toward resolving those; as well as any -20 MR. KATZ: Sorry, one second please. 21

The folks on the phone, somebody has not

muted their line and it's interfering with the conversation. Can you mute your line? If you don't have a mute button, please use star six.

DR. NETON: Okay, I guess I can continue now.

So update on any progress on past issues, and also discuss any emerging issues that are coming out that we think the Board might be interested in hearing.

Unfortunately the Board meets so frequently it seems that I have trouble putting out any ground-breaking progress in between some of these meetings, but we do have some news to report this time.

One thing I would like to discuss is, we've had a number of science issues on the table, and made various progress - progress on various of them over time that I report to the Board. But this year - and I think Larry might have alluded to this in his earlier presentation, we have adopted some specific science goals for this fiscal year.

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These of course are listed here; they are not all of the goals, all the science issues that are out there. The ones that appear here are the ones we've identified as being particularly important to get moving in a rapid timeframe. I've listed them here, we can just go over them briefly.

One is the review of the new solid cancer instance data reported from the Radiation Effects Research Foundation. We are committed to doing that in this fiscal year. These are formally acknowledged and published by NIOSH as our science goals.

As you are aware the new data came out a few months back, and we are going to review those data against what's currently used in the NIOSH IREP model to see if there is any indication that the risk factors might change for certain cancers.

The second issue is in development of the chronic lymphocytic leukemia model. We have committed to that by the third quarter.

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That would be a fiscal year quarter, so sometime in the April to June timeframe we intend to have at least this model developed and out of our shop. This has to go through secretarial office review, so I can't predict how long it will take once it gets through our shop, but it will at least be released for external review by then; or actually internal departmental review by then.

I can report that we solicited the input from three subject matter experts; that's three research hematologists, for the lymphocytic leukemia model that was developed, and we have just received this week the third report. So we are in the process of digesting the reviews that were received, and we will be responding to the comments that we got.

This third one I know has been out there for quite some time and I'm a little bit sheepish to acknowledge that it is still not done. But this is to issue the formal NIOSH position paper on the ingestion and oral-nasal

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breathing issues. I reported several Board meetings ago that we completed the research on that. I know that these are holding up closure on outstanding issues on a number of reviews that SC&A has conducted, and we are committed to getting that done as fast as we can. It merely needs to be written up and formally documented.

fourth bullet The is a formal verification and validation of NIOSH I reported on this I think at calculations. the last meeting. NIOSH IREP was checked for internal consistency and validated by several methods when it was being produced. However, as I think you heard in the public comment session last night, the level of formal rigor behind that review, that is a formal document that we can throw out on the table and say, this exactly is a consolidation of everything we have done, doesn't exist. So we are in the process of compiling that document, and we are committed to having that done by the third

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quarter of this fiscal year. That would be somewhere between April and June of this year.

The fourth one is a new issue that I don't think I've discussed with the Board before, and that is the development of a dose reconstruction methodology for RECA cases. Before I move on to that issue in the next slide, I do want to point out one bullet that is not on this slide, and it's not a science issue, but I think it's important to mention.

If you recall last June, a little over a year ago, we put a report out that was requested by Congress about the presumptive list. asked to cancer NIOSH was make recommendation as what if to cancers should be added to the presumptive cancer list.

We issued that report to Congress, to the Appropriations Committee I think it was that asked for that report. And we indicated that we believe that basal cell carcinoma should be added to the presumptive cancer

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However we did also state that this was risk. an interim report that would be updated when the UNSCEAR, the United Nations Scientific Committee on Atomic Radiation, UNSCEAR, released their forthcoming report. That report has been issued a couple of months ago. We have looked at it. We see nothing in that report that would change our opinion of what we came to in that first report, so we are committed to following up with some type of communication back to Congress to let them know that our original recommendation stands.

Okay, with that aside, I would like to talk a little bit about this new RECA model that I have in my last bullet.

turns out that the amendment t.hat. issued in 2004 to was the EEOTCPA extended Part Ε coverage; that's Department of Labor's part of this program, it extended Part E coverage to uranium millers, miners and ore transporters who worked at facilities covered under Section 5 of RECA.

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This is totally separate from Part B now, so take your Part B hat off and think about Part E.

That coverage is essentially Part E substances. toxic exposure to Well, is radiation is considered a toxic substance by Department of Labor regulations, but they also know that there is a more quantitative way to the probability that that evaluate substance, i.e. radiation, was the cause of endangering health. So when DOL issued their regulation for Part E coverage, they required that NIOSH would perform a dose reconstruction for certain cancer claims that were filed under Part E.

So given that we have engaged in developing some models to determine how we could do dose reconstructions for these RECA cases. We have been engaged in this research for some time now. WE hope to have a draft model done sometime in January that we can start testing and moving some of these cases

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out. These cases have been on the books for awhile, but the good news I think is that we have been informed by Department of Labor that the number of cases expected under this Part E of RECA is small. Right now we're thinking it's somewhere in the vicinity of 50 cases. But one never knows in this program; numbers do have a tendency to change.

it for that's RECA Okay, reconstructions. I would like to just shift gears now and talk about something that is new in our program, and this has to do with the of claimant data sets for coworker use modelings, which will be coming out as 0075. The TIB has been drafted; it's been internally reviewed, and it's undergoing some final tweaking. So we expect it to be issued in the next probably week or so. But I would to bring this up for the Board's attention.

It is interesting that this is somewhat relevant to the discussion that Tim

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Taulbee had yesterday on Savannah River SEC evaluation report. Because he talked about using this type of analysis to reconstruct coworker models for Savannah River. This was not intentional. We did not intentionally collaborate these presentations. But it's actually sort of fortuitous that it came out this way.

Before I get you into this, I would like to actually acknowledge the contributions or the work of Tom LaBone and Janice Watkins of ORAU who conceived of and did the heavy lifting, the work on this project. I'm sure there are others at ORAU, but these are the two technical staff that did much of the work on this, and I'm grateful for their work.

A little bit about the background here. Personal monitoring data are not really available at all sites. There are a few sites out there that have collected large amounts of for instance bioassay samples, but they never computerize them. They are not available

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electronically for us to review, and frankly, there are issues with taking those hard copies and coding them and making them available to our health physicists, primarily because of issues raised by cost and timeliness. How long is it going to take us to go through several hundred boxes, extract the records, code them, do a validation, that sort of thing. So a solution was proposed some time ago that we would rely on the claimant data set.

I have to admit that the first time this issue was broached to me, I was very skeptical. Because I said, how can we prove that the claimant set actually was representative of the general population? that's what. bluow course have t.o be established before we could use these data.

In other words, is the claimant data set an unbiased random sample of the general -- of all the workers at the site. A couple of thoughts come to mind. One is, all

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of our claimants have cancer, so could it be actually a biased high sample, because since they all have cancer, presumably they might have had higher exposures to give them cancers.

I can't think of another argument to the contrary, but there might be another argument that could be postulated why it would represent a sample bias low.

So we wanted to entertain this, but we thought, well, how could we go about doing And this is where Janice Watkins and this? Tom LaBone came in. They proposed a concept where we could use the complete data set that we have for the Y12 workers. It turns out that Y12 has been studied extensively through epidemiologic studies, and in these **CEDR** database, that is the Comprehensive Epidemiological Data Resource database that is managed by the Department of Energy, there is a very large set of electronic data available. And in fact it essentially covers all workers

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at the Y12 plant between 1950 and 1988. That is all monitored workers.

It's a huge data set of the uranium urinanalysis in particular, and it contains for about half million data а bioassay measurements. So it's huge а very comprehensive set of data.

And the idea was, could we use this to somehow test this hypothesis? This is just histogram that shows the frequency distribution of the samples that were actually collected history over the of the Y12 database. As you can see it starts in 1950, pretty low; peaks around 1958 to 1960; drops off in `66, and remains fairly constant. you will note that the heyday of bioassay sample collection, there were upwards of almost 50,000 bioassay samples taken per year.

So this is sort of the universe of all samples. In statistical sampling, this is it. So this is the true value if you will of all samples that we are trying to model.

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So the concept was to test this database -- I did that already -- so decided to do a feasibility analysis to test the claimant data against this database. So we thought, well we can easily develop an annual coworker model using the complete dataset. We have done that already for Y12, and we could establish the 50^{th} and 95^{th} percentiles like we do for our coworker models using the database, which although it had a half a million records, it represented 7,357 workers as you see on the slide. That is still a lot of workers.

We decided to do a worker modeling effort and not an individual sampling effort, because really we are trying to reconstruct does to workers.

So we have a coworker model that we have already established using these 7,000 plus workers. So now we decided, let's develop a coworker model using only the claimant data. Well it turns out that we have

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731 claimants who submitted about 68,000 bioassay samples. So that is roughly 10 percent of the total population. And what would happen if we compared the results of the coworker model developed using the complete set and the claimant data set.

This is a graphic that depicts the agreement at the 50th percentile, and you can see that there is fairly reasonable agreement between these two datasets. The pink line represents the claimant set, and the blue line represents the use of the entire data set.

There are some discrepancies though. If you look early on at 1950 the agreement is not quite so good, and there is also a little bit of a divergence after 1985.

95th the When you go out t.o percentile which are at the extreme ends of the distribution, you expect there to be more fluctuation the of the at extremes distribution; and you in fact do see that. The agreement quite good, is not as

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nonetheless it is fairly consistent across the Board, again, with the exception of the early years around 1950, and the later years after `85.

But then the question remains, well, even though they look like they agree fairly well, how can we demonstrate that it really is a statistically valid comparison. Do they really represent the same population of workers?

And so that is what we set out to do with a technique called a bootstrap analysis. A bootstrap analysis is, you select random sample distributions from the total population. In other words we had 731 workers that we are using for the claimant data set. So let's take 731 samples without replacement, a large number of times; in this case 10,000 times, and develop a distribution of all of those samples.

And this in fact is what this says.

This is a depiction of what the bootstrap

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analysis really is. And we chose to pick the 95th percentile, because if we could get agreement at the 95th percentile, we are pretty sure the 50th percentile would agree, because again, the extremes of the distribution are where you see the largest fluctuations.

So the bootstrap analysis, calculate the 95th percentile of the bioassay results, of the dataset for each randomly draw K workers - in this case that is 731 workers, because that is what the claimant data set had - and without replacement. repeat step two N times. In this case we repeated this step 10,000 times. So we pulled 10,000 sets of 731 people's records, and then we calculated the confidence interval that was generated from those sampling efforts, that is what you see here, which is results for -- this is only for 1953, but I just show this as an example of how the analysis would work. The solid red line in the middle is the -- wait a second -- the

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solid red line is the true 95th percentile. That is of the entire population, the 95th percentile is represented by the red line. The median of the bootstrap analysis is just to the left of that, very close which you would expect, based on that kind of sampling. And then you see the lower 99 percent confidence interval and the upper 99 percent confidence interval identified.

Those are the confidence intervals, of those 10,000 runs, that is the range of the values that came out of that for the $95^{\rm th}$ percentile.

And then the dashed blue line represents the claimant -- $95^{\rm th}$ claimant database reconstructed $95^{\rm th}$ percentile.

So what we have here is the claimant database, the 95th percentile for the claimant database is well within the limits -- is well within the 99 percent confidence intervals for -- using this bootstrap analysis technique. Which gave us confidence, okay,

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| 1 | this is producing some fairly reasonable |
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| 2 | numbers, and this slide summarizes that. The |
| 3 | bootstrap sampling demonstrated that the |
| 4 | results were 36 of 39 years that were |
| 5 | reconstructed - this is for the 95 th percentile |
| 6 | - were within the 99 percent confidence |
| 7 | interval. Yes? |
| 8 | MR. KATZ: Jim, can you pause? Can |
| 9 | people on the telephone hear us? |
| 10 | MEMBER ROESSLER: I can year, yes. |
| 11 | MR. KATZ: Have you been losing your |
| 12 | connection? |
| 13 | MEMBER ROESSLER: No, I haven't lost |
| 14 | it. |
| 15 | MR. KATZ: Okay, thank you. |
| 16 | CHAIR ZIEMER: Okay, go ahead. |
| 17 | DR. NETON: Okay, thank you. |
| 18 | So for those three years that were |
| 19 | outside those 99 percent confidence limits, |
| 20 | that was 1950, `87 and `88, it appears that |
| 21 | the reason was that the claimant dataset |
| 22 | contained less than fewer than 10 percent |

of the total data. So when you have fewer numbers, fewer than 10 percent of the represented samples, then you expect some statistical anomalies to occur.

And in fact there were a few other years that contained less than 10 percent of the data that were fortuitously within the 99 percent confidence interval.

So what we have here is, we have identified that one of the limitations of this technique is, you have to be able to somehow convince yourself you've got at least 10 percent of the real dataset being monitored, which is not exactly trivial in some cases.

So the conclusions of this analysis are that the feasibility did demonstrate that with certain caveats: you have to have at least 10 percent by year of the true data set, at least for the Y12 plant, it would produce a representative coworker model.

I talked about you need to have sufficient sample size. But then we thought,

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| 1 | well, what if you didn't? What if you didn't |
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| 2 | have 10 percent, maybe you could go back and |
| 3 | randomly sample the existing hard copy records |
| 4 | that were there on a random basis, and pull |
| 5 | out and supplement the coworker data until you |
| 6 | got 10 percent, and then reconstruct the |
| 7 | model. |
| 8 | We leave that open as a possibility |
| 9 | for further thought. |
| 10 | And that's it. That is the result |
| 11 | of that analysis, and I would be happy to |
| 12 | answer any questions on the whole presentation |
| 13 | if there are any. |
| 14 | CHAIR ZIEMER: Thank you very much, |
| 15 | Jim. |
| 16 | Let's open the floor for comments |
| 17 | or questions. Dr. Melius. |
| 18 | MEMBER MELIUS: One of the issues |
| 19 | and I think I brought this up in a work group |
| 20 | setting what you are calling a coworker |
| 21 | analysis is really based on all the workers |
| 22 | within a given facility and so forth. It |

doesn't really take into account factors such as job task or job assignment or type of trade or whatever.

So an analysis based on -- even with a 95th percentile -- based on this general group of workers may not be appropriate for a subgroup that has much higher exposures.

And I think there are -- in our discussions we talked about the fact that you have information limitations. So you may not always have the information on the sampling by job task, or job assignment or something like that. Or you may not have adequate numbers to be able to do that.

But I think it's a significant problem, and it's something that you need to think about. Actually it should be that you have to demonstrate that whatever distribution you are putting forward is adequate to capture — to be representative of those groups of workers, or somehow deal with a subgroup in some other way.

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It's not a trivial issue, and it's 1 2 not an easy issue to deal with. DR. NETON: No, I understand. 3 You raise a good point, Dr. Melius. 4 We have talked about this before. And this analysis 5 6 of course presupposes that the workers with 7 the highest potential for exposures indeed monitored in the first place. 8 Well, that's 9 MEMBER MELIUS: my 10 second point. NETON: That's a precondition. 11 And I think we have been held to that standard 12 for a number of sites now. 13 And we have had extensive discussions about -- for instance 14 15 the Y12 site that issue came up. How do you 16 know that the workers who had the highest potential for exposure were indeed the ones 17 that were monitored? So we have gone down 18 19 that path to some degree. Then you have to say well, even the 20 highest exposed workers, is there a set of the 21

highest exposed workers who were unmonitored.

And I agree. We should have to be able to demonstrate that that is indeed true.

MEMBER MELIUS: Then the further complication, is there a particular task or something that were not monitored. The kind of issues we hear about repeatedly at these sites where people are put into very high exposure environments where monitoring deliberately not done or infeasible to do or whatever. And again there is a mathematical side to that. What is the potential exposure they may have experienced in those situations, and how would that compare to that, and does your coworker model, other model, incorporate that potential. Then that is even harder, because you are not measuring anything, so you are having to assume something. Then it is a repeated problem we have run across in these situations.

DR. NETON: I totally agree with you. We have to use extreme care when we determine to whom this coworker model is

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applied. We have to justify that it is appropriate to apply to that particular group.

No disagreement there.

CHAIR ZIEMER: Mark.

MEMBER GRIFFON: Jim, I apologize, I walked in a little late, so I might have missed this in the beginning of your presentation. But I guess my fundamental question is, why are you going forward with this? Is this specifically for Savannah River?

DR. NETON: There are more than just Savannah River sites in mind. I mean there are sites with large amounts of uncoded data, bioassay information in particular that hasn't been coded. It is not necessarily even just for the main radionuclides. For instance, this would be applicable at Savannah River. Tim Taulbee actually mentioned that in his presentation yesterday, it could potentially be useful to reconstruct a coworker model for Savannah River.

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| 1 | But you could also envision |
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| 2 | scenarios where the radionuclides of lesser |
| 3 | usage at the sites that have not been coded |
| 4 | could be brought into play with this type of |
| 5 | analysis. |
| 6 | MEMBER GRIFFON: Then I guess to get |
| 7 | back to the Savannah River question, there was |
| 8 | a lot of data that wasn't uncoded, but I also |
| 9 | see that it seems to me to do this, this is |
| 10 | what I think Tim referred to as TIB XX I |
| 11 | don't know if has a number yet. |
| 12 | DR. NETON: It's TIB-0075. |
| 13 | MEMBER GRIFFON: Oh, it's TIB-0075, |
| 14 | okay. |
| 15 | DR. NETON: It's not been issued. |
| 16 | It's imminent though. |
| 17 | MEMBER GRIFFON: Anyway, you went |
| 18 | through quite a bit of work it seems to me to |
| 19 | enter claimant data here to compare against |
| 20 | the electronic Y12 data. |
| 21 | DR. NETON: No, it was actually |
| 22 | while these are all claimant data, and |

| 1 | claimant data are coded as they come in. |
|----|--|
| 2 | MEMEBR GRIFFON: As they come in? |
| 3 | Okay, so it wasn't extra work for you. |
| 4 | DR. NETON: No, we didn't go and |
| 5 | code 68,000 pieces of claimant data or |
| 6 | 6,800. |
| 7 | MEMBER GRIFFON: Well, I guess the |
| 8 | question is - and I can appreciate that you |
| 9 | don't want to code all the Savannah River data |
| 10 | - but is it a question of, NIOSH doesn't want |
| 11 | to go through and code all that urinanalysis |
| 12 | data for Savannah River? Or that Savannah |
| 13 | River is not making it available to NIOSH? |
| 14 | DR. NETON: No, we have it available |
| 15 | to NIOSH now. It's a matter of resources, |
| 16 | time constraints. The question is, do you |
| 17 | really need to code, say, 600,000 bioassay |
| 18 | samples, or can you demonstrate that if you |
| 19 | have a representative statistical sampling of |
| 20 | those bioassay records and can construct a |
| 21 | coworker model. |

MEMBER GRIFFON: No, I understand.

| 1 | I wanted to make sure that it wasn't an access |
|----|--|
| 2 | issue to the data. |
| 3 | DR. NETON: No, the data are |
| 4 | available. Tim Taulbee assures me that he has |
| 5 | found the records, and we have access to them. |
| 6 | CHAIR ZIEMER: Any comments? Larry |
| 7 | Elliott? |
| 8 | MR. ELLIOTT: Well, I think also in |
| 9 | response to Mark's last question, it also has |
| 10 | to be considered that if we were to go look |
| 11 | for an individual claimant's data in those |
| 12 | boxes, how much time, how many pages would we |
| 13 | have to go through just for each claimant? |
| 14 | So if we have a model that can be |
| 15 | used to address that with perusing through |
| 16 | piles of paper, that was also in this |
| 17 | factored into this consideration. |
| 18 | MEMBER GRIFFON: And does this I |
| 19 | guess I assume we've got to review this TIB |
| 20 | obviously, and it might even be part of the |
| 21 | Savannah River review as well as procedures or |
| 22 | something, I don't know. But I guess I would |

| 1 | assume somewhere in this there is a |
|----|--|
| 2 | justification of why I think this follows |
| 3 | up a little on what Jim was talking about, why |
| 4 | this Y12 analysis proves to me that this is |
| 5 | useful with the Savannah River claim |
| 6 | population. |
| 7 | DR. NETON: I would answer that with |
| 8 | a question: why not? |
| 9 | MEMBER GRIFFON: Well, why not might |
| 10 | be I don't know |
| 11 | DR. NETON: That's why I'm saying, |
| 12 | we thought about this part. |
| 13 | MEMBER GRIFFON: I mean there might |
| 14 | be different percentages of construction |
| 15 | workers applying for claims down there, |
| 16 | because building trades are represented and |
| 17 | the construction workers are not. |
| 18 | DR. NETON: Well, that gets to Dr. |
| 19 | Melius' original question, we have to be |
| 20 | careful to what subset of workers we apply |
| 21 | this coworker model to, I'll grant you that. |
| 22 | Because that is to me a more relevant issue. |

But I think the model itself it has merit, we have done a feasibility. We have demonstrated at least at Y12, that they are not -- this is not a biased population of the overall population at the site. That is all we are saying. Of the overall monitored workers, if you had more than 10 percent of the data in your hand, the claimant population in and of itself is not a biased sample of the overall population.

MEMBER GRIFFON: But all I'm saying is, in this case -- I mean I haven't seen it, so in this case it seems to me that you are comparing a claimant population that might have different sort of demographics than at Savannah River. And this at Y12 it seems that all worker data bounded claimant population data, but is that going to be the case necessarily at other sites.

DR. NETON: I understand what you are saying.

CHAIR ZIEMER: Mark, you are just

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asking if what shows up in the Y12 case, would that show up at a site that had different kinds of demographics and different kinds of jobs and so on.

DR. NETON: Right, because I think if we go around the country, the people more likely to file at different sites might look a little different.

CHAIR ZIEMER: Well, let's ask this follow up question, Dr. Neton, beyond your conclusions, which are on this dataset, what do you see as the next steps in terms of this particular type of analysis? Where do you go from here? Is it the intent to develop a more generalized model that could be used sort of system wide?

DR. NETON: Yes, that was the intent of this entire feasibility analysis was that - - to not go off and just apply this without some sort of statistical analysis. The Y12 database was a convenient set, and I am totally understanding of what Mark was saying

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1 about the transportability of this to other 2 sites. But we thought about this long and hard, and I'm still having trouble figuring 3 out why these demographics would shift from 4 site to site. 5 We are open to any suggestions, and 6 of course this TIB will be I'm sure reviewed 7 by the Board and others. It's out there. 8 haven't used it yet, and we are still thinking 9 10 about it. I mean this TIB does not say go apply this everywhere. 11 It's really almost inappropriate to 12 call this a TIB. This probably should have 13 been a report. 14 CHAIR ZIEMER: Is there another site 15 16 that you could look at to sort of get beyond the one data point to see if --17 DR. NETON: Yes, I suspect 18 19 probably are, but I know this program well enough to know that N equal two, 20 necessarily prove a point either. 21

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CHAIR ZIEMER: I know.

DR. NETON: But it's good for thought. We put it out there. We are not using it. Anybody that can come up with a justifiable reason why it wouldn't transportable, I would love to hear it. mean that's one reason to throw it out there. are not necessarily asking for formal comment from the Board, but this is sort of a summary of where we are.

MEMBER MELIUS: What about -- one of the other factors that comes to my mind would be the sort of monitoring practices at sites, which I've noticed seem to vary over time, and it was resource dependent, and method dependent, and Ι suspect it was dependent on the skills and interest of the health physics team at the site and the amount of resources available to them.

I thought that when we were dealing with the TIB-0052, the construction thing, I seem to recall there were certainly differences there in terms of how they

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approached it. That may be a special population and not appropriate.

The other question I would have is more related to the act. I think you need to -- in my mind, just thinking about this briefly, I think you would need a strong justification to use a method that is based on estimating a person's exposure when there are individual dose records available.

DR. NETON: We would fully use their are available, records when they if the claimants were going to ask for their records. But if we don't get them, we have to have some way of reconstructing them. We are not talking about substituting for their records. just saying, if we don't have your record developed we have а coworker distribution based on the claimant population.

MEMBER GRIFFON: But I think if I heard Larry right, you might be -- you might be, because they'd be so difficult to find, or pull out all those records.

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| 1 | DR. NETON: We always ask the |
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| 2 | Department of Energy for the records, a |
| 3 | person's monitoring records, when we get their |
| 4 | case. And if we don't get them we don't get |
| 5 | them; I mean there is not much more we can do. |
| 6 | MEMBER MELIUS: Yes, but if the |
| 7 | Department of Energy is not bothering to get |
| 8 | them then they are responsible for getting |
| 9 | them, and there is something wrong - |
| 10 | DR. NETON: I understand what you |
| 11 | are saying. This has to do with the uncoded |
| 12 | records - |
| 13 | MEMBER GRIFFON: I don't want to get |
| 14 | into that. |
| 15 | DR. NETON: Again, this is food for |
| 16 | thought. I just throw it out there, and we |
| 17 | can digest it and maybe talk about it at some |
| 18 | other working group meeting or a future Board |
| 19 | meeting. |
| 20 | CHAIR ZIEMER: Any other questions |
| 21 | or comments? Very good, thank you. |
| 22 | Now we are scheduled to have an SEC |

petition update, and this is passed from LaVon 1 2 to Stu to Larry. Larry Elliott is going to give us that update. 3 Welcome back, Larry, to the mike. 4 SEC PETITION UPDATE 5 MR. ELLIOTT: Thank you. I don't 6 7 know if LaVon is on the phone or not, but if you are, LaVon, I hope I do you justice here 8 this afternoon 9 and present your material 10 effectively. this The purpose of special 11 exposure cohort status update is in response 12 13 to the Board's request to present the number of qualified petitions that under 14 are evaluation, and the sites that being 15 are evaluated through the 83.14 process. 16 This assists the Board 17 in understanding that bit of work, and preparing 18 19 and planning for it in their future meetings. The petitions received to date are 20 There are currently 17 petitions in the 135. 21

qualification process. There are 67 petitions

that have qualified for evaluation; nine of 1 2 are still in the evaluation, those progressing through NIOSH evaluation, and 58 3 have been completed. 4 There have been 51 petitions that 5 did not qualify for evaluation. 6 So we'll go through the ones that 7 are still active. 8

The petition, the Special Exposure Cohort petition evaluation reports that are currently with the Advisory Board for recommendation include Chapman Valve. The history of this petition is that the evaluation report was approved and sent to the Advisory Board and the petitioners on August

NIOSH presented its evaluation report at the September, 2006 Advisory Board Meeting. The Advisory Board established a work group to review and evaluate the report at its September, 2006 meeting.

The Work Group then presented its

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31st, 2006.

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findings at the May, 2007 Advisory Meeting, and a decision was made to postpone the recommendation until the 2007 July, Advisory Board Meeting, allowing the petitioners time to review SC&A's report on the NIOSH evaluation report.

The Advisory Board voted six to six on a motion to deny adding the class to the special exposure cohort at its July 2007 meeting, and following this vote the Advisory Board determined they would like to receive a response from the Department of Labor and the Department of Energy concerning potential covered work at the Dean Street facility.

Then prior to the October 2007 Advisory Board Meeting, the Department Labor provided a response to the Advisory Board's questions about the Dean Street facility. The DOE provided an update during the November 2007 advisory Board conference call and at that time they indicated they had completed their investigation the not at

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Department of Energy.

DOE then presented their findings at the January 2008 Advisory Board Meeting, and those findings were that the Dean Street facility should be included as a covered facility, but there was no indication of any additional radiological activities because of the addition.

NIOSH indicated at the January 2008 Advisory Board Meeting that they would revise the Chapman Valve evaluation report based upon the DOE findings. But also we indicated that there would be no changes in our feasibility determination based upon those findings.

NIOSH issued a revised evaluation report on February 5^{th} , 2008.

At the February 2008 Advisory Board conference call the Advisory Board asked SC&A to do a focused review of the new information provided by the Department of Energy, and asked that the information be made available prior to the April Advisory Board Meeting.

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1 SC&A produced a report to the Work Group on March 12th, 2008. 2 Next, NIOSH presented the revision 3 to the evaluation report at the April Board 4 This was also in 2008. 5 meeting. The Advisory Board decided to reconvene the 6 7 working group to discuss a path forward. Work Group met on May 8 2008, and asked NIOSH to send a letter to the 9 10 Department of Energy inquiring about extent of their evaluation. 11 addition NIOSH In agreed 12 to 13 continue looking for the pedigree of the enriched uranium analysis. 14 The Advisory Board voted again on a 15 16 motion to deny adding a class to the SEC at the June 2008 Advisory Board Meeting. 17 The vote ended in a six to six tie. 18 19 The Advisory Board then asked NIOSH to contact the Department of Defense about any 20 radiation related contracts for Chapman Valve 21 to explain the enriched sample. 22

NIOSH has made that contact with the Department of Defense, and DOD responded with no confirmation that Chapman Valve did or did not do work for the Department of Defense.

And we have some additional status to report on that at the conclusion of this presentation if you would like to have it.

Currently the status of the petition and the evaluation report are with the Advisory Board for recommendation.

Blockson Chemical: the evaluation report was initially approved and sent to the Advisory Board and the petitioners on September 5th, 2006. NIOSH presented its at evaluation report the December 2006 Advisory Board Meeting.

then withdrew the evaluation We report after it was determined that it did not address all covered exposures. Αt its 2006 meeting the Advisory December established a work group to review evaluation report for Blockson.

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NIOSH issued a revised evaluation 1 report on July 3rd, 2007. 2 NIOSH presented the revised 3 evaluation report for the Blockson Chemical at 4 the July 2007 Advisory Board Meeting. 5 The Work Group met in Cincinnati on 6 August 28th, 2007. A public meeting was held 7 on September 12th, 2007, to explain the changes 8 that NIOSH had made to the dose reconstruction 9 10 technical approach. A Work Group conference call was 11 held on November 2nd, 2007. At the January, 12 13 2008 Advisory Board Meeting, Dr. Melius indicated he wanted to review the pedigree of 14 the bioassay data, and he wanted to discuss 15 16 the radon model with Mark Griffin. There was no change in the status 17 of the petition, and the evaluation report, at 18 19 the April Advisory Board Meeting. The Work Group planned to meet to discuss a path 20 forward. 21

The Work Group met on June 5th,

2008. The Work Group met again on June 24th and 25th, 2008, to discuss a resolution of the radon issue and any outstanding action items.

The Advisory Board deliberated over the SEC petition at the June 2008 Advisory Board Meeting. The Advisory Board determined that they wanted to see the SC&A radon model in a white paper or a report to moving forward with voting on the SEC.

SC&A issued a draft report on the evaluation of radon levels in buildings 40 on August 12, 2008. The Work Group met again on October $15^{\rm th}$, 2008, to discuss the resolution of issues.

A technical call with NIOSH and SC&A was conducted on December $3^{\rm rd}$, 2008, and a Work Group conference call was conducted on December $12^{\rm th}$, 2008.

Status of this petition and evaluation report are with the Advisory Board for consideration, an update from the working group was given at this meeting.

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| 1 | The Feed Materials Production |
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| 2 | Center, the evaluation report was approved and |
| 3 | sent to the Advisory Board and the petitioners |
| 4 | on November 3 rd , 2006. NIOSH presented its |
| 5 | evaluation report at the February 2007 |
| 6 | Advisory Board Meeting. |
| 7 | At its February 2007 meeting the |
| 8 | Advisory Board established a work group to |
| 9 | review the evaluation report. |
| 10 | In May of 2007 SC&A provided a |
| 11 | draft review of the evaluation report to the |
| 12 | Work Group, petitioners, Advisory Board and |
| 13 | NIOSH. |
| 14 | The Work Group met in Cincinnati on |
| 15 | August 8 th , 2007; again, November 13 th , 2007; |
| 16 | March 26 th , 2008; September 15 th , 2008; and |
| 17 | October 28, 2008. The status of this petition |
| 18 | for Fernald is involved in research and |
| 19 | discussion among the Work Group, SC&A and |
| 20 | NIOSH. |
| 21 | Bethlehem Steel, the evaluation |
| 22 | report was approved and sent to the Advisory |

Board and petitioners on February 27th, 2007.

NIOSH presented the evaluation report at the May, 2007 Advisory Board Meeting.

At that time the Advisory Board determined that it needed further information before making a recommendation on the SEC

The Board tabled its discussion of the Bethlehem Steel SEC evaluation report until the Work Group could look at - which is looking at the use of surrogate data reported back to the Board.

The status of the Bethlehem Steel is that the evaluation report is with the Advisory Board for recommendation.

Hanford, Part 2, which covers all employees during the time period 1947 to 1990. The evaluation report was approved and sent to the Advisory Board, and the petitioners, on September 11th, 2007. NIOSH presented its evaluation report at the October Advisory Board Meeting, and the Advisory Board sent the

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petition.

report to their contractor and the Hanford Work Group for review.

The Advisory Board's contractor issued a white paper questioning whether the additional buildings should be included in the proposed class definition.

In March, 2008 NIOSH issued a revised evaluation report with a modified class definition.

NIOSH presented the revised class definition at the April, 2008 Advisory Board Meeting, and the Advisory Board concurred with NIOSH's recommendation to add a class status. The research and discussion on the petition continues among the Work Group, SC&A and NIOSH.

Nevada test site for the time period 1963 to 1992, this evaluation report was approved and sent to the Advisory Board and petitioners in September of 2007. NIOSH presented the evaluation report at the January 2008Board meeting. The Advisory Board sent

the report to SC&A and the Work Group for review.

The Work Group met on October 29th, 2008 and the status is for the Nevada test site, 1963 to 1992, that the research and discussion on the petition continues among the Work Group, SC&A and NIOSH.

The Mound plant, time period 1949 to present, the evaluation report was approved and sent to the Board and the petitioners in December 2007. The evaluation report was presented at the January 2008 Advisory Board Meeting. The Advisory Board concurred with NIOSH to add a class for the early years, but sent the report to SC&A for review, and established a Mound working group.

The Work Group met on April 1st, 2008, July 14th, 2008 and October 27th, 2008. The status of the Mound petition is that research and discussion continues among the Work Group, SC&A and NIOSH.

Texas City Chemicals for the time

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period 1952 to 1956. The evaluation report was approved and sent to the Board and petitioners on January 18th, 2008. NIOSH presented the evaluation report at the April 2008 Advisory Board Meeting. The Board gave the petition and evaluation report to the Surrogate Data Work Group for review.

SC&A completed a focused review of the Texas City Chemical, Inc. evaluation report in July, 2008. The status for Texas City Chemicals is that the petition and evaluation report are with the Advisory Board for recommendation.

Area four, Santa Susana, field laboratory, time frame, 1955 to 1958. The evaluation report was approved and sent to the Advisory Board and the petitioners on February 15th, 2008. NIOSH presented its evaluation report at the April 2008Board meeting, and the Board indicated they would not take action on this petition until SC&A had completed its review of the site profile.

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SC&A issued their draft review of the Santa Susana site profile on August 5th, 2008. The Work Group had their first meeting on August 26th, 2008. And the status for area four is that the petition and evaluation report are with the Advisory Board for recommendation.

Dow Chemical, 1961 time period through 2006. The addendum two of the evaluation report was approved and sent to the Advisory Board and petitioners on June 3rd, 2008. NIOSH presented addendum two at the June 2008 Advisory Board Meeting. To remind you what addendum two covers, it covers the residual period at Dow.

The Advisory Board asked the Procedures Work Group to review the recently approved dose reconstruction procedure residual contamination. That is TIB, or Technical Information Bulletin 70, assigned the petition evaluation to the Work Group on SEC issues.

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In September of 2008 SC&A completed
a focused review of addendum two to the Dow
Chemical SEC evaluation report.

November 17th, 2008, the Work Group

November 17^{cm}, 2008, the Work Group for SEC issues met and discussed the SC&A report. The general conclusion from the Work Group was that NIOSH's dose model was bounding, but NIOSH needed to verify a couple of numbers for the Work Group.

Status: the evaluation report addendum is with the Work Group for recommendation.

Pantex, time period 1951 through 1991: the evaluation report was approved and sent to the Advisory Board and petitioners on August 8th, 2008. The evaluation report was presented at the Board's September, 2008 meeting, and the status: the petition and evaluation report are with the Advisory Board for recommendation.

General Steel Industries: the evaluation report for this petition was

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approved and sent to the Advisory Board and petitioners on October 3rd, 2008. The petitioner requested that NIOSH delay presentation of this evaluation report until the February, 2009 Advisory Board Meeting which we agreed with.

Linde Ceramics, this covers the residual period at Linde. The evaluation report was approved and sent to the Board and petitioners on November 6th, 2008, and the petitioner requested that NIOSH delay presentation of the evaluation report until the May 2009 Advisory Board Meeting when the petitioner could be present, and we certainly agreed with that.

Savannah River site: the evaluation report was approved and sent to the Board and petitioners on November 18th, 2008. NIOSH presented the evaluation report at this Advisory Board Meeting.

Mallinckrodt, time frame 1958. We presented this evaluation report at this

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meeting, and you have taken your action on it. 1 2 Vitro Manufacturing, the evaluation report was presented at this meeting, and you 3 have taken your action. 4 Metallurgical Laboratory, 5 the evaluation report was provided and presented 6 7 at this meeting, and you have taken your action. 8 SEC petitions currently in 9 10 evaluation process are presented in this side, and you can see that in August we received a 11 petition regarding the Westinghouse Atomic 12 13 Power Development Plant, which covers employees in L and K buildings from the time 14 period January 1, 1942 to December 31st, 1944. 15 16 We expect to be able to deliver an evaluation Westinghouse 17 on Atomic Power Development and present that evaluation at the 18 19 February Advisory Board Meeting in Albuquerque. 20 31st, October of 2007, 21 On received a petition from the Massachusetts 22

Institute of Technology which addressed all locations and all employees, January 1, 1942 to December 31st, 1963. This was - as footnoted, this petition was initiated by a NIOSH finding that we couldn't reconstruct the dose. So it's an 83.14.

In November of 2008 a site visit was conducted - I'm sorry, a site visit was conducted in August of 2008, and NIOSH will present its evaluation report on the Massachusetts Institute of Technology at the February Board meeting.

On April 3rd, 2008 we received a petition regarding the LANL. All service support employees who worked in operational technical areas that had а history of radioactive material use at Los Alamos from January 1, 1976 through December 31st, 2005. Expected completion date is sometime and we have had some difficulties January, with data capture; we have been delayed a little bit there. So we anticipate that we

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will present this evaluation report for LANL support service employees at the February Board meeting.

The Brookhaven National Lab petition came to NIOSH on May 9th, 2008, and we've been working hard on that petition with the support of DOE. All of the Brookhaven records have not been incorporated into any electronic form, and so just finding those records has been a chore.

Our expected completion date is April, 2009, and we hope to present that at the May, 2009Board meeting.

Tyson Valley Powder Farm, which a petition covers all employees who worked in all areas of Tyson Valley Powder in St. Louis, Missouri, during the time period January 1, 1942 through December 31st, 1949. This petition came to NIOSH in June 13th, 2008. We hope to complete this one sometime this month, the rest of this month, and we would present it at your February Board meeting.

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United Nuclear Corp, a petition that covers all site employees who worked in any area of this facility from January 1, 1958, through December 31st, 1969. And in a separate period, January 1, 1970, through July 31st, 2006, which is the residual contamination period. And we expect to provide a completed evaluation report in March of 2009.

Standard Oil petition covers employees who worked in any area Standard Oil development site in Linden, New 1st, from August 1942, through Jersey, December, 31st, 1963. We received this petition September 18th, 2008, and we expect to complete it and present at the May meeting. It will be done in March, and we hope to present it in May.

Just a note: the completion of our evaluation reports were mandated by law to try to achieve completion of these within 180 days, and we have noted that we have missed that mark in several instances. The Savannah

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River site one took a little longer. General Steel Industries, Linde Ceramic Plant, Los Alamos National Lab and Brookhaven National Lab.

We have notified in each instance the Board and the petitioner as to what the delays are and why it's taking us the time it's taking us, and providing a timeline on when we expect to complete the report.

The reasons that we have exceeded 180 days are varied. The Savannah River site petition covered a large site, broad time period, and required a significant data review. And we are still receiving data as you heard from Dr. Taulbee yesterday.

NIOSH was delayed on a number of evaluation reports because we were waiting for DOE to establish the protocol for data captures at Savannah River site, Los Alamos and Brookhaven.

The Linde Ceramics Plant evaluation report was delayed as a result of the

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petitioner changing the basis for the petition 1 2 providing additional supporting and documentation, and that extended the 3 qualification period. 4 The GSI evaluation report exceeded 5 6 180 days by a week as a result of the time 7 required to resolve last minute technical comments from an interview review. 8 So our commitment is that once it's 9 10 apparent to us that the evaluation report timeframe will exceed 180 days we inform the 11 the Advisory 12 petitioners, Board and 13 congressional liaisons. We let them know that you all know that report will not be completed 14 15 within 180 days, the reasons for that and our 16 expected completion date for the report. That concludes the slides, and I'm 17 happy to take any questions. 18 19 CHAIR ZIEMER: Okay, thank you, Let's see if there are any questions. 20 thank LaVon also for the work he 21

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probably did in helping get

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this

together for us. It's always helpful to remind us where we are on each of these SEC petitions.

MR. ELLIOTT: Could I provide you a bit more background on what we have done with the Department of Defense regarding Chapman Valve and make sure that is on the record.

the last Board meeting when Chapman Valve was discussed, NIOSH was asked send an inquiry to the Department Defense, asking them about any work that they may have contracted with Chapman Valve; and secondly, to conduct an additional search to determine if the shipping manifest from the work done by Bechtel contained D&D reference to the enriched uranium sample that was found.

On October 6th, of 2008, the Office of the Secretary of the Department of Health and Human Services sent an email to the White House liaison section, within the office of the Secretary of Defense. It contained an

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explanation of the issue and this request.

Between 1942 and 1987 did the Department of Defense, most likely either the U.S. Air Force or Navy, award any contracts to the Chapman Valve manufacturing company, 1942 to 1958, or the Crane Company, 1959 to 1987, which is located in Indian Orchard, Massachusetts, for any work that would or could have involved the use of presence of enriched uranium.

October 17th, 2008, On the of Health and Human Services Department received a reply from the Department Defense that provided a link to the computer that contained database а listing individual contracts issued by the Department of Defense, and these listings were extracted from a form the Department of Defense uses called a DD-350, a 350 form. The computer database only contained records going back to 1966.

NIOSH reviewed the database and we

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did not find any records of contracts with the Crane Company in Indian Orchard, which is the successor to Chapman Valve.

The nature of these listings, however, are generic, and were not found to be useful for determining the existence of work with radioactive materials. That was not mentioned in the information provided in this database.

For example one contract specified work with nuclear reactors in 1978, but included no mention of the type of work, so it didn't get into detail like we would have hoped.

To view the original contracts would require a time consuming and costly manual search through boxes of records at the National Archives, and we would need the specific Department of Defense agency to help sponsor that search. We have not taken that step to date.

Regarding the Bechtel inquiry piece

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- this goes to the remediation work that was done afterward - NIOSH sent a written request Bechtel asking for worker monitoring to during their 1994-95 records remediation project. In follow up phone calls were made to Bechtel's legal department on September 25^{th} , 2006, October 4^{th} , 2006 and January 19^{th} , 2007. A second letter was sent to Bechtel on 25th, 2007, and NIOSH received April response or acknowledgment to communication efforts that I have mentioned here.

NIOSH located and placed on the Othe certification docket drive for the remedial action performed at the Chapman Valve site in Indian Orchard, Maine. This document written by Bechtel for DOE was at the conclusion of the rededication effort which took place from July to September, 1995. contains a detailed 200 page survey of the site conducted just prior to the remediation Outside references to the enriched work.

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| 1 | sample found in the prior ORAU survey of 1991. |
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| 2 | There is no mention, outside of the |
| 3 | references of the ORAU survey of 1991, there |
| 4 | is no mention of enriched uranium. |
| 5 | There were in fact 11 samples, |
| 6 | isotopically analyzed for uranium, and all |
| 7 | samples appear to be consistent with natural |
| 8 | uranium. |
| 9 | We also identified an additional |
| 10 | lead for Bechtel data in records retrieved |
| 11 | from the Kansas City records center. However |
| 12 | no relevant data has been found. So that has, |
| 13 | I hope, for you a more detailed summary of our |
| 14 | actions that we have taken regarding your |
| 15 | requests to pursue these final threads on |
| 16 | Chapman Valve. |
| 17 | CHAIR ZIEMER: Okay, thank you for |
| 18 | that additional update. |
| 19 | Questions on either the report |
| 20 | itself or the update. Wanda Munn. |
| 21 | MEMBER MUNN: Just as a matter of |
| | II |

interest, whenever we see electronic records

| 1 | stop at a certain point, we know when they |
|----|--|
| 2 | probably started digitizing that information, |
| 3 | is there any way of determining whether there |
| 4 | is any ongoing effort with, for example, the |
| 5 | Department of Defense in this case to go into |
| 6 | the old handwritten records earlier than is |
| 7 | currently - |
| 8 | MR. ELLIOTT: We have not exercised |
| 9 | any effort to do so. And as I indicated |
| 10 | earlier, if we were to do so, we would require |
| 11 | the sponsorship of the particular agency that |
| 12 | we think may have relevance here. They have |
| 13 | to get us into the NARA records. |
| 14 | MEMBER MUNN: This was a much more |
| 15 | broad question than that. |
| 16 | MR. ELLIOTT: I'm sorry. |
| 17 | MEMBER MUNN: The real question is, |
| 18 | do we have any idea whether there is an |
| 19 | ongoing program to begin further |
| 20 | digitalization of records earlier than those |
| 21 | that were undoubtedly done at the time. |

MR. ELLIOTT: I'm not aware of any.

MEMBER MUNN: It's always intriguing to know whether agencies are going to try to transpose their handwritten and typewritten material that they have.

MR. ELLIOTT: I'm not aware of any.
But Tim might have additional information.

MR. KOTSCH: To add to that, we have no indication - we don't know if they are going to computerize those records prior to `66. What Larry mentioned was that records that they have computerized, the information that is coded in those records is not useful. In other words, in their generic context saying, make this valve, do this type of work, it would be unlikely we would find a contractor who would say, and by the way it also included enriched uranium or something to effect. that So the nature of procurements that have been computerized, we feel are not elucidating for the enriched uranium process.

MR. ELLIOTT: Our request to the

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Department of Defense, we can see in the email exchange, went fairly broad and pretty deep. In fact you have met Dr. Paul Blake before who runs the DTRA program, the Defense Threat Reduction Agency program for veterans, he was involved in the response. I talked with him. And he was doubtful that we would be able to find any such contractual record in electronic form that would provide the level of detail that we wanted. He even suggested that we would probably be better off starting at the National Archives and working through that.

But he also offered - I mentioned the speculation that perhaps this was Navy fuel related. And he, as others, have said that he was somewhat skeptical of that because of the amount of enrichment in the sample versus what is in Navy fuel which nobody can talk about. So there - so I don't know where we go from here, but we feel we have done all that we can at this point, unless there is some other thread we need to pursue.

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| 1 | CHAIR ZIEMER: Thank you. Other |
|----|---|
| 2 | comments, questions? |
| 3 | Okay, thank you again, Larry, for |
| 4 | that update. |
| 5 | We are scheduled for a break - a |
| 6 | comment first? |
| 7 | MEMBER MELIUS: Is it possible to |
| 8 | get some of this in a short report or |
| 9 | something from NIOSH, just to - |
| 10 | MR. ELLIOTT: I would be happy to |
| 11 | send this email that was crafted. |
| 12 | MEMBER MELIUS: At least so we have |
| 13 | a record of it as we pursue it. |
| 14 | MR. ELLIOTT: I'll make sure the |
| 15 | Work Group has that. |
| 16 | MEMBER MELIUS: Particularly the |
| 17 | Work Group and the Board, and that way we are |
| 18 | not all hunting through transcripts. |
| 19 | CHAIR ZIEMER: We have a break |
| 20 | scheduled. I think we will go ahead and take |
| 21 | that break, then we will come back. We have |
| 22 | Board work time after that. |

| 1 | Let's take a break. We're |
|----|---|
| 2 | scheduled for 15 minutes; I'm going to give |
| 3 | you 20. |
| 4 | (Whereupon the above-entitled |
| 5 | matter went off the record at 2:26 p.m. and |
| 6 | resumed at 2:47 p.m.) |
| 7 | CHAIR ZIEMER: I believe we are |
| 8 | ready to resume our deliberations. Let me |
| 9 | check the phone lines. |
| 10 | Dr. Roessler, are you on the phone |
| 11 | line? |
| 12 | MEMBER ROESSLER: Gen Roessler here. |
| 13 | CHAIR ZIEMER: Thank you, Gen. |
| 14 | BOARD WORKING TIME |
| 15 | CHAIR ZIEMER: Under Board working |
| 16 | time, we have several issues we can deal with |
| 17 | today. First of all we have some issues |
| 18 | relating to site profile changes at Lawrence |
| 19 | Livermore, and Ted Katz has got some |
| 20 | information for us. |
| 21 | MR. KATZ: Yes, thank you. |
| 22 | So just before Thanksgiving John |

DR. MAURO: called me up to raise sort of a novel issue that SC&A had sort of come across to discuss how to go forward with it.

And the situation is this, and it relates to the discussion yesterday about close outs of site profiles. One of the site profiles as he termed it still sitting on the shelf awaiting the Board to take it up with the Work Group and a comment resolution process is Lawrence Livermore.

And in this case, despite the fact that the Work Group hasn't been formed, and there hasn't been a process resolution yet issue resolution yet - OCAS had taken that site profile review off the shelf and reviewed it and OCAS agreed with some of its findings, and wished to go forward to make changes to the site profile which would affect of course the dose reconstruction done at Lawrence the basis of some of Livermore on findings.

So there are two issues. John

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brought this to me and said, this is a good thing, so OCAS has come to some SC&A staff, some OCAS staff have spoken with some SC&A staff. They want to make some changes to the site profile on the basis of the SC&A review. And they were seeking clarification or what have you from SC&A staff to be able to implement some changes in the site profile.

And John said, and I agreed with him, you know, this is a good thing that this can move forward, even though it hasn't gone through the issue resolution process. But he also, we both recognized that this is sort of an unusual situation because the process is intended to be - have a process resolution, issue resolution with the Board, and the Board has not charged SC&A with doing anything with respect to this.

So this was just before Thanksgiving, so I said to John, well, don't make any significant expenditures on this, don't do any substantial work. Let me contact

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Dr. Ziemer. But I think this is an issue that really needs to come before the Board. is the first time this sort of situation has John pointed out arisen. But as in his presentation yesterday, there is more than one site profile that is actually is in this kind of status. It's been around for awhile, and so the issue could not just be for this case, but also for some other site profiles where OCAS may want to go forward before the Board has gotten to the SC&A review with making improvements to their site profile.

So I contacted Dr. Ziemer, and Dr. Ziemer got back to me after Thanksgiving and agreed that this was an important issue for the Board to take up, and also with a little bit more specificity said, limit what SC&A does at this point to nothing but providing clarifications on the issues it's raised until the Board has had a chance to take this up and give some consideration as to any one of these situations how far if anywhere should SC&A go

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before there has been a work group and so on in interacting with OCAS to make changes just like there are in this instance.

And I don't even know the specifics of the changes. I just know that there were some and they were important for improving the dose reconstructions there.

So that is the situation we have, how it stands with Lawrence Livermore. I don't know far they've gotten, OCAS and SC&A on - well, OCAS in making any changes with the Lawrence Livermore site profile, but I think it's important that the Board decide how it wants to deal with these sort of cases, and how it wants to charge me in terms of managing SC&A as its project officer.

CHAIR ZIEMER: Thank you, Ted. And let me add a couple of comments to that.

Number one, basically in the situation at hand, my instruction was that we were not tasking SC&A to do any new work; that if NIOSH wished to call John and ask what they

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meant by something, a brief exchange of what something meant, that's fine. But I didn't want SC&A doing any tasks outside of the Board's tasking process, number one.

Number two, in principle, let's just talk in general terms, it's conceivable that the Board might not at all agree with what the SC&A recommendation was, and might think that NIOSH's recommendation or procedure or whatever the particular item was the appropriate way to go.

So to go ahead and make changes based on the contractor's review without involving the Board didn't seem to be appropriate either.

So there is two parts to this. One is the tasking issue, and the other is the general principle of the way in which the contractor's report gets used outside of our regular Board process.

So that is where we need some guidance, because in principle this could

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arise in other cases if we haven't gotten to a site profile, but NIOSH has - or a review rather, and NIOSH has said, well, this makes sense, let's make a change. So how do we proceed? So we want to get some input on that.

Phil, Jim? Phil, go ahead.

MEMBER SCHOFIELD: I think maybe on some of these much larger sites that we go ahead and have them take a preliminary look at some of these, just because they are so large and so complex that it takes quite awhile before we get to the point where we can actually sit down both as a work group and as a Board and make any real decisions just because of the sheer volume of records they have to go through in these early stages.

CHAIR ZIEMER: Okay, thank you.

Jim.

MEMBER MELIUS: A couple of points.

I guess NIOSH OCAS certainly can take technical input from wherever. There may be

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an article in a journal that changes a dose reconstruction method, or there may be some other process where they become aware of a potential change. I think there is always a potential at a site meeting or something where they are out doing some outreach that they will learn something new that changes a site profile or something. I don't find that to be problematic at all.

But Ι find it be very problematic that we would have our contractor interfacing with NIOSH and making progress on trying to resolve issues without involvement of the Board at all. And I think it puts us in a very awkward position of having to maybe disapprove of something that our contractor has recommended. But if you go back in time, I've found myself disagreeing with our contractor quite regularly on issues, and I think other members of the Board have also from their perspectives, or certainly seriously questioning their conclusions

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recommendations on a particular point.

So I don't think that is something unusual. Now these may be minor technical issues. They may be something very straightforward. I can't speak to what happened there. But I think it's problematic.

I think we have gotten into a position where our contractor is part of this close out process because of the nature of the way the contract award has been delayed and whatever, been put in the position of trying to rush things through the process. And I think that is wrong and I don't think they should be doing that. I think we need to be very careful about that.

I also think we've even had situations here where the contractor is preparing and presenting reports that not even the Work Group has seen prior to the meeting, and that I think is also inappropriate. They should be working under the direction of a work group, and work group Chair; and there

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should be at least some level of control and so forth, and we need to ensure that their work is independent.

And I have been disturbed, and I think other members of the Board have spoken to me about being disturbed by the sort of the encroaching control over our contractor or attempts to control our contractor by the agency. And I think we have to be very aware of that and very careful.

CHAIR ZIEMER: Other comments?

Here again let me emphasize that in this case that any sort of activity that looks like an issue resolution should not go on without tasking. As I indicated in this case, if NIOSH wished to call John and ask what he meant in a couple of sentences, I don't object to that.

MEMBER MELIUS: Just to clarify, I was not objecting to how you handled it. I thought that was appropriate in bringing it back to the Board and so forth. And again

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that particular question, if NIOSH sees a technical point that they feel they should be changing or addressing, that's fine.

CHAIR ZIEMER: And you are quite right that NIOSH is free to use whatever sources that they see out there that might help them modify or change what they are doing.

Ιt to me, let me propose seems something here, and we can react to it, but one way to handle this would be that if a case arises where NIOSH is looking at a particular issue, and they see something that the contractor, whether it's SC&A or whoever be, that they wish contractor may incorporate, and if it will require a sitting down together to figure out how this works, then we need to be involved, and even if there is not a work group, we can always set up an Ad Hoc Work Group or something to say, yes, okay, here is an issue on this site, and we need to have Board members present.

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set up an ad hoc group or whatever we have. We could do that on short notice if necessary, and I think we would have to leave that up to NIOSH and the contractor, say, something dealing with whatever site. Say it's Lawrence Livermore, and it's, we need to together and hash out the technical details on this. So we can do that. task on short notice. And I think certainly the Board and chair are in a position to do that, either the fallboard or the chair in between, to appoint a group.

But I'd like to get input in some direction from the Board as to how we proceed, so that we make sure that we don't have the situation - that is awkward for all parties if we are out of the loop.

So any other input or suggestions, or if there is a better way to do it I'm open to hearing that. I always like my own ideas, but I'm old enough to recognize that there are two or three people in the world that have

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| 1 | better ideas than I do. |
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| 2 | MEMBER CLAWSON: I understand the |
| 3 | need for this and so forth, but one of the |
| 4 | things I want to make sure, especially as |
| 5 | working chair of some of these groups, that |
| 6 | they especially the member of - the chair of |
| 7 | the working group is involved in it, not just |
| 8 | a member of the Board or whatever like that. |
| 9 | Somebody from the working group or whatever. |
| 10 | Because a lot of these conference calls and so |
| 11 | forth and fact findings or whatever you want |
| 12 | to call them bring us some very important |
| 13 | information. |
| 14 | CHAIR ZIEMER: Well, here we are |
| 15 | talking about cases where there is not a work |
| 16 | group. |
| 17 | MEMBER CLAWSON: Well. |
| 18 | CHAIR ZIEMER: If there is a work |
| 19 | group there is not a problem. |
| 20 | MEMBER CLAWSON: Right, okay. |
| 21 | CHAIR ZIEMER: If the Work Group |
| 22 | gets involved. Here we have Lawrence |

Livermore. We don't have a work group. We have a NIOSH report. We have an SC&A review.

NIOSH has looked at the review. They see something, they have said that's a good idea, let's modify our report. We don't have a work group. That is the situation.

Wanda.

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MUNN: MEMBER There are some similarities in this dilemma to the issues discussed several years ago when contractor could respect to our appropriately represent what was transpiring the Board, for example, before on congressional members. And they were neither a member of the Board nor were they originators of the documents in question sometimes.

But when we don't have a work group, an obvious path through which a Board connection can be made, if we don't notify the Board that this is transpiring, and at least provide an opportunity of a week's time for

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input of some sort, then it seems that we would be getting outside the realm of the structure as we - or at least as I perceived it at the time that we set up our association with our contractor.

Is there any problem with just simply asking both the contractor and the agency to see that situations of this sort are brought to our attention by electronic means and we have an opportunity to respond.

CHAIR ZIEMER: That is not a problem at all, and that can be part of a policy, and actually was done in this case. The question for the chair is, how would you like us to handle that?

MEMBER MUNN: Is this appropriate?

CHAIR ZIEMER: Well, if the Board wishes to sort of develop a policy, and we don't have to do it sitting here. But I want you to be thinking about it. We could even come back tomorrow and adopt something, or you can instruct me to do certain things when such

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1 a situation occurs. 2 But we want to make sure - and I think both NIOSH and the contractor wish to 3 make sure that everybody is in the loop and 4 nothing is happening that we are not aware of. 5 And Joe, you have a sage comment on 6 this? 7 MR. FITZGERALD: Well, just a little 8 I've been working on this 9 more context. 10 Livermore issue, I think, now that I've heard a little bit more, it's the issue I've been 11 working on. And quite frankly, this was a 12 13 data capture strategy that NIOSH was developing for a revisit to revamp the site 14 15 profile. 16 We weren't getting into any issues in terms of recommendations from the site 17 profile. 18 19 CHAIR ZIEMER: In other words how to capture data? 20 MR. FITZGERALD: Well, it was more 21 of a case of for efficiency's sake we covered 22

references. We went to classified databases that the authors of the very first site profile that NIOSH had done four or five years ago had not. So the obvious question when you are going back was, looking at the delta, is there any facilitation that we could give them as to who we had gone to, points of contact, and what databases per se that were missed in the first review that they would then want to take advantage of in this next round review.

So I consider that facilitation; pointing them to the right people, identifying the databases more clearly.

CHAIR ZIEMER: So it wasn't a change in their documents?

MR. FITZGERALD: No, it wasn't a policy exchange at all. It was a facilitation on a data capture. Now the reason I think we wanted to come to you and Ted was simply because it turned out, these data capture strategy plans get pretty detailed and

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| 1 | complex, and once you get beyond the first |
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| 2 | hour or two of facilitation, you start |
| 3 | questioning, well, this is a fairly big job, |
| 4 | we just want to make sure that somebody was |
| 5 | aware, that the Board was aware, that we are |
| 6 | working on this, and it's not simply we are |
| 7 | doing on our own time and doing it off the |
| 8 | premises, that this facilitation is happening. |
| 9 | So I just wanted to make sure that |
| 10 | there was somebody besides just me and John |
| 11 | aware of the fact that we were providing this |
| 12 | kind of support. But it wasn't in the policy |
| 13 | context, it was on the facilitation context. |
| 14 | MEMBER MUNN: Well, it's billable |
| 15 | hours anyway. Right? |
| 16 | MR. FITZGERALD: No, we don't have |
| 17 | an authorization from the Board to work on |
| 18 | Livermore. The Livermore site profile was |
| 19 | completed. |
| 20 | MEMBER MUNN: That's really the |
| 21 | issue here. |
| | |

MR.

FITZGERALD: Well, helping out

is fine in a certain context of time. But once the time gets lengthier, then I think there is more of a concern that at least we alert -

CHAIR ZIEMER: Right.

Jim? Then Ted.

MEMBER MELIUS: In that case, isn't it appropriate to be able to encompass that kind of assistance into one of the tasks. don't remember the tasks well enough that are part of the contract to be able to - if it is adding up to many hours of facilitation, then it should, and there should be some appropriate control from NIOSH and the Board on authorizing that. But it would be more of a generic issue, wouldn't it, Paul?

CHAIR ZIEMER: I think you are right. My concern here would be, if this is a one-time thing, it's kind of a no, never mind. But if it's a facilitate here, facilitate there, pretty soon you are adding up hours, and so is there then a separate task - a

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facilitation task to which we assign a budget.

You can spend up to a certain amount each year facilitating things of this nature, which may not have to do with modifying a report per se, I mean, it may lead to that. But if it's to the extent that our contractor is helping get the job done, and in a sense is helping us by doing that, and helping NIOSH. We do have management money, but I think that's a little different than this. This is - this looks a little different from other things that we have done, doesn't it? John, how does it look to you?

DR. MAURO: The ground rules that I have been using from the very beginning is, once we deliver our report, a site profile review, we do not bill any more time until the Work Group has started and we are authorized to do more work.

Now given that, what could happen that would put us in a position to jeopardize that situation? One, we get a phone call

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from a representative who wants clarification on Pinellas. Explain to me what this OTIP-66 is? I answered that question over the phone.

A perfect example here, let's say we are talking about the situation that Joe just described. Another situation would be documents. We have an obligation to make sure that every document that we use in our report we share with NIOSH. So if we get a call from NIOSH, listen, I see you've cited this, this, this, this document, we have an obligation to make sure that that is available to everyone, so we'll do that.

So there are some what I would consider to be very modest levels of effort which are on the order of an hour or two that very often we encounter, and I do that. I take care of it, and I'll bill that hour. Even though that sort of breaks one of my - I like to put a freeze, because I'll tell you what I do. I like to track how much does it cost us to put the product out, and then from

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then on, from that date, the date on that report, from then on every hour that is billed and every dollar that is billed goes toward what I call close out, so that I could always report back to you how much did it cost to produce the Pinellas report, how much did it cost to support the close out of Pinellas. And I could give you those numbers for every one of these sites.

The way I see it is that this little what I call follow on support to whoever it might be is very very modest, and my general rule of thumb, if it takes an hour or two, that's okay. And that's how I've been running the program.

CHAIR ZIEMER: If you are talking about an hour or two here or there doing describe, things such you which as providing reports, or even what Joe described, sharing your strategy for - or the points of whatever it might be, you contact or easily bill that against the close out

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process, or we can identify it separately, either way. But if it's very modest it's probably not worth tasking separately.

DR. MAURO: That's what we've been

DR. MAURO: That's what we've been doing, yes.

CHAIR ZIEMER: But if in somebody's judgment you are getting - if the thing that Joe is describing now gets very extensive and I don't know, Joe - is he still here? But if you end up having to spend a lot of time facilitating this, and I don't know what might occur.

MR. FITZGERALD: It's a subjective thing. I think it's very profitable for us to help with the data capture on a site we've been at in terms of the feedback we can provide. I think that is a very profitable time. After a couple of hours, you get to the point where you just want to make sure that, you know, that you are aware that we are involved in this.

And of course the other issue that

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| 1 | you raised, which is, if this is something |
|----|---|
| 2 | that would be continuing over time, then |
| 3 | perhaps that is something we need to |
| 4 | benchmark, because it may take more time. |
| 5 | Some sites will be more complicated. And I |
| 6 | think this was a recognition that as perhaps |
| 7 | these sites are revisited, this might be a |
| 8 | continuing role, and therefore we ought to |
| 9 | flag it as something that the Board should be |
| 10 | aware of. |
| 11 | So this was mostly a heads up. |
| 12 | CHAIR ZIEMER: This may be a case |
| 13 | where it is not broken yet. |
| 14 | MR. FITZGERALD: No, I really don't |
| 15 | think so. |
| 16 | DR. MAURO: If you have a task they |
| 17 | can bill it on. |
| 18 | CHAIR ZIEMER: Yes, it sounds like |
| 19 | it's sufficiently modest that it - unless it |
| 20 | reaches a point where in your judgment it is |
| 21 | becoming a significant effort. |
| 22 | MR. FITZGERALD: Well, I just want |

to make sure - we have had instances where the 1 2 Board will say, what are you doing that for? How come we don't know about it? Well, in 3 4 this case we are alerting you that, yes, we are working with NIOSH and looking at the data 5 capture plan for Livermore and providing 6 7 feedback and spending time doing that. So it was more of a heads up, to 8 aware of that, 9 sure you were 10 wouldn't come across as a surprise later on that doing that. But it's not 11 we were crossing over into the issues or policy area, 12 which I think is what I heard earlier, which 13 is not the case. 14 15 MR. KATZ: I just wish you'd use a 16 term other than profitable. (Laughter.) 17 CHAIR ZIEMER: Ted, Wanda, Mike. 18 19 MR. KATZ: I mean just for the record, though, this wasn't made clear to me. 20 So I guess, John, you and I need better 21

this

was

Because

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communication.

not

my

understanding, the characterization that I received of what was the work that needed to be done.

When DR. MAURO: Ι called, Ι realized that we were moving into a mode that we hadn't been into before, where one of our folks might be spending several hours, maybe more, helping out in this capacity, which would be - and I said, let me alert you to this. It's the first time this came up where there possibility that this could was а continue for awhile, and that's the reason I called.

KATZ: Yes, but you explained MR. that they were going to be making improvements the site profile based the to on recommendations you made. I mean that very clearly the characterization. So I did think we were in a mode of making changes based on your recommendations. That's how I understood that.

CHAIR ZIEMER: Well, by the time it

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got to me, the changes were pretty far along.

Okay, that's very helpful. It's

much more clear.

Wanda, do you have some additional
advice?

MEMBER MUNN: Well, I don't know if it's advice. I'm just mulling over here the fact that we came a long way from the original question that was posed. But as long as that question has been posed, it seems reasonable that we should address it at some juncture, in the future. whether now It's orreasonable possibility that might exist with respect to future interactions, especially process has been initiated once any contact, especially contact points.

CHAIR ZIEMER: I think the parties involved are sufficiently sensitive to that that if it reaches that, they certainly have been good about alerting us, even in this case where the threshold actually was pretty low and they still let us know, even though it got

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| 1 | misunderstood a little bit. |
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| 2 | DR. MAURO: I perhaps communicate |
| 3 | too much. |
| 4 | MEMBER MUNN: It's not possible, |
| 5 | John. |
| 6 | DR. MAURO: I have become very |
| 7 | sensitive to the fact that it is very |
| 8 | important for me not to move forward with |
| 9 | anything unless - |
| 10 | CHAIR ZIEMER: Right, and we |
| 11 | appreciate that, so thank you. |
| 12 | Michael, additional comment. |
| 13 | MEMBER GIBSON: there is nothing |
| 14 | wrong with efficiency with the agency and the |
| 15 | contractor trying to work together. But even |
| 16 | on data capture it seems - you know, NIOSH |
| 17 | goes out and takes a look at the puzzle, and |
| 18 | then we want our contractors to take a second |
| 19 | look at that puzzle, to see where someone |
| 20 | might have missed something. And that's how |
| 21 | we get to this resolution process. |

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So it's also - it looks like there

| 1 | could be a way that there may not be |
|----|--|
| 2 | necessarily two complete different views when |
| 3 | you share this kind of information. |
| 4 | CHAIR ZIEMER: Right, but we do as a |
| 5 | practice want to share all the data sources, |
| 6 | have all the folks have access to the - as |
| 7 | much of the information as they are able. |
| 8 | MEMBER GIBSON: But after the fact. |
| 9 | CHAIR ZIEMER: Yes, good point. |
| 10 | Okay, other comments? |
| 11 | Well, it appears to the chair that |
| 12 | we don't need a specific policy at this point, |
| 13 | but we are sensitive to the issue. The |
| 14 | contractor is, NIOSH is, and it appears that |
| 15 | we have a way to deal with it at the moment. |
| 16 | So thank you very much. |
| 17 | Next we have Procedures Work Group. |
| 18 | Let's see how we are in time here? We're |
| 19 | good. |
| 20 | PROCEDURES WORK GROUP |
| 21 | CHAIR ZIEMER: Procedures Work |
| 22 | Group. Wanda, you have a couple of items. |

| 2 | and there was another - oh I think it was the |
|----|--|
| 3 | - wasn't there another one that involved your |
| 4 | work group? |
| 5 | MEMBER MUNN: Yes, there is. There |
| 6 | are a couple of things that we need to report |
| 7 | on. The question is, do you want - from my |
| 8 | perspective, would you like me to give the |
| 9 | Procedures Work Group report now and then we |
| 10 | can discuss the CATI and the other items with |
| 11 | respect to how our tracking system is going? |
| 12 | CHAIR ZIEMER: You might as well do |
| 13 | the whole thing. |
| 14 | MEMBER MUNN: If we have the time to |
| 15 | do that. |
| 16 | CHAIR ZIEMER: How long do you need |
| 17 | - you are going to also discuss the issue that |
| 18 | Dr. Melius raised? |
| 19 | MEMBER MUNN: Yes, if possible. |
| 20 | CHAIR ZIEMER: Okay, let's do that. |
| 21 | MEMBER MUNN: Let's see if we can do |
| 22 | that, and we will try to move through it |

One has to do with the CATI interview issues,

quickly and efficiently - give you a little
CHAIR ZIEMER: Well, I guess
somebody doesn't think you can move through it
quickly.

MEMBER MUNN: It's Christmas time. The Procedures Work Group met in Cincinnati on the 9th of this month. And for the first time approached our data tracking base in a fully digital manner. Of course this database has been worked on for a number of months, and the beautiful contractor has done а job in cooperating with NIOSH to get this running.

And from our perspective as a work group it's working very well. We did go through this fully paperless this time. No one touched a single tree. And we were able to move through the items. We limited ourselves to what we call the third dataset, based on grouping - the last group that we had tasked our contractor with doing, and were able to close quite a few at the time that we

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also started We our day by discussing the new proposed CATI questions that we had all received by email a week or so prior to the meeting. There were as you all know two segments to that, set of one questions for the employee himself or herself, and the other one for survivors.

We had a fairly brisk discussion in the Work Group with respect to whether the questions, especially the questions on the survivor questionnaire were appropriate or even useful; whether they were in fact providing the kind of information that we were attempting to get to.

Having had such a significant amount of feedback from survivor petitioners with respect to their concerns over these questions when they have them, the sort of mistaken feelings that they were sort of being tested, and that they did not have the proper responses; that they were somehow going to

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negatively affect their claims.

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We looked at them fairly closely. I don't think there is any way that you can report in a concentrated form on the real essence of any conclusion that we reached, except that we did understand clearly that the individuals who do these questions try to reassure the claimants at the time that they are carrying out the questionnaire so that the claimant will not be unduly concerned that lack of information on their part creates a negative atmosphere for their claim.

But the letter itself still has contains although it the appropriate information from I believe the viewpoint of most of the Work Group members, it may have too official a tone. This was a finding which one of our previous work groups came to with respect to closing letters that went out to claimants; the information was there, but the perhaps what was not we wanted achieve.

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So we wanted to bring this back to the Board. We took no real action on it, because it was very clear from our discussions that it was likely almost all members of the Board feel strongly about this.

It was our suggestion that we plan to meet toward the end of January again; as a matter of fact we established a date of January 28th in Cincinnati. Prior to that time we would ask that each Board member review that letter and the two questionnaires very carefully, and provide Ted Katz and me with any comment that you have, any suggested wording that you have. We would like to try to, at our next meeting, blend the concerns that are expressed into a cogent response, which we could bring to you at the Albuquerque meeting as a potential for comment.

If anyone has any suggestion as to a move effective way to try to achieve some sort of consensus on the Board with what our response should be in terms of comment period,

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we are certainly open to that.

Yes?

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GRIFFON: Ι just think in MEMBER terms of the comments, the other things that we were talking about on the procedures call is maybe when you through these go questionnaires and the letter, that you differentiate your comments the on questionnaire itself versus the process. Because I think we were finding issues with both parts, and wanted to kind of keep them separate.

CHAIR ZIEMER: And let me ask for clarity. My recollection is that, is it only the questions that the OMB has to approve and not the letter itself? Was that not the case?

MR. ELLIOTT: Is this on? Now it's on. That is correct, the OMB will review the survey instruments and approve those. What you have before you is a cover letter that Wanda has mentioned that goes to the claimant introducing the questionnaire to the claimant,

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giving them an advanced understanding of what is going to be asked of them so they can prepare for the interview process.

And we ask as Mark has clearly identified during discussion if the t.o possible segregate your comments to that of the process versus that of the questions that Because we will need to consider are posed. and react to both sets of types of comments, but the ones that the OMB will be interested in knowing how we - what kind of comment we got on the questions. We wanted to forward those and show how we reacted.

Let me also take this opportunity to say that in the working group discussion I noted for the working group that we had put forward these two revised survey instruments, based upon input that we viewed in the SC&A review of procedure 90, is it? Well, whatever the interview process procedure number is; I'm lost on that right now. But also input from the folks who actually do the interviews, and

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input from the general public during public comment period. We heard loud and clear concerns and frustration and burden that these questions were bringing.

In that light I proposed that there might be another option, rather than this set of questions being posed, we might go at a questionnaire designed specifically to confirm the information that we already have in the claimant file; that is necessary, that appropriate for us to do. But as I said it's short number set of questions that a small would go just to confirming the information at hand with the person. And also - we proposing that this will be done when change contract award situation here, that we will start awaiting the - for DOE claims we would await the arrival of t.he DOE information before we sat down with claimant to do the interview process, so we could cover that with the claimant, address whatever concerns they might have about it not

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containing everything they think it should have, and then there could be another set of questions designed specifically for energy employees, plus a separate design set for survivors that get at information the health physicists feel is essential to doing a dose reconstruction.

And so that might take on a different look than the two examples you have before you that are revised questionnaires based on input. I remind you all -

CHAIR ZIEMER: But that would be down the road, Larry, is that right? Or is that in this one?

MR. ELLIOTT: We are working on this option right now, and if we get a request through the Federal Register notice, it's my intent that we would provide an example of the two that you have got, and this third example, so when we come up with this third example, we are going to share it with the Board as well so you can see what it looks like.

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1 CHAIR ZIEMER: Will that change the 2 reply time in the Federal Register? Or would there be a new separate -3 MR. ELLIOTT: No, it doesn't change 4 the reply time. We have folks working on this 5 6 option that I'm talking about now. Remember 7 at the working group meeting I mentioned that I was awaiting some consensus comments from 8 health physicists that would be factored into 9 10 this option. And so the time won't change, and 11 committed to the working group, your 12 13 comments come in after the suspense date of the public comment period, they will still 14 15 consider the Board consensus comments as we 16 move forward to revise these documents. And we'd account for that in the 17 OMB package that we would send up. 18 19 I would remind you that the review of Procedure 90 and the fact that we 20 have used this same set of questions 21

25,000 cases, probably around 35 to

| 1 | 40,000 claimant interviews. And we are asking |
|----------------------------------|---|
| 2 | ourselves what do we really need. And we are |
| 3 | forced with this OMB requirement to speak |
| 4 | about the burden, and you will see that |
| 5 | mentioned in the Federal Register notice, what |
| 6 | we estimate to be the burden hours based on |
| 7 | the two documents that you have before you. |
| 8 | And I've got to believe that there is a better |
| 9 | type of questionnaire that will put less |
| 10 | burden and less frustration before the |
| 11 | claimants. |
| | |
| 12 | CHAIR ZIEMER: Thank you. |
| 12 13 | CHAIR ZIEMER: Thank you. Now what do the Board members have? |
| | |
| 13 | Now what do the Board members have? |
| 13 | Now what do the Board members have? Do all the Board members have the document |
| 13 14 15 | Now what do the Board members have? Do all the Board members have the document that we saw? |
| 13 14 15 16 | Now what do the Board members have? Do all the Board members have the document that we saw? MEMBER MUNN: Yes, all the Board |
| 13 14 15 16 | Now what do the Board members have? Do all the Board members have the document that we saw? MEMBER MUNN: Yes, all the Board members have the cover letter and the EE and |
| 13 14 15 16 17 | Now what do the Board members have? Do all the Board members have the document that we saw? MEMBER MUNN: Yes, all the Board members have the cover letter and the EE and SV questionnaires. |
| 13 14 15 16 17 18 | Now what do the Board members have? Do all the Board members have the document that we saw? MEMBER MUNN: Yes, all the Board members have the cover letter and the EE and SV questionnaires. CHAIR ZIEMER: So not only the Work |

| 1 | Wanda, is basically that Board members prepare |
|----|--|
| 2 | their comments and provide them to you so the |
| 3 | Work Group can develop from that a |
| 4 | recommendation to bring back to the full Board |
| 5 | at its next meeting; is that correct? |
| 6 | MEMBER MUNN: That is correct. We |
| 7 | are asking that those comments be gotten to |
| 8 | us. |
| 9 | CHAIR ZIEMER: And in that you are |
| 10 | basically asking the Board, is that how you |
| 11 | would like to proceed? |
| 12 | MEMBER MUNN: Exactly. And if that |
| 13 | is the way the Board would like to proceed |
| 14 | then our timeline is very clear. We need your |
| 15 | input in the next couple of weeks. |
| 16 | CHAIR ZIEMER: You will meet in |
| 17 | January? |
| 18 | MEMBER MUNN: We would need your |
| 19 | input in a couple of weeks so that we could |
| 20 | prepare something in the Work Group. |
| 21 | CHAIR ZIEMER: Right, and then |
| 22 | develop a recommendation for the February |

| 1 | meeting, at which time the full Board could |
|----|--|
| 2 | hopefully approve something. |
| 3 | MEMBER MUNN: Either approve or |
| 4 | further deliberate what we bring to you. |
| 5 | CHAIR ZIEMER: Comments or questions |
| 6 | on that? Is that agreeable as a way to |
| 7 | proceed? Mark. |
| 8 | MEMBER GRIFFON: Just a question, if |
| 9 | Larry knows when this third option might be |
| 10 | available and when we might see this. |
| 11 | MR. ELLIOTT: I am giving my |
| 12 | personal thoughts here. The third option may |
| 13 | not even be a viable option once we look at it |
| 14 | and see it. But if it comes out as I think it |
| 15 | may, it's very soon. It's got to be soon, |
| 16 | because the Federal Register notice went out |
| 17 | what last Thursday, is that right? |
| 18 | MEMBER MUNN: That's about right. |
| 19 | CHAIR ZIEMER: And the individuals |
| 20 | have to contact you to get the - |
| 21 | MR. ELLIOTT: Right. So if you as |
| 22 | an individual citizen want to comment in the |

1 public comment period, you send in a request 2 to see --CHAIR ZIEMER: You have to be able 3 to provide this. 4 MR. ELLIOTT: I have to have this 5 ready very soon. Because if we get a call for 6 7 what these exhibits are tomorrow, we have to respond. We are trying to put this option 8 together. 9 10 MEMBER GRIFFON: And again can you this third option little bit? describe a 11 It's got to be a questionnaire that says, you 12 13 know, check off what records we have, and does that agree with your memory of where you 14 worked? 15 16 MR. ELLIOTT: The third option as I tried to explain it before, my concept of this 17 other option is that we take the time to 18 19 confirm what information we have. So that is list of questions, the 20 short critical We have your name, your date of 21 issues.

diagnosis, your work history, this is where

you worked. That would be - if you look at it in three segments, that is the first segment.

The second segment would be for a DOE worker to say, we've received your dose information, and we show that you have been badged these years. You didn't get a badge this year. You didn't have a whole body count. You did provide urine for bioassay on these timeframes. Is that to your knowledge anything different than that? A brief question in that regard.

third segment would be those questions that the health physicists who do dose reconstructions feel are questions that would help them understand best how reconstruct that dose. And I don't know what those are yet. So I'm waiting to see what those are, because they may be no different than the questions we currently have, and maybe a shorter version. But that is where we are trying to go. We are trying to lessen the burden in this third option, and hopefully in

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the two that you had before that we want to lessen the burden. But we also want to lessen the frustration, so that also goes to this process aspect of how to administer the questionnaires.

CHAIR ZIEMER: Josie.

MEMBER BEACH: That third option could be a case by case basis, too. So that sounds like it could be somewhat cumbersome depending on the claimant.

Well, MR. ELLIOTT: I'm not proposing that it be a site-specific kind of questionnaire. The questionnaires have to be standard in the format of questions that are used. And so it goes - the front part, the first segment is probably individual specific, But the questions that come out of that yes. simple. Do we have everything? Is are everything correct? That is the question.

So that is the question we can ask.

It's a little burden, but it is important to verify that we have this information correct.

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MEMBER BEACH: Right, and I wasn't thinking of the first two sections you described. I was thinking of the third one, and maybe I got carried away on my thinking.

MR. ELLIOTT: You know, I've got a problem right now with the two examples that you have before you, and the way we have been using the current questionnaire recently in And that problem is that we already know the question - we already know the answer to many of those questions. So why are we asking the question? You burden the claimant when you ask a question and you already know the answer. And you raise their expectation, and by that you raise their frustration level when you don't do anything that they think you ought to have done with that answer t.hat. they've given you.

So we are coming at it with a very hard critical look at what we need in a questionnaire approach in the CATI interview.

So hopefully we come out of this with less

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burden and less frustration on the claimants. 1 2 That's what we want, but we also want to gain information that will help do the dose 3 reconstruction. So it's a difficult balancing 4 5 act. CHAIR ZIEMER: Thank you. Other 6 Dr. Melius. 7 comments? MEMBER MELIUS: I asked a question 8 yesterday whether this - our comments would 9 10 become part of the docket for this. MR. ELLIOTT: I don't know if you 11 are going to take my word on this, or if you 12 13 want the lawyers to come up and parrot me after this. But again it is not a regulatory 14 15 docket. It is NIOSH's docket, and I believe that I have the ability to insert information 16 into that docket post the comment period. 17 it was a regulatory docket, I am very well 18 19 aware that I cannot do that. So if you don't believe me and you 20 want the lawyers to come up and say that, I 21

quess we can have them do that.

1 MEMBER MELIUS: You say that with more certainty than you did yesterday. I will 2 also trust you that should the lawyers tell 3 you otherwise you'll let us know. 4 MR. ELLIOTT: Thank you. 5 CHAIR ZIEMER: Any further comments? 6 7 Can I take it by consent, Board members, that we will proceed as described and get our 8 comments to Ms. Munn so that the Work Group 9 10 can develop a document for us to look at? objections? 11 MEMBER MUNN: I will send you all an 12 13 email reminding you of what the anticipation is, and so that you will have some specific 14 15 dates in front of you where you can work 16 toward those dates hopefully in getting the information back to us 17 and repeating expectation. 18 CHAIR ZIEMER: Thank you. You have 19 one other issue now you wanted to talk about. 20 MEMBER MUNN: Actually I wanted to 21

touch on one other thing that we had done

during the procedures review process. an opportunity for the first time SC&A's review of OTIP-066, which is a crosscutting procedure regarding the calculation of dose rate of tritides. And we are pleased to see they had two observations and four findings. Unfortunately, the timing is such that NIOSH had not had an opportunity to really take a look at that at all. So that is on our fast track list for requests that NIOSH take a look at that as soon as possible, and we are hoping that we will have some feedback from NIOSH regarding those four findings when we meet again.

As I said, we've gone through the entire third set which dated back to October 29, 2007. And at this point as John Morrow had indicated yesterday in his report, we are approximately halfway through the existing outstanding findings that we have before us.

Now yesterday when we were in other discussions, regarding Savannah River I

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believe, I had made a comment relative to OTIB-0052, and how we were progressing through that, and Jim had asked that I illuminate you further with regard to where we are.

I had thought this might be an opportunity to have you all familiarize yourself with the portion of the O-drive data that we utilize all the time in the procedures activities, because of course the purpose in having this database established the way it is, is for all of you to be able to follow where we are with any one of these procedures.

Are all of you here capable of - I guess Mike may not be right now - but can all of you bring up the O drive on your computer today? If you can, perhaps we can use OTIB-0052 as a very quick lesson in how we move through these in the Procedures Work Group and get a little familiarity under your belt with how to get to information that you might like to have in that regard, if you would like to take few that, I'd minutes to do

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delighted to do so. And I think Steve Marschke who usually does this for us in work group and Stu who - Stu Hinnefeld who handles the NIOSH side of input and upkeep for this particular database is quite willing to throw some of the information up on the Board so that you can see how we do it if you want to do that.

Would you like to do so? Take just five minutes or 10 minutes here to take a look at how we do the procedures things.

Stu has I think some reservations as to how well this is going to go, because apparently things are slow on the net right now. But let's see if we can bring it up. For those of you who are going to try to follow if you can bring up the O drive on your computer and let me know when you are ready.

Okay if you have the O drive up then you will see the shortcut folder there to the procedures, right? Click on that folder.

And wait for a few minutes.

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MEMBER CLAWSON: Which shortcut?
There are several shortcuts.

MEMBER MUNN: It should say shortcut to database. Procedures Database. No? Let me go back and see. I'll start from square one here, and see if I can follow with you.

Now the first thing you need to know about this database when you are working with it is that we have deliberately arranged it alphabetically so that all you need to know is the number of the procedure or the document that you are looking for, and you can just run down - if you notice the finding date, the next thing you see is the title of the - the number of the document, and by looking just at the title of the document alone, and to the column next to it, you can tell how many findings we have on that particular OTIB-0050, and on the third column you will see dash one, dash two, dash three, dash four, that tells you without going any further that on that particular procedure you have four different

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findings, and you can see in the far righthand column what the specific status of each one of those findings is going to be.

Now when we did our work this time, you noticed above the - you see the boxes above the data, the second one from the left, says filter slash sort. What we did when we were working on this group of three, since we knew that the date was crucial for identifying them, we sorted - if you click filter sort, then it will show you three different ways that you sort. We sorted first by date, then second by procedure number, and third by finding.

And then we only looked at open, in progress and abeyance. You see where it says, filter on? We took all of those check marks out except for open, and in progress and abeyance.

And then we clicked on filter on, but I'm not asking you to do that, because that is not the way we want to handle what I'm

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1 trying to tell you to do here. I just wanted 2 you to see how we filtered so that we could bring up only the information that we wanted 3 4 to see. Now if you go back now to what Stu 5 is showing you on the big screen. 6 7 MR. HINNEFELD: Wanda, I'm not showing them anything useful, because this is 8 I'm going to have to restart not working. 9 10 You guys keep working on the phones. MEMBER MUNN: Okay, those of you who 11 have this up on your screen, on the far right 12 13 of your dataset is the move-me-down block, so that as you put your cursor on that block and 14 15 move it down, you are moving through the OTIBs for example by number and if you go down to 16 OTIB-0052, are we following okay, 17 are you getting there, you have OTIB-0052 up? 18 19 MEMBER BEACH: We have several. MEMBER MUNN: Right, so you look in 20 the third column and you will see how many 21

OTIB-0052 findings we have.

22

52, one,

three, four, five six, notice it goes all the way down to 16 before it changes to the next number.

Now looking at those 16 original findings, starting at one, you can go over to the far right and you can see what the status of it is. If we want to go to the detail of this we can find that we already know it is going to be addressed in some other finding.

Now just above the database itself you see the three tabs on the far left in gray: summary, details and procedures. put your cursor first on 52, and then click on will details, it bring for up you the information on that first finding of OTIB-0052. It will show you that it's addressed in findings, in another finding, and it will give you by date on the left what has transpired. It shows what the finding is, and then it shows you what the initial response was from NIOSH on 8/23/2007, and then as you go down those - that form you will see that on the 29th

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of August NIOSH and SC&A discussion, you can see an abbreviation of what was said, what was decided. And then on the 6th an SC&A follow up. If you want to see what that says, the status of this issue is going to be changed. Addressed in finding #16. So the status of this now is addressed in the finding. That is the next one down.

next issue. If you want to click on the next issue, then what comes up is finding #2 and #3. They are both closed. You see what the original finding was, what the NIOSH response was initially, what the discussion was later, and the follow up is that SC&A agrees with the NIOSH initial response. This issue is closed. And the date that it was done.

If you go to the next issue, and you find item #3. It's closed. You can click right on next issue, next issue, next issue, and go all the way down through all 16 of those issues.

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When you are finished looking at the detail, then put your cursor back up on the summary. Statement on the far left of the top of the database. Click on summary again and you are back on the summary sheet.

Was everybody able to follow that It is a little cumbersome reasonably well? the first few times you use it. But once you have become familiar with it, it really and truly is a marvel of complex information one step at a time. So that anytime you have any question with respect to where we are on any one of the procedures that we have given to SC&A to comment upon, you can see what is You can through the filter and sort potential; you can identify how many open items are there; how many are in progress; how It will bring that up for many are closed. you very easily.

Have I puzzled anyone? If you have any questions with respect to this please don't hesitate to address them to the Work

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1 Group. I'm sure Stu would be glad to answer your questions. Steve Marschke is glad to 2 answer your questions with respect to how to 3 handle this database. 4 So once you know how to get into 5 it, it can be a very helpful tool. If you 6 7 have any comments on it please let me know, and were you able to follow any of that Jim? 8 No, I'm sorry about that. 9 Any other questions 10 or comments with respect to the Database? 11 If not then -12 13 CHAIR ZIEMER: Just very quickly, on OTIB-0052 then, somehow my formatting is on 14 black in here, how many of the 16 findings are 15 16 closed now on OTIB-0052? MEMBER MUNN: One, two, three, four, 17 five, six - I count six closed. I count two 18 19 of them have been transferred which to all intents and purposes - no only one has been 20 It essentially closes it for transferred. 21

this particular procedure. There are one,

three, four, five, six that progress, which means they are actively being pursued. And there are - there is one which is in abeyance, which means the action has been agreed to by both the contractor and by the agency, but the action has not yet been completed, and for that reason it is still considered open, because the closing action has not transpired. But it's been agreed to.

Any other question? If not, then that concludes all I have to say with respect to procedures unless someone has some other issue they'd like to bring up.

CHAIR ZIEMER: The only other comment at this point that I would make, and I think Ted, you may have suggested this as well, that on some of the procedures which may have unusually significant impact on what this Board does, you may want to bring them forward separately for closure action rather than aside from the whole set of all the - I forget

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it's over many procedures, but 100 100. But if there are particular around procedures that we feel the Board may want to look at separately, and you would have to identify what those are. Is OTIB-0052 one of those, or if there any high impact are procedures, or in some cases for example, if it's procedure that has particular significance for a particular site. the case of Appendix BB we've moved that out so it's being looked at separately. But I'm just thinking in general terms if there are some that the Work Group identifies as needing special attention, why you might want generate a separate recommendation on those.

MEMBER MUNN: We can certainly do that, and be glad to in the future try to make note of those and make sure that at the very least they are mentioned specifically.

CHAIR ZIEMER: Yes, I think the Work

Group could do this as they proceed and say

this procedure is more than a minor

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| 1 | administrative detail. It has high impact or |
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| 2 | needs some visibility. |
| 3 | MEMBER MUNN: We do have several of |
| 4 | those. |
| 5 | CHAIR ZIEMER: And try do to that as |
| 6 | we proceed. |
| 7 | MEMBER MUNN: We do have several. |
| 8 | CHAIR ZIEMER: So it doesn't get |
| 9 | lost in the details of you know coming to the |
| 10 | Board saying here are 100 procedures that we |
| 11 | are recommending be closed out, it would be a |
| 12 | little like the dose reconstruction group |
| 13 | coming with 100 dose reconstructions and |
| 14 | asking us to approve them all at once, and |
| 15 | that would be difficult. |
| 16 | MEMBER MUNN: We will try to make |
| 17 | sure that those things are brought to your |
| 18 | attention. |
| 19 | And Stu, thank you very much for |
| 20 | getting that up on the screen for us. |
| 21 | MR. HINNEFELD: I'm just waiting for |
| 22 | the electrons to get in the right place. My |

computer is shut down.

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MEMBER MUNN: Well, we appreciate it. Thank you very much. I know it's tough when the process is slow, but I thought it was helpful to have them on the screen for people who might not be able to follow on their computer setup.

Thank you.

CHAIR ZIEMER: Thank you.

We had some space reserved for SEC discussions, and trying remember I'm to whether that for or not was reserved discussing further details on those petitions that we've acted on. Ted? Right.

Now we also have a 4:00 o'clock adjournment time, because we are going to reconvene a little later this evening. We do have some work time, mainly work time, tomorrow morning, so since we do need to have a break and then time for people to have dinner, that is, if your cookies and brownies are all digested by now, then perhaps we will

| 1 | go ahead and recess for this afternoon, and |
|----|--|
| 2 | then reconvene at 7:30 for the public comment |
| 3 | period. |
| 4 | Let me ask if there are any |
| 5 | additional housekeeping items we need to take |
| 6 | care of. |
| 7 | Apparently not. Board members, any |
| 8 | other issues right now? I'm sorry, is |
| 9 | somebody on the line. |
| 10 | So we will reconvene at 7:30 for |
| 11 | the public comment period. 7:30 for public |
| 12 | comment, Eastern Standard Time. |
| 13 | (Whereupon, the above-entitled matter went off |
| 14 | the record at 4:00 p.m. and |
| 15 | resumed at 7:30 p.m.) |

PUBLIC COMMENT

CHAIR ZIEMER: Good evening, everyone, and welcome to the Advisory Board on Radiation and Worker Health.

We have a number of individuals who wish to make public comment this evening, many of whom are not speaking in relation to the local facility, but who agreed to wait until tonight so that the Savannah River folks who were here yesterday would have an opportunity to speak.

And I also want to make sure that there are several folks online, or on the telephone lines, who we've also agreed would be able to address the group.

Let me first see - I want to see if one of our Board members is online. Gen Roessler? She may not be, because her [Identifying information], and she may have had to leave for that.

Terrie Barrie, are you on the line this evening?

| 1 | Okay, John Funk, are you on the |
|----|---|
| 2 | line? |
| 3 | MR. FUNK: Yes, sir. |
| 4 | CHAIR ZIEMER: Okay, standby. |
| 5 | And Dan McKeel, are you on the |
| 6 | line? |
| 7 | MR. McKEEL: Yes, sir. |
| 8 | CHAIR ZIEMER: Thank you, Dan. |
| 9 | Again, Terrie Barrie, are you with |
| 10 | us yet? |
| 11 | (Pause.) |
| 12 | Okay, we'll check back on that. |
| 13 | So let me begin then this evening - |
| 14 | MS. BARRIE: Dr. Ziemer? |
| 15 | CHAIR ZIEMER: Yes. |
| 16 | MS. BARRIE: This is Terrie Barrie. |
| 17 | CHAIR ZIEMER: Good, I was just |
| 18 | checking to see if you were there. Thank you. |
| 19 | And we will get to you and a couple of others |
| 20 | on the line in just a little bit, so please |
| 21 | bear with us. |
| 22 | Before we have the actual comments |

from the members of the public who are assembled here our Designated Federal Official, Ted Katz, will remind us of the redaction rules and related matters.

MR. KATZ: Right. Welcome, everybody, and I'll try to be quick about this, since I think everyone here has probably heard this. But as a requirement we have to repeat this before each public comment session.

So there is a verbatim transcript being made of this session. Ιf you give comments, you have the option of not giving although Ι think all the your name, listed at commentators least want to identified, so that is okay. But if you don't want to you can come up to the mike and speak without giving your name.

If you do give your name it will show up in the transcript. If you give personal information about yourself, such as medical information even, that ordinarily will

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also show up in the transcript. If you give information however about a third party, that information will be redacted; it will be removed from the transcript. It will not show up in the transcript.

And last but not least if there was someone here who wanted to address the Board in private, you could speak to me and we could try to arrange something like that.

Otherwise just to note this policy in all its legal language is laid out where you registered to speak here, and it's also on the NIOSH website with the agenda for this evening.

Thank you, Dr. Ziemer.

CHAIR ZIEMER: And those of you who are here in the assembly, I will call you in the order that you signed up. And since I have to call you by name, your name will appear in the record unless I learn very quickly from you that you do not wish to be so identified.

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But I will proceed with the list as
I have it. First we will hear from Denise
DeGarmo, and Denise will be addressing us in
relation to the Dow Madison facility I
believe.

Denise, thank you. And Denise also has a handout. Board members, I think you should have all received it now. And Denise will also supplement her remarks with some slides here. And I believe those are ready to go as well.

Thank you.

MS. DeGARMO: Thank you very much for allowing me to speak in front of you. a courtesy to the Board I wanted to inform you of some activities that I have taken on behalf of Dow Madison. And in October of 2008 I turned over new research materials Department of Energy, NIOSH and the to Department of Labor, and requested extension of the covered period for Dow Madison to include the years 1954 to extend to the year

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As you know we have several primary documents that establish Dow as an AWE site. Those include the January $4^{\rm th}$, 1956 agreement between Mallinckrodt Chemical Works and Dow Chemical Company for experimental extrusion work.

On the record we also have a FUSRAP document entitled, designation summary for the former Dow Chemical Company in Madison, Illinois.

Document number three consists of the DOE environmental management trip report, May 1988 visit to the Weldon Springs site in Weldon Spring, Missouri.

And the fourth document that established Dow Madison as an AWE site consists of FUSRAP, considered the sites database report on Dow Chemical Company.

In addition to that we have two purchase orders from 1957 and 1958. We have purchase order U-3067-L that was done in

12/05/1957 between Mallinckrodt Chemical Works and Dow Madison; and in addition to that we have a purchase order dated 3/15/1958, U-52990-A which is also another purchase agreement, purchase order agreement between Mallinckrodt and Dow Chemical.

We included - we were able to include thorium into dose reconstructions as of January 8th, 2008, with a letter to Peter Turcic, and I believe you have it on hand, so I did not provide it in an additional copy. But we after several months of additional research in the state of Michigan were able to come up with new evidence presented in the Dow Diamond, which is the Dow Midland or Dow companies corporate journal.

And some of the information we retrieved was pretty revealing. In September, 1954, and you can look at the highlights, they are not very good here, Dow Madison was the first facility built for the mass production of magnesium and magnesium alloys. The sole

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purpose of Dow Madison also was if you read further into this article for the purpose of production and development of magnesium thorium HK31A.

In February 1957 document, we have a secondary document establishing a contract with the Atomic Energy Commission for further development and production of HK31A.

So this relationship between Dow Madison and Mallinckrodt Chemical Works extends beyond just simply Mallinckrodt. We now have reference to a contract which puts Dow Madison into direct contact or in direct agreement with the Atomic Energy Commission for the production and development of this alloy.

In that document that you have in of you it says Dow awarded was contract from the Atomic Energy Commission in 1947 to develop the alloy HK31A. Subsequently company began program develop the a to magnesium thorium alloys in cooperation with

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the Atomic Energy Commission but at its own expense. In 1947 the AEC awarded Dow a contract.

The first rolling of HK31A sheet began in 1953 at Midland, and the following year it was moved to the newly completed Madison, Illinois facility.

The summer of 1963 references Madison as being at the heart of the metals production program for Dow Madison, and you have that in front of you so you can look at that.

Dow Madison was at the heart of the production of magnesium thorium alloys as Dow reported in the Dow Diamond data summer of 1963. Although the company headquarters is at Midland, Michigan, the Madison plant is the heart of the Dow Metal Products Company.

So we already have moved beyond the approved SEC of 1960 as we are beginning to illustrate that the work on HK31, and continuous rolling beyond that 1960 date has

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extended out.

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We have several patents that were given to Dow in regard to magnesium alloy. The first one is dated April 26th, 1960, 2934461, and it has to do with rolling of the magnesium alloy. And we have a copy of that. And I have not submitted that to you, but I am willing to with a copy of that should you want it.

We have another one in `62, again, moving us beyond the 1960 SEC date. This is patent #3039901, anneal from magnesium alloys.

We have another one in 1964 which had to do with the production of fibers, 3121943. And there it is.

And additionally we have found additional references to Dow Madison in the minerals yearbook. 1963 there In specific quote here, and I'm sorry I don't have copies of this, but I was working on this from the car ride down, so I can provide these to you well. The Dow Metal Products as

Company which is out of Madison, Illinois, increased their prices on selected items of alloys, HK31A and HM21A.

In 1964 at the top of the page there is a reference to the fact that the principal domestic producer of magnesium thorium alloys HK31 is again with Dow Madison.

Finally from the Department Energy we were given a document from Lawrence Livermore National Laboratory that links Dow HK31A to a nuclear weapon, and one of the questions that had been raised was whether or not HK31 or Dow Madison had any materials going into a nuclear weapon. And at the beginning of this document, which I have in here somewhere, the first paragraph says, inquiry about thorium regarding your thorium alloyed with magnesium material that were used in weapons parts, and whether Dow Chemical provided the material, I am enclosing the following table.

We have examined the drawings of

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weapon parts that contain thorium in our database. On the drawings the parts in this table specified a particular standard of metal that was produced by Dow Chemical as referenced in the Dow product guide above.

So we believe that we have now made a link between Dow Madison and the Atomic Energy Commission, that and extends relationship with Dow and Mallinckrodt beyond Mallinckrodt. Dow Madison had its own unique relationship with the Atomic Energy Committee that involved magnesium thorium alloy research and development, and in fact this alloy was contracted by the AEC. The thorium and its alloys were used in atomic weapons from 1962 through 1969 as shown in this chart, and given that the magnesium thorium alloys were used in the atomic weapons complex through 1969, and given the development of these alloys, were contracted by the Atomic Energy Commission, not to mention some of the problems with internal and external dosimetry, it appears

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we would have satisfied the statutory 1 2 requirements to extend the covered period to 1954 when Dow Madison began producing and 3 development HK31A through 1972, 4 when actually sold the Madison plant. 5 been contacted by We have the 6 7 Department of Labor who have told us that they have made a draft decision and we should be 8 receiving that in due time, but it is not 9 10 going to be ready for this particular meeting because of the contents of the response; there 11 were quite a few attachments. 12 So thank you very much for allowing 13 me to present this information. 14 15 CHAIR ZIEMER: Thank you very much, 16 Denise. Let me make sure now; you have provided all of your documents to Labor and 17 also to NIOSH and DOE as well? 18 19 MS. DeGARMO: Yes. I do have to now include the mineral yearbook. I need to send 20 off that the Department 21 to of Labor,

Department of Energy and NIOSH, as well as the

| 1 | additional two patents beyond what they have. |
|----|---|
| 2 | CHAIR ZIEMER: And your |
| 3 | understanding from Labor is they are underway |
| 4 | with material? |
| 5 | MS. DeGARMO: I was told they have |
| 6 | the draft decision. |
| 7 | CHAIR ZIEMER: Thank you very much. |
| 8 | MS. DeGARMO: Thank you. |
| 9 | CHAIR ZIEMER: Board members, do you |
| 10 | have any questions from Denise? Very |
| 11 | important development here. Yes, Brad. |
| 12 | MEMBER CLAWSON: I just had one |
| 13 | question. When you submitted this to Labor, I |
| 14 | guess are they going to involve us to let us |
| 15 | know? I know there is kind of a cross between |
| 16 | us and Labor in stuff. |
| 17 | CHAIR ZIEMER: They will certainly |
| 18 | let the NIOSH people - does DOE make any |
| 19 | determination first? I forget the exact |
| 20 | process here. But Larry can you enlighten us |
| 21 | as to what the process here on this is. I |

don't recall the exact sequence of events that

| 1 | this might trigger. |
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| 2 | MR. ELLIOTT: The Department of |
| 3 | Energy is responsible and given authority |
| 4 | under the act to come up with a list of |
| 5 | covered facilities, and they did so early on |
| 6 | in the program, posted that under Federal |
| 7 | Register notice and put it on their website. |
| 8 | The Department of Labor has the |
| 9 | responsibility for setting additional |
| 10 | timeframe limits around those. So this will |
| 11 | be a Department of Labor decision in that |
| 12 | regard, as I understand it, and DOE has |
| 13 | provided input to that DOL examination. |
| 14 | CHAIR ZIEMER: And then once a final |
| 15 | decision is made on that, then NIOSH will take |
| 16 | whatever appropriate action is needed. |
| 17 | Thank you very much. |
| 18 | Next we will hear from Donna Hand. |
| 19 | Welcome, Donna. |
| 20 | MS. HAND: Good morning. I'm Donna |
| 21 | Hand from Pinellas Plant. A worker advocacy. |
| 22 | Can you hear me now? Okay. |

Last night I was talking about a person that had some wounds. hit with classified waste. And then right here with his classified waste at Pinellas Plant, they said that he may have been exposed to photon, electron and neutron radiation. However in the dose reconstruction they will not use electrons, external electrons; they did not only had use neutrons. Не 100 milligrams of dose to him, and he also - it distribution, constant was at the uncertainty was never even ran on the PoC nor in the dose reconstruction.

And airline pilot 200 gets millirems a year, and you are telling me that that janitor person а he was decontaminated the areas, he got cut classified waste. Не was in а neutron generator area, and he only got 100 millirems a year.

These issues were brought up, and that the waste - they said then that the waste

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should have been monitored. The consisted of the treaty on contaminated papers as well as the foams and the metals, all during the trigger testing. They would explode the trigger to test them. And that's what was in 200. And all the waste would be picked up and put in a radioactive bag. that bag it was put into a drum. They'd seal and they'd take it the drum, out to storage area until it's ready to be shipped off to Savannah River.

In 1990 the Tiger Assessment Team came to Pinellas Plant, because in 1986 a report came out saying that they were not following procedures and policies. They were mixing their waste. They were putting radioactive waste into non-radioactive waste. They were putting non-radioactive waste into radioactive waste. And they weren't following any policy or procedures. They were just going ahead and doing processing before policy procedures were in.

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Back to the wound count. According to technical basis 11.5, down at the bottom they say that a wound is defined as any break in the skin. Any wound that occurred in a this is discussing work area and one plutonium - will have plutonium contamination, especially after the event. However, these should monitored, people have been therefore you use the monitoring site, and use

that monitor with an alpha detector, an alpha

detector is the method to use.

There was no alpha detector. There was nothing to detect the wounds. He went to the medical center; they gave him some iodine to put into it, and a Band-Aid. And according to the standards, unless you are going to be exposed, a potential exposure to 500 millrems, they don't put in cuts and wounds. But yet DOL and NIOSH both are requiring that if they be monitored, or they will not use those wounds into it. We have had several people and I'm sure other sites have this internal

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exposure, and it's not being accounted for.

back to the dose Now you go reconstruction; again, like I said, it was only 100 millirems. We have people that have all worked in the same area. They have all had skin cancer. Each one has a different type or distribution. Most of them it's constant, and it's supposed to be log normal because they have been unmonitored, and this is according to their own reports that they wrote in their articles, the 15 articles, is a normal distribution with а mean geometric standard deviation.

They put that its uncertainty, log normal is 1.1. However you run constant at .100, except for one gentleman. In one gentleman his skin cancer was triangular. Why the difference? Why did you change? These are the same workers, same area, same organ; they changed it.

We have an area that in Pinellas Plant that is classified. The workers called

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it Heather. That is in Building 300. In the early years Building 300 was detached, then it became attached. The technical basis document says that there is 300 there. But however there is no other reference to it at all.

In the baseline report that was done by DOE and Lockheed Martin, they have established and given out information that was not classified on 300. Tritium was there. How come the workers that were there in 300, that tritium dose is not attributed at all?

We developed the facts and figures that we got from OSTI, OpenNet, all about the policies and procedures. And the baseline Section back in K lists 28 report radionuclides. DOL and also NIOSH, because DOL health physicists will state that NIOSH is using everything correctly, they completely ignored 27 radionuclides. That was potential exposure to those workers, and they will not use any of those.

Depleted uranium is one of those.

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They had uranium beds. They made the beds.

They refurbished the beds, with the depleted uranium, and their own technical basis document they speak about depleted uranium.

But yet according to them uranium exposure was never to any worker.

get over 2000 BQ's just on that north end, the reason being the pond was given the tritium water. They aerated it, then they sprayed it across the field. That is also where they buried drums. USGA did a survey back in the contamination era. What is this metal thing? In the northwest corner there were drums buried there. They had to pull them up and ship them to Savannah River.

They took another test, and guess what? In the northeast corner there are also drums. They had to go dig them out as well. To this day there is a sign from DOE saying, hazardous area, do not enter.

I wrote to NIOSH to the director

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about the Pinellas Plant doses. I requested a rework of all Pinellas Plant doses. technical basis document in November of 2006 that increased the assigned dose, the increased assigned dose went to .550, so millirem. All the dose reconstructions before, they said, nope, we did it properly. In fact, she stated it was all done by proper methods, and regarding the neutrons we only monitor or we only acknowledge people that had significant exposures. This law says that the dose reconstruction is to be for any worker that may have potential exposure. You do not restrict.

In fact NIOSH in the very beginning of this program got in trouble with Congress because they were being restrictive. This is all doses are to be accounted for, and all potential doses. It doesn't mean if it's a You have got to acknowledge little or a lot. there, those doses were and they potential doses.

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The primary internal dose hazard at Pinellas Plant came from tritium and plutonium. They admit it from Dr. Branch, but however, plutonium is not even used. They just used tritium. their Pinellas On template, it's .930 for photons, it's .930 for tritium. Zero neutrons, zero everything else. And the REMS database in 1990, when you put Lockheed-Martin, it shows there in were neutron doses in those perimeters there.

In 1997, this is after everything has left, because it stopped production in '92, and in '97 the decontamination was completely finished. Again it showed neutron exposure there as well. How come they are not using neutron doses?

Alpha: they refused to use alpha. In Table 6.6 of their own technical basis document it shows there are alpha rays there. It shows Krypton 85, cesium 137, and americium 241, U-238, U-235, U-234, plutonium 238, at 80 percent, plutonium 239 at 20

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percent. This is their own documentation, their own technical basis document, their own information that they are completely ignoring.

list There is а in here equipment that produces this, as whenever they would test and everything they would X-ray that part. The industrial X-rays are not accounted for, and these people were exposed every time they took an X-ray when they tested that part. It is completely - the medical Xrays, unless it shows up in their medical record, they will not give them the medical occupational record.

the However nurse that they interviewed, and she was the one that took it, said I always took two views. It was done every year; two views. That happened all the way up to 1982. From then, guess what, upgrade in her X-ray machine, got an that's when they had exempted and non-exempted employees. And that's when they started the five year, three year, that combination. But

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still some of them had it every year.

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There are more issues. This will be followed up with a formal letter to the Board with all my concerns, and also the documentation.

Thank you for your time.

CHAIR ZIEMER: Thank you. I would just remind you, Donna, that we now have a Pinellas Work Group, so they certainly make note of these issues that you raise as we go forward with Pinellas. Thank you.

Then let's hear from Richard Lee.

MR. LEE: My name is Richard Lee as you just stated, and the only thing I can tell you is, I worked in the Savannah River site from 1980 to 1994. I started off construction as a pipe fitter and a welder, and wound up being management construction.

I can tell you the standard procedures that was done on a daily basis as far as operations goes. I was based out of

central shops. I worked all the shut downs in all the various areas. I had a Q clearance, I could go just about anywhere on the site. I didn't need to be escorted. I did a lot of escorting.

When we went into areas the only time we ever wore a film badge or had a pencil which was a dosimeter was what they called a hot spot or a hot area like a reboiler; then they'd put it on us. Other than that we went into -20, -40, all over the area, never required to wear anything but possibly white coveralls, as far as a worker goes.

When I first started out there from my concern I quested an HP, health physics, guys on a line break we were doing, and pretty well got chewed out and kind of threatened to run out of the area, so I learned real quick, you keep your mouth shut, keep your job, do your job, protect yourself the best you can within the guidelines that they give you.

But that was the norm for

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construction. I was also the gentleman that did the heat exchanges in 100C, the decon. They would assign us, or assign myself either an apprentice or another fitter, but I was the guy that was on it all the time. We nothing there other than plastic suits. We did all the hookup, the flushing. Unhooked it, bolted it back up, called the boilermakers to come to get it, that came from the Ford building and were returned to the building.

We did a line break one time in 100C. Half the time they didn't even know who was in the buildings. There was myself and another fitter. We were bolting up a six-inch flange, 300-pound flange, up in the rooftop. We climbed up there. We were putting the flange up, trying to pin it. We noticed operations had come in in plastic suits below us. Before we knew what was going on, they made a line break; an alarm went off; I got an uptake of tritium, got an uptake of nine,

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before they could get us out. We were never notified or told. My general foreman had to come in. Of course we were already - when the alarm went off we were already trying to get out and get down. But that was just neglect was all that was on their part.

When I went into the Bechtel side as management as a cost engineer, since I had the clearance that I had, I went into all of the areas; even escorted the other engineers for us to do walk downs. We would notify the project manager of the jobs that we needed to do, then they were supposed to set up everything. We'd go in the areas, and I can't remember not one time of ever wearing a film badge or a pencil in all the areas. Ι went in from 700 area in the weapons to all the 100 areas, to the canyons, F&H area, we went all over the site. And that was for a lot of years, and we were never required a wear a film badge. And I kind of thought it a little strange, but I knew in construction we

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| 1 | always had to wear coveralls. I'm kind of |
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| 2 | making a pun, now I'm in a lab coat and |
| 3 | booties, and I thought that was pretty good. |
| 4 | But that was the standard for the day. I mean |
| 5 | there just wasn't what everybody thinks there |
| 6 | was. There wasn't all this protection that |
| 7 | everybody thinks is there, and I was the one |
| 8 | physically on hand. All of us, all the |
| 9 | construction workers, we all know that's how |
| 10 | it operated. And I thought it was important |
| 11 | that you all know that in case you have a |
| 12 | misconception of actually how it went down, |
| 13 | and how it was on the site. |
| 14 | And that's basically all I have. |
| 15 | CHAIR ZIEMER: Thank you very much. |
| 16 | MR. LEE: Thank you. |
| 17 | CHAIR ZIEMER: Let's see, I have |
| 18 | Wayne Knox on the list. I think we heard from |
| 19 | Wayne yesterday. Yes, Wayne, you are back. I |
| 20 | will give you 10 more minutes. |
| 21 | MR. KNOX: One more time. |

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CHAIR ZIEMER: Okay.

1 MR. KNOX: We drove all the way back 2 to Atlanta just to talk to you one more time. CHAIR ZIEMER: Stay close to the mic 3 4 here. MR. KNOX: I'm a wanderer. 5 CHAIR ZIEMER: Well, 6 you can't 7 wander here; we won't have a record of what you say. 8 MR. KNOX: Very good. I will try to 9 10 stay put. Wayne 11 Му name is Knox, and Ι before. addressed you Ι military 12 was а 13 captain in the `60s in radiation physics. Ι major in the military assigned 14 was to 15 Eisenhower Hospital in Augusta, Georgia, 16 nuclear medicine science. And I spent many 15 supporting 17 years, years, the Nuclear Regulatory Commission on the regulatory 18 19 development and compliance side of the house, and doing and operating on inspection teams. 20 have evaluated over 50 percent of U.S. 21

nuclear power plants on behalf of the Nuclear

Regulatory Commission.

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I have even functioned on the DOE Tiger teams, and many special Nuclear Regulatory Commission teams.

I want to talk to you about the real world and this is what a number of people have been trying to tell you: the real world is different from what you think it is.

spent some time in the DOE's environment working up at Hanford. I've worked over here at Savannah River. And I support - of course I can't certify everything that they say, but I have observed many of these things. I am a protege of Dr. Karl Dr. Morgan was almost like a father Morgan. We spent a lot of time together after to me. school working in his little greenhouse and over at his house talking to him about health physics and what went on in the real world. He knew some real world stories. He even told me - and he was almost tearful when he told me about a black man that had an accident up at

Oak Ridge and they brought him in and injected plutonium into his veins. And people - now I can't prove that, but I suspect that knowing Dr. Morgan it was true. And some people up at Oak Ridge might indeed know that.

I say that to say, the real world is different from what you perceive it. I worked as an operational health physicists. We're the knuckle draggers of the group. We have to get the work done, in spite of all the elegant models, in spite of all the weird worded procedures, and even the regulatory requirements. We have to get the work done.

Today I want to address a single point, only a single one, and it concerns regulatory compliance.

My client, I like to call her my client but she is my friend, she is sitting back there. She is the claimant. Mrs. Beulah Lindsay drove back down here from Atlanta for me to address this single point with you.

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We - I reviewed all of her father's data and I found where there was a regulatory shall I say oversight at this point let's not call it a noncompliance; but it is a regulatory noncompliance. This is the letter that was sent to you from her lawyer.

And I quite frankly was expecting some type of reply from you. It was not a threatening letter; it just laid out the facts for your review.

Next she received this letter here, which did not even address the information that I provided to you, and at this point I would like to call it a purely regulatory oversight, because it is if I were looking at it from a Nuclear Regulatory Commission or a DOE perspective, it is not - it will be not in compliance with the regulations.

Basically what I discovered was that DOE, the DOL, directed NIOSH - I think that is the way the procedure flows - to use IREP in order to calculate the Probability of

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Causation of Mr. Lindsay's death from cancer.

NIOSH made a calculation and determined that the Probability of Causation was 32 percent. And it was due to cancer of the stomach that had metastasized to the lymph node, the liver and the lungs.

Based upon your regulations, your regulations, there is no process in which - or shall I say it's unlikely that cancer would move from the stomach to the lungs, that route is closed off, and as such, the lung cancer now becomes a secondary or primary, and based on your regulations when you have an unknown, an unknown primary, it requires you to go back and evaluate all of the possible primaries and select the greatest one of those, and include that in your calculation for the effective Probability of Causation.

That is now Mrs. Lindsay's father

PoC would be calculated based upon two

primaries instead of one. I can show you in

more detail what it's about. I passed around

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to you a little map on how that process works. Basically again based upon your regulation you should have looked at the stomach cancer it had lung cancer metastasis to What you should do is go lungs. to your secondary cancers, lung, and see if there is stomach cancer in that field. Stomach cancer is not mentioned as a place in which - that is the lung is not mentioned as a place in which stomach cancer would move. So therefore you should have considered two primary cancers: stomach and perhaps one of these others. that process is laid out in the procedures; in this description that it's laid out Ι provide to you. And I am requesting that the Board follow the regulatory requirements.

CHAIR ZIEMER: Thank you very much, Wayne. I can't help but making a couple of remarks, the first of which is to make sure that you understand that virtually everyone on this Board has had some considerable real world experience, including two of whom had

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that experience with K.Z. Morgan. So you have a common bond with some of the folks here.

MR. KNOX: Okay.

CHAIR ZIEMER: Number two, I think I should point out that the PoC calculation is actually done by Labor using the NIOSH recommended method. NIOSH does not do the PoC; they do the dose reconstruction, just so that is clear.

Thirdly I wanted to make sure you understand that this Board does not do the dose reconstructions, nor do we review them for claimants. The dose reconstructions are done by dose reconstructors. We do audit random samples of the dose reconstructions, but the important thing here I think for this information needs to feed back to Labor which does some essential determinations of the cancers and so on.

So I want to make sure it gets in the right stream, and there are folks here that can help make sure that the points you

| 1 | are making get to the right people who are |
|----|--|
| 2 | involved in this particular claim, because the |
| 3 | Board cannot under the law get involved in |
| 4 | this particular claim. |
| 5 | But perhaps, Jeff, I don't know if |
| 6 | you can help Mr. Knox or Dr. Knox get to the |
| 7 | right person. We just want to make sure your |
| 8 | information gets to the right place, so that |
| 9 | it can be considered as needed, okay. |
| 10 | MR. KNOX: I do not have a Ph.D. I |
| 11 | have a master's, so you may address me as |
| 12 | Master Knox. |
| 13 | CHAIR ZIEMER: Okay, Master Knox. |
| 14 | Very good. |
| 15 | MR. KNOX: Now who am I going to be |
| 16 | talking to? |
| 17 | CHAIR ZIEMER: Jeff is with the |
| 18 | Department of Labor, and at least make sure |
| 19 | that you get your information into the right |
| 20 | channel so that it can be duly considered. |
| 21 | Now I want to see if Terri Barrie |
| 22 | is still on the line. And Terri, we'd be |

| 1 | pleased to hear from you if you are with us? |
|----|---|
| 2 | MS. BARRIE: Yes, Dr. Ziemer, I'm |
| 3 | still here. |
| 4 | CHAIR ZIEMER: And thank you for |
| 5 | your patience. I know you wanted to speak |
| 6 | yesterday, and by the time we got to you, you |
| 7 | probably had given up on us. |
| 8 | MS. BARRIE: Oh, that's fine. I |
| 9 | wanted to make sure that the claimants from |
| 10 | the Savannah River site had plenty of time to |
| 11 | address the Board. |
| 12 | But I thank you for giving me the |
| 13 | opportunity tonight, and to members of the |
| 14 | Board and Mr. Katz. |
| 15 | What I want to voice my concerns |
| 16 | about is the Ruttenberg Database. And as you |
| 17 | know it contains exposure records for the |
| 18 | Rocky Flats workers. |
| 19 | And I apologize in advance to the |
| 20 | audience if some of these comments are not |
| 21 | self-explanatory. |
| 22 | I understand that NIOSH has finally |

agreed to the state of Colorado's conditions researchers views limit the the compensation program only. Margaret Ruttenberg thought that was the case months ago though, and during the working group teleconference of June 17th, 2008, thought that negotiations were well on their way, and six months ago she had hoped to have the database transferred to NIOSH.

I am trusting that this recent information that NIOSH has sent the agreement letter and everything is being processed promptly. But let's assume that NIOSH and Colorado have finally agreed, and the database is in NIOSH's hands. Who will compare the two databases? I think it's safe to say that the Rocky Flats claimants have no faith whatsoever that NIOSH can conduct this investigation in a fair and impartial manner.

Some of you, the Board members and the audience, are aware that a Rocky Mountain news article reported that a Rocky Flats

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worker was not listed in the NDRC, a project managed by ORAU, a contractor for NIOSH. DOL's requirement for automatic coverage under the Rocky Flats SEC is that a worker be listed in the NDRP or work in building 881. However the Ruttenberg database shows that this worker was exposed to neutron radiation, was not included in the NDRP, nor did he work in building 881.

Are there more workers who should be covered by the Rocky Flats SEC? What is more troublesome is the fact that NIOSH was aware of this database while the Board deliberated the Rocky Flats petition. SEC NIOSH the funding agency for was the Ruttenberg research.

I am appalled that this information was ignored by NIOSH and their contractor, ORAU, when they researched resources for the Rocky Flats site profile and evaluated Rocky Flats SEC petition.

Ignoring this information casts

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serious doubts on the thoroughness of NIOSH's investigation not only of the SEC petition but the dose reconstruction process for the Rocky claim in itself. Flats Is there other information that NIOSH and ORAU overlooked for the Rocky Flats site or for any other site for that matter? Will this database information for the years not included in the SEC that might show that NIOSH's methodology cannot produce a dose estimate with reasonable accuracy?

The Rocky Flats claimants won't know the answers to these questions until the two databases are thoroughly reviewed.

Again I think I can safely say that the Rocky Flats claimants demand an answer, and they demand it now. Contrary to what Dr. 17th, Brant said during the June teleconference, the Rocky Flats special exposure cohort is not a closed case before the Board, and that is the quote. There is nothing in the law that states that the

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Board's recommendation on an SEC petition is final.

There is nothing in the law that prevents the Board from reviewing evidence after the recommendation is submitted to the Secretary of Health and Human Services. There is nothing in the law that prevents the Board to say, hey, after reviewing additional information we amend our recommendation on such and such a petition.

Ι not advocate for another do prolonged and arduous series of meetings. fact I advise against it. The Rocky Flats claimants do not deserve that. However what they do deserve is that every agency follows the law. This process is meant to claimant-friendly. Ignoring evidence is not claimant friendly. Ignoring evidence is not sound science.

I urge the Board of the Rocky Flats working group, in conjunction with SC&A to compare both databases as soon as it is

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received by NIOSH. I understand that 1 2 should not be a long process. Then I also recommend that they 3 4 immediately report those findings to the Rocky of claimants and the members 5 Flats the Colorado Congressional delegation. 6 7 Again, thank you for your time. CHAIR ZIEMER: Thank you, Terry. 8 And I did want to ask, were you on the line 9 10 yesterday when we got the report about the status of that dataset? I believe it's very 11 close - I don't believe NIOSH has it yet as I 12 13 understand it, but they are very close to finalizing whatever agreement is necessary to 14 15 get that data, and then they will move ahead have the Work Group that will be 16 and we monitoring that activity as well. 17 Well, MS. BARRIE: thank 18 you, 19 doctor. CHAIR ZIEMER: I don't believe they 20 have the database yet as I understand it, but 21

they believe that they will be getting it very

| 1 | shortly. So we will keep you apprised of |
|----|---|
| 2 | course on that. |
| 3 | MS. BARRIE: Thank you. |
| 4 | CHAIR ZIEMER: So thank you for your |
| 5 | comments. |
| 6 | Next we will hear from John Funk. |
| 7 | John, are you still on the line? |
| 8 | CHAIR ZIEMER: And John is with the |
| 9 | Nevada test site. Please go ahead. |
| 10 | MR. FUNK: Okay, Dr. Ziemer. Before |
| 11 | I start out I'd like to refer to two |
| 12 | documents, they are reference site profile |
| 13 | documents. One is DOE/RV-317 (REV 1). The |
| 14 | second one is DOE NB/209 (REV 15), and it's |
| 15 | dated December, 2000. As you know that was a |
| 16 | list of all the tests by name and by date. |
| 17 | Okay I will start now. Good |
| 18 | afternoon, Dr. Ziemer, and ladies and |
| 19 | gentlemen of the Presidential Advisory Board |
| 20 | and Designated Federal Officer Mr. Ted Katz. |
| 21 | Thank you for allowing me this |
| 22 | opportunity to once again raise challenges to |

the NTS site profile, TBD documents, the dose reconstruction process. As you know from our you, originally had report sent to I somewhat lengthy PowerPoint presentation with supporting documents and visual aids charts and color like NIOSH uses. Ι have decided to spare you that report, concentrate on only two issues for the moment. However the other issues in the original report are just as important, and I would hope that the next NTS working Board meeting the Board would take the time to review my report in its entirety.

Before I start I would like to mention that the issues I am going to speak on were part of past issues I have raised, and NIOSH will claim that they have responded to these issues, but the fact of the matter is they have not, at least not in the same manner they require claimants to respond, and that is proving their statements with by proof positive documents, require as they the

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claimants to do. Which I now will do with supporting DOE reference documents.

With that I will now get into the issue and explain how the issues are not being properly addressed and expose serious flaws in the reference documents of the site profile TBD documents.

NIOSH using Issue #1, job classifications major factor in as а determining who was or was not exposed to radiation. I would like to mention I have brought up this subject of NIOSH using job classifications up once before on September 19th, 2006Board meeting at the Westin Hotel. And I recent also brought these up in the 180 issues that I sent to the working board.

For unexplained reasons that testimony fell on deaf ears, so I will try presenting this issue once again, hopefully with more success than the last time.

Please bear with me. I will explain how Technical Base Documents,

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reference records, may be wrongly influencing dose reconstructors who are doing dose at NIOSH. And I would like first to call your attention to two reference documents. The documents are DOE/RV-317 (REV 1) and glossary page - glossary section, page 274, under shaft and EG, capital E-G, underground description.

And especially DOE/NV-2009 REV-15 on pages one through pages 151 in tables of shots and date - shots and date and by name. Please note only one shot disseminates the difference between a mine shaft and a drilled shaft, and that is on page 58.

However, the same document does acknowledge there was a difference between mined and drilled shafts in other sections of DOE/NV-209 (REV 15) also in DOE/NV-209 - oh excuse me, I got that number twice. Please bring your attention to by date section page 58 and 59. Ajo is identified as a mined shaft. However on page 92 and 93 it is simply listed as a shaft only.

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And last look under glossary, same document, page 158, under shaft description,

difference between a mine and a drilled shaft.

Now please look in document DOE/NV-317 (REV 1) on page 274, the glossary under shaft, and other crafts - excuse me - is that a page messed up here? Oh, boy. Somehow I got my pages mixed up.

DOE very clearly disseminates, there was a

(Pause.)

I lost track of one of my pages. But essentially what it says, it says in there also, they acknowledge there was a difference between a shaft and a mine shaft and a drilled shaft.

IN the case of the mine shafts, the mines were the predominant labor force on them. In the case of the drilled shafts, the operating engineers, the carpenters, the laborers, the iron workers, carpenters helpers, electricians, they were the predominant crafts on the drilled shafts.

This causes a very big problem because it gives the illusion that only miners worked in all shafts based on language and the job classifications which are not even dated to the period of the testing period, 1962 to 1992. These job classifications go from > 93, `94 and `95. These miner job classifications are very explicit about shafts and tunnels without - are very explicit about shafts and tunnels without disseminating mined or drilled In other words it says they did them shafts. all, of which type of shaft they worked on, just shaft and tunnels and nothing else.

There is however defining no information about other crafts participation underground other than they worked in the tunnels shafts without defining their or did not discriminate duties. But also disseminate the types of shaft importance as well. And because of the way the reference documents for the site profile and TBD documents written, especially are

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documents I have noted, such information the way it is rendered clearly eliminate other crafts as their participation and reentry operations as work practices of other crafts are not disseminated so equally eloquent in their classification descriptions of work practices as miners' job classifications are noted, and such could severely minimize to their importance.

This is only one example of how flaws might happen when using existing reference documents to qualify applicants for exposure based on facts related to jobs classifications or work locations or duties of workers by craft found in the site profile, the TBD reference document, like doi classification and craft responsibilities and duties.

Existing reference to documents related to test information, when using a chart, please see miners' charts. I don't know if Ted got you a copy of that chart,

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CHAIR ZIEMER: Yes, we all have the chart that Ted distributed.

MR. FUNK: Okay, what I did on that, other letter Ι explained in the the mathematical formula I arrived at that. Ι gave the miners 32 mine shaft shots, which I don't even think they had that many. I gave them all the Plowshare. I gave them all the And I gave them all the unspent, along with the tunnels, and even then the whole list only came up to 141 shots of which they were the dominant craft.

And I took the total of the shots and subtracted that, and I percentaged it out, and it came out that the miners only dominated 17-1/3 percent of all the shots on the test site, and the rest of the crafts dominated the 82-2/3 percent of the rest of them.

So that does not - that is not the way NIOSH has showed this. Because I've heard the Board meetings. I've heard miners,

miners, miners, miners did this, miners did that. Nobody else seems to participate in any of the re-entries. They've got miners going to re-entries in the flats. They got them going up in the tunnels. And it seems like the rest of us were out there as tourists or something. I don't know, I got paid for working, and I did an awful lot of re-entries in the flats too.

So I don't know where they are coming up with these figures other than the fact that if you look at these documents, they are either tunnels or they are shafts, and there is only one that identifies it as a mine shaft. The rest of them are simply shafts.

Now there is a lot of difference between the type of shaft they were and the type of shafts they were would determine who worked on them. So that's what we need, to get that document back to DOE, and they need to get the shafts that were mined, identified, and they need to get the shafts that were

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drilled identified in order for people to get a fair consideration.

We also need to get in these job classifications and get these descriptions. For example nowhere does it state - in one place it does say it was a carpenter-welder, but not where he worked. Ιt just general foreman, carpenter, carpenter foreman, general foreman, carpenter welder. It doesn't say anything about in the tunnels if you read the miner-welders' description, it pretty much covers what the carpenters actually did. built the bulk heads. We cut all the steel. That wasn't done by iron workers. Them bulk heads concrete forms; under were the collective bargaining agreement we get all the concrete forms. There was never an iron worker or miner ever touched the bulk head in any of them tunnels, and there was 32 of them coming out of every drift. A drift is a test So that does - that's free shot work. drift. I understand that NIOSH is working toward

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that to eliminate all them people too. So I want to make sure to clear that up.

It took me two years just to get you to admit that we even worked underground. So and I think you will remember that.

Also issue #2, why has the Board done nothing about Mrs. Dorothy Clayton's testimony at the NIOSH Presidential Board meeting at the Westin Hotel September 19th, 2006, at Las Vegas, Nevada, page 176 to page 182 of this particular meeting.

And again when she testified on this exact same issue January $8^{\rm th}$, 2008 at the Presidential Advisory Board meeting held at the Sun Coast Hotel in Las Vegas, Nevada.

I would like to mention, I was recently told that Dorothy Clayton's testimony was in the investigative stage, and the results would be announced sometime soon. However NIOSH very clearly wrote me an email and told me, state in their response, this issue, this report sent to me, I quote:

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Dorothy Clayton's testimony is too complicated and cannot be read by a lay person, and the claimants do not understand the procedures being used. So therefore this subject has been addressed and her testimony will be noted in the records.

Now all you Board members were there during her testimony, and she provided you with the documents that showed they did cook the books up there. And I do not believe that her testimony was so complicated that we had to go get a translator to find out what she said. And when one of those politicians talk, we got to get the news media to tell us what they said, but Dorothy Clayton's testimony was very clear to me.

So at both meetings, Mrs. Clayton provided DOE supporting documents, which show positively that DOE tampered with film badge records, and this has not been properly addressed. And I would like to know where her testimony - and where the results of the

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investigation.

I would also like for those Board members who were not present at the Westin Hotel on September the 19th, 2006, to be provided a copy of Mrs. Dorothy Clayton's presentation and to revisit her testimony at the other Las Vegas meeting on January 2008 so they can see for themselves that widespread film badge tampering did in fact take place, and see for themselves that these film badge records are not as indisputable as NIOSH has stated.

These facts prove that NIOSH stand on documents of DOE and NTS records is fragile at best. NIOSH's position of the stand of the documents records is not so much only fragile but arrogant. In the recent announcement of the Inspector General's office suggesting removing NIOSH from the interview process of EEOICPA is long overdue, and heartily welcomed by NTS claimants as well.

Thank you very much.

| | CHAIR ZIEMER. Inank you, John. |
|----|---|
| 2 | I should point out that in the |
| 3 | packet of material, I think probably you |
| 4 | provided it, but the transcript is included |
| 5 | for Ms. Clayton's testimony, so the Board |
| 6 | members do have a copy of that in case any of |
| 7 | them had forgotten it. So thank you very |
| 8 | much. |
| 9 | MR. FUNK: I didn't know if it all |
| 10 | came through. I had an awful hard time trying |
| 11 | to get it through. |
| 12 | CHAIR ZIEMER: No, we have it, thank |
| 13 | you. |
| 14 | Next we will hear from Dr. McKeel. |
| 15 | Dan, are you still on the line? |
| 16 | DR. McKEEL: Yes, sir, I am. Can |
| 17 | you hear me all right? |
| 18 | CHAIR ZIEMER: Yes, very well. |
| 19 | Thank you for your patience, and please |
| 20 | proceed. |
| 21 | DR. McKEEL: All right. |
| 22 | Good evening to the Board, and the |

| 1 | audience participants. |
|----|--|
| 2 | I'm Dan McKeel, and I serve as an |
| 3 | SEC petitioner for three EEOICPA sites, Dow |
| 4 | Madison, General Steel Industries and Texas |
| 5 | City Chemicals. |
| 6 | My organization, the Southern |
| 7 | Illinois Nuclear Workers, or SINEW, has |
| 8 | proudly joined with the Alliance of Nuclear |
| 9 | Worker Advocate Group, in forwarding |
| 10 | suggestions to President-Elect Obama for |
| 11 | immediate administrative reforms to the act. |
| 12 | ANWAG will also soon be forwarding |
| 13 | copies of its 20 longer term reforms to the |
| 14 | act to President-Elect Obama, and to members |
| 15 | of the new U.S. Congress. |
| 16 | My remarks tonight are surrogates |
| 17 | for questions that would be asked by one of |
| 18 | the missing radiation Board members that the |
| 19 | Bush 43 - |
| 20 | (Pause.) |
| 21 | CHAIR ZIEMER: Did we lose Dr. |
| 22 | McKeel? |

| 1 | MEMBER ROESSLER: The line is still |
|----|--|
| 2 | open. I think he got dropped off. |
| 3 | CHAIR ZIEMER: Dan, I don't know if |
| 4 | you can hear us, but we can't hear you. |
| 5 | Okay, perhaps Dan will call back |
| 6 | in. Dan, are you back on the line? |
| 7 | MR. KATZ: He may not realize that |
| 8 | he is not connected. |
| 9 | CHAIR ZIEMER: Okay, and I don't |
| LO | think I have a phone number for him. |
| 11 | Standby just a minute. Of course |
| L2 | if he is on the phone you won't be able to |
| 13 | reach him anyway. |
| L4 | We have another piece of testimony |
| 15 | to read into the record. So let me ask Ted to |
| L6 | do that, and while we wait for Dan to realize |
| L7 | he's been cut off, and maybe will come back on |
| L8 | the line. |
| L9 | Go ahead, Ted. |
| 20 | MR. KATZ: Okay. Dan? Okay. So |
| 21 | this is unnamed testimony. |
| 22 | In April of 2009 it will be seven |

years since I have filed an EEOICPA claim for survivor benefits. When I filed my claim I did not expect a quick resolution, but I never anticipated that it would take this long, nor did I expect that I would encounter so many challenges along the way.

I consider my claim to be very straightforward. My husband was employed as a health physicist for the SL-1 reactor, by combustion engineering at Idaho National Engineering Laboratory from January 1959 through March 1961.

On the night of January 3rd, 1961, a nuclear excursion occurred at the reactor while a crew of three military men assembling the reactor control rod drive The incident, and the sequence of mechanism. events that occurred that evening and subsequent days are detailed in the IDO report on the nuclear incident at the SL-1 reactor, January 3rd, 1961 at the National Testing Station.

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This report can be accessed through the Department of Energy Idaho operations website.

My husband entered the reactor building twice the night of January 3rd, 1961, in an effort to locate and rescue the three men inside the reactor building. The first time he entered with one other individual in an effort to locate the three men inside and determine if they were alive.

After finding the one survivor they quickly left the building to get assistance to carry him out.

During the second entry my husband entered with three other men. They placed the sole survivor on a stretcher and carried him outside. During this entry my husband's respirator failed, and he had to remove it in order to breathe. This caused him to directly very high concentrations inhale the radioactive material airborne that present in the reactor operating room

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result of the nuclear excursion.

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The reported dose rate inside the reactor was in excess of 1,000 R per hour. Мy husband left INEL in March, 1961, was employed in Germantown, Maryland, bу the Atomic Energy Commission and its predecessor agencies, the Energy Research and Development Agency, and the Department of Energy, AEC, ERDA, DOE, from April 1961 through December, 1988. He was the chief health physicist for the AEC ERDA, DOE, and in this capacity was responsible for the oversight of the radiation health and safety at DOE and DOE contractor facilities.

In that capacity he routinely visited DOE sites and entered radiation areas to perform radiological safety inspections. My husband was diagnosed with multiple myeloma in August of 1994, and died from that cancer on January 30th, 1999.

Based on my knowledge and belief as a former health physicist that my husband's

cancer was caused by the exposure he received at the SL-1 reactor, I filed a claim for the EEOICPA survivor benefits on April 25th, 2002.

On November 13th, 2006, I received a notice of a recommended decision to deny my claim from the DOL Seattle District office. I filed a written object to the denial of my claim on January 7th, 2007, which included supporting documentation. On March 21st, 2007, the final adjudication branch of Jacksonville district DOL conducted a hearing in Orlando, Florida. Subsequently on May 30th, 2007, my claim was remanded back to NIOSH to husband's rework dose reconstruction my report.

Since the remand order I have received two revised dose reconstruction reports, one in July of 2007 and one in September 2008. NIOSH is currently working on a fourth.

I believe that the dose reconstruction reports that I have received to

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date significantly underestimate my husband's combined internal and external exposure.

Among the issues I have repeatedly raised are, my husband entered the SL-1 reactor operating room twice the evening of January 3rd, 1961, yet his report of exposure is nearly identical to individuals who only entered once. It is highly probable that the dosimeter husband was wearing mУ was not readable. IDO 19302 indicates that the extremities of the personnel who entered the reactor operating room were contaminated in excess of 5R per hour.

The probability exists that the film badge was too grossly contaminated to be read. Given the dose rates inside the reactor operating room, it is also very likely that the film badge was blackened to the point that no meaningful information could be obtained from the badge.

Through a Freedom of Information

Act request I obtained all of the occupational

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exposure histories that had been acquired by NIOSH for my husband. Among the external exposure histories provided by INEL, there is a personnel exposure questionnaire which was discussed in t.he dose reconstruction not report. This report indicates that the dosimeter readings for [identifying information] for the period of 12/7/60 through 1/3/61 were lost reading because of the SL-1 incident. That is in quotes.

This would strongly suggest that the dosimeter worn by [identifying information] into the reactor that evening was lost because it could not be read, or that it was literally lost in the confusion of the rescue operations.

husband's internal exposure estimated from cannot be the available records. The estimates in the dose reconstruction report are based only several bioassays that were conducted after the accident, records of the whole body counts

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1 taken at INEL, as well as at Los Alamos 2 National Laboratory immediately after the SL-1 excursion have not been located. 3 4 The dose reconstruction report fails to acknowledge the respirator failure, 5 and obvious internal exposures that would have 6 resulted from that failure. 7 And the dose reconstruction report 8 fails recognize my husband's 28-year 9 to 10 employment history with the AEC or ERDA, DOE. This report states that he worked at INEL 11 from January 1st, 1959 through March 31st, 1961, 12 and from April 24th, 1961, through December 13 16th, This affects 1988. the 14 many of assumptions made in the dose reconstruction 15 report with respect to, "missed dose". 16 The dose reconstruction report does 17 appear to address the "missed dose" 18 not 19 related to the SL-1 accident. my claim for filed survivor 20 benefits under EEOICPA nearly seven years ago. 21

period of time I

In

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that

have promptly

| 1 | responded to every request for information. I |
|----|--|
| 2 | have been engaged in numerous telephone |
| 3 | conversations with both DOL and NIOSH. I have |
| 4 | written letters, and I have participated in |
| 5 | the hearing concerning my claim. I have |
| 6 | provided ample documentation to support the |
| 7 | assertions in my claim. |
| 8 | I believe, however, that the |
| 9 | information that I provided has basically been |
| 10 | ignored. Additionally to date I have not |
| 11 | received a response to the numerous questions |
| 12 | and concerns that I have raised, particularly |
| 13 | those identified in my adjudication hearing. |
| 14 | Thank you for the opportunity to provide this |
| 15 | public comment. I regret that I could not |
| 16 | make them in person. I would welcome any |
| 17 | questions that you have. |
| 18 | CHAIR ZIEMER: Thank you for reading |
| 19 | that into the record, Ted. And I want to |
| 20 | check and see if Dan McKeel got back on line. |
| 21 | DR. McKEEL: Yes, sir. |

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CHAIR ZIEMER: Dan, we lost you very

| 1 | early in your presentation. And I think I am |
|----|--|
| 2 | going to suggest that you start at the top. I |
| 3 | don't know at what point you realized you were |
| 4 | cut off. |
| 5 | DR. McKEEL: Well, the line was |
| 6 | silent, and then it just went, a message from |
| 7 | the operator, if you need to make a call. So |
| 8 | it was cut off. |
| 9 | CHAIR ZIEMER: The others on the |
| 10 | line were not lost; it appeared to be a single |
| 11 | line. So please begin again if you would. |
| 12 | DR. McKEEL: Thank you very much. |
| 13 | So again, good evening. I am Dan |
| 14 | McKeel, and I serve as an SEC petitioner for |
| 15 | three EEOICPA sites, Dow Madison, General |
| 16 | Steel Industries, and Texas City Chemical. |
| 17 | My organization is the Southern |
| 18 | Illinois Nuclear Workers, or SINEW, and we |
| 19 | have proudly joined with the Alliance of |
| 20 | Nuclear Worker Advocate Group in forwarding |
| 21 | suggestions to President-Elect Obama for |
| | I and the second |

immediate administrative reform to the act.

ANWAG will soon be forwarding copies of its 20 longer-term reforms to the act to President-Elect Obama and to members of the new U.S. Congress.

My remarks tonight are surrogates for questions that would be asked by one of the missing Radiation Advisory Board Members, the Bush 43 administration has failed to add to bring the Board to its mandatory full strength of 20 members.

The first comment is, regarding the announcement of CDC's decision who the new Board contractor will be. Mr. Katz said at November 6th conference the call the decision would be announced before Thanksgiving. When it was not I wrote to him as to when the decision would be forthcoming and was told it would be at this meeting in December.

Yesterday the CDC contract officer told us the contractor decision was under policy review, and that he alone was

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responsible for this deadline not being met.

Since none of the current Board members pinned him down why not, I will ask as a surrogate for the missing Board member.

The question would thus be to CDC and the contract office, why specifically has the new Board contractor selection not been made public as the five-year SC&A contract ends today December the 17th, 2008?

My second surrogate missing Board member comment relates to why NIOSH and the Board took 3-1/2 years after the historic Mallinckrodt Destrehan Street SEC petitions were awarded to recognize that Mallinckrodt downtown radiological operation continued well into 1958. The current Mallinckrodt SEC 00133 evaluation report presented for the first time to the Board today in section 4.1, operations description, cites multiple previously known reports to justify this very belated discovery of a heretofore presumably obscure fact.

The cited Mallinckrodt report is

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dated 1994. On page nine of 19 of the 83.14 SEC 133 evaluation report dated 12/1/08 the specific citation is Mallinckrodt 1994, columbium-tantalum plant characterization plant, Mallinckrodt, Inc., St. Louis, Missouri plant; Mallinckrodt, January 10th, 1994, and that is the site research database reference ID 3840.

The missing Board member should have asked two more questions. One, why was this 1994 Mallinckrodt report not reviewed by NIOSH, the Board and SC&A and acted upon in 2005 when the original two MCW Destrehan Street SECs were being examined?

And the corollary question: When did NIOSH first obtain the 1994 report and share it with the Board?

The second question would be: why did it take 3-1/2 years to recognize the 1958 MCW downtown site operations involving the same type of pitchblende-derived raffinate operations that were carried out and discussed

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in detail when the Board wisely voted to overturn NIOSH's recommendation to deny the MCW Destrehan Street SEC-00112-2 that extended only through December 31st, 1957?

As a footnote, there appears to this surrogate missing Board member that from the NIOSH SEC 133 evaluation report, and from the SC&A TR task 10002 report, dated 1/31/2005, there were ample references to 1958 pitchblende raffinate operations at the MCW downtown site.

The references to the SC&A 2005 report include the full citation. Oak Ridge Associated University team, ORAU, technical basis document for the development of an exposure matrix for the Mallinckrodt Chemical Company, St. Louis downtown site, St. Louis, Missouri, period of operation from 1942 to 1958, O-R-A-U-T dot T-K-B-S dash zero zero zero five J-L Westbrook Rev. 00 24 October 2003 D. That's the full citation.

Note the final phrase, period of

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operation, 1942 to 1958. That is included in the SC&A January 31st, 2005 rev zero zero report citation is omitted by NIOSH and ORAU from Joe Guido's December the 1st, 2008 SEC 133 Rev 2 evaluation report citations on page 19 of 19. And that citation reads: ORAU T-K-B-F zero zero five, basis for development of an exposure matrix for the Mallinckrodt Chemical Company, St. Louis downtown site, and the St. Louis airport site, St. Louis, Missouri, rev two, Oak Ridge Associated University ORAU June 14th, 2007, and that is cite Research Database reference I-D 32277.

The missing Board member also would ask a third question, and that is, why would NIOSH and ORAU not cite the full O-R-A-U-T dash T-K-B 0005 report title from rev zero zero issued in 2005 in its rev 02 version issued in 2007?

The surrogate missing Board member would then ask a fourth and final question: Why did SC&A not recognize early in 2005 that

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radiological operations continued at MCW Destrehan Street in 1958 that should have led to an 83.14 SEC or to an extension of the SEC 00012.2 class, to include 1958?

It would seem that the SC&A task one zero zero two report dated 1/31/2005 with at least four allusions to 1958 operations, and urine bioassay samples being taken to include 1958 would have led logically to this conclusion.

listening After to today's I added proceedings, two short comments Board member might have made. First on the Chapman Valve SEC request to Bechtel, Larry Elliot mentioned that his office had three letters and made two phone calls without response. This is the ideal getting any situation to invoke section 73.84W of EEOICPA by asking the Department of Labor to subpoena those Bechtel records. I continue to wonder why this powerful tool is SO underutilized.

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Second, during the Mallinckrodt SEC discussion, NIOSH's Jim Neton was unable to say exactly how many claimants would be covered by the 1958 MCW SEC 00133 class. I believe this type of information should be a standard part of NIOSH SEC presentations. The data should be gotten before NIOSH presents to the Board.

My final comment is that I believe would be improper to redact from transcript of my public comment tonight the officials of the key government names responsible for us not knowing the name of the new Board contractor by today. This delay is causing obvious problems with Board functioning. Those people would be the CDC director, Dr. Julie Gerberding, Acting NIOSH Director Dr. Christine Branch, and the CDC contract officer, Mr. Karl Staudt.

Thank you very much.

CHAIR ZIEMER: Okay, thank you, Dan.

A number of questions for us to ponder. Some

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| 1 | of which we talked about earlier, because we |
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| 2 | asked ourselves how we overlooked certain |
| 3 | things, and it is sometimes difficult to go |
| 4 | back and sort of quarterback those things. |
| 5 | But thank you for your comments nonetheless. |
| 6 | DR. McKEEL: Yes, sir, thank you. |
| 7 | CHAIR ZIEMER: I had another one |
| 8 | here. I missed the second page. Dr. Jack |
| 9 | Bowcord, thank you. |
| 10 | DR. BOWCORD: I will be real brief. |
| 11 | I know this has run over. |
| 12 | Just sitting here tonight listening |
| 13 | to all these people, it sounds to me like |
| 14 | there is a huge dose reconstruction problem. |
| 15 | My Father worked at the Manhattan Project from |
| 16 | 1942 to 1946 and then he worked at Savannah |
| 17 | River for 27 years, and then they come back |
| 18 | and tell me he's got a dose reconstruction of |
| 19 | 11 percent? |
| 20 | Come on. He was all over that |
| 21 | plant, and he was an engineer. He wasn't an |
| 22 | office worker. It goes back what everybody |

said tonight. 1 has said here and 2 reconstruction is based on what you do and not where you work. And I think that is terrible. 3 office building 4 His at Savannah River plant was connected to a reactor that 5 was called Hector, and when that 6 was 7 demolished about eight years ago the office building was taken down as radiation exposed. 8 In 30 years of working for the government 9 10 never wore a dosimeter badge and never wore a film badge ever. When I was in dental school 11 and medical school, I kept asking him, dad, 12 13 why don't you have to wear a badge? You are head all the plant. the 14 over Не was 15 purchasing agent. He was in every division of 16 that plant. I said, we got to wear them at 17 dental school to take dental X-rays. 18 19 don't have to wear them and you are working next to a reactor? 20 think the dose reconstruction 21

plan is a joke is what I think.

1 Thank you. 2 CHAIR ZIEMER: Thank you. to give opportunity 3 want anyone else who may be here who wishes 4 speak but did not have an opportunity to do 5 6 so. Is there anyone else on the phone 7 line that wishes to make comment that didn't 8 have an opportunity to do so? 9 10 (No verbal response.) CHAIR ZIEMER: If not, I thank all 11 participated tonight. of who 12 We 13 appreciate your attendance here. I do want to remind you - oh, okay, I'll do this after we 14 15 are dismissed - want to remind you all that 16 the Board will be meeting again at tomorrow morning. You are welcome to join us. 17 Thank you all and good evening. 18 (Whereupon, the above-entitled matter 19 was concluded at 8:47 p.m.) 20 21

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