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ABRWH BOARD MEETING

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Worker Health held at the Millennium Hotel,

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June 24, 2008

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TRANSCRIPT LEGEND

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- -- (inaudible) / (unintelligible) signifies speaker failure, usually failure to use a microphone.

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PROCEEDINGS

(1:00 p.m.)

WELCOME AND OPENING COMMENTS

DR. PAUL ZIEMER, CHAIR

DR. CHRISTINE BRANCHE, DFO

1 DR. BRANCHE: If someone on the phone could 2 please let me know that you can hear me, I'd 3 appreciate it. Could you let me know? 4 UNIDENTIFIED: We hear you. 5 DR. BRANCHE: Thank you so much. I am formally 6 beginning the Advisory Board on Radiation and 7 Worker Health. This is meeting number 56. 8 are meeting in the lovely St. Louis, Missouri. 9 I'm Dr. Christine Branche and I have the great 10 honor to serve as your Designated Federal 11 Official, also known as Executive Secretary. 12 For those of you in the room, the emergency 13 exits are through this door to my left, and 14 then straight out. Unfortunately, to exit to 15 the street we would need to go up the 16 staircase. We have some persons with 17 disability in the room, which means that if the 18 alarm were to sound we would need some 19 assistance in helping the people with 20 disability -- disabilities out of the room.

1 don't anticipate any problems, but you ought to 2 know where the emergency exits are, should that 3 arise. 4 We do have a redaction policy. That policy is 5 that if -- let me read it formally. 6 If a person making a comment gives his or her 7 name, no attempt will be made to redact your 8 name from the meeting transcript. 9 that at a future date the meeting transcript 10 will be posted on the public web site. 11 NIOSH, the National Institute for Occupational 12 Safety and Health, will take reasonable steps 13 to ensure that everyone who makes a public 14 comment is aware of the fact that your comments 15 will be included. Your name and what you said 16 will appear in the transcript of the meeting 17 and it will be posted. 18 Including the reading of this statement to you 19 today and at the beginning of each public 20 comment period, that's our first attempt to let 21 you know what will happen. This statement also appears at the table, and was posted with the 22 23 agenda and the Federal Register announcement. 24 If an individual, in making a statement, 25 reveals personal information -- for example,

1 medical information -- about themselves, that 2 information will not usually be redacted. 3 NIOSH Freedom of Information Act coordinator 4 will, however, review such revelations in 5 accordance with the Freedom of Information Act 6 and the Federal Advisory Committee Act and, if 7 deemed appropriate, that information will be 8 redacted. And by redacted, I mean removed from 9 the record, blacked out. All disclosures of 10 information concerning third parties will be 11 redacted. 12 And if it comes to my attention that an 13 individual wishes to share information with the 14 Board, but objects to doing so in this public 15 forum, then I will work with you to be able to 16 get the information to the Board without 17 revealing your identity, but you would need to 18 come to me personally. 19 With that, I would announce the names of the 20 Board members for our roll call, and then we'll 21 get started. Dr. Ziemer? 22 DR. ZIEMER: Yes. 23 DR. BRANCHE: Ms. Beach? 24 MS. BEACH: Yes.

DR. BRANCHE: Mr. Clawson?

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1	MR. CLAWSON: Yes.
2	DR. BRANCHE: Mr. Gibson?
3	MR. GIBSON: Yes.
4	DR. BRANCHE: Mr. Griffon is on his way. Dr.
5	Lockey?
6	DR. LOCKEY: Yes.
7	DR. BRANCHE: Dr. Melius had an emergency
8	situation and will be unable to join us in
9	person. He will try to meet with us by phone.
10	Actually Dr. Melius, are you available by phone
11	today or at this time?
12	(No response)
13	Okay. Ms. Munn?
14	MS. MUNN: Here.
15	DR. BRANCHE: Mr. Presley?
16	MR. PRESLEY: Here.
17	DR. BRANCHE: Dr. Poston?
18	DR. POSTON: Here.
19	DR. BRANCHE: Dr. Roessler?
20	DR. ROESSLER: Here.
21	DR. BRANCHE: Mr. Schofield?
22	MR. SCHOFIELD: Here.
23	DR. BRANCHE: We appreciate everyone's
24	participation by phone today and through the
25	duration of this meeting, as well as during the

public comment periods. It is critical that all participants by phone mute their lines. If you do not have a mute button, then please use star-6 to mute your lines. You would then unmute, or use the star-6 to un-mute your lines when you are ready to speak.

If you must temporarily leave the line, please do not put the phone on hold. We would then all be subjected to whatever sound your hold button would have us go through.

Again, it is critical that every person participating by phone mutes their line. Even the slightest sound is picked up by the phone line and then interrupts the ability for all phone participants to hear what's going on in the meeting room. We very much appreciate your cooperation. Thank you so much.

Dr. Ziemer?

DR. ZIEMER: Thank you. It's my privilege to formally call to order the meeting of the Advisory Board on Radiation and Worker Health. We're pleased to be in St. Louis. We've been here several times before. And I want to go on record to tell you that the presence of Ted Drewe's frozen custard has nothing to do with

the fact that the Board chooses St. Louis to meet from time to time.

There are copies of the agenda and related written information on the tables in the back of the auditorium. Please make those available to you if you haven't already done so. Also, as Dr. Branche has indicated, we do ask you to register your attendance with us on the registration form or registration booklet that's out in the corridor.

Also we will have an opportunity for public comment later this afternoon. If you wish to make public comment, please sign up in the public comment book. If you wish to do that and haven't done so already, you can do that during the break. We like to get some idea of how many individuals will be commenting so we can plan for the time accordingly.

I believe that's all the housekeeping items that we have as we open our meeting today. We have a number of topics that we will be discussing, which involve a number of sites around the country, involving various aspects of the -- the compensation program that's operated by Department of Labor and by NIOSH

and Health and Human Services. And we're going to begin -- we'll follow our agenda pretty much as it's indicated. The time's always approximate. If we go over, then we will adjust accordingly; or if we finish something sooner, we will move ahead. So the times are taken to be approximate. We always have to estimate how much time things will take, and sometimes we do that pretty well and sometimes not so well.

Y-12 PLANT (OAK RIDGE, TN) SEC PETITION

In any event, we'll begin with the presentation on the Y-12 Plant from Oak Ridge. We have an SEC petition that will be reviewed by Stuart Hinnefeld from NIOSH, and then we'll have opportunity for the petitioners to comment as well.

MR. HINNEFELD: Thank you, Dr. Ziemer, members of the Board and members of the audience. This first petition evaluation report that I'm presenting pertains to the Y-12 Plant in Oak Ridge, Tennessee. I think it's familiar to everybody. We've done some other work on this site in other petitions.

A little background behind this petition -- I

just do two?

This petition was submitted by a petitioner for whom NIOSH determined we could not complete a sufficiently accurate dose reconstruction. The petition was submitted on September 20th. This is an 83.14 part of the SEC rule, meaning we identified the class and essentially recruited the petitioner to submit the petition. The petition was qualified for evaluation on September 24th, 2007 and we determined that we are unable to complete dose reconstruction with sufficient accuracy for this class of employees at the Y-12 Plant.

Yeah.

Some background behind how we got to this point. You'll recall in July of 2005 the Board recommended the addition of an SEC class from the Y-12 Plant from March 1943 to December 1947. That's the period we're talking about today. The class definition read "employees who worked" -- you know, in part. It was for "employees who worked in the uranium enrichment operations, or other radiological activities" during the specified period. At this time in the history of the program -- remember, this was pretty early on in the SEC process, and at

this time we had not adopted the routine practice of having the Department of Labor review our class definitions for administerability. And so we published this, believing that we had described essentially the people who could be exposed to radiation. And -- but there are a number of ways to interpret these words and so this definition did not really provide for a sufficiently clear path for interpretation --

DR. BRANCHE: Excuse me, Mr. Hinnefeld.

There's a participant by phone we need you to mute your li-- there's a participant by phone we need you to mute your line, please. If you do not have a mute button, then please use star-6. Thank you.

So sorry, Stu.

MR. HINNEFELD: That's okay. So the class definition that was selected didn't provide sufficient clarity or specificity to allow for everybody to agree on how the interpretation of class membership should be applied. So as a result, we saw decisions about membership in the class that we didn't understand, that we --we'd see people who were, in particular,

excluded from the class that we thought -- we didn't really feel like we could -- we could do much of a dose reconstruction for that person, either, so we didn't understand why they weren't included in the class.

And this led to a series of discussions between NIOSH and the Department of Labor about what to do about this situation. And Department of Labor said we're interpreting the words on the page; you know, you wrote the words, we're doing what we can. And so after a series of discussions we determined that the best course of action would be to initiate a new class, add an additional class through 83.14 process, to get the class definition defined more in line with what we expected it to be -- what our interpretation -- we expected our interpretation to be.

A little bit of reminder for the Board and maybe information for the audience for the radiological operations at the Y-12 Plant.

This '43 to '47 period at the Y-12 Plant was when the Y-- where they operated the Calutron devices for electromagnetic separation and enrichment of uranium. These Calutron

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operations involved -- this was -- you know, the primary mission of them -- they're the ones who made the enriched uranium for the enriched uranium bomb at the end of the war. Calutron operations involved production of the feed material -- in other words, you had to prepare the uranium into the proper chemical form and get it so it could be fed to the Calutrons -- conversion of the enriched uranium into the final product. I believe it was enriched with uranium chloride, and then once you had it enriched up to the enrichment you wanted, you didn't want it in the chloride form. I think they probably converted it into an oxide, which then could be made into metal. And then they also had to clean the -- and reclaim uranium from the Calutron internal components, which was quite a large part of the operation because the Calutron didn't deliver all the product possible right to the target collection area and there was material that contaminated the inside of the equipment. Now there were a number of other radiological operations that occurred at the Y-12 Plant, and we -- we see references and some -- and some

general description of that. But we don't really know exactly what buildings those operations occurred in. We don't have a lot of detail about source term and things like that. So after reviewing the information there, we determined that we were -- or we were unable to determine that -- if any specific group of employees was not potentially exposed. Since we didn't really know essentially the extent of the radiological operations, we couldn't really partition the workforce into exposed versus non-exposed.

For available monitoring data that might allow us to do dose reconstructions for internal exposures, internal monitoring data, we have now found a limited number of individual uranium bioassay results from 1944/1945. I don't believe those were available in 2005, I'm not 100 percent sure of that, but it's a pretty limited set of data anyway. It only covers a fairly short period of the operation. We concluded that the available data was too limited to support internal dose reconstruction, but if -- in the event that we had a claimant with their own individual data --

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- you know, one of the pieces of data we had pertained to a particular claimant -- we would attempt to utilize that in a -- in a partial dose reconstruction if it's a non-- a non-SEC cancer case where we have to do a -reconstruct what we can reconstruct. For external monitoring data we have not found any individual external monitoring results. In terms of workplace monitoring data, which we sometimes utilize to inform us about radiation exposures to people, we do have some direct radiation readings and qualitative summaries of those readings, but they're mainly for around the Calutron operations and they -- and a lot of them focus on the X-ray emissions from the Calutron rectifiers, a certain electrical component of the Calutron which emitted its own X-rays. And so -- and there were -- and we know that there were some actions taken to I believe install some shielded glass or something to -- or leaded glass -- in order to worry about those particular X-ray emissions, but we don't really have much information about other potential exposures or kinds of exposures.

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There was an airborne monitoring program that was begun in 1945. There's a -- maybe a few samples here and there, some other times. We have some results. We have some summary descriptions of results. None of these samples appear to be breathing zone samples, however. They seem to be more concerned about production loss type of sampling. You know, where's -where's material getting loose at as opposed to what are people being exposed to. And we've not found any radiological monitoring for any of the other activities -- radiological activities that were going on at Y-12 at the site. This monitoring program was just around the Calutron operation.

We've not obtained sufficient bioassay
information to support internal dose
reconstruction for this class. This is our -our determination of feasibility. The air
monitoring data is -- some air monitoring data
is available, but it's not known enough about
the samples. For instance, sampling strategy
frequency. We don't know if it was
representative of low, average or high exposure
or low, average or high production times, so we

don't feel like that data is sufficient to support dose reconstruction for this class.

We have no bioassay or air sampling results for other radiological operations, because they did do some things besides enriched uranium.

And NIOSH has not obtained any individual external monitoring data -- well, this should actually be in a -- under a bullet called "external dose".

But anyway, continuing on with feasibility of dose reconstruction, we have not obtained any individual external monitoring data during this class period, and we lack the source term information about the non-uranium radiological operations to build a source term model about what the external dose might have been.

We do believe we can reconstruct doses from medical X-rays based on some existing technical -- project technical documents.

The table that we generally provide with these, the summary of feasibility -- again, this is for March 1st, 1943 through December 31st, 1947 -- shows that we believe that, of the possible categories of exposure, we believe we can only reconstruct the medical -- the medical X-ray

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exposures. At least, that's with consistency. As I said before, if we would -- if we have a claimant and that claimant happens to be one of the people who submitted a few of those bio-you know, a bioassay sample or a few of the bioassay samples, we'll make some attempt to assign a dose based on that -- on that sampling. But certainly that would only be a -- we would believe only a portion of the -- of the dose a person might have received, even from that -- even from that mode. With respect to the health endangerment determination -- recall, for a SEC petition we have to -- we must determine first of all is dose reconstruction feasible; and if it's not feasible, then we are to opine on whether there was a health endangerment at the -- to the exposed workers. In -- we did not -- have not found any evidence of a discrete incident that could have resulted in extremely high doses similar to a criticality incident. Recall, this only goes up through 1947. And evidence indicates that workers in a class may have accumulated chronic radiation exposures that we are unable to estimate, so those -- those -- so

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-- and they could have occurred through intakes of radioactive material, or direct exposure to radioactive material. And so consequently we have concluded that the health may have been endangered for those workers covered by this evaluation who were employed for a number of work days aggregating at least 250 work days within the parameters established for this class, or in combination with work days within the parameters established for one or more other classes of employees in the SEC. I think everybody's familiar -- that's kind of our -our boilerplate language about aggregating classes. The rule provides us leeway to either say that the criteria for health endangerment is either presence or 250 days. In this case it looks like presence is not sufficient, so 250 days would be the criterion. The definition of the proposed class that we're proposing for this -- for this action or this evaluation is "employees of the Department of Energy, its predecessor agencies and DOE contractors and subcontractors who worked at the Y-12 Plant in Oak Ridge, Tennessee during the period from March 1st, 1943 through

1 December 31st, 1947 for a number of work days 2 aggregating at least 250 work days, occurring 3 either solely under this employment or in combination with work days within the parameters established for one or more other 5 6 classes of employees in the SEC." 7 Our recommendation to the Board is for the 8 period of March 1st, 1943 through December 9 31st, 1947 we find that radiation doses cannot be reconstructed for -- with sufficient 10 11 accuracy for compensation purposes. Therefore 12 -- here's our table of feasibility and health 13 endangerment findings, our recommendation, and 14 I believe that ends my presentation. DR. ZIEMER: 15 Thank you very much, Stu. For the 16 record, I would like to note that Mr. Presley, 17 Board member, is conflicted on Y-12 and 18 therefore has removed himself from the table 19 for these discussions. 20 And Dr. Poston as well, I'm sorry, also 21 conflicted on Y-12, so both of those Board 22 members have removed themselves from the table 23 for this discussion. 24 Let me open the questions, Stu, with this one. 25 Is my understanding that this new class does

not necessarily replace the previous classes?

MR. HINNEFELD: Well, it sort of -- it sort of subsumes --

DR. ZIEMER: It's intended to be a clarification, but let me ask it a different way. The previous classes were administered in a certain way by Department of Labor.

MR. HINNEFELD: Yes.

DR. ZIEMER: Will those -- will continue to exist -- on paper, at least -- as classes, or do you see this as replacing the previous action?

MR. HINNEFELD: Well, we've had a situation like this before where a later class essentially subsumes the earlier class. I think it happened at Los Alamos with the radioactive lanthanum work, and then the later -- the later addition of a larger Los Alamos class. We didn't -- I don't think we actually made any action to terminate the other class, it still is out there, but the -- I think any claim coming in now and -- would probably be under -- administered under this class since it's broader and anyone who'd be included under the other class would also be included --

1 DR. ZIEMER: Would still be covered by this. 2 MR. HINNEFELD: Yes, yes. 3 DR. ZIEMER: Okay. So in essence this covers 4 the previous actions and covers the issue of 5 concern that NIOSH had --6 MR. HINNEFELD: Correct. 7 DR. ZIEMER: Okay, thank you. Other questions, 8 Board members? 9 MR. GRIFFON: This might be more --10 DR. ZIEMER: Mark Griffon. 11 MR. GRIFFON: -- more to you, Paul. Did we --12 I mean we must have sent a letter regarding the '43 to '47 class, and do you have a -- a copy 13 14 of that? I was wondering (electronic interference) --15 16 DR. ZIEMER: I don't know what that -- we have 17 some noise on the line again. We ask if you're not speaking -- and you shouldn't be, by phone, 18 19 if you're on the line right now -- you should 20 mute your phone. If you do not have a mute 21 button, press star-6. 22 Now in reply to your question, Mark, the 23 wording of the recommendation to the Secretary 24 -- hold on just a moment and I have the letter 25 to the Secretary on Y-12, and with the speed of

1 cyberspace it will finally appear. I'll blame 2 it on the wireless network here, but it doesn't 3 have to be hooked to that. Okay, let me read -4 - I think you're asking me to read the previous 5 class definition? 6 MR. GRIFFON: I -- I quess. I'm assuming we --7 we adopted the same class definition as NIOSH -8 9 Here it is. DR. ZIEMER: 10 MR. GRIFFON: -- but --11 DR. ZIEMER: (Reading) The Board recommends a 12 Special Exposure Cohort status be accorded to 13 all DOE contractors or subcontractors or AWE 14 employees who worked in uranium enrichment 15 operations, or other radiological activities, 16 at the Y-12 facility in Oak Ridge, Tennessee 17 from March '43 through December '47 and who 18 were employed for a number of work days 19 aggregating at least 250 days, occurring either 20 solely under this employment or in combination 21 with work days of other employment occurring 22 with-- within the parameters -- dot, dot, dot -23 - so it's --24 MR. GRIFFON: Did we --

Yeah --

DR. ZIEMER:

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1 MR. GRIFFON: -- the -- the --2 DR. ZIEMER: -- yeah, so the --3 MR. GRIFFON: -- part I'm asking about -- I 4 quess I could pull this up to -- the part I'm 5 asking about is did we put any provision in 6 there to -- 'cause I notice that Stu had 7 forwarded this to the 250-day review workgroup, 8 so did we ask for some provision that -- I know 9 there's one line that we've been adding to some 10 of our recommendations saying that we will --11 will further evaluate whether less than 250-day 12 time frame is warranted -- you know what I'm 13 talking about? 14 DR. ZIEMER: Yes. I mean I don't know if we 15 MR. GRIFFON: 16 included it in that or not, or if it is 17 warranted, can't remember. 18 That is not addressed in this --DR. ZIEMER: 19 MR. GRIFFON: No, it's not. 20 DR. ZIEMER: -- petition. 21 DR. BRANCHE: But that previous decision was 22 from 2005. I think you've gotten a lot more 23 sophisticated with your language since --24 MR. GRIFFON: No, I don't --25 DR. BRANCHE: -- that was done.

1	MR. GRIFFON: think that's that's an
2	issue there, but I mean Stu, you did forward
3	this to the or maybe you forwarded it to the
4	SEC workgroup, maybe I'm confused. It was
5	MR. HINNEFELD: I don't think I sent it to the
6	
7	MR. GRIFFON: It was sent to
8	MR. HINNEFELD: 250-day
9	MR. GRIFFON: to Jim Melius's group, and he
10	has both of those, so I'm I just confused
11	that issue, but
12	MR. RUTHERFORD: Mark, it was sent to Dr.
13	Melius's SEC
14	MR. GRIFFON: SEC workgroup, right.
15	MR. RUTHERFORD: workgroup. We send all the
16	83.14
17	MR. GRIFFON: I gotcha. I was thinking it was
18	on the 250-day issue, but
19	DR. ZIEMER: No, that's simply because it was
20	an 83.14 petition.
21	So the description here is work employees who
22	worked in uranium enrichment operations, or
23	other radiological activities, that was the
24	MR. HINNEFELD: That was the original class
25	description, yes.

1 DR. ZIEMER: Yeah. Other comments or questions 2 for Stu? Now I'm going to ask -- oh, Jim 3 Lockey. 4 DR. LOCKEY: Stu, does this -- does this cover 5 everybody (electronic interference) at that 6 site? 7 MR. HINNEFELD: Yes. 8 DR. ZIEMER: Again we're getting background 9 noise by phone. If your --10 UNIDENTIFIED: Somebody's put us on hold. 11 DR. ZIEMER: -- if your phone is not -- if 12 somebody put us on hold, they're not there to -13 - to mute their phone. 14 MR. GRIFFON: I mean --15 DR. ZIEMER: Yes, Mr. Griffon? 16 MR. GRIFFON: -- I think we're -- overall, you 17 know, this -- this makes sense, the lan-- the 18 proposed language. But one thing in your 19 presentation, Stu, that I wondered about was 20 the -- you said you had -- since we had last 21 talked about this, you've found more urinalysis 22 data? MR. HINNEFELD: A little bit --23 24 MR. GRIFFON: You -- you still --25 MR. HINNEFELD: -- a little bit, found some.

1 MR. GRIFFON: -- didn't think that was... 2 MR. HINNEFELD: I don't think we had -- I'm not 3 4 MR. GRIFFON: When you say --MR. HINNEFELD: -- I'm working from memory 5 6 here. 7 MR. GRIFFON: -- a little bit, I think -- just 8 for consistency purposes, we have -- as Board 9 members, have to sort of understand what a 10 little bit means. I think at other sites I've 11 deemed something to be a little bit and --12 MR. HINNEFELD: Right. 13 MR. GRIFFON: -- dose reconstructions were 14 done, so --15 MR. HINNEFELD: Right, in this -- in this case, I don't know -- LaVon, do you remember the 16 17 number? I was thinking it was maybe 100 or 18 150. 19 MR. RUTHERFORD: Yeah, it was -- it was right 20 around 100, but I want to point out, too, this 21 -- the actual data was right at the end of the actual operational period of the Calutrons, and 22 23 it did not address that period where they went 24 through major cleanup and the dismantlement 25 period of '45, '46 and '47. So it was one --

1 like one run period of -- that they had like 2 100 urine samples in. 3 MR. GRIFFON: But at the end of -- like 1947 it 4 was done? 5 MR. RUTHERFORD: No. 6 MR. GRIFFON: No --7 MR. RUTHERFORD: No, 1945 -- '44/'45. 8 MR. GRIFFON: Oh, it was done before the -- all 9 the cleanup and --10 MR. RUTHERFORD: Right, right, right. 11 MR. GRIFFON: -- other -- okay, all right. 12 DR. ZIEMER: Okay? Other questions? Larry 13 Elliott. 14 MR. ELLIOTT: Just for the Board's 15 understanding I'd like to point out that this 16 issue was really brought to everyone's 17 attention by Denise Brock, who dealt with a 18 series of claimants who were having trouble 19 getting their claims determined for eligibility 20 within this class. And I think our 21 understanding is there's probably 24 or less 22 claims that are so affected by this action, but 23 those 24 certainly deserve all the attention 24 that Denise has brought to this, so I want to thank her for that.

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1 DR. ZIEMER: Thank you, very appropriate. 2 There may be one of the petitioners on the 3 line. Let me ask if -- if a Y-12 petitioner is 4 on the line, and if so, do -- does that person 5 wish to speak? 6 (No response) 7 I had an indication they may or may not be on 8 the line. Apparently not. Thank you. 9 DR. BRANCHE: Make certain you've un-muted your 10 phone. We've given you so many admonishments 11 to mute your phone --12 DR. ZIEMER: Yeah, if you're trying to speak 13 and not getting a response, maybe you are still 14 muted, I guess. 15 (No responses) 16 Okay. 17 DR. BRANCHE: Before we go on, phone 18 participants, I do ask again that if you -- two 19 things. You've heard Dr. Ziemer and me both 20 indicate that we need you to mute your line. 21 would also ask that you resist every notion to 22 put us on hold, begin then the phone line is 23 then obscured by whatever sound your hold 24 system has and so we would have to cut your li-

- we have to go through a lot of labor to cut

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1	you off if you do that. So please, if you need
2	to leave the line, it's better to hang up and
3	dial back in than to put us on hold. Thank you
4	so much for your cooperation.
5	DR. ZIEMER: Thank you. Any other questions or
6	comments for Mr. Hinnefeld?
7	If not, Stu, we thank you very much.
8	DR. BRANCHE: Do we need to read this
9	DR. ZIEMER: Board members
10	DR. BRANCHE: What about this
11	DR. ZIEMER: Okay, we we have received a
12	I believe it was a FAX
13	DR. BRANCHE: No, overnight mail.
14	DR. ZIEMER: Well, whatever yeah, FedEx mail
15	from a petitioner, and this is this
16	copies have been distributed, I understand, or
17	have
18	DR. BRANCHE: Only to the no, only to the
19	DR. ZIEMER: Okay.
20	DR. BRANCHE: court reporter. If this
21	it's fairly complicated. I don't think she's -
22	- the person is not disputing the idea of the
23	class. I think that there was just additional
24	information that the petitioner wanted to bring
25	to light. And I think that the class

1 definition takes care of this --2 DR. ZIEMER: I -- I believe that's --3 DR. BRANCHE: -- person's issues. DR. ZIEMER: -- the case, and this is a very 5 detailed description of an individual case. don't think it would be appropriate for us to 6 7 read this into the --8 DR. BRANCHE: I agree. 9 DR. ZIEMER: -- public record because we would 10 have to redact all of the personal information. 11 So let's simply make copies of this available 12 to the Board members -- includes some pictures 13 and so on of a particular case. 14 understanding, this case would then be covered, 15 anyway, by --DR. BRANCHE: It would. 16 17 DR. ZIEMER: -- by this action, should the 18 Board recommend it and should the Secretary of 19 Health and Human Services so take the 20 recommendation. 21 Board members, it would be in order to have a 22 motion concerning this recommendation from 23 NIOSH. Let me advise you -- and the Chair's 24 willing to hear the motion, if you're so 25 inclined, in simple form rather than in the

detailed language of the -- a memo that would go to the Secretary of Health and Human Services. But if -- if we have such a motion and it passes, then during our work session on Thursday we will provide you with the formal wording of what the action would be as it proceeded to the Secretary.

Ms. Munn, do you wish to make a motion?

Ms. MUNN: Yes, I do. I'd like to move that we accept the recommendation that the SEC be accepted as proposed by NIOSH, and that we subsequently make that recommendation available to the Secretary.

DR. ZIEMER: Okay, you've heard the motion; is there a second?

MR. CLAWSON: Seconded.

DR. ZIEMER: Motion is made and seconded. Is there any discussion on the motion? If you vote for the motion you are voting to recommend to the Secretary that this class of workers be added to the Special Exposure Cohort at Y-12. I add parenthetically it's our understanding from Mr. Hinnefeld that this would become, in effect, the working definition then to, in essence, replace the earlier designations of

1	the classes at Y-12.
2	We'll take a roll call vote, and I should also
3	tell you that the Chair and the Designated
4	Federal Official, under the rules of this
5	Board, will also obtain the vote of Dr. Melius
6	if he's not on the line now. Under our rules
7	we are required to obtain the votes of members
8	who are not present. Mr. Presley and Dr.
9	Poston will be abstaining from voting, so let's
10	proceed with the roll call. Please answer
11	"yes" if you favor the motion or "no" if you're
12	opposed. You may also abstain.
13	DR. BRANCHE: Ms. Beach?
14	MS. BEACH: Yes.
15	DR. BRANCHE: Mr. Clawson?
16	MR. CLAWSON: Yes.
17	DR. BRANCHE: Mr. Gibson?
18	MR. GIBSON: Yes.
19	DR. BRANCHE: Mr. Griffon?
20	MR. GRIFFON: Yes.
21	DR. BRANCHE: Dr. Lockey?
22	DR. LOCKEY: Yes.
23	DR. BRANCHE: Dr. Melius, are you on the line?
24	(No response)
25	Ms. Munn?

1	MS. MUNN: Yes.
2	DR. BRANCHE: Dr. Roessler?
3	DR. ROESSLER: Yes.
4	DR. BRANCHE: Mr. Schofield?
5	MR. SCHOFIELD: Yes.
6	DR. BRANCHE: Dr. Ziemer?
7	DR. ZIEMER: Yes. The motion carries. We will
8	proceed to prepare the formal wording for the
9	Board's perusal later in the meeting. Thank
10	you very much.
11	DR. BRANCHE: There's a participant by phone
12	who would need to mute their line, please.
13	UNIDENTIFIED: Yes, I was cut off my previous
14	phone so I had to call back in.
15	DR. BRANCHE: Then we ask that you not put the
16	phone on hold, please, when you have to leave
17	the line.
18	UNIDENTIFIED: Okay.
19	DR. BRANCHE: Thank you so much. If you do not
20	have a mute button, then if you could please
21	use star-6 to mute your line, we would very
22	much appreciate that.
23	NEVADA TEST SITE WORK GROUP SUMMARY
24	DR. ZIEMER: Okay, let's move on to the next
25	item. We'll have Mr. Presley and Dr. Poston

1 rejoin us, and the next item on our agenda is a 2 report from the Nevada Test Site workgroup, Mr. 3 Presley, chair. MR. PRESLEY: Well, we had hoped to come to 4 5 this meeting and present a pretty set of slides 6 that said that the working group accepts the site profile, as is. But we had a couple of 7 8 issues come to light in a meeting we had 9 yesterday morning, and that's not going to 10 happen. We have asked that SC&A and NIOSH go 11 back and look at these issues that have come to 12 They are These are not new issues. 13 issues that have been discussed in the past, 14 but some people felt like that there was a 15 little bit of difference there so they are 16 going to be looked at, scrutinized and 17 discussed. A recommendation is going to come 18 back to the working group, and hopefully down 19 the road we can make a recommendation on this 20 site -- the NTS site profile. 21 Anybody have any questions? 22 DR. ZIEMER: Okay --23 MR. PRESLEY: Thank you. 24 DR. ZIEMER: -- thank you, Mr. Presley. Board

members, any questions? How -- how many issues

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1 were -- are we talking about here? Is it one 2 or two, or ten or 20? 3 MR. PRESLEY: No, we're talking about two or 4 three. 5 DR. ZIEMER: Two or three issues that will be, hopefully, brought to closure by the next 6 7 meeting --8 MR. PRESLEY: I hope. 9 DR. ZIEMER: -- or by our phone meeting. 10 you. 11 NIOSH PROGRAM UPDATE 12 Well, we're not ready for a break yet. I think we -- we're going to move ahead, Larry, if it's 13 14 agreeable. Mr. Elliott will bring us the program update on the NIOSH program. 15 16 (Pause) 17 MR. GRIFFON: Paul, I -- I did want to note, I 18 -- I have recused -- for Nevada Test Site, it 19 was a shorter presentation than I was 20 envisioning, but for the record, I don't know 21 if you --22 DR. BRANCHE: It was just a -- it was just an 23 update. 24 DR. ZIEMER: Well --25 MR. GRIFFON: It was just an update. I don't

1 know if I had to or (unintelligible) --2 DR. ZIEMER: No, we had no action before us --3 MR. GRIFFON: Okay, right, right, right --DR. ZIEMER: -- so you're okay, yeah. 5 MR. GRIFFON: Okay. 6 DR. ZIEMER: Okay, Larry, proceed. 7 MR. ELLIOTT: Thank you, Mr. Chairman, and 8 members of the Board and members of the public. 9 I certainly appreciate the opportunity to --10 again to bring you a program update on where 11 things stand with regard to NIOSH 12 responsibilities under this compensation 13 program. 14 To date, as of June 16th, NIOSH has received 15 27,367 cases from the Department of Labor for 16 dose reconstruction. We have completed 74 17 percent of those cases and returned 20,089 to 18 the Department of Labor. We break those down 19 into cases that have been submitted with a dose 20 reconstruction report, and that represents 21 There have been 724 cases that have 17,630. 22 been pulled from dose reconstruction by the 23 Department of Labor, and this happens for 24 various reasons, primarily the ineligibility of

There have been 1,735 cases that

the claim.

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are currently pulled from dose reconstruction for SEC class determination and eligibility. The 25 percent of cases that remain at NIOSH represent 6,898, and of those we have completed 789 dose reconstruction reports and those are in the hands of the claimants awaiting return to us indicating that they have no further information to provide. So that's another -- of this 6,898, that's 11 percent that we feel we have finished our work on, awaiting the concurrence of the claimant.

There have been another one percent, or 380 cases, that have been administratively closed in dose reconstruction. And what this is is the fact that we have not received a indication from the claimant that they have no further information to provide and will allow us to move the case back to Department of Labor for a decision. So we're awaiting the return of what we call the OCAS-1, or a form that indicates they have no further information to provide. In this pie chart you can see graphically, I hope, the distribution of these cases that have been completed, pulled for eligibility determination or pulled for SEC class

1 determination, those that have been 2 administratively closed, as well as the active 3 cases. And of those -- also those that are pended. When we say pended, there are a 5 variety of reasons that we would put a case on 6 hold at NIOSH. That primarily results from 7 issues that regard technical approach or a 8 particular site profile that is being held in 9 review and we don't want to proceed on 10 completing those dose reconstructions until we 11 have that particular technical issue resolved, 12 and so we would pend those cases until we see 13 that resolution occur. 14 Of the 17,630 dose reconstructions that we've 15 returned to Department of Labor for 16 adjudication or for a recommended decision, 34 17 percent, or 5,959, have had a probability of 18 causation of greater than 50 percent. 19 leaves 66 percent, or 11,671 cases, that were 20 deemed to have a probability of causation less 21 than 50 percent and found to be non-22 compensable. 23 Just for a point of reference, the early --24 start of this program there were projections 25 made within the government by different

agencies or different entities within the government that dose reconstructed cases would be less than ten percent compensable, so we have considerably moved that bar, as you see in this slide.

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This particular graphic, this bar chart, gives you a sense of the distribution of probability of causation across those claims that we have returned to Department of Labor. It's broken out into deciles or zero to ten percent and on up until you get to the 49 percent bar, and then you see that -- those that are greater than 50 percent on the far right bar. Again, of the 6,898 cases that are still at NIOSH, 2,997 are currently assigned to a health physicist. They're in some state of process of dose reconstruction, with a goal to achieve finality in that part of the process. mentioned this earlier, 789 initial draft dose reconstructions have been provided to the claimants and we're awaiting the return of the OCAS-1; 3,112 cases are not assigned to a health physicist for dose reconstruction. These are probably the ones that are pended, as

well as new ones that have arrived and we're

working to develop what background we need to continue with a dose reconstruction on those claims.

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Of particular note, we now have 4,396 cases, 64 percent of those that we have in an active status, that are older than one year. track that very cautiously and carefully with a lot of attention. If we look at the oldest cases that we have, the first 5,000 that we have received, we continue to monitor our progress on those, and you'll see this broken down -- I think I'll just go to the bottom line here. The most important numbers are shown to We've had 794 of these first you in red. 5,000 cases come back to us, and this is for a variety of rework purposes under our Program Evaluation Reviews, or because the eligibility of the claim or the demographics of the claim changed, which requires us to rework that given claim. The bottom line here is that 37 claims are still awaiting a dose reconstruction. earlier number, 794, has had a dose reconstruction but they've come back. These 37 have never had a dose reconstruction, and we pay particular attention to those because we

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want to get those people, those claimants, a decision as soon as we possibly can. Of these 37 cases -- I'll break them down for you. There are -- they represent multiple individual There are several that are reinstated sites. Y-12 SEC cases, those that you've -- that class that you just took action on are represented in some of these. There are also cases from NUMEC that have come back to us. NUMEC is another class that you've added recently, but we're seeing those come back as not eligible. then we've had -- or NUMEC is -- is a class that was added, but we've not completed our partial dose reconstructions for these -- these particular cases in this 37 that are represented by NUMEC. There's also some Kellex claims here; MIT, which is another class you've added but we are doing partial dose reconstructions on MIT so they're represented here; and Norton. Some are active in this 37 and some are pended -- pended awaiting either a technical approach that we need in order to complete the dose reconstruction or pended because Department of Labor has some action that we're awaiting them to take.

In this line graph you'll see in blue the cases that have been received from the Department of Labor over the history of this program. In green you see the draft dose reconstruction reports that have been provided to claimants, and in red you'll see the final dose reconstruction reports that have been provided to Department of Labor. This is broken out in quarters, and I think we've finally got this abscissa correct, Dr. Poston, so thank you for that correction from last meeting.

In this bar chart we show you the cases that have been completed, by NIOSH tracking number in 1,000 increments. And we break those 1,000 increments down into those that have been completed, by the color blue; those that have been pulled, in the color red; cases that are active are in -- I guess it's mustard; and SEC cases are in light green; cases that are pended are in yellow; and then the admin closed cases are in purple, or lavender.

I mentioned reworks earlier in this presentation, and this slide shows you in trend -- a trend analysis, if you will, the number of reworks that we have been -- we have received

I'd point out for you this series of spikes
later on, starting in the third quarter of
2007. As you know, and I'll talk about in a
moment, we have a number of Program Evaluation
Reviews that were initiated and these red
spikes that you see from third quarter of 2007
up to second quarter of 2008 are a result of
those Program Evaluation Review reworks. I
point out that we have received a total of
7,977 and we've returned 4,583 to the
Department of Labor.

As you know, we -- our first step in -- once we receive a claim from the Department of Labor is we turn to the Department of Energy and seek exposure monitoring information relevant to that claim so that we can proceed with dose reconstruction. We monitor the progress of Department of Energy on their provision of this important information, and we track it. Every 30 days we pulse them and find out where they're at on a given set of requests, and here you see that there are 365 requests that are outstanding, and 96 of those are outstanding greater than 60 days.

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In the procedures working group this morning we talked a little bit about the site profiles for Atomic Weapons Employers that worked with specific types of radionuclides, and in this one, TBD-6000, Technical Basis Document 6000, there are a number of site-specific appendices that were required to be completed. appendices speak about unique exposure scenarios that are not typical to either uranium or thorium, and in this case metals that were worked with, and so we had to come up with a technical approach that spoke to these kinds of unique exposure scenarios. Fifteen of these site-specific appendices have been There are none in review at this completed. time, and one is in -- in development, but that may be -- that may -- appendices may go away. We may find ourselves recommending an 83.14 for that particular site because we've not been able to find any information for that site, so we'll keep you posted as we proceed on that point.

The other Technical Basis Document for Atomic Weapons Employers that dealt with refining uranium and thorium is TBD, or Technical Basis

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Document, 6001. And in the same situation here, we've found some exposure scenarios that needed to be addressed under an appendices, and we have completed six of those appendices and we don't envision any more that will need to be worked up.

Now the Program Evaluation Reports or Reviews. We have had 32 Program Evaluation Reviews that have been issued. You can find these on our web site. These different changes that have been made to our dose reconstruction approaches have resulted from our -- our own efforts to identify a better way to do things, as well as efforts of the -- of the Advisory Board and its contractor in finding issues and resolving issues with us to lead to a better dose reconstruction approach. So you see here 14,217 claims that have been affected across these 32 Program Evaluation Reviews. note for you, however, that that's not -- many of these claims may be duplicated. In other words, a claim may find itself affected by more than one of these PERs, so you can't just rely on the 14,217 and say my goodness, that's a lot. It is a lot, but it's not that total

number. They're counted twice, maybe three times, in this number.

So what has happened here, we -- once we rework a claim against these Program Evaluation Reviews, we see whether or not -- and we do this because there's a potential chance that the dose might increase for a given claim, and so we look at a lot of these claims and we're -- we're thankful when we see one that does increase in dose, and we're very thankful when we see one that crosses the compensation bar and goes to a 50 percent or greater probability of causation. Here you see 249 have switched from non-compensable to compensable.

Now for background, you heard me in -- in the last several meetings I have reported to you that there were 154 that had switched, so now we see an increase here of 92 claims that have been found to be compensable under our Program Evaluation Review. And of those 92, 77 are due to super S; five are due to our Paducah Program Evaluation Review; one is for our LANL Program Evaluation Review; eight are on Blockson and one is on Rocky Flats. Again, I would caution you about the super S because there may be more

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Rocky Flats in that than the one I'm talking about for Rocky Flats. So the numbers are what they are, but they affect claims differently, so just a caution.

That's the -- that's the good news. news is we've done 7,943 in an effort to try to determine whether or not they would be so affected, and there was no change in the compensability of those claims. We have still 6,025 that we are working on as we speak. again, that number may represent, and probably does represent, a lot fewer cases, but many of those cases may be affected by more than one Program Evaluation Review. Hard to get my mind wrapped around it; I'm sure it's difficult to -- for me to express so that you all understand. You're going to hear from LaVon Rutherford later in your agenda on a Special Exposure Cohort class update, but this is just a summary. He will get into more -- greater detail than I'm allowed to here. But to date, as of June 16th, 28 SEC classes have been added since May of 2005. Seventeen of those, or 61 percent, have been -- done so through the 83.13 process or where a petitioner comes forward and

1 petitions for that class. Eleven, or 39 2 percent, have been accomplished through the 3 83.14 process, and that's an instance where we at NIOSH have determined that we cannot 5 reconstruct the dose for a given claim for a site and we establish a class around that 6 7 claim. This represents classes from 22 sites. 8 It represents 1,735 potential claims. Just so 9 you know, there's -- this also represents an 10 increase of three classes, three sites and 170 11 claims from your last meeting that you held. 12 And I think that's it, and with that, I'll take 13 questions. 14 Thank you very much, Larry. Let's DR. ZIEMER: 15 open the floor, Board members, for questions. 16 John Poston. 17 Larry, could you say just a little DR. POSTON: 18 bit more about the cases that were reworked, 19 those high peaks that you pointed out. 20 because of the health physicists or because of 21 changes in procedures or what? 22 MR. ELLIOTT: Program Evaluation Reviews are 23 accommodated in our dose reconstruction 24 regulation, and we are required -- when we make 25 a technical change in our dose reconstruction

1 approach that might lead to an increase in dose 2 estimates, we're required to go back and look 3 at all of those non-compensable claims that 4 might be so affected. And the -- the red bars 5 that you saw on the right-hand side of that 6 chart are really representative of super S, the LANL, the Paducah, the Bethlehem Steel, the 7 8 lymphoma -- these are just to name a few. I 9 can give you the whole list if you want me to 10 run down the list, but there are -- I think I 11 said 32 of those. 12 DR. POSTON: I just want to make sure you're getting quality work from the HPs, that's a --13 14 MR. ELLIOTT: Yes, sir, I think we're -- we're 15 squeezing every bit of sweat they have to get 16 these things done, so ... 17 DR. POSTON: Okay. 18 DR. ZIEMER: Other comments or questions? 19 (No response) 20 Larry, we always appreciate your updates, and 21 it's -- it's good to see them tracking along and the progress that has been achieved, so we 22 23 again thank you very much. 24 SPECIAL SCIENCE JOURNAL PUBLICATION

I'm going to take the Chair's prerogative and

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1 jump ahead here a bit in the agenda, and I'm 2 going to pull an item from tomorrow morning's 3 agenda -- some of the items we're going to try 4 to keep pretty fixed where they involve SEC 5 petitions and people who may be on line, so I'm not going to move those. But we have scheduled 6 7 tomorrow morning a report on a special edition 8 of the Health Physics Journal that has come out 9 within the last week or so, and Jim Neton and 10 others at NIOSH had a big hand in the 11 development of the technical papers in this and 12 he's going to give us an update. 13 members, I think you may have copies of this 14 edition of the Health Physics Journal at your 15 places -- courtesy of NIOSH, I believe. 16 DR. NETON: Yes. Thank you, Dr. Ziemer. 17 You'll have to bear with me a little bit. 18 really did think I was giving this presentation 19 tomorrow so, being the procrastinator that I 20 am, I am -- I guess I'm somewhat prepared but I 21 may -- I may stumble a little bit as I go 22 along, so again I ask your indulgence. 23 I'm here to talk about something that we've 24 been working on for -- oh, probably the last 25 year and a half or so, and that is the special

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edition -- special issue of the *Health Physics Journal* that is entirely devoted to the NIOSH radiation dose reconstruction program.

You know, over the last five or six years we've developed a lot of scientific documents to establish the manner and the methods that we produce these dose reconstructions. And we felt that it was time to put it out into the open literature. NIOSH has formed, a while ago, a scientific steering committee whose mission is to review the state of our science and to figure out what direction we need to go. And it was the consensus of the committee at our first meeting that this was probably the best thing we could do right now, to -- for our program, to get some of our information out into the open literature.

So an overview of the issue -- it's -- it just came out in July, for the July issue. Those of you who are members of the Society would have received their *Journal* a week or so ago. And as Dr. Ziemer pointed out, we provided a copy to each of the Board members for their use and review. I know some of the Board members who are members of the Health Physics Society have

already received a copy, so just consider that
a second copy that you can -- you can put on
your nightstand.

The issue has 15 original articles that cover the science behind the program, and we spent a lot of time looking at what we really wanted to put out there. If you remember, we had, you know, implementation guides for the internal dosimetry, the external dosimetry, those type of documents. And so we tried to -- to capture in this issue those concepts that we thought were key to our program, the efficiency process, those type of things.

It highlights the unique nature of the dose reconstruction for compensation programs, so that's one reason we wanted to get it out there. As you -- as you probably know, dose reconstruction under this program is somewhat different than what you would see for a radiation protection program and also for a -- even a radiological epidemiological study. There are a lot of unique aspects of this program that are driven by the way the law is written.

I would be remiss if I didn't point out that

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this was a joint effort with the ORAU team. Ι commend all the authors who devoted much of their time -- their own time -- to putting these articles together. In particular I'd like to highlight the contribution of Dr. Dade Moeller, who really helped in shepherding this -- this through the process of getting an issue of this magnitude put together. Those of you who've published articles know it's an arduous process to get them published, a lot of backand-forth getting things through the editorial process, and -- and this was sort of magnified by 15 times. We were trying to get these all out at the same time and get people's time commitments organized and on target, and we came pretty close to our target date of getting this out, so I'm pretty proud of what this team has accomplished.

Just for your reference -- now that you have it, this is sort of redundant -- but this is a copy of the cover that came out. I was worried it might not be out in time for the meeting so I just gave you a snapshot of the cover.

You'll see that we chose to put the flow diagram of the efficiency process on there

because we think this is part and parcel of our program, how it's -- how it's operated and how it's actually gained us the efficiencies to be able to process the number of cases that -- that Larry Elliott just mentioned. We couldn't have done the 20-something -- or almost 20,000 dose reconstructions without having this process in place. And it's somewhat unique to the NIOSH program where we have a binary decision, 50 percent or greater or less than 50 percent, unlike some other programs on a global basis that have a sort of a sliding scale that require full-blown dose reconstructions for each -- each case.

The issue is broken into four major sections, as you'll see if you get to look at a copy of it. There's a program background; as you can imagine, it provides the overview of the program, the management issues associated with such a large undertaking. There's an issue of -- it deals with the Advisory Board, authored by Dr. Ziemer. And there's a paper on the scientific issues that we had to deal with in the dose reconstruction program, the large number of issues related to the demographics

1 and the biokinetic models and the 2 Hiroshima/Nagasaki survivors in relationship to 3 occupational exposures -- those sort of things. Data collection supporting studies are in 5 there. We felt it was important to talk about the collection and validation of the data that 6 7 we've done, and also what the role of site 8 profiles was, how they were envisioned and what 9 they ended up being and how we've actually 10 developed some of those. 11 And then in the third section you see dose 12 That's sort of the nuts and reconstruction. bolts of the issues, which you can imagine --13 14 we talk about the internal/external dosimetry 15 reconstruction, environmental, medical. 16 there's a paper in there that deals with 17 bounding analyses in the efficiency process, 18 how we use that to our advantage, and I think 19 that particular example is related to the 20 thorium work at Rocky Flats. 21 And finally there's a section devoted to the 22 probability of causation model IREP that, to my 23 knowledge or my thinking, is probably the best 24 -- the best peer-reviewed publication on IREP 25 to date that's out there. It goes into -- it's fairly extensive. It takes up a good chunk of the Journal, but it's the only place that I'm aware of that presents all of the nuances and ins and outs of what IREP is about, not only the National Institute of Health version of IREP but the NIOSH version of IREP and how there are differences.

I've kind of gone over this, but this just highlights some of the specific issues that we felt were important to include in this issue -- the efficiency process, data hierarchy -- all of these things are included -- either specifically addressed in our regulations on how we do dose reconstructions, or covered in some way, shape or form in the Implementation Guides. So these are all in some way discussed in some detail in the -- in the special edition.

One thing I did fail to mention -- well, let's see, maybe -- oh, no, I didn't, it's coming up. Why did we want to publish this; what were the perceived benefits of getting this out into the open literature? And one thing we felt was extremely important was that these articles would get an independent review of the science

behind the dose reconstructions. That is, we didn't ask any special favors of the Health Physics Journal. We submitted these and they were subjected to the standard blind review process where the Journal would select blind 6 reviewers, they would comment, and then we'd 7 have to negotiate those comments back and forth 8 until there was general agreement among the 9 parties. So an independent review of the 10 science gave us a good feeling that these were not just NIOSH home-brewed science -- science 12 concepts and methods, but they were -- also had 13 some acceptance, at least in the general 14 scientific population. 15 It also helped us to provide citable references 16 that could be used by others for our 17 18

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approaches. It's not uncommon that I get phone calls and e-mails from colleagues who say "I really like what you've done with the medical X-ray stuff, and yeah, you can find it on the web, but how are you going to find it down the line; how can I cite this in my publication?" Well, now it's out there. You know, the medical X-ray -- there's a paper on medical Xrays that's out there. I get a lot of phone

calls on IREP, so that's citable now. And even some of the other issues about data capture and development of site profiles, that sort of thing. So it's citable references that can be used by the general public.

This is somewhat redundant to the previous bullet I just talked about, but we believe that

literature.

bullet I just talked about, but we believe that it adds to the global scientific body of information on dose reconstruction. We've done a lot of work here. We've published thousands of pages of technical documents. It's out there now for -- for historical purposes and there are a number of other programs worldwide -- there's global programs on dose reconstructions that are being formed every day -- not every day; routinely. I mean I was just at a conference out in Colorado where there's a number of countries that are interested in looking at what we've done and adapting it for their specific uses. So we feel it's good to have it out there for their use in the open

And lastly, I think it's important that it assists with communication to the stakeholders on the scientific issues. It would be directly

relevant to those stakeholders who are scientifically oriented, but even for those who aren't, I think -- we hope that it would convey a general sense that these things have been peer reviewed and it's not just, again, NIOSH home-brewed science, scientific concepts, but it is -- has been at least accepted by some peer reviewers that are colleagues of ours in the scientific arena.

We didn't want to hide this publication when it came out. We thought it would be important to let the world know a little bit about it, so prior to the release we had developed a communication strategy. We have purchased 500 copies for distribution, of course 12 of which have been distributed to the Advisory Board. But we intended to provide them to Board members, we have a lot of interest from various Congressional offices who might have some interest in looking at what we've done, stakeholder requests, those sort of things. So we have copies available for distribution for those who -- who would like some.

We've also developed talking points for our staff, particularly our Public Health Advisors

1 who may get phone calls. Word does get around 2 in this compensation program and we prepared 3 our -- we prepared for our Public Health 4 Advisors to be able to discuss -- you know, 5 what does this really mean, what is -- why is NIOSH putting this out, what does it mean to my 6 7 case specifically, that sort of thing. 8 We're also in the process of issuing a press 9 release to notify folks that it's there so they 10 can find it. And we are going to put it on our 11 web site -- not -- we can't put the 12 publication, for copyright reasons, on the web 13 site. But we're going to notice that it has 14 been published. It's on our web site with a 15 summary of the contents and where one might be 16 able to get additional information, reprints or 17 entire copies of Journal. 18 So that's it in a nutshell. I'd be happy to 19 answer any questions, if there are any. 20 you. 21 DR. ZIEMER: Okay, thank you very much, Jim. 22 We appreciate that summary. Larry, you have a 23 comment here? 24 MR. ELLIOTT: I just want to emphasize for the 25 audience and for anybody listening -- anybody

that reads this transcript -- Jim mentioned this but I want to highlight it. The contributing authors to this journal worked on their articles on their own time. They did not take time away from dose reconstruction efforts or site profile development, Advisory Board support, and I commend them for that. But that was one of the ground rules that we set at the very start of this, that we will not sacrifice our work for the claimants just to get this thing out.

DR. ZIEMER: Thank you, good point. John Poston?

DR. POSTON: Well, I'd like to stick my oar in. I think this is a great thing and I commend NIOSH and ORAU for doing this 'cause, as Jim said, it's nice to have this in the citable literature so that scientists can use it all around the world. I think it's a great thing.

DR. ZIEMER: I think one good example of how some of the work is beginning to get noticed is -- and maybe, Larry, you can comment on this -- but my understanding is now that ICRP is looking at formally modeling the class -- or the super S plutonium modeling.

1	DR. BRANCHE: What is ICRP?
2	DR. ZIEMER: International Commission on
3	Radiological Protection.
4	DR. BRANCHE: Thank you.
5	MR. ELLIOTT: Yes, they are, and they are also
6	engaged in several committee work on this
7	relative to aspects of this program. Jim, as
8	associate director for science in OCAS, serves
9	on to advise on one of those committees and
10	your work is on remind me, Jim, it's on
11	DR. NETON: I'm on an NCRP committee.
12	MR. ELLIOTT: NCRP, I'm sorry, that's NCRP, but
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14	DR. NETON: I'd like to be on the ICRP, but
15	NCRP is fine.
16	MR. ELLIOTT: Yeah, we also we're also
17	working with
18	DR. ZIEMER: NCRP, for the record
19	MR. ELLIOTT: that's what I meant to
20	DR. ZIEMER: is National Council on
21	Radiation Protection and Measurements.
22	MR. ELLIOTT: Yeah, we're working with the NCRP
23	on several committee efforts. Jim's on one,
24	but we're also just about ready to commission
25	the NCRP to establish a committee that will

1 evaluate the IREP in great detail. 2 DR. ZIEMER: Any additional comments or 3 questions for Jim? 4 (No responses) 5 Thank you. (Pause) 6 7 BOARD INTERACTIONS WITH CONGRESS 8 Okay. Again I -- I'm pulling another item from 9 tomorrow's agenda, and that has to do with 10 Board interactions with Congress. And Jason 11 Broehm is here and he's -- he's our subject 12 expert on interactions with Congress. 13 welcome. Thank you for being willing to jump 14 ahead. 15 DR. BRANCHE: Actually if I can embellish, Mr. 16 Broehm is -- works in the CDC Washington 17 office, which he'll explain in a moment, and 18 he's an attorney working with our Congressional 19 liaisons and we work primarily with him in that 20 regard. 21 MR. BROEHM: I'm -- I'm not an IT expert so I 22 might need some help here getting my 23 presentation loaded onto the laptop here. 24 (Pause) 25 DR. BRANCHE: No, you don't -- there is no

PowerPoint in your -- in your group from this - for this next group.

(Pause)

MR. BROEHM: Okay. Like Jim Neton, I was expecting to give this presentation tomorrow, so thank you for your patience as I loaded my presentation here. Most of you know me on the I'm Jason Broehm. I work in the CDC Washington office, which does Congressional relations for all of CDC -- I handle NIOSH. my -- I think now about three years of working on this program, I know that periodically we've had Congressional requests. Many of you have interacted in various ways with Congressional staff, and we just thought it would be helpful to sort of provide the overall -- some background, some framework for, you know, what is -- what is Congress, what does it do, what does it need from you. And then -- I don't think a presentation like this has been given before, so...

Anyways, as a -- as an overview just quickly,

I'd like to -- in this presentation provide you

with some background on Congress, tell you a

little bit about my role as CDC Washington

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office representative to -- to NIOSH and to the Board, and then discuss the policies governing the Advisory Board and SC&A interactions with

Congress.

So I'd like to start off with a quick civics lesson -- not to insult anyone's intelligence here, but just to sort of provide the framework for where I'm going. As we all know, and I think learned in elementary school, we have three branches of government: The Executive Branch, for which we work; the Legislative Branch, which is obviously Congress; and then the Judicial Branch, the Supreme Court and the court system, which I'm not going to talk about today but it certainly is an important part of our government. These are three co-equal branches that were set up in our Constitution, and each one was supposed to provide some checks and balances on the others. So in our case, working in the Legisla-- or I'm sorry, in the Executive Branch, we're the federal agencies that are implementing the programs that the government runs and administering programs like EEOICPA. And then

Congress and the Legislative Branch is

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providing a number of important roles that interact with the -- with the Executive Branch agencies like HHS. So quickly, the roles of Congress in this program are -- first of all, Congress passed the legislation that authorized this program to exist. That happened back in 2000. years later it was amended to tweak the law a little bit, add some new requirements.

The other role that Congress has is annually they appropriate the funds that pay for this Advisory Board, for NIOSH to administer the program, for the Department of Labor to do their work, and obviously for the contractors who do their work in this program as well. It comes through that stream, so Congress does have that very fundamental and important role in this program.

And very closely related to that, they conduct oversight of this program, and other programs across the government. The goal in that is to ensure efficiency, make sure things are running the way that they were established to run. And where necessary, to identify the problems that may exist and correct them based on sort of

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spotlighting or highlighting the issues and -and in many cases putting the people who execute the program on the spot and making them motivated to -- to improve the program. So the -- the fourth function the Congress serves then I think a very, very important role, that we certainly can't underestimate, is the assistance they provide in helping our constituents who are claimants in this program or are otherwise interested in navigating this program, and advocating on their behalf. know we have a number of people in that situation here today. I know some of -- some of -- certainly interacted with their members of Congress or their Senators or U.S. Representatives, and it's an important role. These claimants are voters. They elect the Senators and Representatives to -- to do this for them, so -- and these elected officials serve at their pleasure. If they don't do their job, then they may not be re-elected. this is of the utmost importance in -- in their -- their role in helping the people who -- who they serve.

You know, I know that you as Advisory Board

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members hear from unhappy claimants every several months when you have public comment session at one of your meetings. But just to help you understand where Congressional staff are coming from, they're probably hearing from these people on a daily basis, maybe weekly, but it's their job to intervene and do what they can to try to help the process along. So let me run through what Congress needs from -- from NIOSH, from the Advisory Board, really from any program that's run across this government. First and foremost is for the program to be well-run, make -- like I said earlier, to make thing-- make sure that things are working as they were designed to work; and where there are problems, to try to fix those. Next, and very important, is that Congress gets timely information and that that information is responsive to -- to what they've requested, as much as possible. Congress really operates on a different time horizon than -- than most bureaucracies do. Congress really works at a fast pace. Often if you don't get them the information within a few days or, you know, a week, they get impatient. They -- they need

the information more quickly. So bureaucracies don't often respond at this -- at this fast pace, but certainly in my office our job is to try to help that to happen as -- as quickly as possible so that they can get the information they need and proceed with their jobs.

So, you know, basically in preparation for these meetings I notify Congressional staff of the Board meetings, workgroup meetings, let them know, you know, when things are going to come up approximately. Obviously things move around, but try to keep them as informed as possible, share appropriate documents that come from the Advisory Board and SC&A, their contractor.

And then, you know, to the extent that it's possible, one of the things that I'd like you to consider -- as workgroup chairs, in particular -- is when meetings are coming up it is helpful for them to have agendas. I know that that doesn't always happen with every workgroup meeting, but to the extent that you can have an agenda that's -- that's up on the web site in advance or even sent around a day or so in advance, I think that's helpful just

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so that staff who are very -- you know, keep very busy schedules can log onto the call, call in when -- when they have the time and sort of can plan their schedule accordingly. And then the documents that are related to their -- to the sites in question, obviously there are issues here with having to -- to do a lot of work, sort of a lot of this I know happens sort of at the last minute, then it has to go through a Privacy Act review which, you know, we all understand and have to work with. But to the extent that you can sort of plan backward, whether it's, you know, the Board, SC&A, NIOSH, their contractors, and build in the time for the review before the meeting, it's helpful to have those things in -- in their hands. In particular I think for a lot of these workgroup meetings the matrices, these documents sort of lay out the issues that will be discussed in the call. I for one, you know, have been listening to these calls for several years now and -- and still a lot of this just flies right over my head. But if you can imagine a Congressional staff person, many of whom are -- are young and don't have a

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technical background, it's helpful to sort of walk through the issues and at least have something on paper to quide where the Board is going. So I would just say that, as much as possible, please take that into consideration and help them go through this process with you. So my office, the CDC Washington office, as I mentioned earlier, provides Congressional relations support for the whole CDC. really do is sort of a -- sort of a liaison role across CDC with those in the Department of Health and Human Services who we need to coordinate with, and then with Congress. Our job really is to inform Congress of CDC programs and activities, answer questions as they come up. We coordinate any requests for information from Congress. That could take the form of a very simple question that comes up and -- or helping provide information or a status report on a claim for one of their constituents, or it could involve coordinating a briefing to provide information at the staff level, or a -- preparing a witness to testify before Congress at a formal hearing. So I, along with Christine Branche as the

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Designated Federal Official, we're here to -to serve as a resource to help answer questions as they come up and to advise you on how to interact successfully with Congress. And my contact information will be at the end, and certainly you know very well Christine's contact information, but any time you have questions I would encourage you to reach out to her and to me and -- and we can help. So as a preface to this next section which is talking about the policies and guidelines that we operate under, I just wanted to say that Congress serves a very important role in this program and should have access to the Advisory Board members and to the Board's technical support contractor, Sanford Cohen & Associates. So there are sort of two paths that we can go down, in particular for Board members, for how those interactions occur. I would just sort of say, as a starting point, the presumption is that federal employees who speak with Congress do so in their official capacity. You know, if -- if an agency official is asked to provide a briefing on something, we have to go through certain channels of just informing the right

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people, letting them know -- in some cases, you know, tweaks are made, additional people are asked to -- to sit in. But the Advisory Board really is sort of a special case and you are special government employees so you are parttime or not -- not full-time employees in the sense of working 40 hours a week but I know you do put in long hours. But the role of the Board to provide sort of a -- an outside independent voice and -- and review to -- to NIOSH's science is something that, you know, certainly in some cases it may -- may serve Congress better to -- to have those discussions without HHS administration officials present, except for the Board members. And so that -that can happen, but basically in order for an Advisory Board member to speak with Congress in -- in your official capacity as a member of the Advisory Board and a special government employee, you need to follow certain HHS procedures for agency communications with Congress. Having said that, though, of course Advisory Board members may speak with Congress as a

private citizen, providing a different --

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perhaps different inputs and -- and voice, but as I said earlier, the Designated Federal Official and I and my office stand ready to advise and assist you as Congressional requests are received and -- and each one is treated somewhat differently.

So first for appearing as an Advisory Board member, you -- you may speak with Congress in your official capacity following the following When an Advisory Board member receives rules. a request to speak with Congress, he or she should alert the Designated Federal Official, who will familiarize you with the process and coordinate with me in my office and make the necessary arrangements to -- to move forward with -- with whatever's needed. Any written document, whether it's, you know, a single page briefing document that you plan to hand out, or something that's longer and -- and more involved, like hearing testimony, an advisory member needs to share that in advance, and it has to go through a certain clearance process within CDC -- well, NIOSH, CDC and HHS. particular with -- with a hearing, these are more involved and more formalized. Those types

1 of statements, if you're appearing as an 2 Advisory Board member in that capacity, 3 typically what happens is those are cleared at several levels, including -- starting at the 5 NIOSH and -- and/or CDC level. Then it has to 6 qo through an HHS clearance process which goes 7 across the various policy offices of HHS. 8 People have a chance to review and comment and 9 suggest changes. And then the Office of 10 Management and Budget serves a coordinating 11 role across the whole of the federal 12 government, so testimony in those cases would 13 be circulated to our counterparts in the 14 Department of Labor. They would have a chance 15 to comment and -- and provide input on that 16 testimony as well. 17 Now not every document is that sort of reaching 18 in scope in terms of the review. Most -- in 19 most cases a simple one- or two-page document 20 will go through a fairly abbreviated clearance, 21 but it is important to have that reviewed in 22 advance. 23 And then if you appear in this capacity, an HHS 24 representative -- myself, perhaps others --25 would -- would appear with you and -- and

accompany you and get you to the right place at the right time. Then any -- any follow-up information that's requested during the meeting would have to go through the same -- the same clearance and review process as that that was prepared in advance.

So then the other path is appearing as a private citizen. And when speaking with Congress as a private citizen an Advisory Board member really needs to make clear, whether it's in written or oral communications, that he or she -- that you are speaking on your own behalf and not in your capacity as an Advisory Board member, just need to -- need to make that clear.

And then Advisory Board members need to be aware that in this capacity as a private citizen you shouldn't be offering information or opinions about the Advisory Board or other government actions, particularly those that are not public information. Advisory Board members of course I know are constantly reminded by Dr. Branche and others that no information should be released, no documents should be released before consulting with her to be sure that it's

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gone through the Privacy Act review that's -that's necessary to protect individuals' information that shouldn't be shared publicly. And then finally, Advisory Board members should not speak to an opinion or position of the Board unless the Advisory Board has taken a formal and publicly-approved position in accordance with your procedures. So sort of related to this then is meetings that Congress has typically or periodically requested of your contractor, Sanford Cohen & Associates. As we know, they do much of the technical work that supports -- supports your function, and this work and these -- these work products are of keen interest to members of Congress and their staff. And so under the Board's procedures that have been discussed at previous meetings, Congressional offices may speak with SC&A, with or without members of the Advisory Board present. We basically leave that up to the Congressional offices. If they wish to have the meeting with -- with SC&A representatives and not invite a Board member, that is -- is their prerogative. We in the Executive Branch sometimes do have

disagreements with Congress, but we generally try to provide information that they need to -- to serve their role in this process. And so we've -- we've generally provided that access when it's been requested.

So as with an Advisory Board member appearing in -- in their private citizen capacity, SC&A representatives need to speak in that same role and not provide opinions about what the Board is doing or might do or should do. And then the SC&A representatives need to make clear to all parties that they are appearing as employees of a private company, that they are providing their own private opinion and don't represent the positions of the Advisory Board or the -- or of HHS.

And then of course the same proviso, any documents need to be pre-cleared and make sure that -- that they've checked with -- with Dr. Branche to make sure that the Privacy Act review has been done and that those documents are cleared for release before they're shared. And then I would just add -- you know, typically government contractors don't do briefings for Congress. They're sort of

1 providing a support role to the government, and 2 the government officials would do the briefing, 3 occasionally would have a contractor with them 4 to provide a supporting role. But again, as 5 sort of the special case that -- that the 6 Advisory Board has here in terms of providing a 7 -- an independent outside review on NIOSH's 8 science, the -- that's -- the role of SC&A in 9 this process also is to provide that outside 10 scientific voice and we don't want to get in 11 the way in terms of even just providing the 12 appearance of somehow influencing what they -what they say, just by virtue of being in the 13 14 And so it has been the policy to treat room. 15 SC&A as a special case and unique and different 16 from -- from most other government contractors. 17 So that completes what I -- what I had to say 18 to you. I'd be happy to take any questions you 19 may have and --20 DR. BRANCHE: Before you do, I --

MR. BROEHM: Yes?

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DR. BRANCHE: -- have a couple of -- a proviso and some additional information for you, and then please ask -- this is a good opportunity for you to ask as many questions as you wish of

Jason and me and -- as our -- as Jason has explained this information.

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As it concerns the written documents and so forth if you were to appear before Congress -if you were to be asked to appear before Congress in your capacity as an Advisory Board member, understand you're representing the Executive Branch then speaking to the Legislative Branch of government. You can't ask for forgiveness. There are -- this is your -- this is your chance to know that this is the way the procedure is. So to ask for forgiveness later because you did something in a completely -- it would be considered completely inappropriate. You must have all part of your testimony -- proposed testimony cleared by all the levels that Jason just explained.

And Jason, if you can put your slide back up about appearing as a private citizen -- and I will ask Jason to send this PowerPoint slide to me so that we can get this to you. Actually -- MR. BROEHM: And I would just say that Zaida was -- does have the slides on paper and was preparing to copy them for tomorrow, but --

1 DR. BRANCHE: Okay. 2 MR. BROEHM: -- we got -- got ahead of her. 3 DR. BRANCHE: But I think it's fine for them to 4 have them electronically. MR. BROEHM: Yes, we'll get it to --5 6 I think it'll be more helpful to DR. BRANCHE: 7 There's some provisos here that I think 8 are important and I would substitute the word 9 "Congress" with "the press." When you -- many 10 of you interact with the press as it concerns 11 spe-- specifically as it concerns certain sites 12 for which you serve as a workgroup chair. You 13 would be speaking as a private citizen to the 14 press. You'd ha-- I -- I would ask that you 15 make it very clear that you are speaking as a 16 private citizen. You would not be speaking on 17 behalf of the Board. Anything that you would 18 say to the press on behalf of the Board would 19 then have to follow the information that's in 20 Jason's fourth bullet, information that has 21 been formally and publicly approved by the 22 entire Board.

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I have personally been misquoted by the press. It is possible to be misquoted by the press, but I'm not saying that they purposefully do

1 anything wrong, but I think it's always 2 important that you do your part to make certain 3 that they understand that you're speaking as a 4 private citizen and not as a Board member. 5 And with those provisos, I turn it back to 6 Jason to be able to put your last slide back 7 up, and I know that you have several questions. 8 Well, the last slide asks for DR. ZIEMER: 9 questions, so if Board members have questions 10 for Mr. Broehm, this is the time. I think John 11 Poston was first, and then we'll go to Mark. 12 DR. POSTON: I don't have a question for Jason. I do have a comment. We've talked about this 13 14 before. It -- you know, SCA is a contractor who works under the direction of the Board --15 16 this Advisory Board. We establish the tasks 17 that they're going to work on and so forth. 18 And I just want to say that I find the 19 differences between the rules for the Board and 20 the rules for SCA not only ludicrous but 21 hilarious. 22 DR. ZIEMER: Mark? I was -- I was waiting for maybe 23 MR. GRIFFON: 24 a response about that, I don't know if there is 25 any response.

1 DR. ZIEMER: Jason --2 MR. GRIFFON: I know it wasn't really a 3 question, but --4 DR. ZIEMER: -- were you planning to respond to 5 t.hat.? Well, I know that the Board has 6 MR. BROEHM: 7 debated this in the past, and I believe has a 8 written policy that's -- that's been passed, 9 and so I quess that's what I would say to that. 10 DR. ZIEMER: Okay. 11 MR. GRIFFON: I guess that -- that was my 12 question, and maybe we do, but you -- you 13 mentioned that this has been procedure, and we 14 do have an internal procedure on this that 15 covers those bullets? 'Cause I was trying to 16 follow your -- all those bullets and --17 DR. ZIEMER: Well, basically --18 MR. GRIFFON: I was also comparing SC&A versus 19 the Board in my head, but I don't have -- you 20 know, I wondered if we have --21 DR. BRANCHE: It was news to me --22 MR. GRIFFON: -- a written policy or --23 DR. BRANCHE: I was told that --24 DR. ZIEMER: Well, early on --25 DR. BRANCHE: -- there was policy --

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DR. ZIEMER: -- we had a policy written that said that the Board -- the Board preferred to be present at meetings where Congress called on our contractor to give them information.

MR. GRIFFON: Right.

DR. ZIEMER: But we also recognized, based on advice from perhaps your office or at least from the Secretary's office, that we can't dictate -- Congressional offices can call on whoever they want to get information and you cannot invite yourself into their office, so --MR. BROEHM: Right, and that's -- yeah, so I'll clarify that a little bit. The -- when a request comes for a briefing by SC&A, as I understand it, they are then supposed to report that to Dr. Branche and to you, Dr. Ziemer, I believe. That then gets -- sort of makes its rounds to the rest of the Board members. there is one, or maybe there are more Board members, who would like to participate, that offer may be transmitted to the Congressional office that's requesting the briefing. may say fine, any and all comers, we'd be happy to have them. They may not. And just getting back to the sort of Government 101 slide in the

beginning of my presentation, we are two different branches of government with two different needs, and Congress does serve an important role in this program. If they want to invite certain people and not other people, we -- we don't want to get in the way and so that's -- that's the procedure we've been proceeding under.

DR. BRANCHE: If I can just for -- just for -- in my very short experience, every time SC&A has been asked to respond to a Congressional inquiry or participate in a meeting, we've asked if the Board member -- if a Board member can be present, and that has always been honored. And they've even gone to the bother of setting up a conference line so that you can participate by phone. So Jason, my question is, in your experience has there ever been a circumstance where the Congressional member did not wish to have a Board member participate?

Okay, I'm being told oh, yes, okay.

DR. ZIEMER: Early on there were a number -MR. BROEHM: I believe there may be one or two
cases, but I think it's probably more the
exception than the rule. I think generally the

1 Congressional staff would be happy to hear from 2 the different voices who -- both from the Board 3 and from SC&A in such a briefing. DR. POSTON: Dr. Ziemer, I would request --5 since there are at least five new members of 6 the Board and a new Designated Federal Official 7 -- that if there is such a policy that it be 8 distributed to us so we can understand it. I'm 9 10 MR. GRIFFON: Yeah, I agree. 11 DR. POSTON: -- I've taken a poll of the folks 12 that are here that I can speak to and I know 13 that none of us have seen such a policy, or 14 none of us was aware of such a policy. 15 MR. GRIFFON: Well, and speaking as an older 16 member of the Board, I -- I don't -- I remember 17 the discussion about the one issue there, you 18 know, as far as attending meetings with 19 Congress. But I don't remember this being 20 detailed in a policy --21 MR. PRESLEY: -- this detailed. 22 MR. GRIFFON: -- information about when you can 23 provide -- and I've had these discussions with 24 several -- I've had them with Christine, I've 25 had them with Lew Wade about providing opinion,

1	especially as it related to the press, but I've
2	never seen, you know, these detailed bullet
3	points laid out this way and I I think, if
4	it is proceduralized, I'd I'd like to see it
5	as well.
6	DR. ZIEMER: Well, I'm not talking about these
7	bullet points. I'm only talking about
8	MR. GRIFFON: The one
9	DR. ZIEMER: the presence of the Board in a
10	
11	DR. BRANCHE: Yeah.
12	DR. ZIEMER: Congressional request to SC&A.
13	MR. GRIFFON: But I mean then all this other
14	stuff
15	DR. BRANCHE: Right, but as far as the
16	MR. GRIFFON: is this a non is this a
17	proc
18	DR. ZIEMER: Yeah, that one issue.
19	MR. GRIFFON: this a policy being
20	DR. BRANCHE: Well, there's several people
21	speaking at once, I'm
22	MR. GRIFFON: Sorry. I'm asking it if all
23	these things in are they a policy from the
24	agency?
25	DR. BRANCHE: Yes, the other

MR. GRIFFON: And is there -- and is there a policy document -- other than just overheads with bullet points?

MR. BROEHM: Yeah, I don't think that --

DR. BRANCHE: Yes.

MR. BROEHM: -- there is a policy document that states all of this, that I've ever seen. But it's the operating procedures that we as a federal agency, and I think most other federal agencies, follow as parts of the administration, that you go through these clearance processes if you are a federal employee.

DR. BRANCHE: Well --

MR. GRIFFON: So when we -- when we -- as Board members, if we act -- and I've always -- in my communications I always say I'm -- I'm speaking to you as a member -- as a -- as a member of the public, not for the Advisory Board, not for the working group, but -- but would -- would SC&A and the Board members be treated the same under your policy in that regard? Like if you speak to Congress or the press as a member of the public, same rules apply kind of thing, or -- or not?

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MR. BROEHM: Yes, if you're speaking as a private citizen, I think basically the same rules apply.

DR. BRANCHE: As far as the policy issues, I can't speak for any other Department, but let's go back to the clearance issues if you're speaking as a member of the Board and were asked specifically in that capacity, all of the things that Jason explained as far as procedure -- I can't speak for any other Department. I do know that the levels of clearance and so forth apply to the Department of Health and Human Services. I'm not aware that those are written, but those are the procedures that every employee is expected to do. And again, this distinction between the role as a private citizen and that as a member officially of the Board, that's the distinction that I think is important. And I know that, as I said, many of you have been I know interacting with the press. You've made a point to say that you're speaking as a private citizen. My suggestion is that you can't over-emphasize that point, because some of you have been misquoted as having spoken on behalf of the Board when you

1 did not. 2 MR. GRIFFON: But -- but even -- even with that 3 said, I -- I mean I would really like a hard 4 copy, and I know you were preparing for 5 tomorrow, but -- because I think there -- there 6 was one bullet point up there that said as a 7 private citizen you couldn't given an opinion 8 on --9 DR. BRANCHE: Can you put that back up? 10 MR. GRIFFON: -- a Board matter or -- and --11 and I don't know, there's some nuance in there 12 that I want to understand. 13 DR. BRANCHE: Yes, it's the -- it's the --14 MR. GRIFFON: So --15 DR. BRANCHE: Is it the second or the third 16 bullet to which you're referring? 17 MR. GRIFFON: (Unintelligible) one, but that 18 means as in a -- as a Board member, not as a 19 private citizen. 20 DR. BRANCHE: Yeah, this --21 MR. GRIFFON: All right. And where -- where 22 does it talk about us speaking as a private 23 citizen? Is that on the next slide? 24 DR. BRANCHE: All of that applies to you 25 speaking as a private citizen.

1 MR. GRIFFON: So that second bullet applies 2 speaking as a private citizen? I can't offer 3 information or opinions about the Board or --4 or government actions? I mean why do they want 5 to talk to me as a -- as a private citizen if they don't want some information? 6 I think --7 DR. BRANCHE: Well, you have expertise that --8 that brings you to a mem-- as a member of the 9 Board. You have experti--10 MR. GRIFFON: Well -- well, then -- then 11 compare that to the SC&A bullet on private 12 citizen. It said that Congress may seek them out for their opinion. I don't know, I just 13 14 want to understand this better, I guess, before 15 I speak to other people. DR. BRANCHE: Yeah, we -- that -- this is your 16 17 opportunity to clarify that. 18 DR. ZIEMER: I believe what this is saying is 19 if they ask you what the Board's position is on 20 something -- for example, what's the Board's 21 position on -- I don't know, let's pick one out 22 -- Dow Chemical, Madison. Until the Board 23 takes such a position, you cannot --24 MR. GRIFFON: Oh, I agree. 25 DR. ZIEMER: Yeah.

1	MR. GRIFFON: But it wasn't
2	DR. ZIEMER: They could they could ask what
3	a
4	MR. GRIFFON: Board's position. It says
5	if you can read that again
6	DR. ZIEMER: Well, I'm
7	MR. GRIFFON: I'm just worried about the words,
8	you see what I'm saying? If this is a policy
9	document, these overheads are now a policy, I
10	want to understand them.
11	DR. ZIEMER: It was on the other
12	MR. GRIFFON: Back to SC&A, yeah.
13	DR. ZIEMER: on the other slide.
14	DR. BRANCHE: It was the one appearing as a
15	private citizen.
16	DR. ZIEMER: The private citizen slide.
17	MR. GRIFFON: About about Advisory Board or
18	other government actions.
19	MR. BROEHM: That are that are not public
20	information.
21	MR. GRIFFON: That are not public informa
22	okay, so it's okay.
23	MS. MUNN: If they're already public
24	information, if they're already out there, then
25	it does not appear that there's any caveat

1	other than that we make certain we're quoted as
2	private citizens and not
3	MR. GRIFFON: I'm reading through it, that's
4	why I wanted a hard copy, that's
5	DR. ZIEMER: No, you could say that the Board
6	has taken
7	MR. BROEHM: Yeah, and I'm sorry you don't have
8	that in your hands.
9	DR. ZIEMER: this position on something
10	where where the action is public.
11	MS. MUNN: And can even say you disagree with
12	it, as a private citizen.
13	MR. GRIFFON: Right, right, I can say yeah.
14	MS. MUNN: But yeah.
15	MR. BROEHM: Are there other questions?
16	DR. ZIEMER: Brad has a question.
17	MR. CLAWSON: You were mentioning that if we're
18	called in there to talk to Congress, that we're
19	supposed to submit to you what we're going to
20	say. And any time in the past that I've ever
21	talked to them, I don't know what they're going
22	to ask so how am I supposed to submit what I
23	don't know?
24	DR. BRANCHE: He's talking about panel
25	presentations if you were being asked to

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MR. CLAWSON: Okay. Now --

MR. BROEHM: If -- if you are speaking from notes that you're not handing out, you don't need to clear that, although you should -- if you're speaking in your capacity as a Board member -- review that with -- with Christine Branche and go through -- go through those points in advance. But if you're not handing those out as -- as PowerPoint slides or a onepage handout to leave behind, you don't need to go through the whole clearance process that I described. It's a little bit more formalized. And then when speaking of testimony, that sort of takes it to the next level. That's -that's where you go through a long -- weekslong process, in some cases, of reviewing at the varied levels and having comments submitted.

MR. ELLIOTT: But I --

DR. ZIEMER: Larry Elliott.

MR. ELLIOTT: -- I think it's well-advised that if you walk in with notes, you better expect that they're going to want a copy. At least that's been my experience.

MR. CLAWSON: So --

MR. BROEHM: It depends -- it depends what those notes look like. If you have just handwritten notes, it's one thing. If you go in with a full PowerPoint presentation that you're just using on -- for yourself, they may say "Oh, can I have a copy of that?"

MR. CLAWSON: So you're telling me just -MR. BROEHM: So you need to be prepared for
that.

DR. BRANCHE: And then it needs to have been cleared.

DR. ZIEMER: Brad?

MR. CLAWSON: So you're -- you're telling me just shoot from the hip and enjoy it, basically, huh? You know, it's very, very interesting to me that we have so many different policies for so many different groups. I understand the importance of not representing the Board or anything else like that, but why -- why would a member of the Congress or whatever else want to talk to us but to be able to gain our opinion? And it -- it's interesting to me and I realize that they're calling SC&A in to talk to them about

the Board, but they have a totally different -- a totally different process to be able to go through. It's -- it's -- well, it's -- it's the government, I guess.

DR. ZIEMER: Okay. Phil?

MR. SCHOFIELD: I've got one concern, and that's where some Congressional office contacts you out of the blue and has particular questions, maybe about a certain workgroup or issue that's coming up, and wants to know where you're headed on that where -- you know, I'd like a little more clarification on that. Do we stall for time or...

MR. BROEHM: Well, you know, that's something that happens to people in HHS, too. And sometimes it's by design, sometime it's just because they see that as the most direct route or don't know that my office even exists. But if you're caught on the phone and they're asking for information, you could say "Could I call you back?" and alert Christine and -- and/or myself, and we can facilitate that conversation at greater length. If they have very simple questions just about, you know, when's the next workgroup meeting going to be,

you don't need to go through that whole process, I would say. But it -- it does happen and, you know, I wouldn't be overly worried about that. But as a general rule I would try to include the two of us.

DR. BRANCHE: And if it does -- if it does happen, something that simple, then -- and someone just asks you when the next workgroup meeting is, then you could -- I would encourage you to send a courtesy message to me to let me know that that contact has happened.

DR. ZIEMER: But there's a lot of information like that you can answer very simply, and it's public information anyway. People may not have known where to find it on the web site or something like that.

But on the other hand, and I'll just mention this and then we'll call on a couple more folks here, but a couple of weeks ago I got an extensive inquiry from a Senator's office asking the amount of money spent by this panel to investigate issues at certain sites in the country and wondering how that compared with other sites in the country, I think trying to determine whether this particular Congressional

district was getting their fair share of attention -- or whatever. But -- and the office asked for that information back from me in 30 days.

Well, number one, we have a rule that says you

-- we cannot respond to Congressional inquiries

-- the Chair can, nor can the members -without clearing it first with the Board
anyway, so we don't have a way to do that in 30
days. Further, the information was not
information that I could readily get my hands
on. So I turned it over to Christine and -and through their office, maybe working with
Jason, I don't know, was able to get the letter
redone and redirected so the Congressional
office made the inquiry of NIOSH to get the
information. But some of the -- some of the
inquiries are done in good faith, but they are
things we should not get involved in.

MR. GRIFFON: Right.

DR. ZIEMER: Now, Wanda, then Michael.

MS. MUNN: One of the things that continues to be of concern with respect to interactions with Congress is the persistent view that our contractor is not our contractor but rather our

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auditor. In almost every case when I've had occasion to interact, either here in this setting or elsewhere, with Congressional staff, the approach has always been that of "Your auditor has said -- has these findings." It is a concern to -- I -- certainly to me, and I think to others, that that misunderstanding applies. But it certainly seems to be the primary reason -- there are two primary reasons, apparently, why Congressional staff are so eager to speak with SC&A. One is they view them as auditors, and two, they are accessible. They have people available in Washington, D.C. to be able to go to their offices easily. So I rely on you, Jason, to help supply staff with that revised, correct view of what the association is and what findings are being brought to them without having been vetted in this -- in this forum as being preliminary findings, always. It's -there are several things that -- that we look to you as a person to do, and I hope -- we have no way of knowing that you are in fact doing that, so it would be helpful, for me, to hear from you that you do in fact make that effort

because it seems to be such a consistent misunderstanding that we encounter.

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MR. BROEHM: In every communication I have with a Congressional staff person working on this issue -- and in many cases, especially for staff who have worked on this and continue to work on this -- it's a continued conversation over many e-mails and phone calls to continue to explain and re-explain this program. very complex and it's not always intuitive to -- to staff who are coming to it new. I always do my best to explain to them what the Board is and what SC&A's role for the Board is. Now to the extent that there may be misconceptions of what SC&A's role is out there, and I don't hear from those staff, I may not even know that, what their -- what their idea is of SC&A's role. But certainly in phone conversations and e-mails that I have, I put out the information in terms of what -- what role they provide and support they provide to the Board is -- is really what it is. continue to hear the word "auditor" and I don't really know where that came from, but it sort

of has caught hold and, you know, I'll just --

1	all I can say is I'll continue, as I have
2	conversations with staff, to to explain what
3	SC&A's role is.
4	MS. MUNN: That's appreciated. Thank you.
5	MR. BROEHM: Thank you for the question.
6	DR. ZIEMER: Brad? Oh, Mike Mike was first
7	
8	MR. CLAWSON: Go on, Mike.
9	DR. ZIEMER: and then Brad.
10	MR. GIBSON: As far as the proposed testimony,
11	PowerPoint presentations, whatever, what
12	exactly are they going to be reviewed for?
13	DR. BRANCHE: You would be speaking on behalf
14	of the administration, essentially, 'cause you
15	repre in this capacity, you represent the
16	Executive Branch of government in your special
17	as a special government employee.
18	MR. GIBSON: And in my role as a government
19	employee, it's my duty to monitor how HH HHS
20	is implementing this legislation. So if I give
21	draft testimony that I'm asked to give for your
22	review, doesn't that kind of the fox
23	watching the hen house?
24	DR. BRANCHE: When you
25	MR. BROEHM: You I mean you're appointed by

the President of the United States.

MR. GIBSON: Correct.

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MR. BROEHM: You are a government employee.

You work for him. So I'm in the same position. I can't just go up to Congress in my role as a CDC employee and say whatever I want. Usually government employees who do that are considered whistleblowers. You -- again, I explained that you, as Board members, are in a little bit of a special case here and that's why we have the two paths available to you. One is to speak in your role as a Board member. The other is to speak as a private citizen, where perhaps you can be more frank and critical of the program. You know, I don't think in terms of going through the administration review process that if you had something that was critical of -- of NIOSH, or you had recommendations that -- that you thought -- some changes that could be made to improve the program, those would necessarily be taken out. But there are things -- if you put in your testimony, for instance, that the Advisory Board needed \$10 million next year, those are the kinds of things that we can't put in our testimony if it's not in the

1 administration budget. So it's -- it's making 2 sure that -- that administration policy is 3 followed, making sure that you're not asking 4 Congress for vast new resources that aren't in 5 the administration's budget. It's --MR. GIBSON: And actually --6 7 MR. BROEHM: It's again why we have the two 8 paths open to you. And when -- you know, when 9 -- I know an Advisory Board member has spoken 10 in the past and testified to Congress, that it 11 was in that private citizen role and -- that --12 that was one option. MR. GIBSON: 13 But --14 MR. GRIFFON: Was that Dr. Melius? 15 MR. BROEHM: Yes. 16 MR. GRIFFON: He was -- role as a private citi-17 - okay. 18 DR. BRANCHE: And his testimony --19 MR. BROEHM: And he -- he worked --20 DR. BRANCHE: -- was cleared. MR. GRIFFON: Right. 21 22 MR. BROEHM: He wor-- he coordinated very 23 closely with the former Designated Federal 24 Official, Dr. Wade. 25 MR. GRIFFON: That was (unintelligible).

MR. BROEHM: Yeah. So again, it's just an example of whenever this -- this comes up and there's a need for a briefing, a hearing, whatever, it's important just I think to get in touch with Christine very early in the process and then, you know, we can work through the proc-- through what the next steps are from there and what your options are.

MR. GIBSON: But -- well, my options are limited as a private citizen. I can't use information or opinions about Board activities. But if I'm questioned -- asked to give testimony or whatever as a Board member, then I have to have this thing scrubbed, not knowing what'll come out of it. That just doesn't seem...

DR. BRANCHE: Let me just make a distinction, and as -- I think the operative part of that clause is "that are not public information."

As a private citizen, you can speak about Board policy, about Board information that's made public. You can -- you can do that, and you can offer an opinion on that as a private citizen.

MR. GIBSON: Okay.

DR. ZIEMER: Okay, Brad.

MR. CLAWSON: I just wanted to make sure as we're -- as we're discussing this and stuff and we've brought up SC&A, that -- that people understand and realize that John Mauro and the rest of SC&A staff have gone to great lengths to be able to try to involve us in it, and have done an excellent job and we're not -- we're not in any way, shape or form criticizing that. I just wanted to make that distinction.

MR. BROEHM: And as I understand it, when they have a meeting that they then do inform the Board of -- sort of a summary of what happened at the meeting, so from summaries that I've seen, those seem to be fairly detailed and -- and accurate -- not having been in the room, but I mean they seem to not -- not to be too abbreviated or -- or leave things out.

DR. ZIEMER: Okay. Mark, another comment?

MR. GRIFFON: Yeah, the only -- I -- I -- and

I'm going back to Christine's reference there

that -- that are not public information, and

that is reassuring in some ways. The only

concern I have with that is in -- in a role of

a work -- workgroup chair, if -- and that's

1	when you you typ I've typically run into
2	staffers is the the SEC process or whatever.
3	As we all know, things are are often real
4	time, so you know, we always this goes back
5	to this review process and the ability to have
6	documents that are public. If something's
7	discussed on the on a workgroup phone call,
8	I would assume that's public information, even
9	if the transcript's not ready yet. Right?
10	DR. BRANCHE: We make we make
11	MR. GRIFFON: Yeah.
12	DR. BRANCHE: the workgroup meetings
13	available for the public
14	MR. GRIFFON: So they right.
15	DR. BRANCHE: to participate, and
16	Congressional members are often
17	MR. GRIFFON: Right.
18	DR. BRANCHE: on the phone for those.
19	MR. GRIFFON: So to the ext I mean this
20	information was discussed publicly
21	DR. BRANCHE: That's right.
22	MR. GRIFFON: so if someone calls me to
23	follow up on that, I can give my opinion
24	DR. BRANCHE: As a private citizen.
25	MR. GRIFFON: on that as a private citizen,

1 right. 2 DR. ZIEMER: There could be some details in the 3 4 MR. GRIFFON: Yeah. 5 DR. ZIEMER: -- paperwork that the Board is 6 discussing that's redacted --7 DR. BRANCHE: Right. 8 DR. ZIEMER: -- information for the public, so 9 10 DR. BRANCHE: And then that's not --11 DR. ZIEMER: -- so that part could still not be 12 disclosed. 13 DR. BRANCHE: Excellent. 14 MR. GRIFFON: Right. 15 MR. BROEHM: But I -- I would say, Mark, in 16 response to your question, it probably is 17 fairly typical of the Congressional staff, 18 having sat through a whole, you know, hours or 19 three-hours-long call, may come out of that 20 with some questions and --21 MR. GRIFFON: Right, right. 22 MR. BROEHM: -- I think it's -- it's likely you 23 are going to get that kind of question as a 24 workgroup chair. 25 MR. GRIFFON: As what?

1 MR. BROEHM: It is likely that you are going to 2 get that kind of question as a workgroup chair 3 4 MR. GRIFFON: Right. 5 MR. BROEHM: -- of, you know, what -- what --6 MR. GRIFFON: And then it's okay --7 MR. BROEHM: -- what just happened, and 8 explain. 9 MR. GRIFFON: As long as it's okay as a --10 DR. BRANCHE: Right. 11 MR. GRIFFON: -- not representing the 12 workgroup, I always say that --13 DR. BRANCHE: Right. MR. GRIFFON: -- you know, but it's okay to --14 15 to offer my opinion on where -- usually they want to know well, what's the next steps, did I 16 17 miss, you know, something here, you know, or 18 when's the next meeting, sometime -- you know, 19 how's this going to be -- you know. 20 DR. BRANCHE: And the only thing I would offer, 21 again, Jason said to you, it has been helpful -22 - and this really is a protection to you -- to 23 the degree that Jason and/or I can be on the 24 line with you when you speak to that member of

Congress so that we can explain that divide to

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kind of keep -- it's meant to keep you out of trouble in that regard. And there's always a tension -- I just mention this again because we have a lot of people in the audience who've also participated in the -- in some of the workgroup calls -- the Board has put a value on having information as close to real time as possible, which means that you're often discussing documents that have not been Privacy Act reviewed, and there's always going to be that tension of having the latest information that SC&A or NIOSH has provided and you'll end up discussing it without it being yet made available for the public. If you have set your information up in such a way that you have your information available, we can -- and it has been Privacy Act cleared, we do try to get that information on the web site in advance of your meeting. But many of you are presiding over issues where people are working up to the last minute, and you're always going to have to fight that tension, and that's cover that Jason and I can provide for you in your interactions with members of Congress and the public -- and the press.

1	DR. ZIEMER: Okay. Thank you. Jason, thank
2	you again very much.
3	MR. BROEHM: Sure, thank you.
4	DR. ZIEMER: It's been a very fruitful
5	discussion. We're going to take our break now.
6	It's five after 3:00. We actually will yes,
7	let's reconvene in 20 minutes.
8	DR. BRANCHE: Twenty minutes?
9	DR. ZIEMER: Yes.
10	DR. BRANCHE: So at 25 after the hour.
11	DR. ZIEMER: Yeah.
12	DR. BRANCHE: Okay, we'll put the phone on
13	mute.
14	(Whereupon, a recess was taken from 3:05 p.m.
15	to 3:25 p.m.)
16	DR. BRANCHE: Everyone please take your seat,
17	we're about to we're going to start right
18	now.
19	(Pause)
20	We are restarting the meeting after the break.
21	Could someone on the line please let me know
22	that you can hear me?
23	UNIDENTIFIED: We can hear you.
24	DR. BRANCHE: Thank you very much. I
25	appreciate that. Now if you could please mute

your phones, I would appreciate it. If you do not have a mute button, then please use star-6; and when you're ready to speak, please un-mute your phone with the same star-6. And again I ask that if you are participating by phone, it is critical that you mute your lines. Also do not put us on hold. If you feel like you have to leave the line, then please hang up and dial back in, but do not put us on hold. Thank you so much.

Dr. Ziemer?

DR. ZIEMER: Don't you wish you could say that when you call your service provider for help -- do not put me on hold.

DEPARTMENT OF LABOR UPDATE

We're going to jump ahead again on the agenda for a brief time and pull in a presentation that was originally scheduled for tomorrow afternoon, and that is the Department of Labor update. And it's probably good we do that this morning -- this afternoon. Jeff Kotsch from Labor is here and we earlier this afternoon had the update from the -- from NIOSH, so -- and usually we have those kind of next to each other, so it's good we'll get the NIOSH and

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Labor people a little closer together again.

Jeff, welcome back to the podium. We're

pleased to have your report.

MR. KOTSCH: Thank you. Good afternoon. Ι have to apologize. First of all, I'm wading through the back end of a cold and I'm -- so my voice is a little rough. Also since -- I'll --I'll at least take time to put in the caveat that since I was supposed to be up tomorrow, and since I've been kind of under the weather, I haven't really been looking at the presentation that much so we'll work through that, too. And then also I should just say that some of the stuff is -- or the information is redundant, Board meeting to Board meeting. Some of it's background information and that's primarily for the members of the audience that, you know, might be new to the meeting rather than the Board, who constantly gets inundated with this presentation, which is updated number-wise, but -- every -- every couple of months.

And the other caveat I always make is with respect to Larry's numbers and our numbers, we -- we don't agree normally anyway, so as far as

numbers go and some other things --

MR. ELLIOTT: Mine are right.

MR. CLAWSON: Now, kids.

MR. KOTSCH: -- it's all a matter of perspective. But no, the numbers -- we do take snapshots at different points in time, plus -- of our cases and claims. And also obviously things are moving back and forth, it's a dynamic situation between NIOSH and Labor as far as the caseloads go, so it's -- I don't know that we could ever match the numbers, even on a specific day.

Just a little background on the Energy
Employees Occupational Illness Compensation
Program Act. Part B, which is the program -part of the program that we talk about here in
these meetings, became effective on July 31st,
2001. We show 72,273 cases, which encompasses
90,985 claims, have been filed as of June 16th.
The number of claims is always -- I always
mention this, too, then and Larry does, too -the number of claims is always higher than the
number of cases because cases often have more
than one claimant, especially in the -- in the
event of a survivor claim. 40,809 have been

1 cancer cases and 27,289 cases have been 2 referred to NIOSH. 3 Part E, which is the other part of the program, that DOL administers became effective on 5 October 28th, 2004. That was formerly the Part 6 D program that was administered by DOE, and in 7 that part of the program we have 52,458 cases 8 for -- and that includes 72,972 claims. 9 we initiated that program we received from the 10 Department of Energy about 25,000 cases. 11 The -- as far as compensation for the program, as of -- again, I think the 16th is the 12 operative date for most of these slides --13 14 we've compensated a total of about \$3.8 15 billion. About 64 percent of that is Part B 16 claims. That's about \$2.5 billion; \$1.9 17 billion of that is cancer; 287 would be the 18 RECA claims, the -- the miners, millers, ore 19 transporters; and then the remainder of that is 20 tied up with silicosis claims, the beryllium 21 claims -- chronic beryllium disease and 22 beryllium sensitivity type things. 23 \$1.1 billion is for Part E claims. 24 those are the -- Part E is -- in simple terms, 25 is -- are the non-cancer carcin-- I mean

chemical exposure, toxicity types exposures that early on in the program we couldn't deal with and now we can deal with on -- in that part of the program; exposures to asbestos, different chemical -- a lot of degreasers, things like that. And in complement to that, the \$226 million in medical benefits that are paid along with the claims.

As far as Part B benefit overviews -- this is just a quick one -- who's eligible, current and former employees of Department of Energy, it's contractors, subcontractors, Atomic Weapons Employers, beryllium vendors, uranium miners, millers, ore transporters who worked at the facilities covered under Section 5 of the RECA -- of RECA, which is administered by the Department of Justice, and certain family members of deceased workers.

And then quickly again, claim-- claims for Part B can be -- primarily what we're dealing with here are the NIOSH -- the ones where NIOSH gets involved with, which involve primary cancers. There's also chronic beryllium disease, beryllium sensitivity, chronic silicosis and,

again, the RECA Section 5 claims.

The claims filed for cancer under Part B of the Act, potentially any cancers covered under Part B, if it is determined that the covered employee was a member of the SEC and was diagnosed with a specified cancer -- those are the listed cancers in the -- in the Act -- or if it is determined through a dose reconstruction conducted by NIOSH that the covered employee's cancer was at least as likely as not -- which is interpreted as 50 percent or greater -- caused by radiation exposure.

The Part B -- the status under the Act of the Part B cancer claims is 40,809 cases, having 62,900 claims. That's -- have had -- okay, I'm sorry, let's start again. 40,809 cases, with 62,900 claims, 33,118 of those have final decisions. Under the Department of Labor program the case comes in, is developed for medical and employment information. If it's a cancer claim, it goes to NIOSH. They continue to develop -- develop and produce the dose reconstruction report, comes back to us at Labor, and then a recommended decision is made. That's shared with the claimant. They have the

opportunity to object, basically. Whether it's objected to or not, it goes then to the -- to what we call a Final Adjudication Branch, also inside the Department of Labor. They render a final decision, either to compensate or not to compensate, and that's what we're talking about here -- 33,118 cases have final decisions; 1,814 cases have recommended but no final decisions, they're in the process where they're with the Final Adjudication Branch; we are showing 4,192 cases at NIOSH as of June 12th; and 1,685 cases are pending an initial decision, they're in the development process or they have a dose reconstruction but it hasn't been reviewed yet or incorporated into a recommended decision.

This is the standard graphic we often show, the final decisions approved on the left, 13,176.

On the right, the denied final decisions,

19,942. That's the red bar. The other bars,
going across, 32 -- I'm sorry, 3,425 for noncovered employment at facilities that -- or
locations that are not covered under the Act;

11,963 that have probability of causation is -POC's less than 50 percent; 3,074 with

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insufficient medical evidence of a -- of a cancer; 1,100 with non-covered conditions. In the early days -- I mean that's still Part B decisions. In the early days we couldn't do anything with those. Now we can refer those -we work these cases as both Part B and E at the same time, so they would be hopefully covered under the Part E side if they weren't cancers -- or at least if not -- I mean not covered, but at least be -- be looked at under the Part E And 380 cases were denied after determinations of ineligible survivors. And Special Exposure Cohorts -- Larry talked about this -- employment criteria -- the initial ones are in the Act, the three gaseous diffusion plants, certain nuclear tests -prominently up in Alaska at Amchitka, and then of course the new SEC classes that are added -that have been added by the Board. include the specified cancers, the cancers that are listed on the specified cancer list. Causation is presumed, there's no dose reconstructions necessary for inclusion in the SEC. And the process is that HHS recommends SEC designation and if Congress -- the

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Secretary does, and if Congress does not object within 30 days, the facility becomes -- or it gets added as an SEC class.

As of June 12th we're showing, as far as new SEC-related cases, 1,803 cases have been withdrawn from NIOSH for review. Often if they're -- if they're there for dose reconstruction, an SEC class is implemented, then we with -- we compare our lists with NIOSH lists and withdraw those cases to be reviewed as far as being considered under the SEC class. 1,549 have final decisions issued; 128 have recommended but no final decisions; 52 are pending, probably for additional information; and 74 have been closed. So that's 92 percent have final decisions so far of all the cases that are affected by the SEC classes. As far as referral to NIOSH -- again, this is the 16th -- we show 27,264 cases have been referred to NIOSH; 19,618 have been returned from NIOSH. Of those, 17,373 have dose reconstructions. I'm not sure -- it's got to be a bigger number, but 23 being reworked for return to NIOSH -- oh, that's within -- within the Labor hierarchy -- and 2,222 are with --

1 have been withdrawn from NIOSH with no dose 2 reconstruction. 3 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 compensability. 21 22 23 24 25 little off. We corrected this number to match

We're showing 7,646 cases currently at NIOSH, 4,237 of those are initial or original referrals to NIOSH and 3,409 of those are reworks or returns to NIOSH, ones that had a -had an initial dose reconstruction and then -for a number of reasons, like Larry addressed, PERs or -- occasionally -- well, not occasionally, the primary driver other than PERs for -- Performance Evaluation Reports, for our returning cases to NIOSH or dose reconstructions to NIOSH is they're -- the determination that there may be a new cancer, there may be additional employment, things like that that drive us to want to send that -return that case back to have the dose reconstruction looked at again to determine whether that denied case could move towards The dose reconstruction case status -- this slide's showing final decisions for 85 percent of the cases. 17,373 cases have dose reconstructions. I think the slide might be a

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the previous slide. 14,745 dose reconstructed -- dose reconstructed cases have final decisions. 2,152 dose reconstructed cases have a recommended but no final decision -- that means they're somewhere in the -- in our FAB --Final Adjudication process. And 476 dose reconstructed cases have a recommended That's -- again, we have the dose reconstruction from NIOSH. We're just working through the -- the District Offices are just working through the process of creating the recommended decision, so that's... Again, NIOSH case-related compensation is -- is a piece of the larger total compensation. But even at that, we have \$1 billion in compensation for NIOSH-related cases. 10,380 -- I'm sorry, 10,338 payees in 6,722 cases. Of that total, \$810 million have been based on dose-reconstructed cases. That's 7,656 payees covering 5,400 -- 5,420 cases. And another \$193 million has been added due to the SEC classes. That's 2,682 payees in 1,302 cases. This slide is the paid cases under the Act, so there have been -- this is the -- yeah, these are the total numbers, 30,384 paid Part B and E

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20,521 of those have been Part B cases. That included 12,000 -- almost 13,000 cancer case payees, 5,755 RECA case payees. Again, the -- the uranium -- uranium miners, millers and ore transporters. And 1,788 other Part B case payees, primarily the beryllium and the silicosis. And 9,800-plus Part E cases. Again, the toxic exposure type cases. The last time, the Board asked -- and I -- I still want to have a -- or a more of a graph generated, like Larry generates, but I don't know how much that's going to add. But anyway, they had asked about the level of cases that we're getting in, and this still isn't quite what I think we -- you probably want to see, but it's a start anyway. The first -- the upper part is the new Part B cases received by Department of Labor monthly. Just starting recently, in March of this year, 2008, we had 354 cases; then April, 398; May, 381; and 152 in June. So those are Part B cases. I didn't bother with the Part E cases. Again, a lot of cases come in and -- or every case that comes in is considered both under Part B and Part E, but these would be specifically ones that had

cancer -- or at least cancer's a part -- as part of the particular case.

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The Part B cases that were sent to NIOSH is, again, clouded by -- or shows -- it's -- it's not just strictly each month what we forwarded to NIOSH that was a new case. Unfortunately some of these numbers -- and I wasn't able to tickle it out of it yet -- you know, some -- it would include reworks for PERs, SEC things, but primarily the rework numbers so in March of 2008 we sent 677 cases; April, 502; May, 358; and June, 119. So you would expect that number normally to be less than -- if you were just strictly looking at new Part B cases the Depart -- to the Department of Labor and cases that we then forward to NIOSH for dose reconstructions, you should expect those numbers to be smaller than the incoming because cases that come into Labor also are considered for, again, chronic beryllium disease, beryllium sensitivity, silicosis, that kind of stuff. So there would be cases coming in that would be not -- would be more than just cases that we for-- forward to NIOSH. So the beginnings of those ones are ones that are also

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including cases that we're sending back for reworks. But we'll -- we'll work at this. We're try-- I'm trying to get a better indication of what -- but that last number is actually probably not too bad, the 152 -- I always use the rule of thumb it's about 200 cases a month that we've been getting in, been pretty steady as far as Part B cases. we'll try to get a better -- I have to admit, I don't -- I'm not always sure how many cases just strictly go on to NIOSH and it -- there's also a lag there because it may come in one month and get sent to NIOSH the next month as we develop for the medical and employment that they need for -- for the -- for the -- to actually proceed with a dose reconstruction -or for even us to proceed with determining whether that case is one that we can work to a decision.

As is the case, we usually try to -- and there -- there don't -- don't appear to be too many new SEC presentations at this meeting. We usually try to provide some background information on -- on the -- the SEC classes that are up in front of the Board, just for --

for background.

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First there's the Y-12, what I call the early years, '43 to '47, which is a -- it's -- the Y-12 Plant itself has -- has other SECs and it's -- it's been one of the major cl-- you know, sources of cases in the program. Cases, both Part B and E, we've had almost 12,000 from the Y-12 Plant. We've had -- we're indicating about 2,200 NIOSH dose reconstructions and a little over 4,300 Part B final decisions resulting in 2,736 Part B approvals, 2,354 Part E approvals, for a total compensation in both Part B and E of \$50-- I'm sorry, \$531 million. The Dow Madison site, we're showing both Part E and -- Part B and E cases, 357; 3 NIOSH dose reconstructions; 99 Part B decisions; 67 Part B approvals; zero -- it's not a -- it's an AWE site so that's not covered under Part E of the program; and \$9 million in compensation. And at Spencer Chemical we're showing -- again, this slide's got a date of June 17th, we're showing 53 cases from Spencer -- I'm sorry, I guess -- you know, I'm sorry, Dow Madison is both Parts -- Part -- Spencer is both Parts B and E -- Spencer is Part B only. That means

there's no Part E evaluation, 53 cases, two final decisions under Part B and we have not --we have -- there's been no compensation at Spencer Chemical.

And that's it.

DR. ZIEMER: Okay, thank you, Jeff. Phil, do you have a starting question?

MR. SCHOFIELD: Do you have any statistics as to how many of the claimants or payees that there's been a final decision made, that they passed away before they were either notified or paid?

MR. KOTSCH: I -- I have to admit, I don't have -- I know -- I know we have statistics on that. I don't have them with me and I -- and I know that unfortunately that's not an uncommon occurrence, but I don't know -- I don't have the actual numbers. I mean I -- I know that it happens with -- with -- with some frequency that's not, unfortunately, you know, a small frequency, but I don't know how -- how often. But I can -- I can check on that number for you. I know that happens a lot and then we have to -- to proceed with, you know, developing the survivors and then just

processing that -- that, so unfortunately it takes a little bit longer, but those -- those do still get paid.

Okay. Jeff, again, we thank you, as always, for a concise update on the -- the pool of data from Department of Labor, and we look forward to continued interactions with you.

PUBLIC COMMENT

We're going to move in a moment to our public comment period. I'm going to take a brief break in order to get the list of those who wish to participate. If you wish to participate in the public comment session -- and there will be another one tomorrow as well -- but in today's session and have not already signed the paper, we'll give you a couple of minutes to get out there in the corridor and get your name on the list. And in just a moment the list will be brought in and we'll begin that session. So we're going to take about a five-minute brief break here and then we'll resume.

DR. BRANCHE: We'll put the phone on mute.

DR. ZIEMER: And the phone here will go on mute during that period.

(Whereupon, a recess was taken from 3:50 p.m. to 4:00 p.m.)

DR. ZIEMER: We're going to begin the public comment session of the Board meeting. In just a moment I'm going to ask Dr. Branche to read the redaction policy. I also want to alert the speakers that the Board has a 10-minute time limit on public comments. Also that 10-minute is considered an upper limit, not a goal to be achieved. You can think of the difference there.

We generally like to think of the public comment period as just that, comments. It is not generally a question/answer session, although sometimes we do provide -- or try to provide answers if you have certain questions. We try to avoid getting into details of individual cases. NIOSH does have caseworkers available if you have a particular question on a particular case that needs to be answered. So with that, I'm going to ask Dr. Branche to read the redaction policy in connection with the public comments.

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DR. BRANCHE: Thank you, Dr. Ziemer. person making a comment gives his or her name, no attempt will be made to redact the name. an individual, in making a statement, reveals personal information -- for example, medical information -- about themselves, that information will not usually be redacted. NIOSH Freedom of Information coordinator will, however, review such revelations in accordance with the Freedom of Information Act and the Federal Advisory Committee Act and, if deemed appropriate, will redact such in-- redact or remove such information from the transcript that is posted on the public web site. All disclosures of information -- all disclosures of information concerning third parties will be redacted. And again, if you want to bring information, during this forum or in the next public comment period, to the Board but wish not to have your identity revealed, then please speak to me on a break. Thank you. Let me be-- I'm going to take DR. ZIEMER: these speakers in the order that they've signed up. Let me ask if any of the speakers do not wish to have their names identified in the

DR. BRANCHE: If they signed up, they're agreeing to --

DR. ZIEMER: If you've signed up --

DR. BRANCHE: -- they're agreeing to have their

DR. ZIEMER: Okay, by -- just so you know, by signing up here you agree that your name will be in the record, so -- giving you that opportunity if you change your mind on that. Okay, let's begin with John Ramspott. welcome, you may approach the mike.

Testing. I'd like to thank the radiation board and the other organizations and agencies that are here today. The General Steel Industries plant has been near and dear to my heart for the last two and a half years and -- since I first asked the question that -or made a statement that I'd like to find out what actually happened at that plant. done my best, and my wife of course has assisted me in gathering information, with the help of former site experts, family members -everybody we could, includes a couple of physicists who have assisted us, and of course

members of the Board. Just by giving me your - I think courtesy and attendance and really
following what we were trying to do, I believe
we've really come to the crux of what happened
at General Steel.

I know there's probably going to be some dispute and some questions and -- I understand that 'cause not everything's perfect in this world and this is a -- an older site with a unique situation. And being privy to some of the workgroup meetings via telephone conference and listening in and actually attending one today, I can see that this is definitely seriously being taken to heart and looked at. It's a very complex situation.

There are a couple of reasons I'm going to ask for urgency, though, that I think warrant a little special attention. The recent SC&A report actually mentions there are three sites still using these devices today. Two of them happen to be government military sites.

They're noted on the Internet. One's a public site, and I've actually visited that site,

site, and I've actually visited that site,
taken photographs. That's where the operator's
manual came from. I don't think we can wait

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too long to tell these people what's going on with those machines. If we do, there's a problem with that. If there's a real hazard and they don't know it and we don't tell them -- and I can tell you from the site I visited, they think that's like a jukebox, that's -that's safe, walk in, no cooling period. like deja vu General Steel Industries all over. I have photographs. I've talked to the people. They're nice, good, solid people. I felt like I was right back at General Steel. visited that site, too. The new owner actually allowed us to go on site. We now have some video footage of it. And we've tried to share all that and we'll share anything else we have about that site. But that's a real concern. think we need to move on this as soon as we can, as humanly possibly, complex as it is. totally appreciate that, but if there've been changes and those people know about it, maybe they ought to share them with us 'cause we'll find out what they were afraid of and they changed.

Now the other issue -- and this is -- I was listening today, and I'm not going to steal

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anybody's thunder, but there was talk about radiation badges and -- I didn't hear anything about neutrons. And at that site -- and now if you read the SC&A report, there's -- there's definitely neutrons at General Steel. think they're manufactured in four ways, and they're spelled out, and I've confirmed it. The Betatron makes neutrons when it hits the -you're always trying to hit that little platinum target; it makes neutrons. And in the appendix we talk about a photoneutron activation of castings. When the big casting gets hit, that creates neutrons. When we hit the uranium with the 25-million volt Betatron beam, that makes neutrons. No one's denying that. And now one of the physicists is helping me, who is the -- actually it was [identifying information redacted] the gentleman -- the physicist who addressed this Board in Naperville via phone, his old boss, 35-year physicist, Milwaukee School of Engineering, explained to me how the fourth means of neutrons are created. When you make a neutron and that neutron impacts cement or concrete, now all of a sudden you start a chain reaction

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and you get a whole lot more neutrons. wrong, then there's a whole lot of articles I've been reading that are wrong, too. So these badges that we're talking about today, I don't think they measured neutrons. think the survey meters, which we now have a photograph of and now have the man that calibrated the survey meters -- that man was actually at a worker meeting with SC&A, he'll talk to anybody, tell you the same thing he told me, he sent it to me in an e-mail, those survey meters didn't measure any neutrons. They measured -- and they did use cesium whatever to calibrate those survey meters -they did beta and gamma above 50 keV to about 1 million, or 1.3 million. They didn't do anything for 25 million-volt -- 25 million volts. So he said John, those survey meters were a waste of time for those guys. So when they say they walked in there and they used a survey meter and they checked the casting, yeah, I guess you wouldn't get a reading if you didn't have the right tool. So those castings are a lot hotter than those guys thought. So I hope those type of things, when we start

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talking about the -- the badges and -- even at lunch today I quizzed the guys that were there. The production dates, they only had badges for -- what did I hear, '64 to '66? Well, the uranium was gone in '66 -- after '66, and it was winding down to '66. So try and use any information after '66 is a waste of time. There was no uranium there, and that's where a lot of the readings on badges would have come. And then the other thing the guy shared with me at lunch, and I didn't realize it, but they only wore the badges half the time. They wore them when they were in the Betatron. weren't Betatron workers, they were NDT workers, mostly in the Betatron, but then they stepped on the other side of a ribbon door when they were needed while somebody else kept using the Betatron and they're in there working, in Building No. 10 or No. 9. Their badge information, if we did have it, would only be half good. So that's another concern about badges. And if we try to get to the bottom of this based on badges, we're wasting a lot of time.

Now the -- the history that I've put together -

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- and I think it was valid, tried to build a good case for everybody to be able to study. We did a workbook on the site, we gave public comments, we had numerous conference calls, we did outreach meetings, we -- and I tell you what, I do salute Larry. When you said one time you were going to have the Appendix BB, it was almost like hanging a target on your back. Now we had something to shoot at. And we didn't agree with it at first 'cause we wanted all the information out, then let's do an appendix. No, it got it out in the forefront so we really have something to look at now, and we do, and I thank you for it. I thank NIOSH for it. I wrote a critique. I said I think you're wrong. I don't think you got all the information. I got a reply and we got a little differences in there and I think we can work through them. I think they're real. on that critique and a lot of other information, now we have an SC&A report. I'll tell you what, Dr. Anigstein and SC&A putting that report together, I think they really brought it to the forefront what was going on over there, what really happened. And

there's no sense in putting together a report like that unless you're going to do something with it. And I know there's some fine edges that'll be discussed, and NIOSH and SC&A'll go back and forth -- and I hope we're part of that 'cause now we've found new information that we want to present. DOE has provided us with new photographs, big signs on the building which we read when we where there, said don't get within 100 feet of this building. So what's that company do? They built another Betatron building, attached it to the main plant. That's within about 25 feet. So some new photographs actually came out of DOE. have been helping as well, and we definitely thank them.

Now, the SC&A report -- we have it, still trying to understand it all, but now we're going to wait for NIOSH's reply and hopefully we can be part of that review -- would -- I -- I know I'm not the Board, but I'd like to at least look at it, and if I see something and I can put together justifiable proof -- photographs, testimony, whatever, scientific data -- I'd like to share it. And if we can

all come to an agreement, I think we can get some people taken care of.

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And I've had a lot of these workers die recently. You know, it doesn't -- that's -- I heard a question, how many people -- I think Phil asked it, how many people die before they get paid or in the process. And one lady's so close -- I mean her husband died a long time ago and she's so close, and she just died about a month ago. Her husband was only there about 30 years, so he's probably one of the guys that's going to get compensated. So I hope we can wrap this up in a timely manner. I'll do anything I can to help. We've got workers that'll help. If we all head in the same direction, I think it could be done. So if there are any questions -- and again, the -- the neutrons -- I hope I got this right, and SC&A can correct me, neutrons come from Betatrons, they come from interaction with casting, they come from Betatrons hitting uranium and neutrons then hit concrete and more neutrons are made. So we better look at neutrons. I don't think there's any mention of that.

I appreciate your time. If there are any questions, I've got a lot of data -- you guys know I'm a data guy. I've got manuals that's now being scanned. I'll have it for you in CD/DVD, we've got photographs -- I mean from when we visited. I'll give you anything I have that'll help you I hope make a comfortable decision.

DR. ZIEMER: Thank you, John. We appreciate all the input you've given to the Board and to NIOSH over this past year or two, so...
Okay, let's proceed.

DR. BRANCHE: Clarissa.

DR. ZIEMER: Clarissa Eaton, is Clarissa here?
Okay.

MS. EATON: I also just want to add an appre-my appreciativeness of you guys's hard work,
and I hope that you always make a conscious
decision for our Cold War veterans. I think we
owe them more than what is even offered to
them, and we are severely in debt to them. I
am not a claimant, nor am I a beneficiary, but
I am a citizen and these people deserve a lot
more than what's being offered. And I hope you
will use your position to make all remedies

1 available to them. And in any instance that 2 you can expand the program and offer your 3 expertise, because I know all of you are very educated and I'm thankful for that. And I 4 5 would also like to add I -- I have recently 6 submitted a petition for the Hematite site and 7 that I hope you will give them the same 8 consideration in the future. Thank you. 9 DR. ZIEMER: Okay, thank you very much. 10 Then John -- John Dusko (sic) I think is --11 John, welcome. 12 MR. DUTKO: If I use --13 DR. ZIEMER: That's fine. I'll give you a hand 14 here. 15 (Pause) 16 John, spell your last name -- is it D-u-s-k-o? 17 MR. DUTKO: D-u-t-k-o. 18 DR. ZIEMER: --t-k-o. 19 MR. DUTKO: It's one of those. 20 I thank you for letting me speak, sir. 21 privileged -- or one of the guys privileged the 22 last -- since last October to have been working 23 with Dr. Bob Anigstein in researching and in 24 presenting evidence to SC&A. There's three

more fellas here -- Ralph Hersing*, George

1 Luber*, Eddie Brawley -- all were operators, 2 radiographers. They all knew their business. 3 Of the nine guys that gave evidence, I'd like to point this out -- and I didn't run any in-4 5 depth study -- seven of those fellas had cancer 6 of some type or other, either skin cancer, 7 prostate cancer, kidney cancer, yet we've been 8 regularly refused. Now they say we could --9 we can't get a in-depth body cancer from the 10 radiation. 11 Well, us fellows laid on those castings to 12 place it on, stood on it, crawled on them. We 13 did about everything in the world, not 14 realizing at that time -- and had no 15 information that they were activated, along 16 with the donut tube of the machine -- of the 17 compensator. 18 I understand you're going to bring in the film 19 Please do. We didn't trust them then, 20 sir, and we don't -- we won't trust them 21 tomorrow. We wore dosimeters, pencil 22 dosimeters as an instant reference, and trusted 23 them quite a bit more. 24 The biggest part of our time, if we wor-- if we 25 -- let's say we worked a shift in the Betatron,

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we might work a second shift overtime in magnaflux. When we left -- when we left that Betatron, those film badges came off and went on the rack. Now what does that tell us about that beam coming down that railroad track through that ribbon door, and we're 20 feet away working on a tank all -- or a casting of some type, what does that tell us about our dosages from the film badges? Again, I'm not trying to be a smart guy, but I see too many of my fellow workers died, sir, and there's quite a few of them not around anymore. Quite a few. It's -- it's not a laughing matter to us at all. You know, last week one fella came down with prostate cancer and another fella I worked with, Tony Gast*, prostate cancer again. these were Betatron operators and NDT people at Just in short -- in the last couple of weeks, and it seems to keep -- to keep going. And again, I have -- I apologize, I have run no in-depth study, but I think it's an indicator, sir -- it's an indicator. It should be looked into. I'm certainly not a doctor, nowhere near

it. But too many of us people -- too many of

1 us people have come down with the big C. 2 Simply neutrons is a dose introduced by Dr. 3 Anigstein, and rightfully so. We operators, 4 for the first time -- although given bounding 5 doses, your so-called bounding doses by NIOSH, 6 were the first time we saw doses, sir, in In two and a half, three years of a 7 roentgens. 8 short window of time that we had -- for 9 instance -- for instance, with -- with a 10-10 roentgen dose that I had, in two and a half 11 years window of time to three years, I wound up 12 -- by my calculations, by his chart -- with 44-13 roentgen doses. 14 Now, sir, we fired shots three inches steel, 15 nine feet double-A film, with 30 and 40 16 roentgens. And experts tell me that we can't 17 have a deep body cancer? I don't know, I'm not 18 a doctor. I just want to point these things 19 out. 20 But for the first time in 50 years we're 21 starting to find out the truth about these machines, and it's no wonder I see a lot of my 22 23 fellow workingmen not here anymore. 24 thank you very much, sir. 25 DR. ZIEMER: John, thank you very much for your

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Then we'll hear from Bill Hoppe -- Bill? And Bill is with Dow. Thank you, Bill.

MR. HOPPE: I thank you. I was looking through this here questionnaire deal like, and you put a lot of emphasis on what [identifying information redacted] says to say in the pot room they said that they brought the thorium in and they had it under armed quard the whole time, and I can't find anybody that worked in there at that time -- that was a little bit before my time -- that could verify that. Also the way he's got the -- the pots and all that is completely wrong. The -- he said on the back wall there was a workbench. Where the workbench was supposed to be, that's where we had our instruments in that. He also said that they only -- you know, we was around the pots for a short period of time. They had to sit around the pots until they could alloy them. They might be around there for 15, 20, 30 minutes at a time and couldn't get away from it, and all that fumes and that.

They also just checked the area out around seven press in the extrusion department and

1 they ran uranium on the seven press, which is 2 the heavy press, and all around. They ran 3 uranium in the rolling mill -- on the mills and everything else. You know, it's -- it's -there's a lot of confliction (sic) here that we 5 6 can come up with. 7 I've been working on this for seven years, and 8 we've been getting hold of the different 9 people. And seems like every time someone else 10 comes up with some other deal that puts 11 different things together, and we've got a lot 12 of things that comes -- you know, came together 13 that Dow does not give out any information. 14 that's why I was wanting to make sure that that 15 came up. 16 Okay, thank you. I want to ask --DR. ZIEMER: 17 maybe I'll ask Larry. Do we have Bill's input 18 on the Dow information or does someone have 19 this information? Is it new information? 20 MR. ELLIOTT: I think he's speaking about the 21 evaluation report that's going to be presented 22 at this meeting, are you not? 23 MR. HOPPE: (Off microphone) (Unintelligible) 24 MR. ELLIOTT: And I don't know that we've had 25 his particular input on that at this --

1	DR. ZIEMER: Okay
2	MR. ELLIOTT: We're getting that now, so
3	DR. ZIEMER: Well, okay. We we need to make
4	sure that yeah.
5	MR. ELLIOTT: So we have had it
6	DR. ZIEMER: Yeah.
7	MR. ELLIOTT: had some of this since June
8	some of it since June.
9	MR. HOPPE: (Off microphone) (Unintelligible)
10	Silverstein's (unintelligible) came
11	(unintelligible).
12	DR. ZIEMER: Okay, I just wanted to make sure
13	that the information got inputted to the NIOSH
14	folks. Thank you, Bill, for adding that.
15	I need to check and see if there are any folks
16	on the telephone lines that wish to make public
17	comment. They would not of course had an
18	opportunity to sign up. Anyone by phone that
19	wishes to comment?
20	UNIDENTIFIED: Yes, I do.
21	DR. ZIEMER: And identify yourself, please, and
22	
23	MS. KLEA: Yes, I'm Yvonne Klea. I'm the
24	author of Petition No. 93 for E-Tech, which is
25	the Santa Susana Field Laboratory out here in

1 California. 2 DR. ZIEMER: Very good. Please proceed. 3 MS. KLEA: I just kind of have a question comment. In our site profile it mentions that 5 background dose was subtracted from our dose. 6 Now there's no mention up there in the site 7 profile of where they got the background doses, 8 what study they used, and every study that has been written for the facility has been 9 10 questionable on where they picked up their 11 background dose. And the Department of Energy, 12 right now at this time, is working with the EPA to figure out an accept -- acceptable background 13 14 dose. So I have a question about using a 15 background dose to subtract from our estimated 16 doses. 17 DR. ZIEMER: Okay. I think we probably can't 18 answer that question today, but the NIOSH 19 people here are aware of your question and they 20 can follow up on that with you. So thank you 21 very much for that input. 22 MS. KLEA: Yes, thank you. 23 Uh-huh. Is there anyone else on DR. ZIEMER: the line that wishes to comment? 24

(No response)

1 Apparently not. The Chair's aware that Mr. 2 Stephan has joined the assembly and I never 3 want to pass up an opportunity to have you comment if you wish to. He doesn't wish to. 4 5 MR. STEPHAN: Tomorrow. 6 DR. ZIEMER: He's going to comment on not 7 commenting, okay. Thank you. 8 Let me ask if there's anyone else that didn't 9 get a chance to sign up but now has the courage 10 to -- I shouldn't put it that way. It's like, 11 you know, eating powdered milk biscuits; it 12 gives you the courage to get up and do what you 13 need to do. Right. If not, we are recessed until tomorrow morning. 14 15 Thank you very much for your participation 16 today. 17 DR. BRANCHE: At 8:30. 18 DR. ZIEMER: 8:30 tomorrow morning. 19 DR. BRANCHE: We will close the telephone line 20 now, and at 4:45 reconvene for the Blockson --21 for the Blockson workgroup meeting that will be 22 in this room. Thank you. 23 (Whereupon, the meeting adjourned at 4:30 p.m.)

CERTIFICATE OF COURT REPORTER

STATE OF GEORGIA COUNTY OF FULTON

I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of June 24, 2008; and it is a true and accurate transcript of the testimony captioned herein.

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

WITNESS my hand and official seal this the 24th day of July, 2008.

STEVEN RAY GREEN, CCR, CVR-CM, PNSC

CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102