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

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

convenes the

THIRTIETH MEETING

ADVISORY BOARD ON

RADIATION AND WORKER HEALTH

DAY THREE

The verbatim transcript of the Meeting of the Advisory Board on Radiation and Worker Health held at the Crowne Plaza Five Seasons Hotel, Cedar Rapids, Iowa, on April 27, 2005.

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#### TRANSCRIPT LEGEND

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#### PROCEEDINGS

1 (8:15 a.m.)

#### WELCOME AND OPENING COMMENTS

DR. ZIEMER: I'm going to call the assembly to order. This is our third day. I think we all feel like we're sort of on the home stretch, as it were. We -- we are going to continue with our discussions on Mallinckrodt as soon as we clean up the coffee here at the front.

(Pause)

A couple of reminders -- again, as I always do
-- to remind you to register your attendance in
the registration book at the entryway. Again
I'll remind everyone there are copies of the
agenda and related materials on the table in
the back.

Let me give Lew Wade the mike a moment to see - Lew, do you have any additional comments this
morning?

DR. WADE: Well, there was just one comment made by a Board member that a petitioner or two asked last night if we could just very briefly review what happens to a petition once it's been approved by the Board. And I thought I would just walk through that in the broadest possible terms and I would ask Liz to correct

the mistakes that I made, but as you know, when the Board takes an action there are requirements as to the materials it needs to submit to the Secretary. And by your procedures yesterday you've tasked your Chairman with submitting your recommendation within 21 days. Once that submission is made by the Board to the Secretary, it really goes to the NIOSH director, who prepares a proposed decision for the Secretary. And again I remind you that the NIOSH director, in preparation of that proposed decision, will consider the evaluations of NIOSH, the report and recommendations of the

decision.

Board, the information presented or submitted to the Board and the deliberations of the Board. That package makes its way to the Secretary and the Secretary makes the final

A clock starts -- a 30-day clock starts on the day the Secretary receives your recommendation. He has to send a decision to Congress within 30 days. So that clock starts and that will cause a great deal of action to take place within the Executive Branch to prepare the Secretary to

1	see that he makes his decision package
2	available to Congress within 30 days.
3	Liz, anything else that needs to be
4	DR. ZIEMER: Then what happens?
5	DR. WADE: And then Congress acts upon that
6	package as it as it sees fit.
7	DR. ZIEMER: I believe Congress also is
8	mandated to act within is it 30 days?
9	DR. WADE: Thirty days, yes.
10	DR. ZIEMER: Thirty days.
11	DR. WADE: And if they don't act, then the
12	proposed decision becomes effective.
13	MS. MUNN: Thirty calendar days?
14	DR. WADE: Thirty calendar days? I assume.
15	MS. MUNN: If they're not in session, that
16	could be an issue.
17	DR. ZIEMER: One other item I would add
18	parenthetically, Board members, you are aware
19	and we have the letter which we approved.
20	Please understand that I will have to append to
21	the letter an additional paragraph describing a
22	number of attachments which are required, and
23	you'll understand I assume you're
24	comfortable with that. There will be a letter
25	describing that we will attach the minutes of

our meeting, which include the testimony of the petitioners, the -- the documents that are delineated in the rule itself, so there will be that additional paragraph.

#### MALLINCKRODT SPECIAL EXPOSURE COHORT PETITION

DR. ZIEMER: Now we're ready to proceed with our deliberations concerning the Mallinckrodt Special Exposure Cohort petition. We begin this morning with NIOSH presentation, and Larry Elliott is going to present that.

And Larry, let me just also preface your remarks by reminding the assembly of the action that this Board took in -- at the St. Louis meeting, and I'm reading from the minutes relative to that petition -- (Reading) The Board reserves judgment with respect to Mallinckrodt workers employed during the 1949 to 1957 time period until review of newly-located raw data is complete. This material may provide additional pertinent information on monitoring programs and worker exposure for that potential cohort.

And there was discussion on that motion and at the end of that discussion there is an indication -- I don't -- I'm looking for it

here now, but -- oh, here it is. A second motion made, seconded and approved -- (Reading) It is the intent of the Board to make a final determination on this potential cohort at the next full Advisory Board meeting.

So that is the backdrop for where we are today.

### PRESENTATION BY NIOSH

MR. ELLIOTT: Thank you, Dr. Ziemer. Good morning, ladies and gentlemen of the Board and the audience. I'm here again to shift your focus from the discussion you had yesterday afternoon on the site profile for Mallinckrodt chemical workers to again look at the Special Exposure Cohort petition and our evaluation report of that.

As is becoming tradition within this kind of presentation, I will walk you through the petition evaluation process. I know this is probably becoming old hat by now, but I must do this for the record as well as for the benefit of the audience, who may not have heard this type of presentation before. I'll also give an overview and a timeline of this particular petition.

I'll again remind the Board, probably

1 unnecessarily at this point, but I'll remind 2 3 4 5 6 7 8 9 10 11 12 13 14 15 petition. 16 17 18 19 20 21 22 23 24 by rule to determine the completeness of our 25 data resear -- data search. We have to evaluate

you of your responsibilities and your role within this part of the process. I present also a summary of the petition evaluation report, as well as go into the details of the supplement to that evaluation report that was prepared after the February meeting. And I'll conclude with the class definition and the summary findings. Again, the evaluation process is governed by the statute and by the regulation that's listed on this slide, 83.13(c)(1) and 42 CFR 83.13(c)(3). Again, these are the two-pronged tests that must be addressed in evaluating a As well the evaluation process must examine all available data and information obtained. includes the site profile development, information that is pertinent to this facility that may have been addressed at related facilities, the dose reconstructions that may have been attempted or -- and/or completed with regard to the petition. We are also required

the sufficiency of data according to the HP -health physics hierarchy of data that's listed
in our dose reconstruction rule. And we
evaluate the issues of data reliability that's
raised by the petition itself.

As well, if we determine that we cannot reconstruct dose, we then have to address the second prong of the test and whether -- that includes whether health was endangered for this particular class.

Let me speak for a moment here about -- a little bit about -- in reflection of the discussion on the site profile yesterday afternoon, and I'd like to point out that I think there may have -- may be some confusion about site profiles and their -- their content and context as a living document. And in that light, these documents, from their genesis, have been portrayed as living documents. When we -- it goes back to the need to be timely in our efforts for -- for the claimant population in doing dose reconstruction, as well as evaluating petitions here. And so once we feel that we have assembled enough information to process certain types of claims under our

efficiency process, we put that into play and start using those documents, fully recognizing and hopefully trying to make it clear to people that -- with an understanding that as new information comes to light that may inform and enable us to treat other types of cases, we modify those documents.

A revision to a site profile we would consider as giving us the ability to move claims through the system. And as new information we make changes to those site profiles. Certainly we, again, appreciate our colleagues from Sanford Cohen & Associates who bring their perspective to this and call our attention to certain things that we feel and they feel we need to address. And once we address those things, we go back and we look at all claims that were processed under our previous site profile revision and evaluate whether the modifications to a newly-revised site profile would affect the outcome of those claims.

This is a standard operating procedure within my office. It's called a performance evaluation report and any time that a change occurs in any of these documents we go back and

evaluate those claims. If there is a change to a claim, we notify the claimant, we notify the Department of Labor and we work together with both and work through all that.

So I just offer that for the consideration of the Board, as well as the audience.

Let me speak a little bit about the overview and a timeline for this particular petition.

Destrehan Street petition was submitted to us on July 21st last year, and the initial definition for that particular class, as we worked it out with the petitioners, were all employees that worked in the uranium division at the Mallinckrodt Destrehan Street facility in St. Louis, Missouri over the time period of 1942 to 1957.

The submission of this petition met the criteria as outlined in our rule under 42 CFR 83.7 through 83.9, and it was qualified on November 24th, 2004. As I said yesterday, we -- we work diligently with the petitioners to make sure that the basis of the petition addresses all of the requirements within our regulatory process.

The petitioners were notified by letter, and a notice of that submission -- that it had qualified for evaluation -- was published in the Federal Register and put on our web site on December 20th in 2004.

NIOSH evaluated this petition using the guidelines in 83.13, and we submitted a summary of findings and a petition evaluation report to the Board and to the petitioners on February 2nd, 2005. A summary of the evaluation report and finding was then published in the Federal Register on February 3rd, 2005.

At your February 8th meeting in St. Louis we presented the evaluation reports for Mallinckrodt. This eval— the evaluation reports — there were two, if you recall. One report spoke to all DOE, DOE contractors or subcontractors employed by the uranium division of Mallinckrodt during the period from 1942 through 1945. The other report covered the same workers and employees for Mallinckrodt through the period of 1946 through 1948, and then the last part of that report covered 1949 to 1957. We were, as you recall, seeking — for the latter time period we were seeking

advice from the Board concerning the matter of data reliability.

During that meeting a number of additional issues concerning access to data, their reliability and various technical matters that pertained to the time period of 1949 to 1957 were identified during the presentation that we made to the Board. And in response to those issues and others, we submitted a -- or we developed and submitted a supplemental report to our evaluation report that addresses those issues.

The Board sent a recommendation to the Secretary of HHS on March 11th, 2005 recommending that a Special Exposure Cohort class be designated according to all -- accorded to all DOE contractors or subcontractors who worked at the uranium division at Mallinckrodt Destrehan Street during the period of 1942 to 1948. We -- the Board, based upon a recommendation that we made, combined the '42 to '48 period. It was broken down in '42 to '46, if you recall, and they -- or '42 to '45 and then '46 to '48. The Board reserved judgment with respect to workers

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employed during the 1949 to 1957 time period until we -- NIOSH -- had completed its supplemental report.

On April 6th of 2005 the Director of NIOSH sent a recommended decision to the Secretary of HHS that was consistent with the Board's recommendation.

Based upon the considerations of the recommendations, the findings and the deliberations of the Board and the recommendations of the Director of NIOSH, and also the Director of Centers for Disease Control and Prevention, the Secretary of HHS sent his decision to Congress on April 11th, 2005 to add the -- this class to the Special Exposure Cohort. The class definition is (reading) The employees of the Department of Energy or DOE contractors or subcontractors employed by the uranium division of Mallinckrodt Chemical Works Destrehan Street facility during the period of 1942 through 1948, and who were employed for a number of work days aggregating at least 250 work days, either solely under this employment or in combination with work days within the

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parameters, excluding aggregate work day requirements, established by other classes or employees included in the Special Exposure Cohort.

I'm not going to belabor the Board with this, your roles and responsibilities. I think you're very well familiar with them, but for the audience, the Board's responsibilities are governed by statute and by the Special Exposure Cohort rule that was promulgated last May. The main role of the Board is to consider and advise the Secretary of HHS on whether to add a class. Again, you are to consider the petition, the evaluation reports and all available information, and develop and transmit to the Secretary your recommendation. that recommendation, as you know, you must include all of the relevant petitions and any information that is pertinent to that. Lastly, we all are required to adhere to the Privacy Act and control information and prevent unwarranted disclosure of information about the petitioners.

And now I'll go into the summary of the Special Exposure Cohort evaluation report that we

provided for the class with the time period of 1949 to 1957.

Beginning in 1949 Mallinckrodt had established an operational program of radiation monitoring of its employees and its work areas, and this was overseen and directed by the Atomic Energy Commission's Health and Safety Laboratory out of New York. Notwithstanding the data reliability concerns that have been raised, there is sufficient information from the various monitoring activities, together with the information on radiological sources and processes, to reconstruct and validate dose estimates.

After 1949 there are multiple sources of exposure information, and through Dr. Neton's attempt to illustrate in his example yesterday, we feel that we can use the various data sources to complement each other and evaluate any deficiencies or data gaps that might exist in that dataset for Mallinckrodt.

In the SEC petition evaluation report 00012-2, section 7.3, items 2, 3 and 4, you'll find that we address how we might go about doing this.

For example, in item 2 we talk about breath

radon and the limited number of data and the use of that data with regard to the entry of zeroes in that data. We indicate that urinalysis results are the solution for evaluating that kind of a data gap or that type of limitation in the data.

For item 3, lost medical records -- that is an issue that has been raised and one of concern - NIOSH is -- has documented in its search that it did not indicate any loss of medical records, and thus we cannot confirm that medical records were lost, in fact.

With regard to item 4, altered records or conscious cover-up referencing a 1949 dust evaluation which was never finalized, our belief is that the cure for that particular situation is the availability of data from a fully operational health and safety health physics program from 1949 to 1957 that included the oversight of the AEC/HASL.

I won't go over -- this is just another slide of some of the data that was presented in the evaluation report, and you saw this yesterday, as well. And Jim I think did a very good job of summarizing the wealth of information that

exists in the variety of monitoring data that we have.

The purpose of the supplement report was to address a number of issues that were identified but not specifically addressed in the prior NIOSH petition evaluation report. The supplement identifies those issues that were raised at your Board meeting in February in St. Louis, and provides an evaluation of the effect that those issues might have -- might have on the proposed designation of this class from 1949 to 1957.

I'll summarize the issues that were raised.
There were five that are listed here. NIOSH's access to data and reports identified in the 1972 Mason letter as being potentially lost or destroyed. Another issue, items raised in the 1974 Mont Mason notes presented at the February Board meeting. And third, a summary of the content of the six boxes of the Mallinckrodt data that were unevaluated at the time of the February Board meeting. Four, there was concern as to how to calculate exposure from isotopes other than uranium if urine samples were only counted for uranium. And five, the

ability of NIOSH to estimate potential radon -radiation exposures in doses associated with
the blowouts that occurred at Mallinckrodt.
With regard to issue number one, in the
supplemental report we specify that NIOSH has
recovered a record transmittal and receipt form
from the Federal Records Center in St. Louis
which identifies the 22 documents referred to
in the Mont Mason letter, and these documents
were identified as shelf V2161. In that NIOSH
has obtained 21 of the 22 documents, which
consist of reports associated with dust studies
and other facility surveys.

The only document that was not located was entitled "An Annual Report for the Fiscal Year 1950 from the New York Operations Office of Health and Safety Division" which was dated November 13th, 1950. This missing document is an annual report and is -- these kinds of reports are very helpful to us in documenting the reason behind a particular action. But the environmental health and safety data that's summarized in these reports are usually found in other types of documents and other types of reports. So we don't feel that just because

this one report is missing that it leaves us
with a huge data gap. We feel that we can -we can use other information to cover the
issues that might be raised from that lost

document.

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This slide explains the effects of the information that's compiled in the supplemental report, and the data that it's obtained from the documents would not increase the estimated doses calculated from the data that's already in the TBD. It probably, if we obtained additional information -- as I said yesterday for Iowa, additional information would tend to provide more precision in our dose estimates and perhaps even drive them down. The newlyobtained data will be fully analyzed and would -- will be included in the next revision. NIOSH has -- has -- does not find that the records have been lost or destroyed. We feel that most of the data from the documents was found in other sources and was already evaluated and -- and most of it is used in Rev. 1 of the TBD. Those that are not used or not been incorporated will be incorporated.

Item number 2 regarding the 1975 Mont Mason

1 notes as presented in the February Board 2 meeting, and these notes were generated from a 3 trip that Dr. Mason took in 1975 to gather data 4 for an epidemiologic study for Thomas Mancuso. 5 The funding ran out for that study, evidently -- we understand that from other documents --6 7 and it reflects -- this particular document 8 reflects an evaluation in progress. It is not 9 a completed evaluation of the data. 10 Although the study was not finished, the site 11 data have been evaluated and the issues that 12 were identified were revisited by other studies 13 and in other epidemiologic reports. These data 14 were reviewed by NIOSH and ORAU for the 15 development of the Mallinckrodt Technical Basis 16 Document. 17 Based on NIOSH's review of the data, NIOSH does 18 not find any issues that would preclude an 19 ability to reconstruct doses for compensation 20 purposes under this program. 21 With regard to item number 3, this issue being 22 the summary of the content of the six boxes, 23 there were six boxes and two of the boxes 24 contained information from Weldon Spring, so 25 they were collapsed into five total boxes.

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for box one, Weldon Springs, with == about 75 percent of the contents of that box are film badges for 1945 to 1958 for Weldon Springs; 25 percent of that are -- that are left are air sampling data, breath analysis and contamination survey results and radon monitoring for Weldon Springs. And I'm sure that our ORAU contractor will be incorporating that information into Weldon Spring's Technical -- site profile, which I hope we see this week, as Dr. Toohey mentioned last night. Box two also contained Weldon Springs information. About 75 percent of its contents are film badge reports for 1957 to 1961; 25 percent are bioassay data from 1961 to '65. There is breathing zone air sampling data from '63 to '65, contamination survey reports for 1959 and air sampling in the building 301. Box three again contains Weldon Spring's film badge information, and box four contained Mallinckrodt dust studies from plant 4, 6, 6E, 7, shotgun lab and K-65. I would just mention that the distribution of the film badge reports from '57 to '61 have appeared -- appear to us to be pretty random in their order in our

1 evaluation of that particular box. 2 Box five, Mallinckrodt and Weldon Springs 3 information. Approximately 25 percent are for 4 the years of 1946 to '49 and they are film 5 badge records for Mallinckrodt; 75 percent of the box is for Weldon Spring's badge records, 6 7 and approximately half cover 1959 to 1965. 8 other half of that -- for Weldon Springs are 9 bioassay data from '59 to '65 and area 10 monitoring trends. 11 The initial review of these documents indicates 12 that some of the data are duplicate data that 13 we had previously accounted for and 14 incorporated into our Technical Basis 15 Some documents provide additional Documents. 16 data that will also be addressed in a revision 17 of the Technical Basis Document. 18 With regard to item number 4, how will 19 exposures for isotopes other than uranium be 20 calculated if urine samples were counted for 21 uranium only, the inclusion of other isotopes 22 in claimant-favorable assumptions can be made. 23 We can do this through a -- specifically for 24 source materials in which uranium isotopes were 25 predominant, for source materials in which

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radium 226 was predominant, and for source material which concentrated thorium. For air samples which measured gross alpha only, there are also general instructions and assumptions to be followed for conducting dose reconstructions, and those are found in our Technical Basis Document.

The last issue that's spoken to in the supplemental report, and that was raised at the February meeting, was can NIOSH estimate potential radiation exposures and doses associated with the blowouts. A blowout is an event that occurs during the reduction of uranium tetrafluoride to uranium metal. And the existence of a routine urinalysis program after 1948 allows NIOSH to put an upper limit on doses associated with blowouts or other incidences involving internal exposures. can assume uranium excreted by workers on a routine monitoring program is a result of a blowout that occurred immediately after all the workers submitted their previous samples. resultant dose calculation will provide a maximum plausible estimate of the exposure that was incurred. The calculated dose in -- in a -

- all likelihood then would be an overestimate because the urine samples would also include uranium inhaled from routine operations during the work in that plant.

To refine these values, NIOSH can also review information provided by Mallinckrodt workers during their interviews. When such incidents are identified, NIOSH would review the Energy employee's bioassay records for sample -- incident samples and also look at coworker data in that same regard.

The petition evaluation supplemental report addresses the petitioners' concerns regarding reliability and validity of the data. For the years 1949 to 1957, data reliability concerns notwithstanding, NIOSH finds that the radiation dose estimates can be reconstructed and validated for compensation purposes.

NIOSH welcomes the advice of the Board on the weight of evidence determinations involving data reliability for this and future petitions. Specifically, we asked you to consider this in your February meeting and we're looking forward to hearing your deliberations on this point at this meeting.

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In conclusion, the proposed class definition that we offer in our evaluation report is that all DOE, DOE contractors or subcontractors who worked at the uranium division at the Mallinckrodt Destrehan Street facility during the period from 1949 to 1957, we find that for that time period we can do dose reconstruction with sufficient accuracy to achieve either a maximum capping dose or a more precise dose estimate, and therefore we did not have to attend to health endangerment under the Special Exposure Cohort petition. But I would offer that -- and I think we are ready to say that health was certainly endangered to this work force by the type of work that they did and they exposures that they had. Again, I'll remind the Board, for the other two classes which are shown here, '42 to '45, we found it was infeasible to do dose reconstruction with sufficient accuracy, we felt that health was endangered. And for the period of '46 to '48 we also felt it was infeasible -- not feasible to do dose reconstruction, and again that health was endangered for that particular time frame.

That concludes my presentation and I'll stand ready to entertain questions.

#### BOARD DISCUSSION OF MALLINCKRODT SEC PETITION

DR. ZIEMER: Yes, thank you, Larry. Let's open the floor for questions. Dr. Melius?

DR. MELIUS: Got a couple of questions that came up last night that are sort of -- I think are -- think are relevant, though they're not directly related to your presentation. One, is there a petition -- I guess we know from your talk and from what was said last night that for Weldon Springs there's a -- a site profile that's about to go into review with -- at NIOSH coming in from your contractor. And is there also an active -- is there a petition for a Special Exposure Cohort relevant to Weldon Springs?

MR. ELLIOTT: No, I don't believe we have a Weldon Springs petition at this point in time.

DR. MELIUS: Okay. In terms of the work force there, I was a little confused by what was -- the interchange last night. To what extent is there an overlap between the two facilities?

MR. ELLIOTT: Well, as you -- as you heard last night, and I think perhaps Denise or somebody

else is better able to speak to this, there has been a -- there was a large migration of workers from Destrehan Street to Weldon Springs as they ramped up. I don't know what the specific numbers are, but I think I heard from Denise last night probably 50 percent. I don't know if anybody else in the audience can help us here with that, but...

DR. MELIUS: So you've not analyzed that overlap or have information that would -- MR. ELLIOTT: I don't have it at -- at my disposal right now.

DR. MELIUS: Okay. Another question -- this is related to more what you do with site profiles and -- and your -- your analysis of them. Are you -- when you're doing these site profiles and dealing with them in the context of a Special Exposure Cohort petition, do you go -- you're providing sort of a general evaluation. Do you evaluate subgroups at all within that -- within the work force in terms of your ability to do dose reconstruction?

MR. ELLIOTT: Yes. Yes, we do. In fact, I think it's apparent that we do that in this petition, as well as the Iowa petition that we

1 presented to the Board, where we have 2 identified different classes based upon 3 subgroupings -- or -- or events that occurred 4 in that particular facility. 5 DR. MELIUS: I understand events. I don't see 6 subgroupings necessarily in terms of --7 MR. ELLIOTT: Well, the radiographers -- the 8 radiographic technicians for Iowa --9 DR. MELIUS: No, stay in Mallinckrodt, please. 10 I don't --11 MR. ELLIOTT: Stay in Mallinckrodt --12 DR. MELIUS: Yes. 13 MR. ELLIOTT: We do evaluate the different process and the types of tasks that were 14 15 employed in a process, the workers that moved 16 through the process. We're very much 17 interested in -- in not only the highest 18 exposed workers, but those workers that were in 19 a lower or moderate-exposed categories and whether or not we have enough data, enough 20 21 information to evaluate their -- and 22 reconstruct their exposure. 23 DR. MELIUS: Yeah, but -- but -- so -- but do 24 you do any sort of a systematic analysis that's 25 in a -- in the form of a report or... I mean -

1 - I mean you're telling me you generally --2 MR. ELLIOTT: Well, yeah, the --3 DR. MELIUS: -- I mean --4 MR. ELLIOTT: -- the analysis is systematic in 5 the sense that we examine the process and all of the information and all the dose data, the -6 7 - the programs that were put into effect to 8 provide protection to the work force, who was 9 monitored, who was not monitored. Do -- do we 10 report that specifically in a separate report; 11 no, it's all rolled up into our evaluation 12 summaries. Could we do a better job of that; 13 yes, I suppose we certainly could. 14 DR. MELIUS: Okay. That's all my... 15 Okay. Other questions? Yes, Dr. DR. ZIEMER: 16 Anderson. 17 DR. ANDERSON: Yeah, Larry, we saw yesterday 18 the -- kind of the process you go through in 19 the validation, comparing the different data 20 sources to look for discrepancies or gaps or 21 whatever. And my question is, have you done that -- can you say that -- after having done 22 23 that, that you have validated the data, or are 24 we still into you -- you believe you have the

means to try to do that and you'll do that as

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you move along, but it hasn't been done yet.

Is that -- that's my question.

MR. ELLIOTT: Jim, you want to come to the mike and speak to this specifically? This is down in the weeds for me, I'm...

(Pause) DR. NETON: I think we need to be a little careful when we say "validated the data." I think -- it's our position, at least -- that the data that we have, the urine samples and air samples, are -- are -- were properly processed. Now to the extent of this data integrity issue and analyzing that for how we could use the data, we have qualitatively gone through and looked at the data, but we have not yet completed a detailed quantitative analysis to demonstrate that we have reached that conclusion. I think -- I tried to indicate that in my presentation yesterday by the hypothetical example that was discussed last night.

DR. ANDERSON: 'Cause -- 'cause I was just looking at the one slide here, basically, saying that you believe you can reconstruct and validate the dose estimates. My question is

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1 if, when you get into it, you can't, then where 2 are we left? 3 DR. NETON: Right. Well --4 DR. ANDERSON: I mean you gave the --5 DR. NETON: Yeah. DR. ANDERSON: -- the demonstration of when 6 7 there is a discrepancy, then you're sort of 8 left with well, you either have to believe one 9 or the other, or say ever-- all of it's suspect 10 and then --11 DR. NETON: Yeah. 12 DR. ANDERSON: -- then we're sort of into --13 you go from a lot of data to very suspect data 14 and that's --15 DR. NETON: I think what we're trying to speak 16 to here, though, is we have three -- three 17 levels of data, the source term, the air -- the 18 personal monitoring data and the air data. 19 With those three pieces we can do a comparison, 20 as I suggested. But what we're really saying 21 here with this analysis is that, based on the 22 site profile, we believe that not only can we 23 cap the dose, but in almost all instances we 24 can do much better than that. We believe we 25 have sufficient data to do far better than just capping.

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Now if the analysis indicates, for instance, that both air data and bioassay data are suspect -- which I -- I don't want to prejudge, but so far my qualitative analysis doesn't indicate that -- then we're left with source term, and certainly then we can use those data to cap, to put upper maximums, which is relevant for the SEC petition evaluation. I think you have to be careful not to confuse the profile that was designed to do better than maximum dose estimates, to do more detailed analyses, with the requirements of the SEC petition, which is can NIOSH cap the dose. And I think we would assert that given the quantity of the monitoring data we have, plus the knowledge of the source term and the detailed, hundreds of pages of descriptions of the processes, that we can certainly cap that dose. MR. ELLIOTT: To add to that, I think there are various levels and ways of validation. just the fact that in our site profile we identify issues relevant to data is calling recognition to that, and that, in and of itself, is some things that we use to look for

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other data to determine whether or not we need to -- is the -- is the credibility or reliability of the data suitable or unsuitable for use in dose reconstruction, does it prevent us from doing a credible job of dose reconstruction. So we try to bring that to the fore. There are different ways of going through a validation process. Jim's example -maybe it was poorly -- poorly presented or poorly designed yesterday, but it was just an example to illustrate one of those ways. fact that we raised the blank issue in our report is another way. These things all have to be taken into consideration in validating data.

DR. ANDERSON: Yeah, I was just trying to step through our task of, you know, how -- how much -- there seems to be a lot of data. The question is -- and that -- I'm not sure it's been resolved. I mean we have a method for -- or you presented a method to go through how you would go about validating, but one of the things we have to decide is -- is there valid data --

MR. ELLIOTT: Is it feasible to even validate,

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yeah.

DR. ANDERSON: -- is it feasible to validate, and then when you think in terms of Iowa, if you're left with capping, then is the capping process giving you sufficient accuracy. you've shown you can cap, but the question is how meaningful is that -- that cap, and that -that's why I was trying to sort this out. I mean qualitatively it's -- it's good to hear you've looked at it and it -- it seems, but that's kind of where I'm at at this point. DR. ZIEMER: I'd like to pose a question, Larry, or perhaps to Jim. This relates to the missing document from the Health and Safety Lab. I assume there are other annual reports of that type in your data bank or in your collection. Can you characterize the kind of information those reports typically contain? assume they're not necessarily focused on Mallinckrodt. They would have some sort of summaries from across the complex, or what -what can you tell us? I put you on the spot here, but -- but I -- there are -- there are other such annual reports that you've looked at?

1 DR. NETON: Yes, this -- this is the AEC annual 2 report that you're referring to? DR. ZIEMER: Well, it's the one -- the missing 3 4 document that was in -- in the box that was --5 Right. I think we've summarized in DR. NETON: 6 our slide -- maybe Larry -- could you go back 7 to that slide where we talk about what's in 8 there? But in general, these are -- these are 9 broad brush descriptions of the processes, the 10 work force, the general issues that were 11 identified, some summary air sample tables 12 possibly --13 DR. ZIEMER: You would not expect them to 14 contain raw data from Mallinckrodt, necessarily 15 -- or even detailed data from Mallinckrodt. 16 DR. NETON: That's correct, these would be 17 upper-tier documents that would rely on more of 18 the raw air sample data, which we believe we 19 have in most cases. So I don't expect that 20 there would be new information that -- that 21 would not exist somewhere else, if that's what 22 you're referring to. 23 DR. ZIEMER: Yes, that's -- I wanted to clarify 24 that. 25 DR. NETON: Yeah, I'm sorry, I didn't mean to

make a --

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MR. ELLIOTT: These kind of reports summari -- I believe you're right, Dr. Ziemer. These kind of reports summarize issues or activities or problems identified across -- across the complex. They're -- they're a roll-up document. There are several other supporting documents specific from a given site that gets rolled up into this general -- general report. DR. ZIEMER: What I was kind of getting at, if you're familiar with other reports -- for example, would one expect to find a Mallickrodt section in such a report or would the Mallinckrodt data typically be rolled up statistically with other data, anyway? MR. ELLIOTT: I believe the answer to that is yes and no. I think it depends upon the type of incident or the type of issue that's being reported. And if it was critical enough to merit --

DR. ZIEMER: Special attention.

MR. ELLIOTT: -- special attention, it would have said Mallinckrodt had this issue going on, be aware of it for the other sites. And in some cases -- and Jim or Stu can correct me if

1	I'm wrong there would even be listings of
2	summary data where you may not be able to pick
3	out the Mallinckrodt component of that data.
4	DR. NETON: Right, it could be both ways,
5	that's correct.
6	DR. ZIEMER: Thank you. Roy DeHart?
7	DR. DEHART: In February and again last night
8	and today, the issue of quality and validation
9	of the data has been brought up and discussed.
10	Is there evidence, or perhaps even a
11	suggestion, of issues tied to the quality of
12	the data?
13	MR. ELLIOTT: Is there evidence or issues tied
14	to the quality of the data?
15	DR. DEHART: Correct. Is is the data
16	questioned because of of something you have
17	found or something that that is missing that
18	would question the quality of that data that
19	you you're using.
20	MR. ELLIOTT: That's a matter of perspective,
21	of course, but
22	DR. NETON: I think could I maybe chime in?
23	MR. ELLIOTT: Go ahead.
24	DR. NETON: I think you'll find in the 248-page
25	document there's going to be sections that

discuss some of the issues that arose over the monitoring practices. I mean this is a standard -- things that happen when you measure samples. I -- I don't believe, I don't think OCAS believes -- I know OCAS does not believe that these issues that are raised rise to the level that would question the ability to use the data for dose reconstructions.

MR. ELLIOTT: Yeah.

DR. NETON: For example, I'd like to -- this is a good opportunity possibly for me to address an issue that was brought up last night related to the blank samples. There was a -- it was brought up in the Board discussion and also at the evening session that there was some indication in our profile that blank samples were contaminated, and that's true, and I'd just like to read the relevant two sentences, if I may, that are in that profile that were refer-- being referred to.

And I quote (reading) An undated, untitled urinalysis listing found in dose reconstruction project files indicates that closed blank samples were found to have significant levels of uranium in them, indicating contamination in

the laboratory. It was suggested that this might explain the high levels of some of the non-blank, parentheses, worker samples. Thus at least some of the early urinalysis samples must be considered to have been potentially contaminated; i.e., some of the uranium content may have come from the laboratory analyzing them.

So the issue here is that -- this speaks to the quality control, actually, that they were running what I would call in current parlance a method blank. You run a blank with a -- every so many samples and you want to make sure that you're not reporting erroneously positive results because you may have contamination in the lab. And that appear -- that seems to be what's happened here. So if anything, those blank samples would have -- if there was contamination in the laboratory, would have biased the results for the workers high, not low.

So you'll see things like this in here, but again, I've looked through these and I don't believe that any of these rise to the level of really making them unusable for dose

reconstruction.

MR. GRIFFON: Can I just -- Jim, while you're
at the mike --

DR. ZIEMER: Mark?

MR. GRIFFON: While you're at the mike -- I mean I -- I agr-- I -- I misread that yesterday, and so -- so this may be the case of contaminated equipment in this case on a closed blank. This comes -- from what I understand, this is from one -- one memo --

DR. NETON: Right, correct.

MR. GRIFFON: -- and -- and this does speak to my question of feasible to vali-- sort of feasibility, too, feasible to validate. This is one memo about one time period in the lab when they had some high blanks, and -- and they say in that case it would lead to higher -- abnormally high -- it almost might be a memo -- you know, if you want to be a complete cynic about this, it might be a memo sort of explaining away high results. But anyway, aside from that, it's one memo and -- and my question is do you -- do -- do you have a series of quality control reports from those early lab -- I mean this speaks to being able

1 to go back to the raw data --2 DR. NETON: Right. 3 MR. GRIFFON: -- and say okay -- oh, here we 4 have a bunch of low spikes. You know, you 5 might have a situation where all of a sudden for several months they were running spike 6 7 samples and they were all low, and that would 8 throw your data the other way. So this speaks 9 to my question of the reliability of the data. 10 In this ca-- in this one situation, I agree --11 DR. NETON: Right. 12 MR. GRIFFON: -- it would tend to raise your samples higher, but it speaks to the whole 13 14 question of the reliability and the ability to 15 go back to that source term data, and that's --16 that's my question, is that feasible in a -- in 17 a timely fashion? 18 DR. NETON: Well, if the standard is that for -19 - 50 years ago we had every sample with every 20 blank and every calibration curve, the answer's 21 no, we're not going to have that. 22 MR. GRIFFON: I'm not saying that --23 DR. NETON: But -- but that's what you're 24 suggesting that we need to have, and that's the 25 gold standard. We're not going to be able to

1 produce that. We may have some of it. But I 2 would -- I would suggest that in looking 3 through these records, and having run bioassay 4 laboratories and analyzed tens of thousands of 5 samples in accredited laboratories, the methodology employed here is consistent with 6 7 good laboratory practice of running blanks, 8 pursuing erroneously high values based on 9 precipitation and -- and calibration standards 10 that you'll see in here. So it gives a good 11 sense that this laboratory was indeed 12 practicing good laboratory practices for what is actually a fairly standard technique. 13 It's 14 been in use for over 50 years for metric 15 analysis of uranium in urine. So I have a comfort level, OCAS has a comfort 16 17 level with this. We cannot produce every 18 calibration curve in every anal-- you know, 19 reproduce every single result from first 20 principles. That's not going to be possible. 21 MR. GRIFFON: No, I -- I'm -- and -- and you're 22 -- you're --23 DR. NETON: And where do you draw the line --MR. GRIFFON: -- you're right -- you're right 24 25 on that. The flip side of that I guess is this

1 paragraph sort of relies on one memo and one 2 instance, and it draws that general conclusion 3 that therefore all results, if they were 4 biased, they could be biased high. 5 DR. NETON: Right. MR. GRIFFON: So I think that might be stepping 6 7 the other way. You're relying on one memo -- I 8 -- and I -- I'm wrong -- if I suggested that, 9 I'm wrong. I don't suggest that you could, you 10 know, find all the source data, but I'm saying 11 that that -- maybe more than one quality 12 control report might be useful to get a sense 13 of the -- and maybe you have more, I don't 14 know, but one's referenced here. 15 **DR. NETON:** Yeah, we -- yeah, exactly. 16 referenced the ones here that appeared 17 relevant. I mean we try to produce these warts 18 and all so that the Board can look at them and 19 eval -- make their own independent evaluation. 20 From my -- from my experience in running 21 laboratories such as this, it's -- it's --22 contamination's almost always a problem and, if 23 anything, you end up biasing the results high. 24 But that's just been my experience. 25 MR. ELLIOTT: I think what we are learning here

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is that we need to speak to the other source documents as well in the same series, how many did we look at, perhaps. And if this is the only one that raised our awareness and our atten-- draw our attention, that's important for -- for the public and important for this Board to understand, that it's not just one document with this one issue that's brought -and, you know, been brought to attention, that there's a series of documents that we looked at. Maybe we didn't see the whole universe, but what I'm learning from this dialogue is that we need to be a little more comprehensive in our reporting about -- about this kind of an issue. I do know that we looked at more than just this one quality control document.

DR. ZIEMER: Jim Melius.

DR. MELIUS: Yeah. First of all I just would concur with Larry's conclusion. Based on that I think it really would be helpful for us to get a better sense of, you know, what's your overall, you know, support for a particular statement or for a particular process or, you know, whatever it may be -- some sort of exposure monitoring system so -- so we

1 understand the depth of that and so forth. 2 My ques-- follow-up question has to do -- sort 3 of a follow-up with what I asked earlier and 4 what Henry -- Henry asked about, and maybe I 5 can ask it more clearly, but as we discovered 6 with Iowa yesterday, there are -- questions 7 come up about sub-- saying -- called subgroups 8 within the cohort or within the -- in the work 9 site based on department, building or job title 10 or whatever, and as to the quality of the data 11 for -- availability of monitoring data for 12 them. And I guess my question specifically for 13 Mallinckrodt is have you gone through a 14 systematic effort to look -- look at the population there and determine whether you have 15 16 adequate data for, you know, sig-- significant 17 subgroups within the -- the population in order 18 to be able to complete dose reconstructions. 19 MR. ELLIOTT: Well, personally I have not gone 20 through the data, Dr. Melius, but --21 DR. MELIUS: No -- no, I'm not asking you --22 MR. ELLIOTT: -- but I -- I believe -- I do 23 believe that -- that the ORAU team and my staff 24 have gone through the Mallinckrodt data to the 25 point that they understand the different job

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categories, the different tasks that were involved there, and the data that speaks to not only a department but also to those different jobs and tasks and job categories.

DR. ZIEMER: Jim Neton -- add to this.

DR. NETON: Maybe I can amplify a little bit on that. We -- we have, as we discussed yesterday and as Larry presented today, monitoring data on the majority of the workers after 1948, so in case -- almost all workers in the later years, so we have that to start with. Now that doesn't mean that that -- those data themselves are going to be able to reconstruct doses for some of the processes such as the filter cake operators and that sort of thing. But we do have air monitoring data to supplement that in many buildings -- in most of the buildings during those time periods. And Dr. Melius, you pointed out yesterday, some of the data the end/N\* may be small. That may be true and I've asked for that answer. I'm still waiting -hopefully I'll have that sometime later this morning.

But what happens when you do a dose reconstruction is you -- you look at the case

under evaluation, and we say is this -- where did this person work, what did they do. You look at -- do they have urine sample data, do you have air monitoring data, and then you -- you try to make a determination -- for instance, if the air monitoring data we have -- admittedly, if it's sparse -- but if it's enough for that particular case to put that case over 50 percent, then that is sufficient to do a dose reconstruction.

Now on the other hand, that end/N\* may be small, but at the same time you may say well, let me not just use that one building. Let me take the maximum value in all the buildings in that year and assume the worker inhaled that, and if that value comes out very small, low PC, we have some pretty good confidence that we're forwarding the Department of Labor a dose reconstruction that is useful to them.

So you have to take this in context of how the dose reconstruction process works. In that context, though, we have looked at all classes of workers around the building -- around the site and made a determination that as far as we could tell, looking through all these

1 individual scenarios, we can do a dose 2 reconstruction for the workers at Mallinckrodt 3 in this time period. 4 DR. MELIUS: Yeah, and -- and just to follow 5 up, is that available in a report? Is that --DR. NETON: Well, I think it's -- in general 6 7 terms, it's in the SEC evaluation report where 8 we -- where we speak to -- we have the process 9 knowledge, we have air monitoring, we have all 10 those pieces of data that allow us to do that 11 type of analysis when we do a dose 12 reconstruction. 13 DR. MELIUS: Yeah, but the problem, Jim, and some of it has to do with we're marrying one 14 15 process that's the site profile --16 DR. NETON: Right. DR. MELIUS: -- the individual dose 17 18 reconstruction process with an SEC, so not sort 19 of faulting you personally or the --20 No, I understand. DR. NETON: 21 DR. MELIUS: -- the organization, but the 22 problems -- you make very general statements 23 about this in -- when it's presented, either an 24 overview of the site profile or in the SEC 25 evaluation, and -- and we're trying to look

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behind that a little bit for the SEC evaluation process and -- and it's hard to do when all we get are general statements. I mean they may be true and they may be fine, but it would be nice to have some document or some documentation that backs that up that's specifically relevant to an SEC evaluation as opposed to the ongoing -- going -- going process, and maybe that's asking too much. But -- but I think it would be -- it would be nice to have and -- and -and the problem is that when we probe behind it, you -- you -- and it may --it may be correct in terms of the site profile process. You know, you -- don't have an answer to that because you've looked at it generally. May-maybe somebody at ORAU has, you know, looked at that who's -- who's done the document and under-- understands the details better, but it's not something that -- that we can see and readily reach conclusions on. It's much the same problem we had with Iowa, though --

DR. NETON: Right.

DR. MELIUS: -- fortunately Iowa was, I think, a simpler situation to deal with.

DR. NETON: Right, but short of having done all

the dose reconstructions and demonstrated that we can do every single one, I'm not sure how we could demonstrate that to you other than to provide you these analyses that talk about the quantity of the monitoring data, the quality of it and -- and the process knowledge and source term knowledge that all speak to doing a dose reconstruction with sufficient accuracy. We're trying to flesh out the picture, and I think we've made a very good case that we know the picture very well.

DR. MELIUS: Well, I --

DR. NETON: And outside of --

DR. MELIUS: -- I disagree with that in the sense that -- and I'll give you a specific example, and I know you've already been criticized for this, but the -- this hypothetical example you present isn't helpful. It would have been a lot more helpful -- I'd rather have seen five slides showing specific examples where one would expect or reasonably expect there might be difficulty, you know, because of missing data or the years involved or the department or, you know, sparse data, whatever, that you -- this is how you would --

1	would have done it in those in those cases
2	and do that, and I don't you know,
3	something like that, rather than hypothetical
4	this is how we're going to going to do it.
5	And again, that's not sort of appropriate for
6	the site profile, but it is appropriate for SEC
7	evaluation.
8	DR. NETON: I don't disagree with you. It
9	would have been better to have that. But
10	again, short of doing that for all 311 cases
11	DR. MELIUS: Well, but
12	DR. NETON: it can't be done.
13	DR. MELIUS: I gave tried to give a
14	reasonable number, five, so
15	MR. ELLIOTT: But with with that said
16	with that said, we have the data that we could
17	use to do that.
18	DR. NETON: That was the point
19	MR. ELLIOTT: The data exists.
20	DR. NETON: I was getting at. The data
21	MR. ELLIOTT: That's the point
22	DR. NETON: do exist.
23	MR. ELLIOTT: of the illustration.
24	DR. NETON: I agree it would be better to have
25	it done up front. I mean we're not

DR. ZIEMER: Let me offer, also, that another report, for example, if it's sorted by job title or whatever, would still be a general report. It's simply another way of crosscutting what we already have. So aside from individual discrete examples which certainly were helpful when we went through the original process of describing the efficiency process and so on where you took specific cases, that's helpful for illustrating. But whether that -- a specific case necessarily says you can do it for all cases, that question is always out there.

It seems to me that the information that we're talking about for all these jobs is already contained in what material we have. If -- if -- yes, perhaps if you wanted to look at it a different way and cross-cut it differently, you'd do a different sort on this. But in my mind, the information at least is there.

Mike, I think you're next.

MR. GIBSON: With respect to this information you've found that showed that there was a -- a blank sample with contamination at a -- was there additional information that told you what

1 they did to correct the problem? 2 DR. NETON: I have not personally reviewed that 3 memo, so I can't tell you that. But again, if 4 there were -- it was a problem, the indications 5 are that the worker results would have been biased high, so --6 7 MR. GIBSON: Well --8 DR. NETON: -- I mean that --9 MR. GIBSON: -- I think that depends on what 10 they did with the lab. They also could have 11 very well have increased the minimum detectable 12 activity of the units that read the bioassay 13 samples, which would have masked exposures to 14 employees. DR. NETON: But in the way we --15 16 MR. GIBSON: Is that true? 17 DR. NETON: Yeah, a contaminated laboratory 18 would increase the MDA, you're absolutely 19 correct. But in the way we do our work, if -an increased minimum detectable activity would 20 21 have increased the missed dose that we use for 22 the calculation purposes. The higher -- the 23 higher the minimum detectable activity, the 24 more the missed dose calculation using the

internal dose models will be in our -- in our

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1 analytical process. But I have not read the 2 memo so I can't speculate as to what they did 3 to correct it is the direct answer to your 4 question. 5 Thank you. Henry, did you have another comment? 6 7 DR. ANDERSON: Just more of a general -- I -- I 8 guess I'm looking as we move forward on these 9 and we look at others that the issue of data 10 validation I think is something you need to 11 think about. And there may be -- you know, you 12 can't do it today, but as you look at others to bring forward to us, I think -- and that's sort 13 14 of what I think you were referring to. I think 15 there are some issues that are separate from 16 looking at the site profiles that in the SEC 17 petitions it would just give us more to point 18 to if we had that -- some examples or what has 19 been done -- to detail that, I think that'd be 20 helpful. 21 MR. ELLIOTT: I agree, I think there are many 22 lessons learned here in these first two SEC 23 efforts that we've brought to you. 24 DR. ZIEMER: I saw Dr. Toohey approaching --25 approaching the mike. Perhaps you have some

1 additional comments on this issue. 2 DR. TOOHEY: Well, I have the answer to a 3 question that was asked. We have in the files 4 316 Mallinckrodt cases, 200 Weldon Spring 5 cases. Of those, 110 claimed employment at both sites. 6 7 DR. ZIEMER: Okay, 110 employed both 8 Mallinckrodt and Weldon Springs. 9 MR. ELLIOTT: That's our case population now. 10 That doesn't speak to the overall employee 11 population for Mallinckrodt and Weldon Spring. 12 That's just our case population. Right, Dick? 13 DR. TOOHEY: That -- that is correct, that's 14 just pulled out of NOCTS, and it's all claims, 15 so it would include some that have already been 16 completed. 17 MR. ELLIOTT: I hate to ask another question, 18 but just so we're all clear, we -- you know, 19 the follow-up question to that would be how 20 many of the Mallinckrodt cases worked at other 21 sites besides Weldon Springs, because we probably have a few of those -- a minority, I'm 22 23 sure, but --24 DR. TOOHEY: Give me about ten minutes and I'll 25 let you know.

1 DR. MELIUS: Well, while you're at it, can --2 just someone tell how many of the Mallinckrodt 3 have been processed already, how many are 4 outstanding? 5 DR. ZIEMER: We'll -- we'll get that information. I think Mark is next and then --6 7 DR. MELIUS: And of those -- of those -- excuse 8 me, of those, how -- how many -- what's the 9 overlap? I mean 'cause I assume there's some -10 - some of the --11 DR. TOOHEY: Overlap on the sites? 12 DR. MELIUS: No, in terms of being processed 13 through the sys-- I assume the 110 haven't been 14 because Mallinckrodt -- 'cause Weldon Springs -15 - well, I guess you have done some Weldon 16 Springs, so yeah, we'd want to know the 17 overlap, also. 18 DR. ZIEMER: Okay. Mark Griffon. 19 MR. GRIFFON: Yeah -- yeah, not to be a broken 20 record on this issue, but I -- I think the --21 the question of the data's there, that -- that 22 raises questions with me and -- and harping on 23 the validation issue, I know I'm harping on it 24 probably, but I -- I -- I think it's important

to remember -- you know, one reason NIOSH is

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involved in all this is that a lot of people in the public, a lot of claimants right now, have concerns or issues with DOE's data, with database data that's been used for previous studies, and -- and you know, one of my fears coming through this whole process has been that we don't want to just regurgitate the same data without independently -- and this is this NIOSH independence -- independently going in and validating it.

Now I -- I know that -- that, you know, Jim and Larry indicated it was more than one memo regarding quality. I have to trust them -- you know, 'cause we -- we just haven't had any more depth on that issue, but I think that is -- that -- that's why I raise it so often, because this is database data from the CER database used for epidemiological studies. Without going back one step, you know -- you know, it may be very -- it may be perfectly valid, but we -- you know, I think that was one reason that NIOSH was put in this role as -- to sort of have that independent look back to the raw data, at least -- and I'm not saying all raw data, but at least to the extent that you can

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validate those database data that you're using. The second thing I wanted to point out that -that I -- I am very happy, actually, in this profile and Jim's discussion of -- of -- of --'cause I pushed for this often, this concept of using air sampling and the urinalysis data to sort of check and see if urine intakes sort of are in the same ball park. That kind of a process is a good one, so I do appreciate that effort and there's a lot of data there. guess the -- the time -- the problem that I'm having is that, you know, where we're at today is, you know, can we -- can -- can I make a judgment on -- on the validity of -- of the data being used, and that's -- that's where I'm sitting here today and it's -- it's different than reviewing the site profile, that we're in a petition evaluation process, so that's what I'm stuck on and I don't know that I have all the facts to answer that, which is making it difficult.

MR. ELLIOTT: Can I make a comment on Mark's first point? We -- we do take that very seriously, that -- that -- we see that as a -- not only a responsibility but an obligation to

provide an independent review. And I'd just offer this, Mark, as you -- as I think the Board knows you know, we're not just relying on CER data. Each case that comes in, we go back to DOE and we ask for the original badge or urinalysis data. We don't accept annual summary data. We don't accept roll-up data. We want to see the original badge and urinalysis or whatever source readings that we can get, and we use those. We do not -- you know, we don't trust the -- the roll-up, the cumulative summary, the annual reporting data. That's -- just so you know.

MR. GRIFFON: (Off microphone) Yeah, and Dick mentioned (unintelligible).

DR. ZIEMER: Jim Neton?

DR. NETON: Well, I wasn't going to address what Larry was, but I was going to say that on top of that, though, we have -- we're developing -- you know, we're doing a couple hundred sites at the same time, and -- and it turns out that the Health and Safety Laboratory was -- was intimately involved in many of the AWE or DOE type early operations, and so we are developing some knowledge base about the

processes and methodologies that were used. And I can speak directly to the air sampling analysis. That was questioned early on in the Mallinckrodt review. We have gone back and done some very extensive digging into that. AEC -- HASL -- was involved in the air sampling program there, so we're -- we're developing a comfort level with the techniques, the Whatman 41 filtration, the correction for selfabsorption, the -- you know, the quantity of the air samples through the filters, breathing zone versus general area, so we're developing a very good picture of what was done in those areas. So we have the air sampling data. believe we have a very good picture of how some of the samples were analyzed.

Now -- I see Mark is thumbing through the profile -- I'm sure there's some early indications in the '40's of some ion plate measurements that may or may not be valid, but once you get to the '49 time frame, you're really talking about typical scintillation-type counter measurements, a HASL counter, Cassidy counter some of you may know it by, so we have those data. So again, we can bracket that,

bounce that up against the urine data that we have, and even the fluorometric methods were done at some time periods in the '50's at HASL and so we -- we can have some comfort that they were providing guidance from the University of Rochester as well as the Barnes Hospital data. And so, yeah, we're trying to do that to the -- as best extent we can.

MR. GRIFFON: I -- I was scolded so I didn't get to the table I wanted to --

DR. NETON: I'm sorry, I didn't mean to censor
your -- censor --

MR. GRIFFON: That's okay, I won't -- I won't (unintelligible). This is more of a follow-up 'cause -- 'cause Dick Toohey did mention yesterday that a lot of the individual DRs you have their individual data. Do you have any -- and probably not at your fingertips, I -- I understand -- any idea of the percentage -- and I guess for '49 to '57 is our time period of interest -- how much of that raw -- the raw records are you getting in these -- in your requests 'cause I think that's important to us...

DR. TOOHEY: Yeah, off -- off the top of my

1	head, I do not know the answer to that
2	question. What our procedure is, when we get
3	the DOE submittal and it appears to be
4	incomplete which may be all the way from
5	monitoring for a lot of years but not for a
6	couple, or no data at all then we go to the
7	CER database and search on name, Social
8	Security number, whatever, for data that's in
9	our database to see if the previous
10	epidemiology study captured that data and we
11	can plug it in. But what
12	MR. GRIFFON: I guess that that was sort of
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14	DR. TOOHEY: what fraction of those that
15	represents, I don't know.
16	MR. GRIFFON: Okay.
17	DR. TOOHEY: My my I I know in general
18	terms what it is, and it's about 20 percent or
19	so where we need to go into another source of
20	monitoring data and attempt to capture it.
<ul><li>20</li><li>21</li></ul>	monitoring data and attempt to capture it.  MR. GRIFFON: That's general terms for
21	MR. GRIFFON: That's general terms for
21 22	MR. GRIFFON: That's general terms for Mallinckrodt or

1 quote, smaller sites. 2 MR. ELLIOTT: Does that include AWEs? 3 DR. TOOHEY: Yes. Well --4 MR. ELLIOTT: See, AWEs are a --5 DR. TOOHEY: -- generally AWEs --MR. ELLIOTT: -- whole different bag of worms. 6 7 DR. TOOHEY: -- we get no data. 8 MR. ELLIOTT: You get no data for AWEs --9 DR. TOOHEY: Exactly. 10 MR. ELLIOTT: -- by and large. 11 DR. TOOHEY: But what we've then been able to 12 do in data capture trips where we've gone to a 13 health and safety lab or records repositor and 14 found some of this data, we've now scanned that 15 in and entered it and linked it to the 16 claimant. 17 MR. ELLIOTT: Let me be clear on what I just said. For AWEs, DOE doesn't help us out. We 18 19 get no feedback for AWEs from DOE, so we're 20 left on our own devices and the data that we do 21 get for AWEs is, as Dick says, through our own capture efforts, through our own search and 22 retrieval efforts. 23 24 MR. GRIFFON: Well, I -- I guess that's --25 that's sort of an important question in my

1 If you had, you know, 80 percent of --2 of the people for this '49 to '55 time period 3 had individual -- along with the CE-- you know, 4 you could -- you could mostly rely on the individual data, and only 20 percent you had to 5 go ba-- you know, that -- I think that would 6 7 play an important role in this -- in our 8 discussions -- deliberations. I know that -- I 9 know you don't have that answer, but... 10 MR. ELLIOTT: We don't stop when we don't get a 11 response from DO-- or when we -- DOE says we 12 don't have it, we go back and we push them 13 again until we are satisfied they can't find 14 it, they don't have it or it's lost. 15 DR. TOOHEY: Well, again with the caution, to 16 the best of my knowledge and belief at this 17 point in time, I don't think we have any cases 18 of claim files missing data where we did not 19 have something on that file in the CER 20 database. The CER database I think is more 21 complete than the DOE submittals we're getting, 22 claim by claim. 23 DR. ZIEMER: Thank you. Gen Roessler? 24 DR. ROESSLER: You've mentioned the HASL 25 laboratories, not only the importance in this

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project but -- or this site but others, have you had -- I'm trying to do a little mental arithmetic here to think of how old these people would be now who were responsible back at that time, and I think they probably are still around. Have you had access to any of the people who were setting up the procedures? Have you been able to talk to them and ask them any questions with regard to their goals and --DR. NETON: Yes, we've -- we attempt, where we can, to do that. And I think I presented some indication at the Bethlehem Steel presentation that we discussed this with Naomi Harley, Dr. Harley, who was -- must have started when she was five years old there because she's still very much young, but she has provided us some -- some input. She actually measured the -- the air samples at many of these early facilities and gave us a pretty good description of the quality control practices of those measurements.

We're trying to contact Al Breslin, who some of you may know. He's still living in New Jersey somewhere, but we've not been able to contact - he was the architect of the early AEC/HASL

air sampling program. We have indications of write-ups by him in the '60's speaking to what the process was early on, but we felt it would be better to get it directly from his mouth. Some of you may know Sue Pazanne\*. She was not there directly in those time periods, but she is a well-respected radiochemist who has quite a bit of institutional knowledge about past activities at HASL.

We have been there. We visited the laboratory. I have sort of a unique situation that when I was at New York University obtaining my doctorate, there was a very excellent collaboration between New York University and the Health and Safety Laboratory in lower Manhattan, so we -- we communicate fairly regularly with those folks.

MR. ELLIOTT: We actually had Naomi write up a description of the procedures and the practices as she could recall and we presented that doc-- I think that's -- that was for Bethlehem Steel, but it's applicable for this situation.

DR. NETON: Right, and we've also been to the HASL facility twice now for -- for data capture efforts, to collect records and also to look

for procedures. Some of the early procedures are actually still available, but not many.

DR. ROESSLER: I think from my perspective in health physics, HASL has been around for a long time, has a very, very good reputation. I can't say specifically on this site, but a very good reputation for running reputable, credible laboratories.

DR. NETON: I will have to qualify that, though, and say that HASL does have that reputation, but they did not do all the analyses at Mallinckrodt, just to be -- to be -- let the record show that. But I would say that one reason we have some comfort is after '49 much of the health physics work at Mallinckrodt was done under the tutelage of the HASL folks, which gives us a little more comfort there.

## **PETITIONERS**

DR. ZIEMER: I think we perhaps are ready to move on and hear from the petitioners, and I'd like to thank you, Larry, for your presentation. I'd like to turn the podium over to Denise Brock at this time, representing the petitioners.

1 MS. BROCK: I would like to thank the Advisory 2 Board for again affording me time on another 3 4 5 6 7 8 9 10 11 12 13 14 process. 15 16 17 18 19 20 21 22 23 24 notebook paper, so kind of bear with me with 25 some of this.

busy agenda, and I'd also like to commend the Board on their decision and expeditious recommendations to the Honorable Secretary Leavitt on the SEC petition regarding Mallinckrodt years of 1942 to '48. I'd also like to thank Dr. Lew Wade, Dr. John Howard, the OCAS staff, NIOSH staff, Department of Labor, as well as Senators Bond, Talent, Harkin and Grassley, along with all the members of the Missouri and Iowa Congressional delegation who have been so helpful in this SEC I'd also like to state that I wrote this statement a few days before I even came here, and since we were here and we've heard statements from other people in the audience and myself, a lot of this is going to be repetitious, so please excuse me for that, but it's going to be too hard for me to kind of pick and choose through it. I've also interjected some notes, even on

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Twenty-seven years ago yesterday while I was at school, my father died in my brother's arms. remember waiting for the school bus that morning. I yelled goodbye, but my dad didn't answer. And I decided, for whatever reason, I was going to wake him up so I walked back to his room. I didn't really care if I missed the school bus. I was probably actually hoping I would, and I -- I walked back to his room. oxygen mask was on and the room was cold and kind of damp. It always felt like that, even if the sun would come in through the windows. I always remember that feeling. So I said goodbye to him again out loud and I still didn't get any response. So I leaned down and I shook him and he woke up, startled. said bye, Dad, I love you. And for the last time in my life my father looked me in the eyes and said goodbye, my girl, I love you. hours later, while I was at school, my brother came to my classroom and told me that my father had died. Sorry. I find it so ironic standing here before you

today so many years later asking for justice

for my father's coworkers. I've heard that we

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are all put on this earth for a reason. I
don't think I'm here by coincidence. I don't
think it was just happenstance that in 2001 I
heard about this program. I was blessed. I
feel like, for whatever reason, God let this be
my purpose, and I just hope my dad can see me
and I hope that he's proud.

As I stated in my previous statement in St. Louis, my father worked at Mallinckrodt from 1945 until 1958. He died at age 52, but not before years of pain and suffering from his radiation-induced cancer. My mother, thankfully, has been compensated under this program. And I'm here today, as I was in St. Louis, not asking you for compensation for my family or for myself, but for those workers and survivors who cannot, for reasons of health or finances or who I've stated before cannot begin to fathom the complexity of this program. I'm here to honor the memory of my father and the coworkers who will unfortunately never see or hear an admission of guilt, nor receive an apology or payment. I'm here to ask the Board to consider my statements, my findings and my

pleas in the decision that they must render

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regarding the remaining years of 1949 to 1957 requested in my SEC petition, 00012.

And I'm not here to deny that NIOSH has a collection of Mallinckrodt data. I am here to state that what data exists possibly has major credibility issues. I'm also here to say that quantity of data does not necessarily mean quality of data.

Mallinckrodt was the first U.S. feed material processor for the United States atomic weapons program for World War II. Their work began in 1942 by producing a ton of pure uranium a day. Mallinckrodt remained a prime contractor for processing until 1957 when the last Mallinckrodt plant closed and the last of its employees in those operations were terminated. Mallinckrodt hired at least 3,500 employees, and they were assigned to these secretive operations for a varying length of time, having multiple job titles and descriptions during this time period. The workers at Mallinckrodt who helped build this atomic pile helped win the cold war, and they gave their lives for their country, and they're known to be among the most highly exposed workers to internal and

This is

1 external radiation in the entire history of the 2 United States Atomic Energy Commission. 3 By NIOSH's admission, many of these workers 4 worked long enough at appalling concentrations 5 of alpha-emitting dust to accumulate more than 6 a permissible lifetime inhalation exposure. 7 would like to restate again for the record that 8 I respectfully disagree with the NIOSH 9 conclusion that it is feasible to estimate dose 10 for this time period. 11 NIOSH's position is that HASL did its own 12 monitoring, and this means that there is verification on the Mallinckrodt data that was 13 14 not in place prior to 1949. Again I'm 15 unpersuaded that the post-1949 data is any more 16 credible than that of Mallinckrodt's. 17 SC&A's audit report for Rev. 0 notes that there 18 were dramatically different results for 19 monitoring by Mallinckrodt and HASL of the same 20 exposures. HASL data is higher in 15 cases and 21 lower than Mallinckrodt in 12 cases. 22 according to the chart in the SC&A audit report 23 Rev. 0. If I understood correctly, NIOSH is 24 actually using the lower on HASL than -- than 25 the higher ones on Mallinckrodt, if I

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understood that correctly. And on page 77 of Rev. 1 on the TBD, '50 to '54 it says it's unclear who actually did the urinalysis, whether it was Barnes or the AEC. So we cannot answer the question on who has reliable data or whether it's reliable at all. And this does not change the fact that there is evidence to doubt the credibility of any of this data due to the huge liability concerns of all involved. I'd also like to restate from the February meeting that there is no isotope-specific monitoring for raffinates. There was frequent exposure in plant six to these raffinates whose pathways for uptakes are not well understood. And I don't want to go into again -- Richard Miller touched on that yesterday about the lime and the exothermic reaction that happened, so I think you've all heard that I just don't feel the need to go back into all that. But I do think that because of this reason and being no isotopic-specific urinalysis to quantify any raffinate uptakes, and the burden of proof on NIOSH is so high to establish internal dose, that this is the circumstances that Congress created the SEC, at least part of it.

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As noted above, the TBD does not address internal and external radiation dose from open wounds and burns, which I stated in the past workers have testified to and stated that it was quite prevalent. I was present at an SC&A interview with workers and I did videotape The workers discussed these burns, these acid burns and the open wounds, as well as numerous other things and -- I mean it wasn't limited to incidents or occurrences. talked about a lot of things. I actually have a lot of these tapes and have offered those to NIOSH to share with their staff and the Board. In the SEC petition evaluation report, SEC 00012-2, page 3, NIOSH is seeking the advice of the Board for the time period of 1949 to 1957. It also states that any documentation that raises questions concerning the integrity of data management or reporting at Mallinckrodt helps sustain the lack of credibility accorded by the Mallinckrodt claimant population to the government concerning the employees' radiological exposures at Mallinckrodt and concerning the dose reconstruction program

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Evaluation Reports.

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One of the issues brought to the Board's attention by NIOSH at the St. Louis meeting was an August 1975 memo titled "Notes and Summary of Visit by M.E. Mason, August 1975." according to NIOSH, disputed a 1972 memo from Mont Mason to Dr. Thomas Mancuso in where Mason states his concerns about the possible destruction of records identified as shelf list V2161. NIOSH further believed that the August 1975 memo confirmed the conclusion that these records were not only found, but complete. even if NIOSH would be in possession of these records, it still does not speak to the credibility nor the completeness. Since that February meeting, after NIOSH told the Board and the petitioners and the public that this particular memo was referencing to the records with the shelf list of V2161 and that they were in possession of such, they've now stated that their initial beliefs were This information can be found in the

This obviously raised another red flag with me.

supplement to Mallinckrodt SEC Petition

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The memo was withheld until NIOSH was compelled at the February meeting to produce it. was so much commotion over this revelation over this particular memo, and it was trying to discredit the memo that was in my petition. The initial findings of NIOSH were obviously an assumption and have now been changed. now claims to have since recovered a records transmittal and receipt form from the Federal Records Center which appears to identify the contents of shelf list V2161. It actually says "appears". What does appears mean, exactly? Is this V2161? I've noticed it says 21, 22, 23, 24. I mean if it is, how do we know it's How do I know? How does NIOSH know? complete? How do we know it hasn't been altered or tampered with? And how do we know that NIOSH is not going to come back in a month from now and state that this wasn't it or that there's extra -- extra information or extra data? NIOSH also states in the supplement that because this records inventory form did not include actual exposure or process information of possible relevance to the development of the Mallinckrodt site profile, it was not grouped

with other records that were identified as relevant for that purpose. NIOSH also claimed that this memo somehow proves that the latter years of the record/data keeping at Mallinckrodt are above reproach because HASL or the AEC was involved.

This document, in my mind, does just the opposite. It confirms my views that there are numerous problems and inconsistencies. It also confirms that there remains much uncertainty about the validity and completeness of data for not just the '42 to '48 time frame, but the remaining years. And as far as the AEC, as I stated at the February meeting, they allowed this operation to continue with unacceptably-high levels and saw it as an opportunity to experiment on worker population and not as a moral outrage.

My first thought on this memo is that it's not authored by just one person. The first page speaks about Mason in the third party in three separate statements. The first statements says, and I quote, I can make a study without the AEC NYO documents but they are authentic references above the level of contractor bias

or Mason shortcomings, end quote. Number two, I quote, Becher-Mason conflict about usefulness of Mallinckrodt U.U. shortcomings, unquote -- I'm sorry, numbers as an indicator of body burden, end quote. Number three, Becher argues that I am contradicting myself in that at Weldon Spring Mason used U.U. values to report exposure to AEC in compliance with regulations of 100 CFR 20.

This not only shows that Mason clearly is not the author at the beginning of this memo, but clearly speaks to the contractor bias.

NIOSH again is assuming that Mason authored these notes. NIOSH also states in their supplemental issue response that Dr. Mancuso ran out of funding and his study for the Mallinckrodt populations workers (sic) was not completed or released.

Dr. Mancuso was terminated. He was blackballed. His findings or conclusions on the studies he did were inconvenient for the AEC. When they hired him I understand that they were so sure that he was going to come back and be government-friendly, they actually referred to it as Mancuso's Folly. When his

conclusions were not as they wished, Mason actually had to take his findings and move those to other places. Many years later I believe he was exonerated, but I believe that he had years of problems with the Atomic Energy Commission, and that was due to, again, the conclusions that he had that were inconvenient or not AEC-acceptable.

NIOSH states that based on their review of the

1975 Mont Mason notes, the context of those notes and the review of the data, NIOSH found no issues that would prevent dose reconstruction. I wonder if I'm reading the same notes, because this memo solidifies and confirms the many uncertainties that exist regarding the Mallinckrodt population and any data which exists.

On the first page of the August 1975 memo there are several different opinions on what numbers or data are even useful. On the second page the author states that the broad subject of measurement of internal depositions, the value of urine bioassay in assessing body burden, the relative worth of body gamma counts versus urine and fecal bioassay is as old as the art

of health physics, and older in industrial hygiene.

On page 77 of Rev. 1, the top paragraph again states it's even unclear who did the urinalysis on Barnes and Mallinckrodt. I also have a 1951 memo that was attached in my petition that actually tells Mallinckrodt to stop sending urine samples for a certain time period because the lab was overworked.

The 1975 memo also indicates that there are large numbers of these employees whose records did not list them by Social Security number and then later had to be manually cross-matched, still leaving some out. One must account for human error. We have no assurance that there is complete or accurate data for any individual Mallinckrodt employee, and neither does NIOSH. There is serious question as to the credibility and completeness of this entire Mallinckrodt data.

My daughter had a call from a worker who told her he was having a difficult time getting his records from Mallinckrodt or the Department of Energy, wanting to see individual records or whatever he had. Come to find out, he found an

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old badge that he had been assigned, and the badge had his Social Security number on it, but actually had the wrong middle initial. remembered talking to his supervisor at Mallinckrodt and they had actually discussed some of his records being mixed up with either his brother's or somebody else's. And we've noticed through other workers a lot of times their records would say, for example, R. Jones -- and maybe that would be Ralph, and then you'd have a Richard Jones. And when you have to cross-match these things, there was concern that perhaps people's Social Security numbers with the name, with the data, were all being confused. And if anybody's interested, my daughter had written something up, has a copy of that gentleman's badge and all the information that was needed and it's actually been notarized. NIOSH provided a summary of what was in the five boxes that they have retrieved. Box one is approximately 75 percent film badge readings, many of which -- according to NIOSH -

- are duplicative and already have been

addressed in the TBD. Data from '46 to '49 has

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already been incorporated. They state that some additional documents that support dose reconstruction can be incorporated into the TBD -- can be incorporated -- but that '53 to '58 time period is not yet incorporated, which this information is largely specific to particular operations. It can be incorporated into the next revision of the TBD. How many times are we going to revise this? Again, I understand it's a living document, but these workers are dying. They don't have years to wait for this. The remaining 25 percent is air sampling data, breath analysis, et cetera -- but for what years and for whom, and is this already Box two and three are insignificant to this petition because it retains Weldon Spring Box four is Mallinckrodt dust studies and annual uranium urine results for '48 to '58.

annual uranium urine results for '48 to '58.

To that I must again state garbage in is garbage out. How do we know the credibility of this data? We already know that zeroes were recorded for dust and urine when no tests were even taken. And checking for uranium in the

1 urine does not account for other things such as 2 thorium or plutonium. 3 And as far as the plutonium -- I'm actually 4 looking for something; give me a minute here. 5 (Pause) In Rev. -- it looks like Rev. 0, page 23 of 32, 6 7 and I'll just take a part of this out of here -8 - there may have also been trace radiological 9 impurities supplied, uranyl nitrate, 10 hexahydrate, recovered from separations 11 operations at Hanford and sent to either -- to K-65 or Mallinckrodt for processing. 12 review of Rev. 0, pages 43 to 44. NIOSH has 13 14 not found any data on these operations. Page 15 And I'm wondering if NIOSH has actually 16 investigated that to see if in fact there was 17 plutonium. 18 As I stated, I have several workers that feel 19 that there was plutonium at the Destrehan 20 Street site. 21 Box five is 75 percent Weldon Spring film badge 22 and the remaining 25 percent is film badge for 23 Destrehan during '46 to '49. This is largely 24 not applicable. I already have a cohort from 25 '42 to '48.

I'm curious if there's been any detailed discussion or explanation on what is in these boxes. Does NIOSH findings -- are they the same as with the TBD. And I -- I don't know if it's been assessed -- I think Larry may have just discussed this -- and by who or if there's been any individual assessing of this -- independent.

Back to the August 1974 memo, page 6 continues to talk about disagreements and information not matching. Page 8 of this memo refers to the J factor. In the same paragraph it states, and I quote, other adjusting factors were used in calculations to produce an I index value for each person where I equals 100 was presumed to equate to a potential lung dose of 600 rem from alpha dust.

Well, no wonder the company and the AEC had liability concerns. In my petition there is substantiating documentation as to these concerns and in graphic explanation as to allowing these employees to get to levels sometimes in excess of 1,000 rem to the lung. I understand that the annual limit at the time was 15 rem, a standard set by the AEC. But yet

by their own admission they allowed these workers to be exposed to over 40 times the set amount before even addressing the problem.

In my SEC petition I submitted a memo dated 10/3/1972 to Dr. Mancuso from Mont Mason. The third paragraph reads both Mallinckrodt and the AEC were mindful of the sensitive human relation problems and the health department bent over backwards to gain and hold the confidence of rank and file, as well as union representatives.

It goes on to refer to a 1949 dust study and subsequent removal of a number of over-exposed workers, and it then reads, and I quote, every action, every statement by management was carefully thought through. Carefully drafted explanations and responses were prepared in advance of announcing the transfer of people. Managers, supervisors, medical staff and health department staff were all coached and coordinated. As part of caution and on the advice of attorney, a formal report was never prepared on this study. Thus there is no document to subpoena, only a list of names and numbers and work sheets. There was no lengthy

description of the basis for calculations to be pulled apart by the scientific community, with the possibility that such controversy would undermine confident -- employee confidence in the company's safety measure, end quote. Is not NIOSH and ORAU a scientific community? And are we to believe that after all of the lies, the cover-up, the mishandling of data, the liability concerns, that just because HASL has oversight that the surge of conscience and transparency and honesty just happened all of a sudden for these latter years? This was a joint effort to keep these workers in the dark and to quell any liability issues at that time or in the future. The mind-set of Mallinckrodt and the AEC was horrific.

As one reads through the remaining pages of the August 1975 memo that NIOSH has found, it appears repeatedly that the data are much in question. It seems as though the data has been reworked consistently. On page 23 there's reference to either Becher or Mason authoring at least this and the following pages, but I've yet to ascertain which one of those gentlemen actually did that. This entire document

illustrates the multi-faceted problems in collecting data, recording it at different times in different ways using different practices, and then trying to keep it all together. It confirms that there are multiple opinions as to the quality and completeness of

any of this.

memo.

I don't want to insult anybody's intelligence by reading aloud from that or to con-- and continue quoting from the memo. I know that you've all probably had an opportunity to read it, and I'm quite sure that everyone has taken note of the numerous issues that I have. I would, though, like to call your attention to a memo of January 31st, 1951 from Eisenbud to W.E. Kelley -- or from Eisenbud to W.E. Kelley states about a year ago you asked if we -- if it would be possible for us to estimate our potential liability among the long-term Mallinckrodt employees. As I explained at that time, you presented a rather knotty problem, one which, in the present state of knowledge, would not be answered even to a first approximation. This memo again was a 1951

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I understand that SC&A have only been able to
do a partial review for Rev. 1, so it's
impossible for me as a petitioner to prepare my
case and have further quality arguments and
reference to a site profile when our auditors

haven't been given a chance to complete the

work.

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It puts me at a distinct advantage (sic). can't not (sic) argue or comment without reading and reviewing these TBD shortcomings. I realize that SC&A did the best they could, but because of this revelation of Iowa's cohort being withdrawn from them and it had to be reviewed again, it put me and my claimants at a disadvantage, as well. It was kind of put on a back burner, and SC&A were -- were trying to look at these at the same time, so it was rather difficult. It put us all kind of under the gun, so it made it really difficult for me. I was finally able to take a look, obviously, at the partial report. But I actually have a to-do list that I want to talk about in a little bit that NIOSH has.

I have another concern about the sperry cake and the issue of the dermal contact. I believe

1 that there were no specific numbers in 2 reference to that. We know that dose 3 consequences from exposure raffinates are 4 significant. Routine inhalation of even a 5 milligram quantity of sperry cake, one 6 milligram per month over a few years, has a 7 potential for significant internal radiation 8 dose, notably to the bone surfaces and the 9 Thorium 227, the main decay product of 10 actinium 227, is a potential concern for the 11 lung dose, as well. 12 Dose from the radionuclides has not been 13 evaluated in any of the documents I've seen, 14 and I don't see anything in Rev. 0 to help answer that, nor -- I don't think there's 15 16 anything in Rev. 1. 17 I don't really understand a lot about the 95 18 percent confidence level, but I want to know if 19 I understood correctly that workers who were 20 marginally exposed were averaged down. 21 looked like the time-weighted averages discriminated against them, and that's a 22 23 concern for me, as well. 24 And I obviously have no background in health 25 physics, but I can read. And I know that Janet

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Westbrook from the beginning has stated that NIOSH could dose reconstruct the entire population of Mallinckrodt workers. opposed this SEC from the beginning. know that in Rev. 1 from the beginning she's stating that dose can be performed from '42 to '48, and I guess I just didn't understand that. I understand now that it's been addressed. But something as simple as work hours is incorrect, at least the way I look at it in Rev. 1, it looks as though these workers were working maybe five days a week, maybe six days a week at -- at 40 hours or 44 hours. That is incorrect. My workers have told or the survivors' workers basically state that they worked seven days a week, 40 plus hours. might be one of the few things that survivors know, but one of the few things. And the workers that are living always state to that fact.

I'd like to talk about the splitting of this petition again. I know we brought this up or I brought it up in February, but by breaking this up into these subclasses has serious equity issues. For example, if I have a worker that

worked in 1947 and worked his 250 days and he has one of the 22 SEC cancers, he has a coworker that started in 1948, but he's missing that 250-day mark, I would have to go back to these workers and try to explain to them how their coworker was able to have that same cancer and job and be compensated in an expeditious manner, and they're not covered. There just -- it does not seem equitable to me and it's very difficult to have to try to explain it. I don't think there is an explanation.

For the record, I would also like to talk about coworker data and survivor claims. I also wanted to state, in reference to that, that when I sent in the original claim for my mother's claim I happened upon something called a Leo Goodman incident report. It's been so long I don't remember exactly what was in that, but there was some information, probably two or three things, in reference to incidents or occurrences at the Mallinckrodt site. I noticed when I was reading through NIOSH's things that they had never heard or found anything in reference to any incident at

Mallinckrodt, and I just don't think that's accurate.

I've sat through telephone interviews with these survivors -- and granted, they don't know much. You're talking about people that are in their seventies or eighties who have years and years ago heard from a spouse that they were involved in some sort of accident, and that's usually how they refer to it, an accident. during this telephone interview the NIOSH CATI question is raised to the survivor and claim -you know, claimants in reference to this accident or occurrence. And if by some stroke of luck a survivor can remember their spouse telling them about a situation, they're asked specifics. This is a monumental task for a spouse or a child or even sometimes a grandchild to hurdle. They're asked if the worker ever had chelation therapy. Every claimant I've had asked me what is chelation ther -- I don't know what chelation therapy is. They're expected to know details that they received second or third-hand, or maybe not at all, and sometimes as back (sic) as the 1940's. They were also required to have two witnesses

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to this accident.

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Well, you know what, my dad was involved in an accident. He worked there before I was born. My mom was 78 when we filed her claim. hadn't really remembered the accident, but my sister did. So we had to rely on that memory of -- of my sister triggering my mom. My uncle was also involved in an accident there, and I -- I think I've brought this up before. When my aunt went through her telephone interview, I was with her like I'm with any of the claimants that ask for my help, and I was on the -- the phone, the extension, and she was going through the questions through the interview. Typically everybody we have from NIOSH or ORAU are very polite. They're helpful. They're empathetic. But every once in a while -- you know, when you get that volume of people, you get somebody that's maybe not so nice or has had a bad day. This was horrible, because this woman began to ask my aunt questions about this accident, and she wanted to know if my uncle would -- had chelation therapy and my aunt didn't know what that was. She then asked her about being

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shows it on a monthly basis. As of April 18th -- again, last week -- we had sent -- or week before last, I guess -- we had sent out 8,537 cases. Again, the month of April not being completed yet, that number of 193 doesn't reflect the full month.

Draft modifications is portrayed here. just so you know, draft modifications refer to the changes that occur in a draft dose reconstruction report while it's in the hands of the claimants, before that report has been sent on to the Department of Labor. A DR draft is considered modified when DOL provides NIOSH with new information that would affect the case, such as a new cancer, new employment information or something of that sort, or a change in the survivorship. Modified dose reconstructions are illustrated in blue and represent the draft dose reconstruction reports that have been modified due to the receipt of the new information from Labor. And the modified DRs that are sent back to Labor after they've been changed are shown in red. The final dose reconstruction reports that have

The final dose reconstruction reports that have been sent to DOL are depicted in this slide on

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a monthly basis. I'd be pleased to note for you that as of Monday this week, we went over the 8,000 mark. This slide was prepared last week, as of April 18th, and at that point there was 7,851. We have now gone over the 8,000 mark back to DOL.

With regard to our request for exposure information from the Department of Energy, we have sent 18,543 requests. We have received 18,053 responses to those requests. The age of the outstanding requests are shown, those greater than 60 days being 74; 18 for greater than 90 days; 30 requests we've been waiting on for 120 days; and 50 requests we have been waiting on for over 150 days. I can speak -- I know this question's going to come from Dr. Melius, so I'm going to speak to it before he has to ask it. I'm going to save him a little time and it'll save us all a little -- little energy and just give you a quick summary. Albuquerque operations office, particularly with regard to the site for Lawrence Livermore, we have seven requests that are over 60 days past due, and we have one request that's over 120 days past due.

1 Amarillo operations office, Pantex being the 2 site, three requests over 60 days. 3 Chicago operations office, that would include 4 Argonne East and Argonne West as the sites in 5 question, five requests over 60 days, two requests over 90 days and four requests over 6 7 120 days. 8 The General Electric facility in Vallecitos, 9 California, we have 12 requests over 150 days 10 and two that are over 120 days. 11 Let me just run -- Richland operations office -12 - this is Hanford and PNNL -- we have 29 requests over 60, 13 over 90, 20 over 120, and 13 14 25 over 150. 15 The remainder are from Savannah River, Fernald, 16 Nevada, Honeywell, and they vary in the dates. 17 We follow up on these on a monthly basis. 18 numbers do change and we are digging into 19 specifically now what's holding up any case 20 beyond 90 days, and we will be making some 21 determinations as to whether or not there is 22 ever any information that's going to be 23 forthcoming. So we have renewed our vigor in 24 following up on those cases, besides just a 30-25 day follow-up period.

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Cases here that are shown are telephone interview statistics. These are the CATIs, the Computer Assisted Telephone Interviews. Cases for which at least one interview has been completed, 18,130. This is -- these numbers are, as you -- as you recall, a case can have more than one survivor and so we interview everybody associated with a case. So these numbers represent actual interviews beyond the -- just the single cases that we have. Interview summary reports that have been sent to claimants, around 24,000 -- or 20 -- close to 25,000, and the number of interviews that have yet to be conducted for the claimants right now, or as of April 18th, were 270. Since January 1st of 2005 we have been conducting between 300 to 400 interviews per month, and you can see that on this slide -shows how -- how the CATI process has -- has, I think, been a very successful aspect of our program, at least in showing some completion here. And these folks that do the Computer Assisted Telephone Interviews are also tasked with doing what is called the closeout interview, and that's not captured in this kind

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of -- in this slide. These are only the interviews for the dose reconstruction that are shown here. Any closeout interviews -everybody gets a closeout interview to make sure that they understand the report and we can answer any questions that they might have, or encourage them to sign the OCAS-1 or to provide additional information that might better inform the dose reconstruction on their claim. We have -- these statistics are -- basically can be found on our web site in the claim information page and shows the process of handling a claim. And as of April 18th, 2005 we had 8,952 cases that were in a pre-dose reconstruction assignment development part of the process where information is being collected, interviews had been done, reports were being exchanged on those interviews with claimants, et cetera. Cases that have been assigned to a DR-ist (sic), a dose reconstructionist, 1,197. We have 477 draft dose reconstruction reports that had been sent to the claimants, were in the

April 18th. And we had sent, as of that date,

7,851 cases -- completed cases to Department of Labor.

This graphic illustrates a cumulative figure of cases received by NIOSH in 1,000-block intervals or increments, with a breakdown of the overall cases by tracking number that have been completed. That's the number indicated on the top. And represented in red, the cases by tracking number that have been completed prior to January 1, 2005 -- that would be these (indicating), and then those that are represented in blue are the cases by tracking number that have been completed since January of 2005. This graphic's intended to give you a perspective on how we're doing with regard to the oldest cases, how we're doing within each one of these 1,000 incremental blocks, how we've done since January 1st of this year. There are pulled cases that are accounted for in these figures. A pulled case has been a case that DOL has retrieved from us for many reasons, perhaps -- the most -- it could be that -- that -- there have been cases sent to us that were CLL, and at this point in time we're -- CLL's the only cancer that's not

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adjudicated under this program. It could be that -- the most unfortunate and most disturbing, to me, reason would be that the Energy employee or the only survivor of that Energy employee has deceased, and those are not -- those are very few, but I do watch those on a -- on a close -- very close basis. They have my attention.

As mentioned at the previous Board meeting, we

As mentioned at the previous Board meeting, we are focusing on completion of the first 5,000 claims. We have been somewhat successful at reaching this goal. Since January 1st, 2005 we've completed 120 cases in the 1,000 block, 117 cases in the 2,000 block, 81 cases in the 3,000 block, 87 cases in the 4,000 block, and 103 cases in the 5,000 block.

It's our -- we have a concerted effort underway and have set a goal that by June 30th of this year we will have attended to all of the remaining cases in that first 5,000, either through dose reconstruction or assignment to a Special Exposure Cohort class or a -- by completed dose reconstruction it could be actually a dose reconstruction sent to the Department of Labor or one that has been

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drafted and placed in the hands of the claimants. So ORAU has some incentives to do this and we're working very closely with them to make sure that this goal is achieved. I think it's very important that we attend to this. This is part of our timeliness concern that we're facing.

Administratively closed records are shown in this graphic by month, since we first started tracking these. And what happens here is where we have a situation where, for whatever reason, the claimant decides not to sign the OCAS-1 form, not to return it to us, has perhaps got to the point of a height of frustration with the program or with us or whatever and they -they've just shut off communications with us. We do go back to them. We try to talk them through the process, try to encourage them to file the OCAS-1. We encourage them to do that. We explain to them that if they have another cancer, if they have additional employment, we could follow up on that together. We explain that the signing of the OCAS-1 is not an agreement that they sign saying they are in full agreement with our dose reconstruction,

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but it is simply that they have no further information to provide and it allows the case to move forward so that a decision can be proffered from DOL. And as you can see, we're dealing with a few of these but not very many. Reworks. I talked about modifications earlier. Well, reworks are a little bit different breed here. A rework refers to a process that occurs at the final dose reconstruction report stage, meaning that the case has been in the hands of the Department of Labor. And in this situation, something has been identified by the Department of Labor or by the claimant that would require us to go back and revise our dose reconstruction report, redo the reconstruction or add new -- because of new information, add to that report. These -- these revisions can be initiated because of additional employment, additional cancer, new information that the claimant identified after they had signed the OCAS-1 and sent it on to -- and we had sent it on to DOL. So there's a variety of reasons as to why these are -- are kicked back to us from the Department of Labor.

We maintain a high level of contact with the

claimant population through phone calls and correspondence and e-mails, and that's shown on this slide. Our contractor receives the bulk of the phone calls, and I believe this also includes the CATIs, as well as the -- this 150,000 number of phone calls to ORAU includes CATIs and closeout interviews.

Pardon me?

DR. WADE: (Off microphone) (Unintelligible)
finish up (unintelligible).

MR. ELLIOTT: Okay. Pick it up -- pick it up, he says.

Last time we had some interest in hearing more about the compensation rates by cancer model, and this slide presents some of the caveats associated with the slides that I'm going to show after this. These results that I'm about to present to you are through April 20th of this year. They're based on claims which NIOSH received notice from the Department of Labor of a compensation decision, so there's a number of claims that won't be included in this that are still at -- are at DOL, but we haven't learned from DOL what the final decision was. These rates may be skewed by the DR efficiency

1 process. The rates may not be predictive of 2 any future results. And unless otherwise 3 noted, the rates reflect claims with only one 4 reported primary cancer. Does not include 5 secondaries or multiples, unless so indicated. 6 As you might expect, lung tops the list here at 7 almost 70 percent. Leukemia's a close second 8 at 67 and 61 and 56, then we see liver and 9 other types of leukemia, endocrine gland and 10 other respiratory. 11 DR. MELIUS: Lew, if you and Paul could try not 12 to get your heads together because --13 DR. WADE: We will not get our heads together 14 again. 15 Thank you. I thought that was MR. ELLIOTT: 16 just a bump in the road trying to get me to 17 hurry up. 18 But non-melanoma skin is shown here, and 19 several other cancers. I'm not going to go 20 through this. You have it -- have these 21 slides, as well, in your briefing packet and 22 they're on the table for the public. 23 think it's indicative of the radiogenicity of 24 the cancer, as we suspected, based upon the 25 scientific knowledge and understanding of

cancer causation.

Nothing much here I want to say about this slide. You can peruse the numbers as you can.

Trying to pick up the pace here, this is -this is where I need to make notice for you
that at least one secondary cancer, primary
cancer unknown, we see about a 71 percent
compensation rate. For cases where we have
multiple primary cancers, we show about 42
percent.

The percent of -- there are -- for those cancers where we have 30 or more claims and we have not seen anything -- any -- any of those claims compensated are listed at the bottom of this slide. I do know that there is one female genitalia that I think Labor has in their hands right now that may be compensable. We have to wait and see what happens with that one, a very high dose.

Petitions received, we're moving on to a different topic here, but petitions that we have received, 26 total. We have 20 active in our hands. HHS decision has been made on one, that would be Mallinckrodt for the years '42 through '48, as you know. There have been six

petitions administratively closed for lack of
basis.

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Our -- briefly, our accomplishments since the last Board meeting, we have exceeded sending 8,000 dose reconstruction reports to DOL. We've seen Secretary Leavitt's final decision to add a class of workers for Mallinckrodt. That was sent to Congress on April 11, 2005. We have participated in 33 meetings at 13 sites since March 1st. This is a huge commitment of staff effort. We send not one, but we send several people -- PHA, a public health -- or a health physicist, public health analyst, a communications specialist -- typically to these meetings, and it's been a very resourceintensive effort. But I think it's been very beneficial to us and to the people who show up at those meetings. We can answer questions. You may have seen a different presence here at this meeting for NIOSH. We've had public availability sessions at this meeting where we've asked claimants -- we noticed all of the Iowa claimants that we would be here. I've had mixed results on this -- from why'd you tell me that you were going to be here when you can't

1 tell me that anything's changed, to hey, it's 2 good to see a face associated with my claim. 3 So we're evaluating our presence and our -- how 4 we represent ourselves at these meetings, and I 5 would appreciate hearing any thoughts or comments that the Board members have, on an 6 7 individual basis, about that. 8 Finally, we would lay claim that 12 Technical 9 Basis Documents have been approved since 10 January, and seven Technical Information 11 Bulletins have been approved. 12 I think that will conclude mine and I'll gladly 13 answer questions, and I'm sorry if I was too 14 quick and I glossed over things that you wanted 15 to hear more about. 16 DR. ZIEMER: We'll have time for a couple of 17 questions, and I should tell you why Lew and I 18 had our heads together. We are eliminating 19 Jeff from the program. He's aware of it. You 20 have the Labor report in your document, so that 21 will help us with the time a little bit. 22 Jim, and then Leon. 23 DR. MELIUS: Larry, I thought you --24 DR. ZIEMER: Larry already answered your 25 question, Jim.

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DR. MELIUS: Yes, I know, and I was saying he stymied me, but I've got to come up with some new questions or a new way of asking 'cause I'll give you credit, the cases completed by tracking number -- my usual question is how many have you -- what have you done for me lately, and you know, you've got that on there, so that's -- that is helpful.

I have a comment and a question. My comment is sort of the half-full, half-empty sort of argument, but if my calculations are right -and they're certainly at least ballpark -- it still leaves you within the first 5,000 with about 2,400 of those first 5,000 to be completed; 3,200 of the next 5,000 to be completed, and so on. And at the completion rate that you're going at now, assuming even no new cases, I believe that's at least two years to catch up with the backlog and presumably with the current rate of increase -- of claims coming in, I think that stretches it out to about three to four years to -- or closer to four years to catch up with the backlog. Now that's assuming no SECs and making lots of other assumptions. So I guess I'd like to have

a little bit more information. One, on what you're doing with the backlog to catch up, and secondly, how we're dealing with -- who is that

MR. ELLIOTT: Who is that backlog?

DR. MELIUS: Yeah, what kinds of ca-- I believe in the past I've asked for information on -can you tell us the -- why -- why are we having problems with these? Is it -- what --

MR. ELLIOTT: Well, since there's two -- over 200 sites represented in our caseload, as you Several of those are AWE sites. Several -- we have several individual cases where just one or two cases from a given site. We won't have a site profile for those. We're looking at what we can do for those on an individual basis. We have employed an overestimation approach, a new Technical Information Bulletin that provides an overestimation approach, and if the case is compensable, we move it on. it's not and we have to refine our dose reconstruction or find information to process

I think you're going to -- from this meeting to the next meeting you're going to see a dramatic

increase in our rate of production, given some of the things you just alluded to, that SEC classes are going to help us out with reducing our backlog. This overestimation approach that we've approved for the ORAU folks to use is going to help. We're -- we've talked to them about adding staff where we need to add staff to get this work done. There's a variety of efforts underway. Now without, you know, going into greater detail -- that I don't have at my fingertips right now -- I can't -- I can't add any more.

DR. MELIUS: My recollection is that ORAU was working on some evaluations or reports related to some of this -- these backlog issues -- again, specifying who was rep-- who was included in that backlog by type of case or site or something.

MR. ELLIOTT: That's right.

DR. MELIUS: Could those be made available to the Board, at least, if you don't have time to...

MR. ELLIOTT: No, I don't think they're -they're in a shape or form that I'm happy with
yet. We're working with the contract-- our

1 contractor to get to a point of understanding 2 on how we're approaching our backlog issue in a 3 variety of ways, and I'm not at a juncture 4 where I'm ready to commit publicly to how --5 the different approaches that we're applying. 6 We're working on that, and as soon as we have 7 something that I can make publicly available, I 8 will. 9 DR. MELIUS: Is there a time frame on that? 10 MR. ELLIOTT: I hope we'll be able to say more 11 about that at the next meeting in July. 12 DR. MELIUS: I mean I'd just point out -- I 13 mean this -- there's a lot of frustration which 14 we've heard today from Mallinckrodt, and we 15 heard yesterday with Iowa, about the length of 16 time it's taking for cases to be processed 17 through this program. And I mean I think a 18 much more -- I mean I think a plan needs to be 19 developed and it needs to be shared with the 20 public --21 MR. ELLIOTT: I agree --22 DR. MELIUS: -- as soon as possible. 23 MR. ELLIOTT: -- I agree 100 percent. 24 let's keep things in perspective here. 25 we -- we sent letters out to the Iowa claimants

1 once we were dealing with the revision to their 2 site profile and indicated that we were no 3 longer processing dose reconstructions until we 4 had a revised site profile. Now that we have a 5 decision -- a recommendation for a class, we're 6 watching what's going to happen with that 7 decision on that class. Mallinckrodt, we have a revised site profile. 8 9 We are dealing with a petition. We have --10 have no recommendation. We will go back and 11 start processing claims under Mallinckrodt Rev. 12 1 site profile immediately, and we're going to 13 start moving those claims through as we await 14 this Board's deliberation on -- on 15 Mallinckrodt, and then finally a decision on 16 Mallinckrodt as an SEC, so I'd just offer that. 17 I agree with you 100 percent. We need a cogent 18 plan and we're working toward that end. 19 DR. MELIUS: Well, I think we need to re-20 evaluate the approach entirely. I find it 21 increasingly unacceptable that people are 22 having to wait this long. 23 DR. ZIEMER: Leon Owens. 24 MR. OWENS: Larry, I had an opportunity to 25 attend three of the DOL outreach meetings in

Paducah, and I would like to compliment Mr. Stu Hinnefeld and Ms. Heidi Deep. Particularly -- Stu fielded some very tough questions. We had a large -- large group of workers in attendance, and so I do think it adds a lot of benefit, a lot of value to have NIOSH representatives present at those meetings.

MR. ELLIOTT: I thank you for that, Leon, and it's kind words like that that keep us going with these difficult challenges. I would like to add, though, that the 33 and the 13, the statistics in those outreach meetings, do not include our worker outreach, our worker input. Those are just the town hall meetings that DOL has sponsored. So while we're doing that, we're also going out and doing our own worker outreach. As you know, we spent a session down in Paducah during this time period since the last meeting, so...

DR. ZIEMER: Thank you. Henry and then Mark.

DR. ANDERSON: Just quickly, you mentioned the

-- we're now going to be getting into some of
the site profile updates. What -- what's the
process? I think you mentioned you would
potentially go back through those that have

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already been adjudicated to see if it'd make a change? How -- how's that happen? I mean --

MR. ELLIOTT: Well, we do that --

DR. ANDERSON: -- that could get to be very
onerous.

Oh, it is resource-intensive, as MR. ELLIOTT: everything we do here in this program. consumes us. But we have a standard operating procedure, one of those procedures which has been reviewed in task three, which is called program evaluation reporting where if a change is made -- and we've made numerous changes, numerous revisions in site profiles, as you may know. And each time one of those is -- is completed, one of those revisions is completed, we look back through the cases that were dose reconstructed under that particular version, and the evaluation must include whether or not the modification that was made or a change in the site profile that was made would affect the compensability outcome of the case. So if the case was already found compensable, we disregard it.

If it's found non-compensable under the prior version, we evaluate the change and whether it

1 would make a difference in the compensability. 2 If it does, we reprocess that dose 3 reconstruction, notify the Department of Labor 4 and the claimant that we're doing so and we 5 send that revised dose reconstruction back. To date, I do not believe -- Jim could correct 6 7 me if I'm wrong, if he's still in the room, but 8 I do not believe that we have made any changes 9 -- we've not seen any non-compensables turn 10 into compensables based upon revisions that 11 we've made. 12 DR. ANDERSON: That was my next question, yeah. 13 Thanks. 14 DR. ZIEMER: Mark. 15 MR. GRIFFON: Just -- just a quick one, Larry. 16 I -- I think it might have been last meeting 17 you mentioned that -- or it was -- I think the 18 question was raised and you had mentioned that 19 ORAU was doing a report for you on self-20 identified SEC classes, and I wondered if that 21 was completed yet. And if it is, if the Board 22 can get a --23 MR. ELLIOTT: We -- we -- it's not -- I've sent 24 it back because I wasn't -- I wasn't fully 25 satisfied in how that was developed. I want to

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see more detail in it and more effort put toward that end, and we're awaiting a revision of that report. I can tell you that some of the -- some of the sites that involve early years we're really focusing our attention on that. I've asked specifically, because I know -- we've got Linde site profile in our hands, and if you've looked at it, you know that it is reserved for the early years, and I'm saying that right there, in my opinion, is a potential class and so we should look at that. looking at Linde, we're looking at NUMEC, there's a variety of those sites that have -in the early years where their data is nonexistent to very minimal, or the monitoring program was not what we would hope it should have been, we would like to have seen it. MR. GRIFFON: That can be -- when it -- when it's completed, it can be provided to the Board?

MR. ELLIOTT: Yep.

DR. ZIEMER: Thank you. Okay. Thank you very much, Larry, for the update. I think we're ready to move on. We have one item on our afternoon agenda that may have some substantial

1	discussion, and that's the first 20 dose
2	reconstruction case wrap-up, and we have the
3	subcommittee met earlier this week was that
4	this week? Monday Monday morning, and most
5	of you were actually here for that, but we have
6	the materials that come to us as a
7	recommendation from the subcommittee. I think
8	it would be helpful if the Chair identified to
9	you again what those materials are.
10	MR. GRIFFON: Paul, I
11	DR. ZIEMER: Mark, can you help us?
12	MR. GRIFFON: I was just going to ask, does
13	it make sense I know Jim said he had a
14	motion, and then we have a task order for SEC
15	task to consider.
16	DR. ZIEMER: We have
17	MR. GRIFFON: And I this is going to be a
18	lengthy process, so I'm just afraid that we
19	might lose
20	DR. ZIEMER: Well, it's an issue of what should
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22	MR. GRIFFON: (unintelligible) numbers
23	DR. ZIEMER: go first. Actually the
24	MR. GRIFFON: I know.
25	DR. ZIEMER: the task order for SEC was

1	we had a draft of that, I think, in our
2	telephone
3	DR. MELIUS: And we said we would take
4	action
5	DR. ZIEMER: And we would act on it here.
6	Perhaps
7	DR. MELIUS: I have a copy if we need
8	DR. ZIEMER: Perhaps we can act on that quickly
9	since
10	DR. WADE: (Off microphone) (Unintelligible)
11	have it.
12	DR. ZIEMER: that was developed. We've had
13	that for a month or so. Would you like to do
14	that next? It is the next item.
15	MR. GRIFFON: I think tho yeah, those two.
16	DR. MELIUS: And with with the Chair's
17	permission, can I hand out the issue related to
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19	DR. ZIEMER: Yes.
20	DR. MELIUS: transparen so these people
21	can glance at it.
22	SEC PETITION REVIEW TASK ORDER
23	DR. ZIEMER: Okay, let's let's get the
24	document before us. It's called Special
25	Exposure Cohort petition review task order.

1 The draft does not have a date on it. I guess, 2 for convenience, put today's date on it. 3 (Pause) This would be task five for our contractor. 4 5 MR. GRIFFON: And we --DR. ZIEMER: Jim -- Jim is passing out an item 6 7 which is a different issue, so that doesn't --8 this is not part of the item before us here. 9 MR. GRIFFON: We had a -- I mean we had a 10 fairly lengthy discussion on this on the 11 previous Advisory Board phone call meeting, and I -- I think -- well, I think we just need to 12 13 consider it now, and there might be --14 DR. ZIEMER: Sure. MR. GRIFFON: -- possible amendments to it that 15 16 we want to consider, I don't know, but... 17 **DR. ZIEMER:** Does everybody have a copy? 18 actually trying to recall whether this came out 19 of the subcommittee. Lew, can you help me --20 MR. GRIFFON: It was --21 DR. WADE: Yes, it was prepared in -- really 22 during and after our subcommittee meeting in 23 Cincinnati, and then it was placed before the 24 Board on its phone call on -- I think it was 25 April 11th, and there was discussion of the

1 full Board, reserving for this meeting action 2 on the -- on the task. 3 I think there's also the possibility that if 4 this task document is agreeable to the -- to 5 the Board, the Board might instruct me to undertake the independent government cost 6 7 estimate, which would expedite the matter. But 8 I think first there needs to be an intellectual 9 agreement on this document. 10 DR. ZIEMER: I'll interpret the document as 11 coming from the subcommittee. It constitutes a 12 motion, does not require a second. It's open 13 for discussion. Basically what we have before 14 us is a motion to approve a task concerning 15 Special Exposure Cohort petition reviews. 16 Wanda Munn. 17 MS. MUNN: I don't know whether my note on this 18 -- the copy that I have -- is something that I 19 made during the time we were discussing it and 20 whether it was something we agreed to, or 21 whether it's just a note that I've made to 22 myself. 23 Item number three, (reading) the contractor 24 will be required to review -- I have inserted 25 "up to" --

1 DR. ZIEMER: That's correct, we --2 MS. MUNN: That was a part of our discussion, 3 wasn't it? 4 DR. ZIEMER: We basically agreed that we 5 weren't guaranteeing that there would be eight, 6 so the scope would be up to eight. I think we 7 had agreed on that. I don't see that in this, 8 but in paragraph three, without objection, add 9 the words "up to" eight SEC petitions. 10 scopes an upper limit for purposes of the cost 11 estimate, not guaranteeing to the contractor 12 that there would necessarily be this many petitions. We don't know how many we would 13 14 need reviewed. Thank you. 15 MS. MUNN: And I have --16 DR. ZIEMER: Another? 17 MS. MUNN: -- one other issue. Although there 18 is nothing in this particular item which would 19 point to my concern, nevertheless it's a 20 concern I want to raise. 21 When we first began looking at task orders for 22 our contractor, we -- I was under the 23 impression that the contractor was going to be 24 providing -- specifically -- technical, 25 scientific expertise that we were not able to

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provide here in our own group. And I -- I had no -- at that time, we reviewed some of the credentials of virtually all of the people that was my understanding would be the primary actors in what SC&A would be doing for us. On our last telephone call we had a considerable amount of input from a policy individual that I questioned afterwards and was told that that individual had become a part of our -- our SC&A task force. I was a little taken aback by that because I did not -- I was unaware of the fact that -- that we were providing our -- our contractor with instructions to follow our -- our directive by way of inviting policy makers on the team. I guess -- because we're now in the process of putting together another task, I guess -although I don't see that it's the Board's requirement to review the credentials of every person that our contractor chooses to -- to take on, I do have to question that particular item as -- with respect to credentials. DR. ZIEMER: Thank you. This may be an issue we would have to discuss with the contractor in

that particular case. Lew, do you have any

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comments on that or can you shed any light on that?

DR. MELIUS: I have some comments when --

DR. ZIEMER: I was not on the phone call. To address this?

DR. MELIUS: Yeah, my recollection is that -if I'm guessing right at the individual you're referring to -- is that he was listed as a member of the team when we approved the initial contract, and so it was not a surprise to me to see him and hear that he was involved in -- in doing this and was listed in a -- I thought at the time when we approved it -- in an appropriate fashion and in an appropriate role, and -- again, if I'm guessing right at who you're referring to 'cause I was not part of this more recent phone call -- was that he was also -- has a considerable knowledge about the DOE complex and I believe has been -- if he's -- to the extent that he's played a role, and I don't think it's been a major one, has been based on his historical knowledge of information throughout the -- the complex, particularly on specific sites.

DR. ZIEMER: Not necessarily technical, but

1 information on or knowledge of --2 DR. MELIUS: Availability of technical 3 information. I mean --4 DR. ZIEMER: -- availability of technical 5 information. DR. MELIUS: -- the particular individual is 6 7 fairly knowledgeable technically, but -- again, 8 my sense is he's -- and recollection at the 9 time is he's being used -- was to be used and 10 involved in an appropriate way, given his 11 knowledge and background. 12 DR. ZIEMER: Wanda? 13 MS. MUNN: If that is in fact the case, then I 14 must have been completely blind at the time 15 that we were -- time I thought I was paying 16 attention. I am not comfortable with that. 17 am not at all comfortable with individuals 18 other than the technical expertise I clearly 19 understood we were seeking from our contractor. 20 DR. ZIEMER: Thank you. And again, that may be 21 an issue we would explore, perhaps with our --22 with our contractor. Whether or not that 23 impacts directly on this motion, I'm not sure. 24 DR. MELIUS: Can I just -- I would hope, given 25 some of our past history, that if we're going

to try to deal with issues like that, that we would do it with full involvement of the Board.

DR. ZIEMER: Thank you. Any other comments on
-- on this particular document?

DR. WADE: Could I ask a clarifying question,
Mark, since -- I assume that you're listing
here a number of tasks, any or all of which
could be engaged, depending upon the Board's
wishes. So it's not that all of these things
will be done, it's up to the Board as to what
would be done. Is that correct?

MR. GRIFFON: Yeah, I guess we -- I mean we had -- you mentioned on the conference call that we had, and I don't know if we want to amend this to give an option at the discretion of the Board to make a decision as to whether we want a more limited scope for certain reviews or the full scope. And that mi-- you know, that might be something we want to consider.

DR. ZIEMER: It would seem to me that if we had a task that was sort of encompassing, then we could use that for a particular issue, or portions of the task, could we not? Is it not better to have a broad task under which work can be done rather than a very narrow one that

1 confines you so you cannot do certain things? 2 DR. MELIUS: Yeah, my sense was also that 3 number two here would be developing the 4 procedures that were going to be used and very 5 well that one of those procedures would be the option to sort of focus on different --6 7 DR. ZIEMER: Procedures would --8 DR. MELIUS: -- evaluations as we --9 DR. ZIEMER: -- spell that out. 10 DR. MELIUS: -- go --11 DR. ZIEMER: Right. 12 DR. MELIUS: -- along, and -- and again, I don't know if we need to include it here, but 13 14 certainly as we operationalize this that we 15 should take that into account 'cause it may 16 very well be appropriate, and to make these 17 more timely, also, I think. 18 DR. WADE: I am comfortable with that. I just 19 wanted to make sure that was the sense of -- of 20 the Board. I think this gives me that 21 flexibility, but I wanted to clarify that 22 sense. 23 DR. MELIUS: And as I said on the conference 24 call, I am uncomfortable with trying to do this 25 piecemeal, one question at a time, or two

1	questions, 'cause I think it's just going to
2	tie us up and we're going to lose consistency
3	over time.
4	DR. ZIEMER: Thank you. Other comments?
5	(No responses)
6	Okay. Are you ready to vote on this document
7	then? Let us vote.
8	All in favor will say aye?
9	(Affirmative responses)
10	Any opposed, no?
11	(No responses)
12	Any abstentions?
13	(No responses)
14	Motion carries. Thank you.
15	DR. WADE: And then a clarification. Do I have
16	the permission of the Board to undertake
17	developing an independent government cost
18	estimate to this task?
19	DR. ZIEMER: Any objections to have our
20	DR. WADE: Thank you.
21	DR. ZIEMER: Okay. Our official will proceed
22	with that.
23	DR. MELIUS: I actually think the sooner we can
24	get this in place, the better given the
25	numbers Larry just showed.

MR. GRIFFON: Just a logistic question. I think you have the most electronic version, minus the "up to" eight ca-- so if you can just -- I don't need to send any revised versions -- DR. WADE: No, I'm fine with that. Thank you. DR. ZIEMER: Okay. We have a proposed motion from Jim Melius. We also have -- which may or may not be a long issue, but I believe the -- the case report -- the 20-case wrap-up is really the important issue that we need to address before we lose a quorum.

MR. GRIFFON: I don't know, I think if we want to -- you want to -- I would -- I would defer to this motion first and then -- I think that makes more sense.

DR. ZIEMER: Okay. The Chair will recognize Jim for the purpose of making a motion.

## DISCUSSION, LEGAL OPINION FROM DOJ

DR. MELIUS: Yeah. This motion cames out (sic)
-- (unintelligible) from our discussions
yesterday and -- or maybe it's out lack of
discussions or ability to discuss the reported
legal opinion from the Department of Justice.
And I'm not sure whether we want to task NIOSH
with this or whether it would be better to be a

letter to the Secretary asking -- 'cause this is a response 'cause at least it's implied to us that the Office of General Counsel for the Department is where -- at least this was communicated for -- this opinion was communicated from, so let -- why don't I read the motion into the record.

The Board has serious concerns about the reported Department of Justice legal opinion regarding the handling of classified information as the basis for decisions within the Special Exposure Cohort program. While fully supporting the need for preventing the release of classified information, the Board also recognizes the importance of transparency to the EEOICPA program. Due to the long history of secrecy at DOE nuclear facilities, former workers are very suspicious of secrecy related to any health-related information used as the basis for their claims.

The Board respectfully requests the following information: One, who requested this legal opinion and what was the rationale for the request; number two, what agencies were involved in the discussion of this legal

1 opinion and to what extent does this opinion 2 apply to programs in those agencies; number 3 three, requesting a copy of the legal opinion 4 and a presentation by an attorney familiar with 5 the basis for the opinion at our next meeting. The Board believes this information is critical 6 7 for the Board to properly and fully carry out 8 our responsibilities under EEOICPA. 9 DR. ZIEMER: You've heard the motion. Is there 10 a second? 11 MR. ESPINOSA: Second. 12 DR. ZIEMER: The motion has been seconded. 13 It's open for discussion. Wanda? 14 I feel I have to repeat the question MS. MUNN: 15 that I asked yesterday. I don't understand why 16 bullets one and two are of any consequence. 17 Who requested it, why it was requested, doesn't 18 seem to be an issue. The Department of Justice 19 is within its prerogative to do that. 20 for a copy of the legal opinion and an attorney 21 familiar with the basis is, in my view, certainly within our prerogative and we made 22 23 the statement in the first paragraph that 24 workers are suspicious and that concerns us. 25 really do not see that bullets one and two are,

1 frankly, germane. 2 DR. ZIEMER: Are you making a formal motion to 3 exclude them or are you just raising the 4 question? 5 I would move that we exclude bullets MS. MUNN: one and two. 6 Okay. There's a motion to exclude 7 DR. ZIEMER: 8 bullets one and two. Is there a second? 9 DR. DEHART: I second. 10 DR. ZIEMER: And seconded. Now what is open 11 for discussion is the removal of bullets one 12 and two. You may speak in favor of the motion, 13 you may speak in opposition to the motion. 14 Indicate what your -- Roy. DR. DEHART: I'm speaking in favor of the 15 16 motion. I'm not certain that that information 17 has any bearing on the action of the Board. 18 The impact of the legal opinion does, but who 19 may have instituted the inquiry or whether or not it was done solely from the Department of 20 21 Justice I think is immaterial. 22 I would also ask the question, have we seen the 23 full opinion, so I think that needs to be put 24 in the motion. DR. ZIEMER: The motion asks for a copy of the 25

1 opinion, I believe, yeah. 2 DR. DEHART: Yes. 3 DR. ZIEMER: Bullet three --4 DR. DEHART: Well, how can we have a concern 5 when we haven't seen the full opinion? 6 DR. ZIEMER: We've seen a slide depicting the 7 opinion. 8 I would put that in there, we have DR. DEHART: 9 not seen the opinion. 10 DR. MELIUS: Well, I tried to address that by 11 saying the reported Department of Justice legal 12 opinion, in the first sentence. That was what 13 I was trying to get at. 14 To speak --15 MS. MUNN: Alleged. 16 DR. MELIUS: To speak against the motion to 17 remove --18 DR. ZIEMER: Go ahead. 19 DR. MELIUS: -- I'd like to explain -- was -- I 20 thought it would be helpful to understand the 21 context for this opinion, and that would be 22 knowing who requested it, what was the 23 rationale for that request, as well as in 24 bullet two, what other agencies were involved. 25 For example -- and I guess we didn't get to

hear this presentation today, but I'm aware that the Department of Labor is developing their regulations regarding Subtitle E, and so issues of classification versus due process and so forth may be something they're wrestling with at the time. That would sort of put this in context and I think help to -- help us to understand what was involved. Also to know how this was being applied, was there a related opinion that was being developed for Department of Labor's program or for some other part of this program, and it would just, I think, more fully inform us about this opinion and be able to understand it better. And that was the rationale for bullets one and two.

DR. ZIEMER: Henry?

DR. ANDERSON: Yeah, I guess I'm speaking against. I -- I would agree, I -- I guess what I'd like to know is was this done specifically, narrowly and for our program or what's often -- it could be that this is a general opinion related to classified information that then sweeps us up in the late notification as somebody looked at this and said this may impact you, when in fact it was developed for

other programs. And that places our issues in context and might then result of a need for a more detailed review or comment on specifically our program. It's for -- if it's for our program and it was apparent nobody here knew about it, that is also important to know. So you know, I think it would be -- the context of it and how broad-sweeping it is -- we may learn that when we get a copy of it, and it may all be in that copy as to who it's being sent to, but I just think it would be helpful to know how specific is it to they reviewed our program versus other issues.

DR. ZIEMER: Okay. Wanda, do you wish to speak for your motion?

MS. MUNN: It still doesn't matter whether it's just our program or whether it's the whole wide world. The Department of Justice is within their prerogative to do that. As a matter of fact, it's their responsibility to do that. And so since they're doing it, from our perspective our only concern needs to be how it affects our program.

DR. ZIEMER: Okay. Others who wish to speak in
-- okay, Michael?

MR. GIBSON: I think it's important to leave the first two bullets in there only because we've tried to -- been open and honest and work in good faith with NIOSH and Department of Labor and everyone else in this program. I think it's just -- it'd be a good road map and a good history to determine how this was -- was -- what the genesis was for this, since no one seemed to be able to tell us earlier.

DR. ZIEMER: Thank you. Henry, did you have another comment?

DR. ANDERSON: No.

DR. ZIEMER: Anyone else wish to speak for or against the motion?

(No responses)

Are you ready to vote? This is a motion to delete bullets one and two. If you vote yes, you are voting to delete those two bullets. A no vote is a vote to retain them. Okay? Those who favor deleting bullets one and two, raise your hand, we'll get a count here -- one, two, three and the Chair will vote, four. And those who oppose deleting the bullets -- one, two, three, four, five, six, seven, so the bullet -- the motion loses and the bullets will

1 remain in. 2 We are now back to the original motion with all 3 bullets in place. Now this may pose a dilemma 4 for those who voted against the bullets because 5 they may favor the main motion but be concerned 6 about the bullets. That's the nature of what 7 happens when things are amended or not amended. 8 Sometimes you take the good with the bad. 9 But let me ask if there's any further 10 discussion, in which case we will vote for the 11 main motion as unamended. Are you ready to 12 vote? 13 Okay. All who favor the motion, say aye -- or 14 let me ask for a show of hands. Okay, one, 15 two, three, four, five, six, seven, eight, 16 nine, ten. 17 Opposing? One. And no abstentions? And the 18 motion thereby carries. 19 It appears to the Chair that this should be a 20 request to the Secretary. 21 DR. MELIUS: I believe so, yes. 22 DR. ZIEMER: And if that's the case -- and I 23 may need some help here. 24 Our normal role is to advise the Chair -- or to 25 advise the Secretary. Lew, is there any

problem if we simply -- we are asking the Secretary if this -- if this information can be provided. Is that appropriate or should -- should we go through NIOSH on this.

DR. WADE: I don't think there's any problem with it, in my opinion.

DR. ZIEMER: Okay. With that added comment, then, the Chair will proceed -- do I have to do this in 21 days? We will -- we will send this forth as soon as possible. Thank you. Mark.

DR. WADE: Could I take up an issue before
Mark, just -- the future schedule?

DR. ZIEMER: We're going to run out of time.

## FUTURE SCHEDULE

DR. WADE: It'll take two minutes. But would someone get Cori in the room as I start to do this? I would refer you to this piece of paper you have, a future schedule. The only reason I do it, there is a key decision that needs to be made, triggered by this piece of paper. Very briefly, all I'm trying to do with this is at every meeting to look two meetings out and give you a sense of what might be coming downstream. I think it is terribly important that we coordinate, for example, comments back from

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your contractor on the TBD as we might be contemplating an SEC petition, for example.

And I think there are many things that will come from this.

If you'll look at the July meeting, there are some changes already. You've asked that we take some of the task three work and move it from the October meeting to the July meeting, and I'll take that as an instruction from the Board. We need to add the Mallinckrodt SEC work now to the July meeting. You have just asked in your letter to the Secretary for a transparency issue briefing at the July meeting. I am operating on the assumption --Larry, correct me if I'm wrong -- that the IAAP -- IAAP SEC rad workers issue might not have to be worked, given the action with regard to the SEC, or does it still need to be worked? MR. ELLIOTT: I think it still needs to be worked up and given to the Board as an evaluation report because the Board took separate action on that. They're awaiting our evaluation report.

DR. WADE: Okay. Thank you. The only issue that requires work is you'll notice that we

1 would expect to discuss a Y-12 SEC petition in 2 July. The Y-12 site profile review is on your 3 contractor's list. Right now it's late in 4 their scheduling. I would like you to consider 5 elevating it in the scheduling so that they can begin to work on it now, so you would have as 6 7 much possible benefit of their review as 8 possible when we come together in July. 9 ask for the sense of the Board that that would 10 be acceptable to you. If it is -- I've already 11 discussed this with Joe and I think he's ready 12 to proceed. I didn't want to take that action 13 without consulting with the Board. 14 DR. ZIEMER: Any -- any comments by the Board 15 members or --16 DR. MELIUS: I think that'd be good. 17 Can I have one other comment -- agenda comment, 18 though? 19 DR. ZIEMER: Yes. 20 The Bethlehem TBD or the DR. MELIUS: Yeah. 21 review of it and so forth, I believe -- well, I 22 actually know, because of -- I was copied on 23 some correspondence. The Congressional 24 delegation in western New York and our two 25 senators had -- had requested clarification

1 from DOL. I think it's -- the issue is were 2 there additional runs at the -- Bethlehem 3 Steel. And last I heard, which was a couple of 4 weeks ago, was that there had been no response 5 from Department of Labor on that issue. think it'd be worth exploring on the part of 6 7 NIOSH to -- just as a scheduling issue, 'cause 8 potentially the DOL response could change the 9 site profile one way or the other, and I'd hate 10 to have us deal with it, particularly given the 11 long history there, and then have -- suddenly 12 have a -- some change come down from DOL. It may very well be that it doesn't affect it, but 13 14 15 DR. WADE: Okay, I understand, and I will take 16 on that action. 17 DR. ZIEMER: Okay. 18 DR. WADE: Now we have a Board meeting 19 scheduled in early July. Cori, the tentative 20 dates are? 21 MR. GIBSON: Sixth, 7th and 8th. 22 MS. MUNN: Sixth, 7th and 8th. 23 DR. WADE: The 6th, 7th and 8th? We were 24 contemplating meeting in Oak Ridge. We now

have this competing need for a St. Louis

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meeting. I don't know that we can resolve that today. I know, Mr. Presley, you had comments you wanted to --

DR. MELIUS: Paducah, half way.

MR. PRESLEY: I've got a real concern. meet in Oak Ridge July -- or anywhere else, especially Oak Ridge -- Oak Ridge has -- for the last 20 years we in Oak Ridge, including myself, we get two days holiday. This year it lie-- it goes for a Monday and Tuesday. A lot of people in Oak Ridge are off and take vacation that whole total week, 'cause that gives them nine days. I do not feel like that that would be fair to the people in Oak Ridge if we have a meeting on a holiday weekend. I also do not feel like that it would be fair to people in -- anywhere else if we have a meeting on a holiday weekend. We get enough concerns and bounce-backs about some of the other things that we do, without having meetings on holiday weekends and have to travel on weekends ourselves and things like this. That's my concern.

DR. WADE: Thank you. I don't know that we're going to be able to work through this right

now. I think I -- I understand that concern. If you would leave it to myself and staff to try to work through these issues, realize that there are -- there are definitely competing demands on us and we will try again with the wisdom of Solomon to work through this.

MR. PRESLEY: With the wisdom of Solomon, if we can work through this, can you do it as early as possible? I hate to get down a week or two before a meeting and then have to start making arrangements. I think that goes for all of us.

DR. WADE: Understood.

MR. GRIFFON: Can I --

DR. ZIEMER: Thank you, Lew. Yes, Mark?

MR. GRIFFON: Just -- just one question along the lines of our future agenda here, and this is a -- we spoke -- or we questioned Larry earlier on his backlog. This is quickly becoming our backlog. I'm looking at the Savannah River profile on there. We've had that for a while. The task three procedures review, we've had that for a while. And I wonder if we -- I think we committed in -- in the subcommittee meeting to working on the procedures review in between these next two

1	meetings via workgroup, subcommittee, I'm not
2	sure what that process is going to be. But I
3	wonder if we can do the same thing for the
4	Savannah River my concern is I mean,
5	quite frankly, this week has been a challenge
6	for everyone to get through. And if we load up
7	this meeting again with five four or five
8	major items like this, we're not it's not
9	going to work, and
10	DR. WADE: Understood. The Sava for the
11	record, the Savannah River profile I think
12	we've just received or are about to receive?
13	UNIDENTIFIED: (Off microphone) It's been
14	submitted.
15	DR. WADE: It's been okay. But your
16	UNIDENTIFIED: (Off microphone)
17	(Unintelligible)
18	DR. WADE: Okay, your point is well made.
19	Thank you.
20	(Whereupon, several Board members commented,
21	off microphone and simultaneously.)
22	DR. ZIEMER: We have it. Yeah, Mike.
23	MR. GIBSON: It seemed, you know, here a few
24	months back, a year ago, we met more often than
25	four times a year. And it just seems to me,

with -- with the addition of the SEC process, the dose reconstructions and everything else that four times a year is not enough for this Board to meet, and I just wonder why we can't pick up the pace and go back to maybe every six weeks or so like we did before.

DR. WADE: Certainly.

DR. ZIEMER: Yeah, we -- actually -- you have four meetings here. We had a meeting in February, and we had one or two telephone meetings plus the subcommittee meeting, so we actually have ended up having about six or more meetings a year right now. A few -- a couple of those didn't involve everybody, but it actually has -- it has seemed to me to be a fairly rigorous pace, but --

MR. GIBSON: I quess --

DR. ZIEMER: -- but if we have business we need to conduct, we'll -- we can do that.

MR. GIBSON: I'm just -- I was just agreeing with Mark that, you know, the pace is picking up and even if we have six a year and it's not enough, you know -- I mean I think it's our duty, that's what we were appointed to do is to do this business. And so, you know, this has

been a pretty taxing week...

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DR. ZIEMER: Henry?

DR. ANDERSON: Yeah, I was going to say that's one thing we may be able to do is the subcommittee meetings, maybe those could be done by phone, and that would give us an extra half -- I mean we've already started -- they were two-day meetings. It's fairly easy to find two days together, but when you move that out to find three days, and then a travel day as well, it starts to get problematic. think a two-day is a lot easier to find, and if we're going to do three days, I would rather pick up an extra half-day by doing the subcommittee on an alternative maybe phone schedule issues, and deal with the -- these issues in a close -- by the -- the number two is -- is -- I'd like to get October dates as soon as possible 'cause there's a number of key meetings that are in October already, so before we start to try to get three days and not have them be a Friday, Monday and Tuesday.

DR. ZIEMER: Right. Right.

MR. ESPINOSA: I'm in the --

DR. ZIEMER: Rich.

MR. ESPINOSA: I'm in the same lines with Henry, just a little bit different. I schedule my agenda -- I schedule my schedule around these meetings, so if I don't know in advance and the meeting changes, like we talked about the July meeting changing, you know, if I have the notification now, I might be able to address it. But if I don't have the information till later, I'm looking at missing a meeting or days of that meeting.

DR. WADE: I'll commit to trying to put out dates for the July and October meeting the next week.

DR. ZIEMER: Jim?

DR. MELIUS: Can I suggest since we're relooking at this issue that we also re-look at
this issue of the subcommittee as a whole, that
-- that we may really have to split up into -have more than one subcommittee, and that way
it spreads the work out a little bit more
clearly and it would also I think make the
scheduling of some of these meetings -subcommittee meetings easier -- easier to do.
And I don't know whether something -- whether
you want to work on, Paul, or whether you want

to work on it with a workgroup, but -- but I think for our July meeting I think we should -- I think we have a -- we have a sense of what -- well, SEC petition reviews are very hard to do other than as a committee, so those are going to have to be (unintelligible). Site profile, dealing with dose reconstruction, some of these other issues, I think that we can -- dose reconstruction reviews, I think we can do better as subcommittees. Again reporting back to the Board, but I think it may be a more efficient process and --

UNIDENTIFIED: (Off microphone) Workgroups.

DR. ZIEMER: We can do workgroups, but remember that if it's a sort of regular process, then we get into the subcommittee type action.

DR. MELIUS: Our charter is up the end of
August, and so a July -- make a decision on
this at our July meeting and it would be
appropriate 'cause we could then amend the
charter to deal with the subcommittees, and I -

DR. ZIEMER: Well, the subcommittee does not have to meet as a committee of the whole, obviously. For example, there could be a

1 subcommittee meeting between April and July if 2 -- for a group of four or five is all that we 3 need, actually. In fact, that was the original 4 intent. We named everybody to the subcommittee 5 in order that we could choose any four or five who were available at a given time. 6 7 DR. MELIUS: But -- but as you pointed out, you 8 -- you were -- it seems like you feel like 9 you're meeting all the time, and that's because 10 you chair the subcommittee and therefore at 11 every meeting you have to be at. And again, I 12 -- maybe that's not practical and -- not taking 13 away from the amount of effort you're putting 14 into it or anything, but -- but again, let's 15 iust --DR. ZIEMER: Well, I was --16 17 DR. MELIUS: -- let's just --18 DR. ZIEMER: -- I was a little protective of 19 the -- of the subcommittee's work during the 20 early days, but they've matured and actually 21 they get along better without me sometimes, but 22 we can certainly do that and -- and think 23 about, for example, having a separate 24 subcommittee for dose reconstruction and a 25 separate one for the site profiles, for

1	example. Right now it's a combined thing, but
2	understood. And if necessary, Lew, we
3	certainly can I think the ones who are sort
4	of the main ones on that initial subcommittee
5	could meet, with or without the existing Chair,
6	in between our next or before our next
7	meeting.
8	DR. WADE: Right, I think I think we'll push
9	to see a subcommittee meeting before our next
10	meeting, with the task three issues on it at a
11	minimum.
12	DR. ZIEMER: Now we've just been avoiding
13	getting to these final 20 cases, Mark, but
14	MR. GRIFFON: I got to do it now?
15	DR. ZIEMER: the time has come.
16	MR. GRIFFON: All the crowd's gone, it wor
17	no.
18	DR. ZIEMER: Let's make sure that we all have
19	the documents.
20	MR. GRIFFON: That's what I was going to ask,
21	can we take like a 10-minute and I can hand
22	out
23	DR. ZIEMER: Oh, okay, Mark, you
24	MR. GRIFFON: One more delay.
25	DR. ZIEMER: Yes, we will take a 10-minute

1 break and then reconvene. 2 (Whereupon, a recess was taken from 3:05 p.m. 3 to 3:15 p.m.) 4 DR. ZIEMER: We're ready to resume 5 deliberations. We have already lost one person 6 -- we didn't lose him; he had to leave. 7 Espinosa had to leave. We still have a quorum. 8 Mark is all set to go, but the Chair discovered 9 that there is yet another item, Mark. 10 REVIEW AND APPROVAL OF DRAFT MINUTES 11 We actually did not approve the minutes. 12 have two sets of minutes. The first, the 13 minutes of the subcommittee, of our last 14 meeting. I'd like to ask if there are any 15 additions or corrections to the minutes of the 16 subcommittee. 17 Motion to approve the minutes? 18 MR. PRESLEY: So moved. 19 DR. ZIEMER: Second? 20 MR. OWENS: Second. 21 DR. ZIEMER: All in favor, aye? 22 (Affirmative responses) 23 Opposed, no? 24 (No responses) 25 Minutes are approved for the subcommittee.

1 Now the minutes of the Board itself, this is 2 the February meeting in St. Louis. I'm going 3 to first point out a couple of changes. 4 Executive Summary, page 6 -- Executive Summaries don't have to have a lot of detail. 5 I'm going to ask that the sentence under 6 Tuesday, February 8th, where Ziemer announces 7 8 to members of the public to utilize the 9 microphones and so on, I don't think that needs 10 to be in there. If there's no objection, we'll 11 delete that from the Executive Summary. 12 I assume that you all have looked at your 13 There's one place where we talk about places. 14 Mark Griffon's motion, and I'm -- I have a 15 feeling it was a motion, not notion. Mark, you didn't discover that? 16 17 MR. GRIFFON: I have no notion. 18 DR. ZIEMER: You have no notions. 19 MS. MUNN: Sometimes it's right. 20 DR. ZIEMER: On -- there are actually several 21 versions. I'm looking -- in my version it's 22 page 15. It's probably 15 on yours -- broad 23 heading: Site Profile Modifications and 24 Schedule, Status Report. Is that on page 15 25 for you? Go down under Mr. Kenoyer, second

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MS. MUNN:

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paragraph, the 19 site profile cases, last sentence, minor change but change it to read "teams remain intact and are moving ahead" -- teams remain intact and are moving ahead -- minor grammatical change.

On page 16 -- and I need some help on this.

This was the issue of holding the vote open. The last sentence is fairly -- that whole paragraph, which is one sentence, is long and convoluted. I would like the Board's permission to break it into pieces and to explain -- by adding some words such as "the vote was held open so that the votes of Dr. Anderson and Dr. Andrade could be obtained, " and then in parentheses -- and here's where I need help -- I was going to add "Dr. Anderson voted for the motion" as we have the vote but then we don't indicate -- the vote was held open, but then what, so we would indicate Dr. Anderson voted for the motion. And then we've got to say something about Dr. Andrade. going to say Dr. Andrade -- Andrade's untimely death on February 10th precluded his participation. Does that sound too crass or --

No. No, that's appropriate. That's

1 to clarify. 2 DR. ZIEMER: He was unavailable? 3 MS. MUNN: No. Demise. Demise, yeah. 4 DR. ZIEMER: Is untimely death okay? 5 MS. MUNN: Untimely death or demise. 6 DR. ZIEMER: Okay. 7 DR. ROESSLER: No, untimely death. It's 8 factual and --9 DR. ZIEMER: So I will give Cori words to that 10 effect, but that's agreeable. I want to be 11 sure to show the open vote and close that loop 12 on that one. Okay. Thank you. Without objection, I'll make that change. 13 14 UNIDENTIFIED: (Off microphone) No objections. 15 UNIDENTIFIED: (Off microphone) 16 (Unintelligible) give all these to Cori? 17 DR. ZIEMER: Yes, I will. On page 17, Board 18 Working -- Board Discussion/Working Sessions. 19 It starts out (reading) Returning to the issue 20 of the SEC evaluation. Dr. Melius, I didn't 21 understand that at all, and I wondered if you 22 did. (Reading) Dr. Melius offered he didn't 23 feel it would have been more helpful than 24 having the site --25 DR. MELIUS: I was just noticing the same

1 thing. 2 DR. ZIEMER: Could you help us -- if it's 3 agreeable with the group, we'll ask Dr. Melius 4 to tell us what that means. No, maybe we can 5 work on this afterwards. I don't think the sentence makes much sense, as written. 6 7 not clear to me, at all. 8 DR. MELIUS: I'll give you a change. 9 DR. ZIEMER: Yes. On page 20, Board 10 Discussion, where it says (reading) Dr. Ziemer 11 cited the Board -- I'm suggesting we just omit the words "the Board to" and just say Dr. 12 Ziemer cited the section. It's the section 13 14 being cited. 15 There's a spot that -- oh, here it is, page 38. 16 Mr. Griffon, his motion was -- is this a motion 17 or a notion? This is what I was asking about, 18 Mark. 19 MR. GRIFFON: I think it is notion, right? 20 Yeah. 21 DR. ZIEMER: I wasn't sure whether that was 22 referring to a previous motion that you had 23 made --24 MR. GRIFFON: No. 25 DR. ZIEMER: Okay. So it is notion then?

1 MR. GRIFFON: Yes. 2 DR. ZIEMER: Okay. Very good. 3 MR. GRIFFON: Great notion. 4 DR. ZIEMER: So the notion will remain. 5 page 55, the second paragraph from the end, it says (reading) An unidentified member of the 6 7 audience indicated Congress had been aware dose 8 reconstruction wouldn't happen overnight. 9 I remember that particular part of our meeting 10 and in fact the unidentified member was Tom 11 Horgan, and I think the minutes should so note 12 that, and in fact I pointed that out to Tom and 13 he indeed would like that statement to be 14 attributed to him. 15 UNIDENTIFIED: Where is that? 16 UNIDENTIFIED: Which one? 17 DR. ZIEMER: That's on page 55, at least in the copy I'm looking at. Second to the last 18 19 paragraph where it refers to an unidentified 20 member of the audience. The unidentified 21 member has now been identified. 22 MS. MUNN: Who was it? 23 DR. ZIEMER: Tom Horgan from Senator Bond's 24 staff. 25 Are there any other corrections or additions to

the minutes?
(No responses)
Then with those changes, I can have a motion
for approval, as amended.
MS. MUNN: So moved.
DR. ZIEMER: Second?
MR. PRESLEY: Second.
DR. ZIEMER: All in favor, aye?
(Affirmative responses)
Opposed, no?
(No responses)
Thank you very much, the minutes are approved.
Now we're ready does anyone else have
anything that we can put in here before we get
to Mark?
The documents that you need now
DR. MELIUS: Quit stalling, Mark, let's get
going.
DISCUSSION OF FIRST 20 DOSE RECONSTRUCTION
REVIEWS
DR. ZIEMER: I think the latest version
there should be an April 27th draft. It's a
it's a mark-up draft called individual dose
reconstruction case review progress report,
first 20 cases. You should also have a summary

1	of findings matrix, cases 1 through 20. And do
2	we need to have the SC&A checklist?
3	MR. GRIFFON: Yeah, they they actually just
4	gave me a a revised checklist, so I don't
5	have copies of that and I should have
6	DR. ZIEMER: Perhaps everybody
7	MR. GRIFFON: It's only two pages.
8	DR. ZIEMER: Everyone has their original
9	checklist. You can are there many changes
10	in it?
11	MR. GRIFFON: Yes, it's it's kind of
12	different. Number one they they did
13	totals on those deficiencies now.
14	DR. ZIEMER: Okay.
15	MR. GRIFFON: And the second thing, they made a
16	new column for unknown or uncertain, so it
17	looks a little cleaner.
18	DR. ZIEMER: Okay. Are you able to proceed,
19	though, Mark, with what we have at hand and
20	MR. GRIFFON: Yeah, yeah
21	DR. ZIEMER: So let's proceed with
22	MR. GRIFFON: let's try this then.
23	DR. ZIEMER: what we have here.
24	MR. GRIFFON: Okay. I guess it makes sense to
25	start off with the text.

1 DR. ZIEMER: Yes. 2 MR. GRIFFON: The summary report that we 3 discussed in the subcommittee meeting. 4 DR. ZIEMER: Yes, and incidentally, the 5 subcommittee approved this document conceptually for the full Board, so this 6 7 constitutes a motion. It is before us for 8 action. 9 MR. GRIFFON: Right. 10 DR. ZIEMER: There were some places that needed 11 to be filled in, some numbers and other things like that, but this is for formal action, and 12 13 most of you have already seen it in 14 subcommittee session. Thank you. 15 MR. GRIFFON: Yeah, and -- and there still are 16 some places where numbers need to be filled in, 17 but we've made a lot of progress these last 18 couple of days, and I thank Kathy and Hans 19 Behling from SC&A. They helped me pull some 20 pieces together. 21 The third paragraph -- let's see, actually -- I guess the -- the first changes that really take 22 23 place in this draft are -- don't occur until 24 the second page, the second paragraph. 25 MS. MUNN: The aforementioned SCA?

24

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MR. GRIFFON: Yeah, and it says (reading) The aforementioned SCA report includes a summary checklist -- I identified that as a checklist now -- with findings in behalf of -- that's the way it reads on -- in the report -- 20 case reviews, instead of the 15. And then I left --I left the numbers in here. I know we had had some discussion about not reporting the numbers, but I -- I tried to include their summary numbers from their checklist, which we're getting copies of right now, and it indicates that a total of 69 identified in the 20 cases -- SCA considered the majority of deficiencies, 49 out of 69, to be low level deficiencies, with four scored as medium level deficiencies. The question here is, someone's going to add that up and see that it doesn't equal 69. The rest are unknown, and I wasn't sure exactly how to integrate that into -- into this summary report. I also feel a little uneasy submitting a report to the Secretary where we have 16 unknown that we can't -- you know, unknown ranked findings. That's a little interesting position.

DR. ZIEMER: Yeah, maybe we can talk a little

1 bit about what that means --2 MR. GRIFFON: Right. 3 DR. ZIEMER: -- in terms of unknown. Can you 4 describe that --5 MR. GRIFFON: Well, yeah --6 **DR. ZIEMER:** -- whole thing? 7 MR. GRIFFON: -- if -- I think for the most 8 part, if we look at the matrix, it -- it -- it 9 -- most of these unknowns come on on the AWE 10 cases and -- which are the first one through five in our listing, and the -- if you notice 11 in the last column, the Board action, number 12 13 six, that means -- and NIOSH's resolution 14 actually is written there so you can see it. 15 These have been deferred to site profile 16 reviews, so you know, SC&A basically said well, 17 since we've deferred discussion on these issues 18 and resolution of these issues, we don't know 19 how significantly it could affect the case at hand, and that -- that's sort of why they 20 21 ranked it as unknown, I guess. DR. ZIEMER: I wonder if -- terminology-wise, 22 23 if we used words such as "the impact of which 24 has not yet been determined" or something, so

that it's clear that it has -- it is yet to be

25

1 resolved, as opposed to we just don't know. 2 What we need is some wording to that effect, I 3 suppose. 4 MR. GRIFFON: Yeah, "yet unresolved" or 5 something to that ef-- I agree with your notion 6 -- notion. 7 MS. MUNN: Could that be incorporated in the --8 DR. ZIEMER: Should I make my notion into a 9 motion? 10 What -- do you want to suggest particular words 11 right now or do you just want to ponder that? 12 Maybe -- maybe we can try to solve that. 13 mean we need something here to act on. 14 MR. GRIFFON: Yes, I mean if someone can come 15 up with -- we can insert a sentence right after 16 that, where we talk about the numbers, saying 17 that 16 deficiencies are -- are -- they're --18 the potential -- the potential significance of 19 16 deficiencies remains to be determined, or --20 or is -- is as yet undetermined. Or has yet to 21 be resolved, yeah. 22 DR. ZIEMER: The potential impact of the other 23 16 deficiencies --24 MS. MUNN: Why don't you just say the remaining 25 deficiencies?

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              DR. ZIEMER: -- has not yet been resolved or --
2
              or determined?
3
              MS. MUNN: Or still in resolution.
4
              DR. ZIEMER: Yes.
5
              MR. GRIFFON: Potential impact of the remaining
              deficiencies --
6
7
              DR. ZIEMER: Of the remaining 16 -- is it 16?
8
              MR. GRIFFON: If you want to put -- yeah.
9
              MS. MUNN: Of the remaining deficiencies.
10
              DR. ZIEMER: How many is it?
11
              MR. GRIFFON: It is 16, yeah.
12
              DR. ZIEMER: We have 40 --
13
              MR. GRIFFON: It is 16.
14
              DR. ZIEMER: Of the remaining 16 deficiencies -
15
16
              MR. GRIFFON: Is yet --
17
              DR. ZIEMER: -- is yet or remains to be resol--
18
              is yet -- has not yet been resolved? Or not
19
              yet been determined?
20
              MS. MUNN: I would say are still in resolution.
21
              DR. DEHART: (Off microphone) (Unintelligible)
22
              which implies --
23
              DR. ZIEMER: Or is -- is --
24
              MR. GRIFFON: Still in resolution?
25
              DR. ZIEMER: Yeah, are still under review?
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1 DR. DEHART: (Off microphone) Under review, 2 something sounds (unintelligible) --3 DR. ZIEMER: Let's see, potential impact is 4 singular -- is still under review. Right? 5 MR. GRIFFON: Is still under review, okay. 6 DR. ZIEMER: Potential impact of the remaining 7 16 deficiencies is still under review. 8 MR. GRIFFON: Potential -- I said potential 9 significance, otherwise, the same thing. 10 DR. ZIEMER: Significance. 11 MR. GRIFFON: 'Cause that's the way the 12 checklist is labeled, potential significance. 13 DR. ZIEMER: Yes, thank you. Of the remaining 14 -- is still under review. So that will take 15 care of the ambiguity of the unknown. 16 MR. GRIFFON: Right. Okay, thank you. 17 And then the last sentence, (reading) It is 18 noted by the -- by SC&A that the sum -- and 19 this is an addition in the bottom of the 20 checklist which has just been handed out to --21 this is just a clarification of how to 22 interpret these numbers, which we discussed in 23 the subcommittee. 24 DR. ZIEMER: Yes, right. 25 MR. GRIFFON: (Reading) It is noted by SCA that

1 the sum of the deficiencies for these 20 cases 2 should not be used to gauge the impact on 3 individual cases since several low level 4 deficiencies for one individual case may raise 5 the potential significance of (sic) that case. 6 DR. ZIEMER: Very good. MR. GRIFFON: 7 Is that --8 DR. ZIEMER: Uh-huh. 9 MR. GRIFFON: Okay. 10 DR. ZIEMER: Let's ask if that is agreeable. 11 Conceptually we agreed that that sentence 12 should be added. Is everybody comfortable with 13 that? 14 Appears to be, okay. 15 The next paragraph, the only -- I MR. GRIFFON: 16 just added parenthetically case ranking and 17 site/program-wide ranking in there to better 18 define -- as Wanda pointed out, this -- this --19 you know, we have SC&A's checklist, then we 20 have our matrix, and it's still going to be a -21 - may be a bit difficult to walk through, but I 22 tried to better define the Board's names for 23 these things, case ranking and site/program-24 wide ranking.

And then the next paragraph, toward the bottom

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I -- I did some editing to say (reading) case rankings are the same as those listed in the aforementioned SCA summary checklist.

This -- we spent quite a bit of time -- I spent quite a bit of time with Kathy Behling and she really did the grunt (sic) of this work trying to match up the checklist against the matrix. In the future I think -- the -- the good part of this was in the future we're going to have SC&A develop the checklist and matrix so that we won't have this -- this merge issue. Right now, as it stands -- and this -- I don't know if this is going to create confusion, but right now, as it stands, the number of items on the matrix does not equal the 69 mentioned in the checklist, but that's because Kathy pointed out -- and -- and where in the matrix -- like under 1.1 we put in parentheses that G.2 refers -- in the summary of the findings, G.2 refers back to that checklist. We wanted to have some way to tie them together. And what -- what happened here is that in the discussions -- my matrix was developed from the early discussions of all the findings, and I think that when they

did the checklist they sometimes rolled two of

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these findings into one deficiency, so there -there's more of -- more of these in the matrix
than there are in the checklist. I think it's
like 80 to 69 -- you know, it's slightly
different.

That won't happen the next time. I think that would be a lot better if it just matches up neatly.

DR. ZIEMER: Yes.

MR. GRIFFON: Yeah. Anyway, moving on to that sentence, (reading) Case rankings are the same as those listed in the aforementioned SCA summary matrix -- summary checklist. site/program-wide ranking considered the broader potential impacts of the findings and resulted in 49 -- I don't know if I need parentheses there -- 49 low level deficiencies, 35 -- that number is actually wrong. I just -and this is -- this is what we really have to go through the matrix and -- and discuss these, the site/program rank, low level, medium and high level findings -- 49, 35 and three high level deficiencies. And that -- that number, 35, actually has gone down by -- just looking through as I'm editing. Not making apologies,

1 but doing this at 5:30 in the morning, you 2 know, I made some mistakes, so we -- we should 3 go through these and make sure that -- I think 4 it's more like 32 now, something around 32, but 5 I'll -- I'll sum those up again after we go 6 through our discussions of the individual 7 findings, anyway, so... 8 MS. MUNN: (Off microphone) So this is the same 9 49 (unintelligible). 10 DR. ZIEMER: Did you have a question, Wanda? 11 -- we -- need to use the mike there. 12 MS. MUNN: I don't really know what I'm asking. Is that the same 49 where -- I'm --13 14 MR. GRIFFON: No, yeah, it -- this --15 MS. MUNN: A different 49. MR. GRIFFON: Yes. It's a different 49 'cause 16 17 the total number of findings on our matrix -it doesn't match up, so to make that happen 18 19 we'd have to -- we'd have to go back and -- and it might be worth it, just for clarity, 20 21 although I don't look forward to the -- to the 22 task. It doesn't match up one-to-one, so 23 that's -- might be confusing. 24 DR. ZIEMER: Well, as long as we know what --25 we're using it -- I suppose you're concerned

1 about if it goes forward in some form to the 2 Secretary, that that may raise confusion. 3 could add a sentence to explain that, I 4 suppose. I'm not sure which would be the 5 better route. MR. GRIFFON: 6 Right. 7 DR. ZIEMER: To try to match it up one-to-one 8 or simply indicate that there was this -- would 9 you call it an overlap or something of that 10 sort? 11 MR. GRIFFON: Yeah. Yeah, I -- you know, I 12 think it's -- it's -- it's an over-- they 13 rolled -- they rolled some of the findings 14 under discussion into one -- one deficiency in 15 their checklist, so that was -- that's how that 16 hap-- that's how it happened. 17 DR. ZIEMER: Right. 18 MR. GRIFFON: But there's no -- there's no 19 different findings. I should say that. You 20 know, they're all the same set of findings, 21 it's just that sometimes they rolled them into 22 one finding as opposed to keeping them 23 separated out. 24 DR. ZIEMER: Yeah. 25 Anyway, I'm -- I'm willing to do MR. GRIFFON:

1	it either way you want there, if we want to add
2	some clarification or if we want to go back and
3	make it match, either way. I can work with
4	SC&A on that and if we need to.
5	DR. DEHART: Mark, I hate to suggest it, but I
6	think for clarification for ourself
7	MR. GRIFFON: Yeah, I know, I know.
8	DR. DEHART: that it's that you'll
9	probably need to do that or somebody do
10	that.
11	DR. ZIEMER: Wouldn't a single sentence,
12	though, rather than have him go back and try to
13	re-do that whole matrix you're talking about
14	re-doing the matrix otherwise, are you not?
15	MR. GRIFFON: I think I'd have to re re-do
16	the matrix anyway.
17	DR. DEHART: Yeah.
18	MR. GRIFFON: So that that but that's
19	fine. I think we need to. And like I said,
20	this next the next round, with SC&A
21	developing the matrix as they go, this won't
22	happen. You know
23	DR. ZIEMER: It won't happen
24	MR. GRIFFON: we won't have this match-up
25	issue, right. So I I will I will do

1 that. I still think we -- we could step 2 through --3 DR. ZIEMER: Sure, let's --4 MR. GRIFFON: Yeah. 5 DR. ZIEMER: -- do that. 6 MR. GRIFFON: The next paragraph was also 7 modified. (Reading) SC&A concluded that 19 of 8 20 dose reconstructions reviewed during this 9 initial basic audit were considered to be 10 sufficient for the purposes of determining 11 probability of causation -- parentheses, case 6 12 DR may be -- may not be sufficient -- for the 13 specific cases reviewed; however, concerns were 14 identified which could have a broader impact on 15 the overall dose reconstruction program. 16 That's a little -- that might still need some 17 editing 'cause it's not really a "however" 18 anymore. 19 MS. MUNN: (Off microphone) That 20 (unintelligible) be taken out (unintelligible). 21 MR. GRIFFON: Yeah. 22 Is there -- it's not clear to me DR. ZIEMER: 23 why that one sentence -- Case 6 DR may not be 24 sufficient -- is in parentheses. Isn't that 25 simply the next statement?

1 MS. MUNN: Yeah. 2 DR. MELIUS: Yeah. 3 MS. MUNN: It's a stand-alone sentence. 4 MR. GRIFFON: All right. 5 DR. ZIEMER: In fact, you might say "However, case 6 -- the case 6 dose reconstruction may 6 7 not be sufficient." Is that what --8 MR. GRIFFON: Yeah. 9 DR. ZIEMER: -- basically what is being said? 10 MR. GRIFFON: Yeah, okay. 11 DR. WADE: Is there any chance of that being 12 resolved? I mean it would seem to me if -- if we knew that issue, it would be more complete, 13 14 or will we not know that issue? 15 DR. ZIEMER: I think that goes back to -- we 16 may not even need to know that. 17 DR. MELIUS: Yeah. 18 DR. ZIEMER: That -- that becomes a NIOSH 19 issue, I believe. This is -- this is simply a 20 finding. There's one case where the DR may not 21 have been sufficient. We're not asking that 22 that -- that that be resolved for this report. 23 MR. GRIFFON: Right. 24 DR. ZIEMER: We're simply reporting it --25 MR. GRIFFON: I think that --

1 DR. ZIEMER: It seems to me it goes back to 2 NIOSH. Right here it's just one de-identified 3 case. NIOSH will know what case it is. It's -4 - the burden on them would be to take action. 5 I don't think the purpose of this is for us to 6 resolve issues on cases individually. 7 Is that the understanding of the group? Yes. 8 Oh, I ended up -- however, Mark -- with two 9 "howevers". 10 MR. GRIFFON: Yeah, I was going to say we might 11 want to have that sentence read, after the 12 parentheses with POC -- finish it up to say 13 "for the specific cases reviewed," period. 14 Then say "However, case 6 dose reconstruction 15 may not be sufficient, " period. 16 DR. ZIEMER: Period. 17 MR. GRIFFON: And then "Additionally, concerns 18 were identified which could have a broader 19 impact on the overall dose reconstruction 20 program." That's a little better. 21 DR. ZIEMER: Did everybody get that? Doesn't change the concept -- a little bit of 22 23 wordsmithing. Okay. 24 MR. GRIFFON: No. 25 DR. ZIEMER: We'll agree that that's

1	acceptable. Proceed.
2	MR. GRIFFON: Okay. I actually don't think
3	there were any other there was a an
4	editorial change somewhere in the last
5	paragraph, but that was that was it. Oh
6	DR. ZIEMER: The last paragraph on the
7	conclusions, or that section?
8	MR. GRIFFON: The last paragraph on page four,
9	there was just a change editorial to where
10	DR. ZIEMER: Ongoing concerns?
11	MR. GRIFFON: Where yes, consistency of
12	cases.
13	DR. ZIEMER: And the change is just that
14	deletion?
15	MR. GRIFFON: Just there are "there are"
16	instead of "you have" similar
17	DR. ZIEMER: Oh, yes, that was just an
18	editorial
19	MR. GRIFFON: That's all yeah. That was
20	the only other I believe the only other
21	changes.
22	DR. ZIEMER: Okay, any questions or comments on
23	this this
24	MR. GRIFFON: I also one thing one other
25	thing I should point out, on page two it says

1 insert table with one through 20 and sites, 2 POC, et cetera. Stu Hinnefeld did provide 3 this. I don't know if he handed or --4 DR. ZIEMER: Yes, that was distributed and 5 should be at your place. 6 MR. GRIFFON: So we'll probably just ask to get 7 this in electronic form and insert it in there 8 and --9 DR. ZIEMER: Right, so this is Table 1. 10 MR. GRIFFON: Right. 11 DR. ZIEMER: Which lists the case number, using 12 the pseudo-number 1 through 20, it gives the 13 probability of causation, the IREP cancer 14 model, the location, working years and work 15 decade. Which is what we had when we did the 16 selection, so that gets inserted. Okay? 17 MR. GRIFFON: Right. 18 DR. ZIEMER: And then attached to this would be 19 the scorecard and the matrix. 20 The checklist and the matrix. MR. GRIFFON: 21 think we were -- did we refer to the Board --22 or the Board methodology I think we referred 23 to, also, did we not? 24 DR. DEHART: You intended to put that in there. 25 I don't know, may... MR. GRIFFON:

1	MR. PRESLEY: (Off microphone) On on Table
2	1, (unintelligible) get that?
3	MS. MUNN: Yeah.
4	MR. PRESLEY: Third one down, you need to
5	change that from 1040 to 1940.
6	MR. GRIFFON: Oh, yeah.
7	DR. ZIEMER: Yeah, that was one of our earliest
8	earliest work decades.
9	MR. GRIFFON: Yeah. We refer to this the
10	Board has developed a methodology, attachment
11	2, so this would be this this other text.
12	DR. ZIEMER: Okay, Attachment 2 will be the
13	methodology previously approved. Right?
14	MR. GRIFFON: Did we previously approve that?
15	DR. ZIEMER: Did we approve that? I thought we
16	had or did we?
17	DR. DEHART: I don't think so. We talked about
18	it.
19	MR. GRIFFON: I don't know, 'cause we added
20	I added on these these ranking or not
21	rankings, but these action action numbers,
22	six different actions may be taken, and that's
23	the last column in the matrix now.
24	DR. ZIEMER: Right.
25	MR. GRIFFON: So we should look at that.

1 DR. ZIEMER: So we do need to look at that. 2 MR. GRIFFON: Yeah, I think we can look at that 3 along with the matrix. That would make sense. 4 DR. ZIEMER: Now that document is entitled 5 Methodology for Categorizing and Ranking DR 6 Case Review Findings. That would be Attachment 7 2. 8 So Mark, are you ready for us to review that 9 then? 10 MR. GRIFFON: Sure, yep. 11 DR. ZIEMER: Okay. There's an introductory 12 paragraph. Does anyone have any issues to 13 raise with that? 14 (No responses) 15 The rankings -- it simply describes a graded 16 approach and gives the bullets -- criteria, 17 basically. 18 MR. GRIFFON: Yeah, the difference -- one 19 difference from before is that we -- we had one 20 -- I had one ranking before and now we've got 21 these -- these case-specific and site/program-22 wide rankings, so that paragraph on the ranking 23 of findings changed a little bit. 24 DR. ZIEMER: Yeah, how --25 MR. GRIFFON: The bullets -- the criteria are

1 similar, though. 2 DR. ZIEMER: Right, how -- how have you 3 changed? Do we have the latest copy? 4 MR. GRIFFON: Of the matrix? 5 DR. ZIEMER: My copy still says there's a 1 to 6 5 ranking system. Do I have --MR. GRIFFON: Oh, you have the right copy and I 7 8 didn't cha -- I changed the matrix, but I didn't 9 change the numbers on there. I have low, 10 medium, high now on the... 11 DR. ZIEMER: Okay, so that first paragraph 12 under Ranking the Findings, let's -- let's look 13 at how that should be worded. 14 (Reading) The graded approach should be based 15 on the importance of the identified finding, 16 other cases at the facility or other cases 17 program-wide. Two separate rankings will be 18 assigned, case ranking and site profile --19 site/program-wide ranking. 20 And then we would say what, a low, medium, high 21 ranking system? Or --22 MR. GRIFFON: Are based on a low, medium, high 23 ranking system? To rank -- two separate 24 rankings will be assigned based on a low, 25 medium, high ranking system -- low, medium,

1 high qualitative ranking system? I don't know. 2 DR. ZIEMER: So in parentheses, rather than a 1 3 to 5 ranking system, we would say a low, 4 medium, high ranking system. 5 MR. GRIFFON: That's fine. 6 DR. ZIEMER: We don't have to explain that; 7 it's self-explanatory then. Is that agreeable? 8 (No responses) 9 And then you would have your bullets saying 10 what the rankings are based on. Any comments 11 on those? They remain the same. 12 (No responses) 13 Anything on the categorization? We have 14 actually now the -- the six categories that 15 Mark has suggested, I think these we need to 16 agree on, that would be used in the summary ma-17 - matrix. Those are on that second page, 18 options for Board action, 1 through 6. 19 Let me ask if there's any changes or 20 modifications. 21 MR. GRIFFON: And -- and number 4 and 5, I 22 should just note that, you know, NIOSH 23 disagreed; Board and NIOSH reach compromise. 24 Sometimes in -- in the -- in the matrix notes 25 you'll see, you know, SC&A and NIOSH are in

1	agreement. I think SC you know, it really is
2	the Board SC&A on our behalf, I guess, so
3	DR. ZIEMER: Let me make sure I understand
4	number 5. This would be one where NIOSH says
5	that they disagree, and basically the Board
6	says okay, we're not going to do anything about
7	that. In essence, we are accepting the
8	disagreement if we do nothing.
9	MR. GRIFFON: That's correct, yeah. Maybe I
10	should say Board accept
11	DR. ZIEMER: I don't know, I'm asking
12	MR. GRIFFON: That was the
13	DR. ZIEMER: how you wish to characterize
14	MR. GRIFFON: That was the intent.
15	DR. ZIEMER: I think the I think the effect
16	is, if we drop the matter, the Board is
17	accepting NIOSH's
18	DR. MELIUS: Yeah.
19	DR. ZIEMER: disagreement.
20	MR. GRIFFON: Right.
21	DR. MELIUS: And I apologize 'cause I haven't
22	been involved in the subcommittee meetings so I
23	may misunderstand, but the prior point that you
24	brought up, Paul, regarding SCA and NIOSH
25	excuse me, SCA and the Board being the same

1	entity, my understanding of the process is that
2	there's a resolution process that goes on
3	between SCA and NIOSH prior to our involvement,
4	and so I guess I'm a little concerned that it
5	sort of somehow implies that the Board has
6	approved of whatever SCA's
7	DR. ZIEMER: This is the final wrap-up
8	DR. MELIUS: So it just applies
9	DR. ZIEMER: where
10	DR. MELIUS: to the final.
11	DR. ZIEMER: the final thing that would go
12	in the right-hand column after all of the back-
13	and-forth iterations have occurred, how the
14	Board finally disposes of all issues.
15	DR. MELIUS: Okay.
16	DR. ZIEMER: For example, if if where
17	where it says NIOSH
18	DR. MELIUS: No, I I understand that.
19	DR. ZIEMER: NIOSH agrees, they'd be
20	agreeing with SC&A and accept that that closes
21	it.
22	DR. MELIUS: Okay, I may have misunderstood
23	Mark then 'cause I thought he said Mark was
24	saying that S was implying that the Board and
25	SC&A were equivalent and

1	MR. GRIFFON: No, no, I
2	DR. ZIEMER: No, the Board is ultimately taking
3	an action.
4	MR. GRIFFON: Yeah, I meant I meant as far
5	as it applies to this
6	DR. MELIUS: Yeah.
7	MR. GRIFFON: Right. I'm sorry.
8	DR. MELIUS: So so it should say the Board.
9	MR. GRIFFON: Yeah.
10	DR. ZIEMER: Yes.
11	DR. MELIUS: Okay.
12	DR. ZIEMER: So number 5 would say, rather than
13	the Board drops it, the Board accepts? But
14	what what that's NIOSH disagrees; the
15	Board accepts
16	DR. MELIUS: The Board concurs with NIOSH, I
17	think.
18	DR. ZIEMER: Board concurs, 'cause that's the
19	effect.
20	MR. GRIFFON: Yes. Yes.
21	DR. ZIEMER: And then any others?
22	(No responses)
23	We may find as we proceed that some of these
24	are not useful or we may need others, and of
25	course the Board can change these at any time.

1 I think Mark is suggesting this may be a good 2 starting point, and it enables us to come to 3 closure, at least on this first set of 20. 4 MS. MUNN: Sorry. 5 MR. GRIFFON: I'm sorry, we had a sidebar. 6 DR. ZIEMER: I'm just saying, Mark, it seems to 7 me that once we put these into use, if we find 8 that some of these are not useful or we need 9 others, we can always modify this, but --10 MR. GRIFFON: We can always revise that, right. 11 DR. ZIEMER: -- as a start, this seemed --12 based on your work on the matrix -- seemed to be a useful way to come to closure on the 13 14 issues that have been identified. 15 MR. GRIFFON: Right. I tried. 16 DR. ZIEMER: Do I take it by consent then that, 17 with that minor change, these six categories 18 are agreeable? 19 (No responses) 20 Stu? 21 MR. HINNEFELD: I didn't want to interrupt, but 22 I was just curious who said --23 DR. ZIEMER: Yes, you did. 24 MR. HINNEFELD: I have, okay, sure. Who should 25 we send Table -- the electronic version of

1	Table 1 with the changed date in it?
2	DR. ZIEMER: I think get it to Mark so he can
3	insert it. He has the electronic
4	MR. GRIFFON: (Unintelligible) forward it to
5	yeah.
6	DR. ZIEMER: Or Cori? Who?
7	MR. HINNEFELD: E-mail's easy, I can send it to
8	everybody.
9	MR. GRIFFON: I guess send it to me first and
10	then I'll send the dra
11	DR. ZIEMER: So he can incorporate it, yeah.
12	MR. GRIFFON: I'll make these changes and
13	forward the draft to everyone.
14	MR. HINNEFELD: Okay.
15	DR. ZIEMER: Send it to Mark.
16	MR. HINNEFELD: That's fine.
17	DR. DEHART: Could I ask that the next time we
18	see it it's been a while since I've seen a
19	clean text if we could see one without
20	lines.
21	MR. GRIFFON: This was for full transparency of
22	the changes I was making along the way, so it
23	makes for difficult reading, I agree.
24	DR. ZIEMER: Actually you could send bo you

1 MR. GRIFFON: Yeah. 2 DR. ZIEMER: -- a red ver-- or a mark-out 3 version plus a clean version. 4 MR. GRIFFON: Okay, yeah. 5 DR. ZIEMER: Yeah. Are we agreeable on this Attachment 2 then? 6 7 Jim, another comment? 8 DR. MELIUS: Not yet. 9 MR. GRIFFON: He's thinking of one. 10 DR. MELIUS: No, I don't have any right now. 11 DR. ZIEMER: All right --12 DR. MELIUS: Or I actually do have a comment. 13 DR. ZIEMER: Yeah, go ahead. 14 DR. MELIUS: When I read this summary of 15 findings and where this code is put as the 16 Board action, I just -- the heading on the 17 table that are NIOSH resolution, I'm -- I find 18 it a little bit confusing 'cause sometimes it 19 says NIOSH and SC&A agree and sometimes it 20 doesn't, and it's -- it's unclear. I just 21 think, for future -- not to make you go back 22 through and change this -- that we ought to be 23 -- some consistency in the language that we use 24 'cause that's what confused me --25 DR. ZIEMER: Actually I believe we may want

1	both. This this tells about the resolution
2	going on with our contractor, but the final
3	column is the Board's handling of everything up
4	to that point. I believe that's the intent.
5	Mark?
6	MR. GRIFFON: That is the intent, yeah. Yeah.
7	DR. ZIEMER: I believe I believe this is
8	intentional that he
9	MR. GRIFFON: So it's probably
10	DR. ZIEMER: SC&A and NIO that that
11	tracks what happens until we finally take a
12	final action, which shows up in the last
13	column.
14	MR. GRIFFON: But still there's probably
15	inconsistencies, I would I would
16	DR. ZIEMER: But that intent is to have those
17	two entities
18	DR. MELIUS: Okay, then label it more clearly -
19	-
20	DR. ZIEMER: as you go across, and then the
21	Board action. Okay?
22	MR. GRIFFON: Label them more clear yeah.
23	DR. ZIEMER: Gen Roessler.
24	DR. ROESSLER: Since this final column is the
25	most important one, I think a footnote on the

1 bottom of the page on this document would help 2 to put down what number 1, 2, 3, 4 and 5 and 6 3 mean. 4 MR. GRIFFON: Yeah, and I --5 DR. ROESSLER: We could footnote everything, 6 but I think it -- that one thing would help. 7 MR. GRIFFON: Yeah. 8 DR. MELIUS: Well, I would even suggest that 9 we, where necessary, make that column a little 10 bit wider so that if there's -- if we've gone 11 beyond just a simple 1 through 6 in terms of 12 what action the Board has taken, that we sort 13 of write that out a little bit. 14 MR. GRIFFON: Maybe I'll spread this onto legal 15 paper and just write out that they -- yeah. 16 were trying to fit it on one page, too. 17 DR. ZIEMER: Dr. Roessler, did you get your comment in? 18 19 DR. ROESSLER: Yes. 20 DR. ZIEMER: Are we ready to look at the matrix 21 itself then? And do we have -- oh, we do have 22 the -- we do have the checklist, and that would 23 -- Mark, is there anything else we need to say 24 on the checklist? You've already pretty well 25 explained it. That would be inserted in the

1 packet, too. 2 MR. GRIFFON: Yeah, the -- I don't think I got 3 -- did I get... 4 DR. ZIEMER: It -- it now has the 69 --5 MR. GRIFFON: Yeah. DR. ZIEMER: -- deficiencies, the 49 lows, the 6 7 four mediums and the unknown --8 MR. GRIFFON: Which we can label --9 DR. ZIEMER: -- which I think we probably are 10 going to call that something else, SC&A folks, 11 un-- not yet resolved or -- yeah. 12 understand what it is so we're -- we're okay on 13 it. I think we're going to call it something 14 else here. It's not as if we don't know what -15 16 MR. GRIFFON: Right. 17 **DR. ZIEMER:** -- the deficiencies are. 18 going to be addressed in a different way. 19 MR. GRIFFON: The one thing I would ask, and I 20 -- I told SC&A I would -- I would ask this of 21 the Board, this has been revised to include 22 some sections, especially -- I think Section G 23 was revised -- to accommodate the use of the 24 single checklist for DOE and AWE sites, so this

is slightly different than the one you've seen

25

1 before, and they asked if we could review and 2 approve this because they're planning on using 3 it for the next 18 they're already -- you know, 4 so they wanted us to take a close -- closer 5 look --6 DR. ZIEMER: Mark, you've already seen this, so 7 what is your advice to us on that? 8 MR. GRIFFON: My -- my advice is -- is that 9 they -- they made some changes to the 10 footnotes, which I think were important, 11 especially the one -- the sum of the 12 deficiencies, which is similar to the language 13 we put in the summary report, that you 14 shouldn't pay attention to those percentages 15 too much, too closely. 16 DR. ZIEMER: And then Section G --17 MR. GRIFFON: And I -- I think overall, Section 18 G seemed -- seemed appropriate. I still -- you 19 know, my generic concern -- and we'll get into 20 that when we get into the matrix -- is this 21 unknown column or yet-to-be-determined column. 22 I -- I think -- I'm not sure how we can handle 23 that. But as you'll notice when we look in the 24 matrix, a lot of these key issues that we raise 25 in this review have been deferred --

DR. ZIEMER: Right.
MR. GRIFFON: to site profile reviews, and
so how complete can our report be on you
know, and I and I understand this is an
ongoing
DR. ZIEMER: But but at the moment
MR. GRIFFON: Right.
DR. ZIEMER: you feel that it's appropriate
that we have the contractor proceed with this
new format?
MR. GRIFFON: I think so, yes.
DR. ZIEMER: Is the is the committee or
the Board agreeable to that?
(No responses)
Appears to be no objections. Without
objection, we'll consider that that has been
approved for use in the next round.
Now let's proceed to the matrix. We actually -
- Mark, I think you actually need to
individually look at each of these items, do we
not, and make a judgment on them?
MR. GRIFFON: Yeah, yeah.
DR. ZIEMER: And there
MR. GRIFFON: Well, and it to complicate
this matter of one just a little bit, that

1	last column, which Jim rightly pointed out is -
2	- is not I guess it's really SC&A and
3	NIOSH's resolution, but it was provided by
4	NIOSH and SC&A hasn't reviewed that, I don't
5	believe. So so I think
6	DR. ZIEMER: You're talking about
7	MR. GRIFFON: we're not clear if that's
8	DR. ZIEMER: the column
9	MR. GRIFFON: a final resolution
10	DR. ZIEMER: called NIOSH resolution?
11	MR. GRIFFON: Yes. Yes.
12	DR. ZIEMER: So SC&A has not yet seen that and
13	agreed that particularly in those cases
14	where it says that they both agree?
15	MR. GRIFFON: Right.
16	DR. MELIUS: Need to resolve the resolution.
17	MR. GRIFFON: We need to resolve the
18	resolution.
19	DR. ZIEMER: As far as the this Board's
20	action has to be with respect to the last
21	column, and there are a number of items in the
22	list for example, if if the last column
23	is number 1 or is item 1, basically the
24	Board really has to do nothing. I believe
25	that's correct.

1 MR. GRIFFON: Right. 2 DR. ZIEMER: If the designation is 2, the Board 3 really has to do nothing. I mean we can 4 approve. If it's number 3, we would -- we 5 would have to take a specific action. Likewise for 4. Likewise for 5. And 6, perhaps we 6 7 would have to agree that that's what's going to 8 happen. So there's a number of these where --9 MR. GRIFFON: Yeah, and --10 DR. ZIEMER: -- we would have to say yes, that 11 is what --12 MR. GRIFFON: I should point out 3 actually --13 you know, it says "unless the Board recommends 14 action through..." so we may recommend, if it's 15 a 3. I think that's a -- you know. 16 DR. ZIEMER: Right, in the absence of action, 3 17 remains, but we would have to look at 3s to determine whether we want to take action. 18 19 MR. GRIFFON: Right. 20 DR. ZIEMER: Now my question is, does the Board 21 -- does this require SC&A to actually review 22 this before we take action? I mean we can --23 we can approve -- we can approve these 24 documents and the matrix as a -- as a -- as a 25 document format-wise and content-wise, with the

1	exception of approving the last column in terms
2	of the actions. Or particularly if it
3	requires SC&A to do some review before we
4	finalize.
5	MR. GRIFFON: I I guess the the the
6	question on the NIOSH resolution column is that
7	and I've pointed out a few that they've
8	as I was working with them, they found a few
9	where they 10.1 is one example where they
10	indicated that it says (reading) SC&A
11	concurred with the assigned medical dose in the
12	February, 2005 report.
13	And they they indicated to me that they did
14	not agree with that, so I don't want to have a
15	misstatement of facts in this matrix as we move
16	it forward, either, you know.
17	DR. ZIEMER: Well, it appears to me, although
18	we're close to closure, there may be another
19	small step
20	MR. GRIFFON: Yeah.
21	DR. ZIEMER: that has to occur before we are
22	ready to actually close on this. Is that am
23	I correct on that?
24	MR. GRIFFON: I think it's probably yes,
25	yes.

1 DR. ZIEMER: Then what I'm going to suggest is 2 a motion that we accept all of the documents 3 and attachments, including the matrix, except 4 for the Board -- well, except for the -- the 5 rankings --MR. GRIFFON: I don't know that we can --6 7 DR. ZIEMER: What -- what I'm trying to say is 8 that we accept the matrix as a vehicle for 9 doing this, but we're not yet agreeing to 10 either the rankings or the Board actions until 11 SC&A has an opportunity to review the NIOSH 12 resolution column. That would be a motion that 13 would seem to me to be in order, if someone 14 would wish to make it. 15 MR. OWENS: Dr. Ziemer, I'll make a motion that 16 the Board accept all the documents that we have 17 reviewed and that we accept the matrix in 18 principle, awaiting final resolution of a 19 column -- I guess it's NIOSH resolution --20 where SC&A would be involved. 21 DR. ZIEMER: And our Board action, therefore. 22 MR. OWENS: Yes. 23 MR. PRESLEY: I'll second that, but I'll ask a 24 question, also. Do we want to make it the last 25 two columns where --

1 DR. ZIEMER: Yeah, it's the NIOSH resolution 2 column which SC&A would review, plus the Board 3 action column. 4 MR. PRESLEY: Right. 5 DR. DEHART: Yes. 6 DR. ZIEMER: Now I'm going -- since officially 7 this was on the floor as a motion from the 8 subcommittee, I will regard the current motion 9 as a substitute motion that replaced the 10 original one if there's no objection. 11 similar motion, but it is more specific on what 12 will happen here. 13 (No responses) 14 Now, comments on that motion? 15 This is a question I think is DR. MELIUS: 16 relevant. In regard -- again, I was not at the 17 subcommittee meetings or discussions of this, 18 but in regard to the number 3s where -- has 19 there been discussion among the subcommittee as 20 to which ones we would want to move forward in 21 some way? 22 MR. GRIFFON: 23 DR. MELIUS: No. 24 MR. GRIFFON: We didn't get that fa-- I don't 25 think we had this form of the matrix available,

1 did we? In the subcommittee meeting? 2 DR. ZIEMER: Not -- not fully and --3 MR. GRIFFON: Right. 4 DR. ZIEMER: -- and you can look down through 5 here and there are some 3s in here currently, and you'll see the nature of them. They are --6 7 they are valid cases where there's bona fide 8 disagreements between NIOSH and our contractor 9 as to how one might approach things. 10 DR. MELIUS: Well, I would just -- if we're 11 going to close this out, do we want to put out 12 a -- a document that is sort of our final report where we haven't resolved those 3s? 13 14 DR. ZIEMER: No. 15 MR. GRIFFON: No. 16 DR. ZIEMER: No, we're only approving this as a 17 -- as an instrument that still remains to have 18 that last step occur before it's closed out. 19 DR. MELIUS: Okay. And just --20 DR. ZIEMER: We're approving the text, the 21 types of attachments that would go with this 22 report, the matrix, the form of the matrix, the 23 content --24 DR. MELIUS: Uh-huh. 25 DR. ZIEMER: -- with the exception of those

1 last two columns. 2 DR. MELIUS: And again, this is a question --3 does this mean this Board action coding, are we 4 really going to move toward something that 5 would be a 3A and a 3B or something -- 3A where 6 we've taken -- at some point taken a step and 7 recommended to the Secretary that some change 8 be made, or 3B where we did not, or does -- or 9 are we intending to then change the code? I'm 10 just confused by sort of the --11 DR. ZIEMER: Well, I don't know the answer to that. Mark, did -- I don't know if the 12 13 subcommittee addressed that, per se. 14 MR. GRIFFON: We didn't get that far, no. I 15 mean I'm fine with that. That seems like a 16 reasonable approach, to me, that we need to 17 know whether we did or did not send any 18 recommendation to the Secretary on that certain 19 finding, and 3A and 3B is just as good a system 20 as -- you know, does that make sense? 21 but that -- but --22 DR. ZIEMER: Well --23 MR. GRIFFON: -- that isn't done until we go 24 through all the findings. 25 I don't know, I -- I can't pre-DR. ZIEMER:

1 judge what -- which of -- whether that would be 2 better to break it out right now or just to go 3 through it and if there's a specific action for 4 a -- for a 3 item, we just act on it. 5 DR. MELIUS: Uh-huh. DR. ZIEMER: It definitely requires some 6 action. 7 8 This is Bob Presley. I believe MR. PRESLEY: 9 you're going to have to act on it and then the 10 final -- you're going to have to change that 11 code to something you can send to the 12 Secretary. 13 DR. MELIUS: Exactly. 14 MR. PRESLEY: Or if not, you're -- you've still 15 got an open-ended problem. 16 DR. MELIUS: Yeah. 17 DR. ZIEMER: Okay. 18 MR. GRIFFON: All right. 19 MR. OWENS: Dr. Ziemer, the motion was just to accept the -- you know, the documents and --20 21 and the matrix in principle, and whatever 22 revisions or changes might need to be made 23 before we finalize it, I think we could do 24 that, hopefully.

MR. GRIFFON: Yeah.

25

1 DR. ZIEMER: Actually if -- if -- in case of a 2 number 3 finding or closure on a report, there 3 -- there will -- the Board would have to take 4 an action. Whether we call it 3A and 3B or 5 not, there would be an action sort of up or 6 down as to whether you go forward. 7 MR. PRESLEY: You might want to call it 3-1 or 8 3-6. I mean it's something you'd have to show 9 closure on. 10 DR. ZIEMER: Right. 11 MR. GRIFFON: That's fine. 12 DR. ZIEMER: Yes, Wanda. 13 MS. MUNN: Just a question that puzzles me a 14 little, whether -- I assume we want to have as 15 many of the last two columns complete as 16 possible before we send this away. I'm 17 wondering on 12.9 whether that can be one of 18 those that can disappear by putting the 19 response in the resolution column. I can 20 understand not having anything when you have an 21 item that hasn't --22 MR. GRIFFON: Yeah. 23 MS. MUNN: -- been discussed, but --24 MR. GRIFFON: Here -- here's another -- another 25 reason for -- that -- that we still need some

1 work on this table. The NIOSH resolution 2 column came to me in one file and SC&A sent me 3 another file that had additional findings that 4 weren't in the matrix that NIOSH was reviewing, 5 and I merged the two. So where you see blank on NIOSH resolution, it's often that they --6 7 they hadn't considered that one at all, so they 8 need to re-look at this, as well -- if that 9 made any sense. So -- so this is still not a -10 - quite ready for prime time, obviously. 11 needs -- needs to be edited. 12 DR. ZIEMER: The motion is not -- will not 13 preclude doing that. 14 MR. GRIFFON: Right. Right. 15 In fact it would mandate following DR. ZIEMER: 16 up and coming to closure. So there may be some 17 of these that NIOSH also needs to look at. that what you're saying? 18 19 MR. GRIFFON: Yes, yes, probably only --20 probably only four or five that they hadn't --21 that weren't in the version that they were 22 reviewing. This was in real time, as we know. 23 DR. ZIEMER: Yes. Okay. 24 MR. GRIFFON: So... 25 DR. ZIEMER: Other questions or comments?

1 (No responses) 2 So we will vote on this, and if approved we 3 recognize that we have not yet come to closure 4 on the first 20 case. We're getting closer and 5 closer. MR. GRIFFON: Believe it or not. 6 7 DR. ZIEMER: Are we ready to vote? I do want 8 to thank Mark especially for a lot of time and 9 effort put into developing and filling out in 10 the matrix, together with our contractors and -11 - and NIOSH folks who helped pull this 12 together. It's been a good process for us to 13 develop a methodology for handling our -- our 14 reviews. 15 MR. GRIFFON: The only thing I'm comforted by 16 is that I think going forward we have a much 17 cleaner system. 18 DR. ZIEMER: Yes. 19 MR. GRIFFON: 'Cause I'm not looking forward to 20 editing this matrix to make -- to making the 21 numbers match up. We went through two days of this, Kathy Behling and I --22 23 DR. ZIEMER: Right. 24 MR. GRIFFON: -- and it -- but going forward, 25 it'll be much cleaner with --

1	DR. ZIEMER: Yes.
2	MR. GRIFFON: SC&A filling in the matrix.
3	DR. ZIEMER: Very good. Are we ready to vote
4	then?
5	MS. MUNN: Yes.
6	DR. ZIEMER: Okay, let's vote. All in favor of
7	accepting the document under the terms
8	indicated, please say aye.
9	(Affirmative responses)
10	Any opposed, no?
11	(No responses)
12	Abstentions?
13	(No responses)
14	Thank you very much. The motion carries and we
15	are close to closure on the first 20 dose
16	reconstruction reviews.
17	PUBLIC COMMENT
18	We have on our agenda at 4:15 a public comment
19	period. I've received requests from a couple
20	of individuals to address the assembly. First,
21	Delbert Moore I believe from Iowa. Is
22	Delbert in the assembly?
23	(No responses)
24	Does not appear to be. Also well, this is
25	Dan McKeel, it's but he's addressed us

1	already. It says after the after Denise and
2	the SEC, which he's already done, so this
3	this one's already been covered.
4	Were there any other members of the public who
5	had a desire to address the assembly? Please
6	approach the mike and you can identify
7	yourself, please.
8	MR. RUBY: Hello, I'm Doug Ruby.
9	DR. ZIEMER: Doug.
10	MR. RUBY: I'm here representing my dad, John
11	W. Ruby.
12	DR. ZIEMER: Uh-huh.
13	MR. RUBY: I had a quite a lengthy little
14	thing I was going to read, but I noticed in the
15	paper articles that a lot has happened since we
16	came up Sunday to meet with NIOSH, and that you
17	guys apparently have come to some resolution on
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19	DR. ZIEMER: Are you are you from the Iowa
20	group?
21	MR. RUBY: Yes, my father worked at IAAP.
22	DR. ZIEMER: Yes, then you may be aware that
23	the Board has has made a recommendation to
24	the Secretary to approve
25	MR. RUBY: I just read

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DR. ZIEMER: -- Special Cohort status --MR. RUBY: -- that article two minutes ago, so I'd come up and I had a nice long spiel for y'all, and I would like to say that Silas Mason was the contractor my father worked for and that I don't think that -- we were pretty upset when NIOSH wanted to review the declassified DOE information they told us about Sunday. See, they originally denied my father, and I had to appeal it based on the ground water -it was quite alarming a year ago to find out that NIOSH did not take ground water contamination into consideration on that first dose in construction (sic) at that meeting, and then subsequently they've done the right thing. I just want to thank you guys for, you know, being stand-up on this. You've got to realize that a contractor -- DOE may hire somebody to do something, but back then what did we know about these kind of hazards, you know. second off, you know, contractors have been known to fudge on the rules, so to speak. they have some -- some blame in this, but they also were ignorant, as the whole country was, as -- you know, the dangers of working with

1 this kind of stuff. But after eight years of 2 frustration, I just want to thank every one of 3 you guys for being stand-up and doing the right 4 thing. 5 To me, NIOSH was not our friend. But I -- you 6 know, I just felt like they were working 7 against us from the start, but you guys have 8 pretty well shot my statement all to heck, so I 9 guess that's all I wanted to say. Thank you --10 DR. ZIEMER: Okay, thank --11 MR. RUBY: -- very much. 12 DR. ZIEMER: -- you for coming, in spite of the 13 change in what you were going to say. I think probably you're more comfortable than you would 14 15 otherwise have been, so -- but we're glad 16 you're here. 17 Are there any other folks -- yes, please 18 approach the mike. 19 MS. WILBURN-YOAKUM: Yes, my name's Linda Anne 20 Wilburn-Yoakum and my father died 29 years ago 21 May 7th this year. This is my mother in the 22 pink. She'll be 92. I've been five years 23 fighting this and I -- I agree with the 24 gentleman. It's been a bad fight. My father 25 also worked for Silas Mason, and records is

something that just wasn't -- and the government, also. They just said no, we didn't do anything there. Oh, yes we did. And no, we don't have any records.

I've got a couple of questions. I'm sure you've heard lots of crabbing from everybody. You basically approved the (unintelligible) variation on what the radiation levels were going to be. I was unable to make it here Monday and Tuesday. I'm sorry, I know I missed a lot. I did the St. Louis. I just couldn't get my mother and be here.

My father's levels are unknown because of records. He was a steam-fitter and a pipe welder. He worked all over the plant. He didn't work on Line 1, he didn't work on Line 6. He worked all over the place, and I guess I've -- I've got a couple of questions. Iowa City at the University Hospitals told him two weeks before his death that his death was due to his work, and that was what he did at Burlington in the Armory. It's -- it's known and accepted that my father had a rate of -- diagnosis of cancer during the period when they said it was. He had the thyroid cancer.

He had the bladder. It -- it's not a problem there, but the problem is they say he only had 12 and a half percent. Well, who are they gauging these records against and where are they getting their records? What are they made from, other plants, other places? I mean I know you've heard it all.

But my biggest question is, I pushed the button on everything I can. Deb McCurran\*,

Congressman Leach's associate in Ottumwa, has been wonderful the last two years, but it's been very flustrating (sic) for both of us.

Now I have -- I have taken and filed, my mother's been denied. We appealed. I told them they had to come here. My mother wasn't well enough to go, so they came here and they said unless you've got new information, Ms.

Hill said you're just going to be automatically denied, and we were. So I had to get a request in a certain amount of time to keep her case alive, so to speak. Now where does that put her since they only give him a 12 and a half

percent? And I mean like he -- he told us all

hush-hush secret you didn't talk about. Had he

kinds of things before he died. That was a

1 not been dying, he wouldn't have told us 2 nothing. 3 DR. ZIEMER: Perhaps one of the NIOSH people can answer that, but if -- if in fact he was a 4 5 member of IAAP during the designated time period and if in fact the action that this 6 7 Board took yesterday proceeds through Congress 8 -- which is the ultimate step -- then I would 9 assume -- and NIOSH, you can help me out here -10 - I would assume that all of these folks are 11 part of that cohort, are they not? 12 MS. WILBURN-YOAKUM: Are they accepted in this 13 even though they've been denied for like --14 they told us she -- they told her she -- he had 15 to have 50 percent to qualify. 16 DR. ZIEMER: I suspect we may have to have one 17 of the individuals look at the dates and so on 18 to confirm that the -- there are some criteria 19 in terms of dates and numbers of working days, 20 but --21 MS. WILBURN-YOAKUM: The dates coincided with 22 the dates. 23 DR. ZIEMER: -- that -- I think we can have 24 that done and should be done privately, not in 25 open session, since there are privacy issues.

1	But please be aware that the action already
2	taken earlier this week was to recommend
3	Special Cohort status
4	MS. WILBURN-YOAKUM: I read that.
5	DR. ZIEMER: for the Iowa group, so that may
6	indeed change your situation.
7	MS. WILBURN-YOAKUM: Okay. I thank you for
8	your time. I know the Board has a lot to do.
9	DR. ZIEMER: Thank you very much.
10	MS. WILBURN-YOAKUM: Appreciate it.
11	MR. RUBY: (Off microphone) Can I ask one more
12	question?
13	DR. ZIEMER: Yes, please.
14	MR. RUBY: This Special Cohort did pass. Okay,
15	the articles in the paper my question is
16	related to this. It says that it's going to
17	Mike Leavitt next and they're actually the
18	article says they are in 60 days they may be
19	sending checks.
20	DR. ZIEMER: Well
21	MR. RUBY: Because in 2000 it already went to
22	Congress. It still does have to go to Congress
23	after Mike Leavitt?
24	DR. ZIEMER: Keep in mind that this Board is
25	advisory. We we have we are advising the

1 Secretary of Health and Human Services that 2 this class be added to the Special Exposure 3 Cohort. 4 MR. RUBY: Okay. 5 That -- the Secretary of Health and Human Services has yet to take our advice 6 7 and do something with it. MR. RUBY: Right. Now, see, the paper says in 8 9 2000, though, this -- that it wouldn't have to 10 go to Congress this time because in 2000 they 11 passed something it refers to in the newspaper. 12 Are they incorrect? DR. ZIEMER: Well, the initial legislation of 13 14 course is in place, but Congress has to approve 15 addition of classes. Is that right, Jim --DR. MELIUS: Well, I think Congress -- be --16 17 correct, Congress has a chance to turn down --18 DR. ZIEMER: Yes, if they don't act on it 19 within 30 days to turn it down, it's --20 automatically becomes part of the class, so in 21 that sense --22 MR. RUBY: That's all they need. 23 DR. ZIEMER: -- yes, they -- they can --24 Congress can turn it down. If they do not, 25 then it --

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MR. RUBY: Okay. Thank you.

DR. MELIUS: Essentially the time frame would be 21 days our letter will get up to the Secretary. Secretary has 30 days then to make a recommendation. If the Secretary agrees with our recommendation, then Congress has 30 days to act. If Congress doesn't act to stop it, then DOL would be able to process the...

DR. ZIEMER: We're -- we're not in a position to say the check is in the mail, but -- other -- other members of the public who wish to address the assembly? Thank you very much. If not, I just want to double-check. We may have some odds and ends here. Lew, help me out. What have we not yet covered?

## SC&A, INC. CONTRACT UPDATE STATUS

DR. WADE: The only thing that we've not covered at this point, and I don't know that we would do it other than to make mention of it, is that -- you know, the SEC (sic) contract was originally awarded for \$3 million for five years. That was two years ago. We're coming to the expenditure of that \$3 million for the work that's currently on the books. This Board will need to make a decision as to what work it

1 wants SC&A to do next year and will need to 2 build that into an estimate of cost, and then 3 I'll need to proceed to try and secure that 4 money. So at the July meeting we'll need to 5 have a discussion of what you would like to see 6 your contractor do next calendar year. 7 DR. ZIEMER: And that would involve identifying 8 perhaps numbers of dose reconstruction cases, 9 numbers of site profile reviews and any 10 assistance with petitions that the Board may 11 wish to identify. 12 DR. WADE: Correct. Is that correct? 13 DR. ZIEMER: We --14 MR. GRIFFON: Do we have to do a cost estimate 15 at that meeting? 16 DR. WADE: No. We have -- we have cost 17 figures. Now just for you to start to think 18 about -- it costs approximately \$200,000 for 19 your contractor to review a site profile and 20 approximately \$350,000 to review 20 dose 21 reconstructions. So with those kinds of 22 multipliers, you can begin to estimate what you 23 would like to see done. I have no figure to 24 offer you in terms of the SEC task. 25 Now it's well possible that these numbers will

be high estimates because we are developing -SC&A is developing more efficient procedures,
but I only have the numbers available that -that are real at this point. Again, to
complete the record on this, it -- those
numbers are about twice what was originally
estimated when you looked at this contract.
But I must quickly point out that the Board has
really more than doubled the work of the
contractor through its six-step process. So I
think all of this is in order.

I think you need to think about what you would like to see done and then I have to try and secure the funding for that. That is not a given. I'm very supportive of the process and will work very hard to secure that funding.

MR. PRESLEY: Lew --

DR. ZIEMER: Robert?

MR. PRESLEY: -- Robert Presley. Before our next meeting can you come up with a list of the task that are on the board -- I mean that would be on the books, the possible tasks for us so we will all be playing on a -- on a same sheet of music, and also the ones where you have an estimate for cost, could you please put that in

1 -- estimate for -- per job so that we can look 2 at that before we go to our meetings so we've 3 got some idea of what to talk to before we get 4 there, please? 5 DR. WADE: I understand. What I'll try to do 6 is -- I think it's appropriate that I would 7 write to the Board and provide you with this 8 information before the next meeting. 9 information would be public at the next 10 meeting, obviously. 11 DR. ZIEMER: Okay. Wanda? 12 MS. MUNN: That was the question I was going to 13 ask, this will be a public meeting? 14 DR. WADE: Yes. I think when we discuss 15 numbers at this level, without getting into the 16 details of the labor rates of the contractor, I 17 think we can have these deliberations in 18 public. 19 MR. PRESLEY: We can have them. 20 DR. ZIEMER: Yeah, I think -- and -- and Lew is 21 sensitive to -- to the level at which you can 22 discuss the numbers. We can't discuss 23 individual hourly rates and those kinds of 24 things, but we can discuss costs of total 25 contracts.

1	Yes, Arjun, did you have a
2	DR. MAKHIJANI: Yes, I had a question
3	DR. ZIEMER: comment or question?
4	DR. MAKHIJANI: For Dr. Wade or or you, Dr.
5	Ziemer, I presume in view of the motion that
6	was passed in regard to the SEC task order that
7	you would be expecting a response from SCA in
8	the form of a proposal and cost estimates by a
9	certain date.
10	DR. ZIEMER: Lew is going to touch base with
11	John on that
12	DR. WADE: I'll have to go to the contracting
13	officer and then we'll approach SC&A.
14	DR. MAKHIJANI: Okay, fine. Thank you.
15	DR. ZIEMER: You don't have to do anything at
16	this point.
17	DR. MAKHIJANI: Right, I just I just
18	since I'm not personally familiar with it, I
19	just wanted to be clear about the process.
20	Thank you.
21	DR. WADE: I think that's all, Paul.
22	DR. ZIEMER: Let me ask if there are any other
23	items to come before the Board today?
24	(No responses)
25	Thank you very much. It seemed a little bit

1 like a marathon --2 MR. PRESLEY: Can I say -- can I say one thing 3 before --4 DR. ZIEMER: You bet. 5 MR. PRESLEY: Now that it's all over and deliberations are all over with, Mark and I 6 7 went to Germantown last week and met with Sanford Cohen & Associates, went through a lot 8 9 of -- tremendous amount of paperwork up there. 10 I want to thank Sanford & Cohen (sic) for going 11 up there with us, allowing us to look with 12 them, and also thanking -- Larry Elliott's 13 people did a fabulous job on getting that stuff 14 ready for us to look at in a timely manner. 15 They did a very, very good job. I want it on 16 record that they did. 17 DR. ZIEMER: Thank you very much. Mark, you 18 want to add to that? 19 MR. GRIFFON: Yeah, and I would just add I was 20 impressed that you got the clearances as 21 quickly as you did, so that trip worked out 22 pretty nicely. 23 DR. WADE: Yeah, I mean when we last met by 24 phone, or even the last time the subcommittee 25 met, I wasn't sure it would all come together,

1	but it did. And there are many, many people to
2	be thanked for that, but I think good process
3	was followed.
4	DR. ZIEMER: Thank you very much. We are
5	adjourned.
6	(Whereupon, the meeting adjourned at 4:35 p.m.)
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## CERTIFICATE OF COURT REPORTER

## STATE OF GEORGIA COUNTY OF FULTON

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I, Steven Ray Green, Certified Merit Court Reporter, do hereby certify that I reported the above and foregoing on the day of April 27, 2005; 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 27th day of May, 2005.

STEVEN RAY GREEN, CCR

CERTIFIED MERIT COURT REPORTER

CERTIFICATE NUMBER: A-2102