### U.S. DEPARTMENT OF LABOR

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# ADVISORY BOARD ON TOXIC SUBSTANCES AND WORKER HEALTH

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MEETING

THURSDAY
APRIL 25, 2019

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The Board convened in the Lamar Ballroom at the Augusta Marriott at the Convention Center located at 2 Tenth Street, Augusta, Georgia, at 8:30 a.m. Eastern Daylight Time, Steven Markowitz, Chair, presiding.

#### PRESENT:

SCIENTIFIC COMMUNITY

JOHN DEMENT
GEORGE FRIEDMAN-JIMENEZ
MAREK MIKULSKI
KENNETH SILVER

MEDICAL COMMUNITY

MANIJEH BERENJI STEVEN MARKOWITZ, Chair CARRIE REDLICH CLAIMANT COMMUNITY

KIRK DOMINA RON MAHS DURONDA POPE CALIN TEBAY

DESIGNATED FEDERAL OFFICIAL

DOUG FITZGERALD

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## 1 P-R-O-C-E-E-D-I-N-G-S 8:42 a.m. 2 MR. FITZGERALD: Good morning, 3 4 everyone. My name is Doug Fitzgerald, and I'd like to welcome you to the second day of this meeting 5 6 of the Advisory Board on Toxic Substances and Worker 7 Health. I'm the Board's designated federal officer, or DFO. 8 9 Before we convene, just wanted to go 10 over some general housekeeping and remind people that should there be an emergency, the exits are 11 to the back of the room. Please exit and find your 12 way outside if there is an emergency. 13 14 Also, there are restrooms located out the back and to the left down the hall, and there 15 16 are water fountains there by the restrooms as well. Our agenda today will take us to around 17 the noon hour. We'll do our best to try to adhere 18 19 to that schedule. There will be no public comment 20 period today. And with that, Mr. Chairman, I will turn 21 it over to you. 22

1 CHAIR MARKOWITZ: Thank you. Unless someone requested, I think we might skip the 2 introductions today, because everybody in the room 3 4 here at least was present yesterday. So unless there's an objection? Okay. 5 6 So Ι thought we'd continue the 7 discussion about the claims review from yesterday. If there were any additional claims that people 8 Silver? 9 wanted to discuss? Dr. 10 MEMBER SILVER: An interesting Parkinson's claim was for a Y-12 machinist who hired 11 on in the 1950s and spent 45 years mostly as a 12 13 He had a couple of managerial positions machinist. towards the end of his career in 1995. 14 His primary care doctor diagnosed him 15 16 with Parkinson's at age 82. The family had already interacted with the EEOICPA program for squamous 17 cell carcinoma. Under Part B he did not get 18

compensation because the probability of causation

remember correctly, people keep coming back as they

get additional skin cancers, and the IREP model

was never more than about 15 percent.

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But if I

sometimes comes out in their favor, but it did not in his case.

His primary care doctor made the diagnosis of Parkinson's and said, I feel it's due to job exposures, citing welding fumes, carbon steel, and stainless steel. The primary care doctor didn't say that the worker was a welder, so it's ambiguous as to whether he had bystander exposure or direct exposure to welding fumes.

I don't know if this helped or hurt, but the primary care doctor also said that the worker's spouse could really use some help with in home care in that diagnostic letter.

So there were two industrial hygiene reviews, one out of Jacksonville. A deficiency there is that they ignored the two types of steel that are in the site exposure matrix.

The DC industrial hygiene group did a better review and pointed out that there was a direct disease link work process for Y-12 stainless steel, carbon steel, and Parkinson's.

A deficiency of the DC review was that they didn't

seem too curious about the percent manganese in the metals that he was working with, and that baffles me.

When an agent has been associated with a specific outcome, why wouldn't they put it any brain power into estimating the percent manganese in the ore or the metal over that period of time?

When it came to levels of exposure, they characterized them as low, and when the CMC looked over the file in the IH report, the CMC concluded that none of the exposures could have resulted in this gentleman's Parkinson's disease.

The CMC report I referred to yesterday has appeared to have a lot of cut and pasted boilerplate; in particular, a sentence about Parkinson's under 50 being associated with genetic factors. Well, this man was 82 when he was diagnosed, so that's kind of irrelevant, but it was in there.

I learned a lot about Parkinson's and secondary Parkinsonism yesterday, but a chapter by Robert Feldman, the late, great neurologist from

Boston University who trained lot of occupational, environmental, and neurotoxic specialists, asserted that PET, positron emission tomography, was helpful for the differential diagnosis of idiopathic Parkinson's disease versus toxic Parkinson's disease. Berenji said there was I know Dr. mixed evidence on that, but I thought that was a missing piece of information from his file. So the claim was denied -- oh, one other relevant fact is that the primary care doctor mentioned that L-DOPA was slowing the progression of disease, and although there's a lot of nuance to this, kind of heuristic, is that if L-DOPA works,

So despite my problems with the exposure assessment in the CMC report, I came down on agreeing with the determination, as the onset was 82 years old, L-DOPA was working, and in my gut, I felt it was probably idiopathic Parkinson's disease.

it may not be a toxic agent that's causing it.

CHAIR MARKOWITZ: Ouestions or

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#### 1 | comments?

MEMBER SILVER: And I'm not heartless.

I also want you to know that he also developed

D-cell lymphoma and they got Part B compensation

and survivor's compensation.

CHAIR MARKOWITZ: Dr.

Friedman-Jimenez?

MEMBER FRIEDMAN-JIMENEZ: A couple of comments: first, rather than speculate about the percent of manganese in the various steels, which does vary quite a bit, is there any way to get actual information data from the SEM or -- some of this may actually be classified. Is there a way to find out what the percent manganese in the various steels that were used in the different plants, or is that just futile?

MEMBER SILVER: Well, if I had been asked to work on this case, I would have gone to a couple of reference sources. I'm at a school without a lot of library resources, so we inadvertently have a great historical collection of the Kirk-Othmer Encyclopedia of Chemical

1	Technology, 2nd Edition, from the era of the Cold
2	War.
3	It often has chapters written by
4	engineers and chemists who worked in the DOE
5	complex. So that is one reference source that I
6	would have checked on typical manganese
7	concentrations.
8	I think it might be available in the
9	open literature, but in curiosity, we've seen,
10	on the part of the CMCs and the IHs tells that you
11	never take that kind of deep dive into the
12	literature.
13	MEMBER FRIEDMAN-JIMENEZ: Okay. It
14	sounds kind of difficult to determine with any
15	confidence for a particular job, not to mention
16	a whole career, but I think manganese is a pretty
17	common alloy in most of the stainless steels ever
18	used. Is that right, John?
19	MEMBER DEMENT: Yes.
20	MEMBER FRIEDMAN-JIMENEZ: Yes. So
21	the second question: I don't know that the test,
22	for example, the globus pallidus versus the

substantion diagram finding are really accurate 1 enough to distinguish with confidence idiopathic 2 Parkinson's from manganism. 3 4 Typically, manganism shows findings in the globus pallidus, but they are not very 5 6 sensitive, and they're not very specific either. 7 So I don't know that we would expect that that would be used to distinguish. 8 I think a default presumption that 9 10 Parkinson's and manganism are not clinically confidence 11 distinguishable with would be Marek, do you think that's accurate? 12 reasonable. 13 MEMBER MIKULSKI: I think that's a fair 14 comparison. I was looking at this last night actually, and this is exactly what you have been 15 16 saying. There's really -- those are not sensitive enough to use diagnostic testing in order to be 17 able to use it as definite diagnosis. 18 19 CHATR MARKOWITZ: This is Steve 20 Markowitz. I think the Procedure Manual actually recognizes that -- I'll try to find the section 21 -- but I think they aggregate the various relevant 22

1	ICD diagnoses and consider them to be subject to
2	the issue, so Mr. Nelson, do you have a comment
3	you want to make?
4	MR. NELSON: Just a point of
5	clarification. In the Procedure Manual in Appendix
6	1, Chapter 15-4.16 of the Procedure Manual says
7	tells CEs to develop claims for Parkinsonism,
8	Parkinson's disease, and any reasonable alias in
9	the same manner.
10	CHAIR MARKOWITZ: Okay. Thank you.
11	That was the section I was referring to, yes. So
12	that's a healthy approach. Dr. Redlich?
13	MEMBER REDLICH: I think that case also
14	illustrates one other aspect of Parkinson's, which
15	is the incidence does increase with age, and he
16	was 82 years old, and age is considered a risk
17	factor. It's also more common in men than women,
18	especially in the older age group.
19	MEMBER MIKULSKI: Do we know at what
20	age he was diagnosed?
21	MEMBER SILVER: Eighty-two.
22	MEMBER REDLICH: That's right.

MEMBER SILVER: Is it at all relevant that when he retired, he had no neuro symptoms at all? At age 64? If it had been the manganese, would you have expected some neurotoxicity before the later onset of full-blown Parkinson's? Or is that not a predictable case representation?

MEMBER MIKULSKI: I don't know.

MEMBER FRIEDMAN-JIMENEZ: I think that's a valid question. I don't think anyone here knows the answer, but what is the latency period for manganese-induced Parkinsonian-type symptoms? It's something that may be studied or may not have been studied.

But these kinds of facts or factors, I think, are useful in distinguishing, but even with that, with the age, with the response to L-DOPA, with the imaging findings, it's still quite difficult to distinguish -- and I don't know that it's a reasonable goal, and I think the Procedure Manual really has it as well as we can formulate it -- to consider them indistinguishable, clinically.

1 CHAIR MARKOWITZ: Steve Markowitz. Ι think, Marek, the issue of latency maybe 2 something that, when we move forward on providing 3 4 some advice to the Department, we should -- if there are data on that issue. 5 6 Any other questions or comments on this 7 Is there another claim that people want to review? Dr. Dement. 8 This is a fairly brief 9 MEMBER DEMENT: 10 This is an interesting case. This is an individual who worked at Sandia as a metallurgist, 11 a materials scientist from '78 through 2008. 12 13 He developed symptoms of Parkinson's at age about 60 and submitted a claim. 14 The claim was originally denied, or the recommended decision 15 16 was to deny it based on lack of finding exposure information on the SEM. 17 His authorized representative, who was 18 19 his spouse, asked for an appeal at the time the recommended decision was made and stated it was 20 difficult to get information on his exposure from 21

the cause of classification.

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was

case

The

1	recommended for further development.
2	So the interesting thing about this is,
3	this individual published a number of articles that
4	specifically talked about his work with these metal
5	alloys, and as they produced a whole list of
6	publications that came along with it, peer-review
7	publications that demonstrated it.
8	The claims examiner went back and made
9	the direct disease link between manganese alloys
10	and Parkinson's, and it was awarded. So that's
11	a good story.
12	CHAIR MARKOWITZ: So it is useful to
13	publish. Is that the lesson?
14	MEMBER DEMENT: Yes, sometimes people
15	read them for different reasons. I looked at the
16	publications. They were really complex
17	publications, quite detailed.
18	MEMBER MIKULSKI: What was the site
19	that he worked on?
20	MEMBER DEMENT: Sandia.
21	CHAIR MARKOWITZ: Questions or
22	comments on this case? Are there additional cases?

1	Dr. Silver?
2	MEMBER SILVER: Are we ready for some
3	more COPD?
4	CHAIR MARKOWITZ: Yes, we can move back
5	to COPD, that's fine.
6	MEMBER SILVER: The claimant was a
7	laborer
8	CHAIR MARKOWITZ: Is this an accept or
9	a denial? Just to
10	MEMBER SILVER: This was a denial.
11	CHAIR MARKOWITZ: Okay.
12	MEMBER SILVER: Employed at the Nevada
13	test site for eight years, from 1980 through 1987.
14	There were several sources of exposure
15	information. The SEM was the main one relied upon
16	by the IH and the CMC.
17	But the employer also had hazard
18	profiles for some of the work areas that were
19	included in the file, and I was distressed that
20	the DOL people did not rely very much on the
21	information provided by the employer.
22	But the main exposures considered were

silica, asbestos, certain metals, lead, diesel exhaust as well, wood dust, welding fumes, and cement.

She was not diagnosed with COPD until age 73, after having left the Nevada test site for a couple of decades. Early in the assembling of facts for her, it was referenced to six years of heavy exposure to asbestos when she started at the Nevada test site, but that dropped out of the documentation as the claim progressed, and I'm not sure why.

Missing from her files were the actual pulmonary function test data. Her primary care doctor diagnosed her with COPD, referred to the six years of heavy asbestos exposure and the other vaporous gases, dust, and fumes that I've mentioned. He didn't use that phrase, but we all know diesel, cement, silica dioxide fall into that category.

She reported smoking only one to two cigarettes per day, and no one ever questioned that in the documentation that I saw. But when it

reached the CMC, the CMC saw fit to include a paragraph about how 80 to 90 percent of COPD cases are due to smoking, and it seemed like cut and pasted boilerplate with no reference to her actual smoking habits over the years.

She also developed Lewy body dementia while the claim was being processed and relied on her authorized representative, a family member, to advocate for her.

So the claim was denied, and then on the question of whether I agree with that, I just looked at the Board's presumption that it's not yet been accepted by DOL and saw that she had five years of exposure to asbestos, diesel, cement, other vapors, gases, dust, and fumes. With that as my guiding light, I felt it was an unfair denial.

She had not been through the former worker program; I think that would have helped her, since I know they do some fine grains, characterization of people's work histories for the Nevada test site program.

I think DOL ignored the fact that she

1 spent time at Area 51. I'm not the conspiracy theorist I used to be, but we do know that they 2 had a lot of exotic materials being incinerated 3 4 out there. They also overlooked information about 5 6 her work with molten asphalt rubber to fill cracks 7 in the road. That information was provided by her employer, and other information provided by her 8 employer said she spent time around a tank farm, 9 10 which would have resulted in gas and vapor exposure. So that's my take on it. Any insights 11 on Claim Number 5427? 12 So I reviewed that 13 CHAIR MARKOWITZ: claim also, and my take on this is that one doesn't 14 need to resort to the approach of vapors, gas, dust, 15 16 and fumes, but actually for a laborer in the SEM, the Nevada test site, if you look at how labor and 17 COPD overlap, what the exposures are, which 18 19 includes cement, diesel exhaust, and the like, she 20 would appear to have had those exposures. In fact, my puzzle on this case -- the

CMC just followed what the IH said, so I ultimately

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said there wasn't significant exposure, and the CMC followed.

But let me just read from the IH report briefly. Conclusion, quote, in the absence of compelling data to the contrary, it is highly likely that the Claimant, in her capacity as a laborer at the Nevada test site, was significantly exposed to cement, diesel exhaust, lead, mercury, crystalline silicon dioxide, welding fumes, and wood dust.

Any exposure to these agents that she might have received would have been incidental in nature, parentheses, occurring in passing only, end parentheses, and not significant.

So there's a direct contradiction within that statement, and I don't know that that was ever corrected or not, but it was puzzling. The other aspect is the references that the IH provides are the usual references; meaning, the site exposure matrix, facility database, and then a number of textbooks, which clearly don't have the detailed kind of information that would allow a person to

1 make these conclusions. So clearly, they were relying on their 2 expertise understanding of the site and industrial 3 4 hygiene. So that's where I found this case to be puzzling. 5 6 CHAIR MARKOWITZ: Comments? 7 Questions? MEMBER POPE: I also see that -- I 8 reviewed that case as well, and I concur with Ken, 9 10 and as well with you, Dr. Markowitz. 11 seemed like there was a trend or commonality with the IH starting out saying there is a connection, 12 13 and then having that contradicting language It says there is a connection, and it 14 following. says no, I think it's environmental related. 15 16 CHAIR MARKOWITZ: Yes, Mr. Domina. I quess a couple of 17 MEMBER DOMINA: questions. Some of these claims -- I apologize, 18 19 I haven't had a lot of time to review, but some 20 of them are still open, and we have questions about -- there's got to be a mechanism for us to talk 21

to DOL about them, to get more information or see

where some it is going, in my opinion.

Then the other part where we always see below regulatory limits; who is that? Who is the regulatory limits that they're referring to? What agency is that, and are they all citing the same agency?

And then an observation from me: You look at all of these sites that have SECs for the Part B for radiation; if you don't have any radiation data, you don't have any data for anything else either, because that always came first.

We need to come up with a mechanism -just me saying out loud -- if there's an SEC, they
need to look at stuff different for the chemical
exposure or any other exposures, because you know
there is no monitoring, because a bunch of us lived
it.

And to put the onus on the worker to come up with stuff when the claim is not properly adjudicated, just like the one John just talked about -- well, the guy's a metallurgist. Then you're taking somebody who is obviously at a lot

1	lower level, trying to tell them they weren't
2	exposed. I mean, that's criminal.
3	CHAIR MARKOWITZ: Any other comments
4	or questions?
5	MEMBER MAHS: Yes, I just had a look,
6	and I think that four out of the six that I reviewed
7	had that same contradictory statement. It says
8	substantially exposed, but to low or very low or
9	in-passing exposure.
LO	CHAIR MARKOWITZ: By the way, just to
L1	follow up on Mr. Domina's question: Mr. Vance,
L2	are you on the if you're on the phone, the
L3	question is whether any of the claims we were
L4	provided for review are still open cases. I would
L5	have thought they would be closed.
L6	MR. VANCE: Hi, good morning, Dr.
L7	Markowitz and the Board. Yes, John Vance. That
L8	was not a criteria we were looking for, so we were
L9	just looking for cases that met the requirements
20	for the pool, which was Parkinson's denied,
21	Parkinson's accept.
22	So there could very well be cases in

1 that sample size that have other ongoing issues or are currently in some sort of appeal. 2 CHAIR MARKOWITZ: Okay, Okay, thank you. 3 4 Dr. Silver? 5 MEMBER SILVER: What a great idea, 6 Kirk. If a site has an SEC, it's an admission by 7 government that radiation, kind of the main act at the site, is not being adequately monitored. 8 So it's not a great stretch to infer that chemicals 9 10 were not being well monitored in that time frame. I don't know how we get that recognized 11 administratively and legally, 12 but it's just exploding with common sense. 13 Steve Markowitz. 14 CHAIR MARKOWITZ: In any of the claims that people looked at, in the 15 16 pre-1995 period, was there any evidence that the industrial hygienist actually used monitoring data 17 from the site in order to influence their decision? 18 19 I didn't see any reference to any 20 monitoring data in the decision-making, and that's not a criticism, that's just an observation 21 22 reinforcing what you're saying; that there is

1 minimal highly relevant monitoring information for decision-making on a claim. 2 Hi, this is Mani MEMBER BERENJI: 3 4 Berenji. So to answer Dr. Markowitz' question, I actually did review a claim, COPD, that was 5 6 approved. ID is 017, Date of Birth, Case 7 (Redacted). This was an individual who worked at 8 9 multiple locations within the Rocky Flats Plant, 10 so this individual worked as a radiation monitor, machinist, tool maker, construction millwright, 11 12 as well as a supervisor. He was involved in construction and welding inspections, and he worked 13 for multiple subcontractors over multiple periods 14 of time. 15 16 So his work history, at least with DOE was fragmented. From what I was able to gather 17 from the record it appears that he worked from 1962 18 19 to 1967, then there was a two-year lag, and then he worked from 1969 to 1973, and then from 1999 20 to 2003. 21

This is an issue I had with some of the

claims that I reviewed, especially with the folks who have had fragmented work histories. So I'm not really sure what he was doing between 1974 and 1998. I wasn't able to gather if he was still engaged in some sort of other type of activity that could have introduced him to additional exposures. That wasn't very clear to me.

But at least from that I was able to review, I thought that first and foremost, there were actually the occupational medicine reports, so this individual had multiple injuries at the Rocky Flats Plant, and there was actually good occupational medicine records with respect to the injuries that he had.

With respect to actual exposure data, again, I just did a brief review of this case, so I'd probably have to go back to get some more detail, but at least from what I was able to see, there were some sampling reports done by industrial hygiene at the Rocky Flats Plant.

This is addition to SEM, as well as the fact that this individual was in the Former Worker

Medical Screening Program.

I think it was just in this particular worker's interests that he was close to National Jewish, which is a very renowned health care system, and they actually have some of the best pulmonologists in the country. So he actually did have good surveillance screening program.

They actually did screen for beryllium, and he actually had some additional screening surveillance, monitoring for metals.

So I think this individual, by the fact he was at Rocky Flats, had good access to medical care, good access to screening protocols at National Jewish, and at least the industrial hygiene report that was issued on June 19th, 2018 did incorporate both the SEM as well as some of the sampling reports.

But again, I always run into this issue as well, with respect to the exposures that were listed in the industrial hygiene report, because I do find that there is still a discrepancy.

The exposures that were listed were

cement, diesel 1 asbestos, engine exhaust, silicone dioxide, welding 2 endotoxins, fumes, ammonia. 3 But I think we've all mentioned this 4 at some point during this meeting, but the low to 5 6 moderate, low to very low exposures in terms of 7 the way they categorize the exposures, I feel like this is an underlying thing that we've come across 8 day in and day out. 9 10 Despite the fact that this individual on the DOE Former Worker's Medical Screening 11 Program actually had some sampling data -- I'd like 12 13 to get John Dement's input on that at some point -- I feel like there's not a comprehensive way to 14 incorporate all those data points. 15 I feel that 16 the industrial hygienists and DOL still resort to SEM at least 90 to 95 percent of the time. 17 CHAIR MARKOWITZ: I'm sorry, I didn't 18 19 catch that last point. They still resort to 90, 20 95 percent what? MEMBER BERENJI: The SEM. I feel 21 that's their main go-to. Again, I feel that if 22

1	you get some industrial hygiene input from some
2	of our colleagues here, but despite the fact that
3	there was actual sampling data, I feel that the
4	DOL still resorts to the SEM. I feel that that's
5	an unfair protocol.
6	CHAIR MARKOWITZ: This is, oh yeah I'm
7	sorry, Calin Tebay.
8	MEMBER TEBAY: Calin Tebay.
9	Oftentimes we only see at the HWEC when folks come
10	in with claims. The IH data that's in the file
11	is submitted by the claimant.
12	MEMBER BERENJI: That wasn't made
13	clear to me, because I wasn't there. But I'm not
14	sure if DOL actually
15	MEMBER TEBAY: That's what my question
16	was going to be.
17	MEMBER BERENJI: Yes, that I don't
18	know.
19	MEMBER TEBAY: I've personally never
20	seen and to be honest with you, a year ago, almost
21	to the day, we met with the DOL, the district office
22	in Seattle with DOE and the HWEC. And this the

1 topic we discussed for two hours, this boilerplate language of not significant or significant but not 2 above OELs and PELs, and where they're getting their 3 4 information for this boilerplate language. So this has been going on for quite some 5 6 time, and the only time we've seen IH data that 7 could potentially back up those kind of statements is when the claimant themselves had mined some kind 8 of IH data from their own site and then submitted 9 10 it themselves. MEMBER BERENJI: Yes, I'm not sure 11 about this particular case. 12 It wasn't made exactly 13 clear to me whether this was submitted by the 14 claimant or DOL. CHAIR MARKOWITZ: You know -- Steve 15 16 Markowitz -- the other aspect of this is from the medical end. If you receive an IH report that lists 17 some exposures and then ranks them frequent or 18 19 occasional, low, very low, or frequent, I would expect there to be significant variation among the 20 CMCs on how to interpret that information. 21

I could easily see that one CMC would

1 say that a frequent low exposure to X is not significant because its low; whereas, another one 2 would say it's frequent even though it's low, and 3 4 therefore it is relevant to the person's disease. Consistency is very important, and I 5 6 don't know that that's been looked at or how you 7 develop a system that would be consistent so the claimants are treated equitably. 8 9 MEMBER MAHS: Kirk asked about any open 10 One of mine definitely is, 5227, a lady cases. They had asked for a review 11 from Savannah River. due to them not giving her the final recommendation, 12 13 just the final decision, and they didn't use the testimony of two of her co-workers. 14 15 CHAIR MARKOWITZ: A question for Mr. 16 just following Domina's Vance, uр Mr. on So if there are cases that are in some 17 suggestion. and Board members want to submit 18 sense open, 19 comments on those cases that might be useful in the review of those case, how should we handle that? 20 Should we submit them to the Department? 21

Before you answer that question, I'm a little

1 uncomfortable with, in any sense, setting up an expectation that all the claims we look at and that 2 all Board members would be obligated to do this, 3 4 because that's not our role. However, if we find issues that the 5 6 Department would want that feedback on, then there 7 should be an avenue to do that. MR. VANCE: Yes, this is John Vance. 8 Yes, I think that would have to be a conversation 9 10 between the Board and the DFO and the program, as far as how that mechanism would want to work. 11 I'm just not sure how we want to do that. 12 13 I mean, some of these cases were in some process of adjudication for a variety of things, so what 14 their status is now or what it will be in the future 15 16 is hard to tell. So I think that's a conversation between all three parties. 17 This 18 MR. FITZGERALD: is Doug 19 Fitzgerald, DFO. I wouldn't want to make a decision on the fly here without looking at this 20 a little more closely, but I think the charge of 21

the Board should be one that looks at more general

1	sort of application of the statute and the laws
2	and how the program is conducting its business,
3	rather than getting into individual cases.
4	I would hate to set up that expectation
5	that the Board is going to weigh in and actually
6	weigh in on individual cases. That's not to say
7	that if things that are found in the normal course
8	of business appear to be egregious, that the program
9	should be made aware of, I don't want to cut off
10	that opportunity either.
11	But to set up a sort of formalized
12	process where the Board weighs in on individual
13	cases I think might be problematic.
14	MEMBER SILVER: Dr.
15	Friedman-Jimenez?
16	MEMBER FRIEDMAN-JIMENEZ: So, I
17	understand your point, and I agree. I think,
18	though, it's very useful for us to communicate in
19	some formal way our opinions on various good and
20	bad things that we've discovered in these reviews.
21	So maybe we should aggregate our
22	findings in our reviews in a systematic way that

we can then transmit to the DOL that will hopefully be useful in changing some of the problematic things and reinforcing positively the good aspects of the reviews.

CHAIR MARKOWITZ: Steve Markowitz. You know, we do, this segues into the larger issue of, how do we move forward on claims review, and what do we do about it? We've been given 20 claims; we've had some time review them. I'm sure we haven't reviewed all of them. I'm sure we haven't been able to spend sufficient time on many of them to be able to weigh in properly.

Nonetheless, it's been a very useful exercise for us to understand the claims process and how the various pieces of information are used.

We do have an outstanding request to the Department for an additional 80 claims to review, including -- and this is from the December 10th request, just to remind you -- including 20 chronic beryllium disease claims, 20 sarcoidosis claims, 20 interstitial lung disease claims, and 20 asthma claims. So that's an outstanding request

already.

The second issue above and beyond that is, how do we want to move forward with getting to what I think Dr. Friedman-Jimenez was talking about, which was being able to do our Task number 4 of our charter, which is to evaluate the industrial hygienists and the physicians for objectivity, consistency, and quality of their input into the process. So if we could discuss that, how to move forward.

The floor is open for ideas. Dr. Dement?

MEMBER DEMENT: I think, as we walk on through these claims, and as we continue to go through the claims, I think there's some themes that recur in some of the claims across the board, and that's -- I really think that's the areas we ought to concentrate on.

Some of them have to do with the issues of consistency between IH assessments and CMC assessments. So I think, rather than concentrate on any particular IH or CMC or any particular claim,

1 I think it might be beneficial, as we sort of wind down, and we go through a set of claims, COPD, 2 Parkinson's, asthma, or whatever, that each one 3 4 of us sort of synopsize our observations. After you've taken a look at all of 5 6 these claims, what are the major points that you 7 have seen with regard to the positive aspects of how the process works and perhaps those that need 8 to be have some attention paid to. 9 10 And after we've had a chance to do that sort of by category, I think maybe if we reconvene 11 and sort of compare notes, if you will, and we see 12 some commonality in observations across the claims 13 that might have some areas that could be addressed. 14 I don't know how else to bring this to 15 16 a reasonable conclusion. I think the discussion that we've had in the last day or so has been 17 helpful, and I think we have seen some emerging 18 19 things, but I think we may see more as we dive more 20 deeply into the process. Silver? 21 CHAIR MARKOWITZ: Dr. What is the status of 22 MEMBER SILVER:

1	the Board's request for an outside contractor to
2	assist with claim reviews? I agree with Dr.
3	Dement's approach, but the phrase, after we've
4	reviewed all these claims, gives me pause, because
5	most of us have day jobs.
6	I love doing this stuff, but it would
7	be really helpful to have an outside contractor.
8	The first version of this board thought our
9	colleagues at the Association of Occupational and
10	Environmental Clinics could help us get this done
11	if resources were available.
12	CHAIR MARKOWITZ: Steve Markowitz.
13	The short answer is, we have no outstanding request
14	to the Department for resources to do any work.
15	In other words, there was a request of the first
16	board. That board's term has expired; this board
17	has not made that request.
18	MEMBER SILVER: I think we should move
19	to make that request again, and maybe we could add
20	some more specificity to it as we sort of move
21	forward.
22	CHAIR MARKOWITZ: So that's not a

1 formal motion, but we can discuss that and formulate a formal motion. Dr. Friedman-Jimenez? 2 MEMBER FRIEDMAN-JIMENEZ: I agree with 3 4 both Dr. Dement's proposal of synopsizing our findings and requesting additional resources. 5 6 And to make things maybe a little bit 7 more difficult, I think it's also important to look lung cancer, mesothelioma, some of 8 at cancers: the leukemias, bladder cancer, cancers that are 9 10 likely to be caused by chemical carcinogens. These are a different framework for the causal 11 inference, and I think it's important for us to 12 look at the cancer cases as well. I don't know 13 14 how many there are, but I wouldn't want to completely ignore the cancers. 15 16 CHAIR MARKOWITZ: Well, it is true --Steve Markowitz -- it is true that we only looked 17 at claims -- this board -- for two conditions: 18 19 COPD and Parkinson's disease, and it may well be 20 that the approach that the industrial hygienists and physicians take is somewhat different by 21

different condition because of availability of

information, their own working assumptions, or whatever.

So I would agree that, not just cancer, but we need to figure out, not necessarily today, but we need to figure out the portfolio of different types of claims that we think should be looked at so we end up with a credible set of conclusions, recommendations, that's based on a broad look at the program. Dr. Redlich?

MEMBER REDLICH: I think in that regard having a better sense of which are the most common claims, under which categories, and also what the trends have been, because I gather that there may be more of X disease and less of Y in terms of where to focus efforts.

CHAIR MARKOWITZ: Well --- Steve Markowitz -- in our December 10th data request we asked for that information for lung diseases, for the most common Part E conditions in general, for neurologic conditions, for cancers, and for kidney disorders. So that information has been requested.

1 MEMBER REDLICH: Yes, it would be very 2 helpful to get that data to best focus efforts. MEMBER BERENJI: This is Mani Berenji. 3 4 I do agree with that. I think we should be able 5 to compile all the statistics and then at least 6 try to have some spreadsheet where we can actually 7 kind of tease out these particular cases, approved and denied. 8 And then there are obviously some other 9 extenuating circumstances with some of 10 claims, but I feel like we should have a systematic 11 approach, and we should hopefully be able to at 12 13 least get some quality data. So at least from my experience on this 14 board so far, I feel like we've never actually 15 16 gotten an actual handout or spreadsheet just looking at how many claims they process per year, 17 what percentage are denied, what percentage are 18 19 approved. I feel like, at least for me, it's been 20 a struggle. CHAIR MARKOWITZ: Steve Markowitz. 21 We 22 did get some data on Parkinson's disorders, but

no data on the rest of the conditions, at least this board.

But think the use of the word Ι systematic is key, because so far what we have are for two conditions. claims We have, relatively short time for review, our impressions, our initial impressions about what these claims show, and we're not going to draw any conclusions from those initial impressions because it wouldn't be appropriate. What we need is a more systematic approach to examine the appropriate set of claims.

What we've done so far has been very useful because it does allow us to send out some preliminary categories of concern; issues that we would raise with industrial hygiene evaluation with the medical evaluation and the like, so it helps us design that kind of systematic evaluation.

But that's what's needed in order to understand the issues, because we have a taste of it, but we don't have a full understanding, and we couldn't credibly represent to the Department that we had any particular recommendations or, I

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	should say, not any particular recommendations,
2	but a set of recommendations.
3	That wasn't meant to be a summary
4	statement. Dr. Friedman-Jimenez?
5	MEMBER FRIEDMAN-JIMENEZ: Two other
6	potential issues that I want to ask whether we
7	should raise them. One is injuries like
8	chemical-induced injuries, chemical burns,
9	dermatoses; the second is impairment, disability
LO	and impairment. In other words, time lost from
L1	work. Do we want to get involved in those two
L2	issues?
L3	Because also those are involved in
L4	occupational medicine decision-making and would
_5	be related to chemical toxic substance exposure.
_6	So I'm raising it as a question; I'm not advocating
_7	for doing that.
L8	CHAIR MARKOWITZ: Steve Markowitz.
L9	Well, the first question is, is it within our
20	domain? Is it within our charter to address those
21	issues?
22	And I would say to the extent that Task

1 4 is looking at the objectivity, consistency, and 2 quality of the industrial hygiene and the medical input into the program, that impairment 3 and 4 chemical-induced injuries, which are just another outcome, would fall within what we're -- the advice 5 6 that DOL has asked us to produce. I'm not asking 7 for a bigger set of issues, by the way, but I don't see how they wouldn't be conceived as being within 8 the domain. Dr. Dement? 9 10 MEMBER DEMENT: Sort of by definition 11 of what our charter is on that part to look at issues 12 of claim in of objectivity, across terms 13 consistency; there's no other way to get at that without looking at claims in rather great detail. 14 definition, 15 that, by is And timeand 16 effort-intensive process. So I think we're all interested in 17 spending time on these claims, but each one of us 18 19 has limitations; we have other jobs to do. so I think we do need some other hands to take a 20

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I don't know quite what that looks like

look at this and help us with the process.

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1	yet; where do we find that expertise, but I think
2	it's appropriate that we request some assistance
3	to get to the point that they've asked us to get
4	to.
5	CHAIR MARKOWITZ: And that statement
6	was in reference to claims review in general,
7	perhaps including impairment, including chemical
8	injuries and the like; is that right?
9	MEMBER DEMENT: Yes, broadly speaking.
10	CHAIR MARKOWITZ: Well, it seems the
11	sense is, and correct me if I'm wrong Dr. Redlich,
12	you wanted to say something?
13	MEMBER REDLICH: I mean, I think
14	there's a Workers Comp system for acute events like
15	a person is actively working. I don't think that
16	it's part of our task. I think if there were a
17	chronic, long-term sequellae of that acute exposure
18	event that then resulted in a chronic condition,
19	that that would be.
20	CHAIR MARKOWITZ: Steve Markowitz.
21	It's hard to believe that we could rely on the
22	excellent review by state workers' comp system of

chemical-induced injuries. It's possible, but it's not a default conclusion. And to me, it's just another outcome that could be within the ballpark of what's looked at. This is MR. FITZGERALD: Doug Fitzgerald, DFO. I've been listening to your back-and-forth on this, and there's some unknowns right now in terms of whether or not there will more resources, or if we have the capability of doing that. I think you could probably make all sorts of connections and linkages between what

I think you could probably make all sorts of connections and linkages between what you're asking, as others have said, to look at. But is the link really a strong one? If you have limited resources, and you all have limited time and other jobs and that sort of thing, what is the work that this board should be focusing on and prioritizing? Is that going to spread us too thin in the absence of other resources?

CHAIR MARKOWITZ: So I think the sense of what I've heard so far is that the Board would request additional resources in order to conduct

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an appropriate systematic evaluation of an appropriate number and variety of claims in order to weigh in our task of contributing to an assessment of the objectivity, consistency, and quality of the industrial hygiene and medical input into the program. Is that an appropriate summary?

MEMBER DEMENT: It's an excellent summary of where we are.

CHAIR MARKOWITZ: I think we should formulate that or some version of that as a recommendation. I'm looking to Kevin because I'm hoping we can get it on the board, hoping he's remembered exactly what I said.

(Laughter.)

MR. FITZGERALD: This is Doug Fitzgerald again, DFO. I just want to raise the issue that what the Board may not be familiar with is our procurement process. Even if we, as an agency, agreed with the request and thought it was a valid one, the procurement process within the federal government is lengthy, and it requires a lot of -- it will take time.

So like I said, in the absence of any additional resources coming in, this Board still has to pursue its mission. So I'm just cautioning you in trying to manage expectations that even if there was an agreement immediately that we should provide the Board more resources that you're asking, it would be some time before those resources were available.

Thank you for that CHAIR MARKOWITZ: advice. To me, what that translates into is that we move in parallel on this issue, meaning that we make our recommendation about a request for resources to do that systematic evaluation, even as we continue to review claims, aggregate our concerns as I think Dr. Friedman-Jimenez mentioned, Dr. Dement said, from the claims we have now, perhaps from a limited number of additional claims for different -- not necessarily 80, but a limited number of additional claims for different outcomes that we're able to design that systematic evaluation with greater specificity within a reasonable period of time.

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1 But a recommendation that asks for those 2 resources, at least puts it on the table that frankly, to weigh in on the issues that we need 3 4 to weigh in on Task 4, we can do so, but it's limited unless we have additional resources. 5 But to move 6 in parallel, because we know it will take time to 7 get those resources. Ms. Pope? MEMBER POPE: Yes. I think it's very 8 9 important for us to -- it's great for us to 10 acknowledge and identify the different concerns have about these claims 11 that we that we're reviewing, but is there a way that we can, and 12 13 especially for the benefit of the new members on the Board, to find out the recommendations that 14 we did submit that have been approved and accepted 15 16 by DOL? CHAIR MARKOWITZ: What the status is? 17 MEMBER POPE: Yes. 18 19 MEMBER BERENJI: This is Mani Berenji. 20 So yes, I think that would be great to have some sort of dashboard in terms of questions that we've 21

brought up, the DOL's responses, what percentage

2 outstanding. I think that way we have some sense of what's happening, and we can hold ourselves 3 4 accountable, but we can also hold DOL accountable 5 for what's being done and what's still outstanding. 6 So honestly, I'm happy to put that 7 together; that's not hard. We'll just have to get a spreadsheet and create a dashboard. 8 9 CHAIR MARKOWITZ: That would be great, 10 and I think we'll make that as an action item. 11 In fact, Ms. Leiton yesterday volunteered to bring us up to date on the status of the interview of 12 13 the claimant by the industrial hygienist, which they agreed to, as long as the claims examiner was 14 involved. 15 16 She didn't quite know the status, but volunteered to -- so that applied to the other 17 recommendations, we will ask them for, and I'm sure 18 19 they will accept Dr. Berenji's assistance in 20 organizing that. So we will do that, thank you. This is Kirk Domina. MEMBER DOMINA: 21 You know, yesterday when Ms. Leiton was talking, 22

have been implemented, what percentage are still

they just came out with Procedure Manual 3.0, and she said it's soon going to be 3.1. I'm just curious if something they're doing there affects anything that we could be doing, so that we're not doing something that we've got to redo.

MEMBER BERENJI: That's where the dashboard would come in. So like I said, at least we can have some sort of working document we can add on to it over time, but at least we have some way to kind of keep ourselves accountable.

Because at least from my perspective,

I find this kind difficult to be able to track,

so it would be good to have some systematic way

of tracking what we're doing, what they've been

able to accomplish, what's still outstanding, and

then really trying to advocate for more concrete

data.

I know we keep asking for specific numbers, but I would love to be able to see some bar graphs starting from the date of implementation of Part E from, I believe it would be, what, 2004, 2005 to the present.

I'm not sure why it's difficult to get that concrete data just in terms of how many claims come in on a yearly basis, what percentage are approved, what percentage are denied. I mean, at least we'll have a better sense of what's happening, because at this point, I still don't know.

I'm not sure if you folks have additional data that I don't have access to; I still don't know.

CHAIR MARKOWITZ: To follow up on your comment, Mr. Domina, we've raised this issue before about Board input into the policy-making process, and I think I recall that the Department's position is that it is not the Board's role to review policy changes that are under consideration, except for the initial Board's weigh-in on the that official rule. But it's not our role to weigh in on changes as they are under consideration.

And that's understandable. I must say, it gets awkward sometimes, because the new Procedure Manual 3.0 removed the section about asbestos exposure from 1986 to 1985, which is a

topic that the Board, in general, has spent a lot of time on, asbestos consumption.

That section was removed unbeknownst

to us, and then we discussed it initially yesterday:
what does that mean? And there will be some
back-and-forth between the Board and the Department
about that. Maybe the next version will have it
restored, and maybe not.

So it's awkward, because that's a topic that we spent a fair amount of time on, and a section is gone that's obviously relevant. Now, after the fact, we're going to provide our opinion about that, but it's an awkward process, let me just say that.

So I think Kevin's put the agenda back on, thinking that we should probably move on. But I would like to go back to that recommendation.

So the recommendation is that the Board requests resources in order to conduct a systematic evaluation of an appropriate number and variety of claims in order to assess the objectivity, quality, and consistency of the industrial hygiene and medical evaluations that are part of the claims

1	process, parentheses, Task No. 4.
2	Okay. So how should that be modified,
3	now that we see something in writing? To conduct
4	a timely systematic evaluation? Okay. So take
5	out the, and put in a timely.
6	MEMBER BERENJI: I think we need to
7	specify what exactly we mean by resources. Are
8	we talking about manpower? Are we talking about
9	technological? I think we need to specify that.
10	CHAIR MARKOWITZ: Okay, so let's
11	suggest some words.
12	MEMBER BERENJI: I would probably put
13	in parentheses, personnel. I mean, do we have a
14	specific number of folks I mean, we could
15	probably put a range.
16	CHAIR MARKOWITZ: Well, we clearly have
17	to flesh out some details, but I don't think we're
18	capable of doing that right now. I think we should
19	flesh it out over the next four to six weeks so
20	that they have something real to go on.
21	But I don't think we need to do that today.
22	MEMBER BERENJI: Okay.

1 CHAIR MARKOWITZ: But the categories, 2 I agree with. So personnel and IT MEMBER BERENJI: 3 4 support; I would probably at least put that in And then in terms of timely, I think we 5 there. 6 need to specify that too. Within what, a six-month 7 time frame, a one-year time frame? CHAIR MARKOWITZ: Well, there are two 8 time frames: one is receiving resources, and the 9 10 other time frame is actually conducting evaluations. 11 Of course receiving resources, we want 12 13 to make that as short a time period as possible, and finishing the evaluations, we want to be 14 realistic. 15 16 MEMBER BERENJI: I feel like we really need to be specific with these folks, because I 17 feel like a lot of these recommendations are very 18 19 vague, and these folks need to be told, like, we want this, this and this, and we need to be very 20 specific. At least that's been my experience so 21

I'm not sure if you folks agree, but --

far.

1	CHAIR MARKOWITZ: I agree with the need
2	for specificity; I don't think the recommendations
3	have been all that vague, but that's another issue.
4	Dr. Silver?
5	MEMBER SILVER: I'm concerned that the
6	phrase, personnel and IT support could be
7	misinterpreted to mean that DOL would reassign
8	their personnel on an in-kind basis to assist us.
9	I think what we really want is what the NIOSH
10	radiation board has, which is an external
11	contractor.
12	CHAIR MARKOWITZ: Now it says, in order
13	to conduct that could be interpreted that we
14	set out the general framework of that evaluation,
15	but then don't necessarily oversee that evaluation.
16	So the question is, do we need to be, in this
17	request, more specific than simply to say, to
18	conduct? For instance, we could say, to design.
19	MEMBER SILVER: Design and direct?
20	CHATE MADIANTEE
20	CHAIR MARKOWITZ: Dr.
21	Friedman-Jimenez?

1	say design, I would I like the word conduct.
2	I mean, there are several epidemiologists,
3	industrial hygienists on this Board that can help
4	with the design. That's not where the
5	labor-intensive part of it is.
6	But actually doing the record
7	organizing, selecting the records; we could design
8	what kind of sampling we want, what diagnoses we
9	want, but the actual work involved is beyond our
10	capacity.
11	CHAIR MARKOWITZ: I see. So resources
12	to support the Board to is that what you're
13	getting at? So after, the resources would be to
14	support the Board in order to conduct?
15	MEMBER BERENJI: I think this is
16	getting too wordy already. I mean, I feel like
17	this needs to be pretty concise and succinct.
18	CHAIR MARKOWITZ: So this is what we're
19	looking at. Either, does it need any additional
20	wording or, for that matter, should any wording
21	be deleted to reflect what we're after?
22	MEMBER MIKULSKI: Take out the second

1	in order?
2	MEMBER BERENJI: Yes, I think that's
3	a little too much wordiness.
4	CHAIR MARKOWITZ: I'm sorry; take out
5	in order?
6	MEMBER BERENJI: In order
7	MEMBER MIKULSKI: The second one. And
8	then the variety of claims to assess the
9	objectivity, quality, consistency
10	CHAIR MARKOWITZ: Yes, that's fine.
11	I don't know whether, Doug, as the designated
12	federal official, whether you see any areas of that
13	request that are so vague that it wouldn't transmit
14	the intended request?
15	Obviously, there are going to be details
16	about numbers of claims, types of claims, and all
17	that. That, we will provide. But at this level,
18	is there anything additional in specificity
19	MR. FITZGERALD: If you're trying to
20	kind of create a placeholder for a more refined
21	request later, I don't know that you need to be
22	more specific than this. But you were going to

1	have more discussions, I think, about the wording
2	of this, so
3	MEMBER REDLICH: Did we want to specify
4	an external contractor?
5	CHAIR MARKOWITZ: Questions?
6	Comments?
7	MEMBER SILVER: If this is just a
8	placeholder, I don't think we're going to bring
9	on the wrath of government procurement specialists
LO	if we mention the would-be contractor by name.
L1	I mean, we're going to refine this, so
L2	if this is a statement of our sentiments, then I
L3	would propose we put in the Association of
L4	Occupational and Environmental Clinics as the first
L5	board discussed.
L6	CHAIR MARKOWITZ: Mr. Mahs?
L7	MEMBER MAHS: It may be just me, but
L8	you have the support to support, but would it be
L9	better to replace the support with, assist the
20	Board, instead of two supports there?
21	CHAIR MARKOWITZ: All right. That's
22	good. Speaking about I don't think we should

1 name a particular -- at this point, I don't think we should name a particular -- it could be limiting, 2 if actually, because what that particular 3 4 organization doesn't want to do it? 5 But my concern is that it's possible 6 that a blend of internal and external resources 7 might be able to make this happen. For instance, the claims need to be 8 organized, indexed, and it's possible that there's 9 10 internal support that could do that in preparation 11 for, ultimately some -- we need some physician time and industrial hygiene time to evaluate these 12 13 claims, and we wouldn't want that from inside the Department, because of conflict of 14 interest, essentially. So it could be some blend of internal 15 16 and external. MEMBER REDLICH: That was what I was 17 asking, not knowing what resources are available 18 19 internally, it seems that that request should just 20 be as open as possible, the point being, request 21 resources.

MR. FITZGERALD:

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I don't want to speak

1	officially for the energy program, but I think,
2	in general, the availability of federal personnel
3	is going to be very limited.
4	CHAIR MARKOWITZ: Is what?
5	MR. FITZGERALD: Very limited.
6	CHAIR MARKOWITZ: Very limited. Okay.
7	So we could leave it, then, as external contractor,
8	and if some limited internal resources are
9	provided, as long as there's no conflict of interest
10	or whatever, that would be fine with us.
11	MEMBER REDLICH: It could be such as
12	
13	CHAIR MARKOWITZ: Such as an external
14	contractor? This is part of the reason why many
15	chapters in medical industrial hygiene texts have
16	single authors; otherwise, they'd never get done.
17	MEMBER BERENJI: This is Mani Berenji.
18	So I'm thinking the way we should probably break
19	this down is maybe have some bullet points, so that
20	way it's a little easier to read.
21	So, the Board requests resources (such
22	as external contractor to provide personnel and

1	IT support) to assist the Board with the following:
2	and then just literally bullet points, so at least
3	it's easier to read.
4	Then we can just list by bullet point
5	what we want. In order to get the colon in there,
6	and then we just bullet-point, just go for it.
7	We just add whatever we want. At least it will
8	be easier to read.
9	CHAIR MARKOWITZ: I wouldn't
10	underestimate the ability of the Department of
11	Labor to read these requests.
12	MEMBER BERENJI: I don't think so, at
13	least from my
14	CHAIR MARKOWITZ: I would read bullet
15	points if they were multiple parallel tasks, right?
16	So conduct this, to assess that, to provide that:
17	three equivalent parallel paths. But this is just
18	a single function, which is to allow us to conduct
19	a systematic evaluation of X, Y, and Z.
20	Why don't we do this? Why don't we go
21	with bullet points and see what they sense. But
22	while we do that

1 MEMBER REDLICH: It's a related concept, so -- I'm normally in favor or short 2 bullets and short sentences, but I think in this 3 4 case they're linked. CHAIR MARKOWITZ: Right. Because the 5 6 conduct and evaluate -- to assess is the function 7 of the evaluation, right? So it's, they're linked. So other comments, questions? This is 8 now a motion, or someone needs to make a motion 9 10 to accept this recommendation. I make a motion to MEMBER BERENJI: 11 accept this recommendation. 12 13 MEMBER REDLICH: Second. 14 CHAIR MARKOWITZ: So the floor is open. The motion is to accept the recommendation that 15 16 the Board requests resources (such as an external contractor to provide personnel and IT support) 17 to assist the Board in order to conduct a systematic 18 19 evaluation of an appropriate number and variety of claims to assess the objectivity, quality, and 20 consistency of the industrial hygiene and medical 21

evaluations that are part of the claims process.

1	I would make a friendly amendment that
2	the phrase, to provide personnel and IT support
3	and additional resources as required, just to leave
4	open that we may have forgotten something.
5	MEMBER REDLICH: Yes.
6	CHAIR MARKOWITZ: Okay. It's open for
7	comments. I'm just looking for the Board charter
8	to make sure that this language is entirely
9	consistent. So I would actually make another
10	friendly amendment. Where it says in the third
11	line, to assess, I would say, to assess and to
12	ensure. I add that because that's what the charter
13	says.
14	Okay. Are there additional comments
15	on this, because the floor is open. Otherwise,
16	we need to take a vote.
17	MEMBER BERENJI: So this is just a
18	placeholder, correct? I mean, we're going to
19	refine this over time.
20	CHAIR MARKOWITZ: Well, this is a
21	request to transmit to DOL, and yes, we need, over
22	a relatively short period of time, to start to fill

1	out exactly what that would look like.
2	So let's take a vote. How do we take
3	a vote? I can't remember.
4	MR. FITZGERALD: We'll run down the
5	list and get everyone's
6	CHAIR MARKOWITZ: Okay. Okay.
7	MR. FITZGERALD: And then just go with
8	the names as they're represented here. Dr. Dement?
9	MEMBER DEMENT: Yes.
10	MR. FITZGERALD: Dr. Dement is a yes.
11	Dr. Friedman-Jimenez?
12	MEMBER FRIEDMAN-JIMENEZ: Yes.
13	MR. FITZGERALD: Dr. Mikulski?
14	MEMBER MIKULSKI: Yes.
15	MR. FITZGERALD: Dr. Silver?
16	MEMBER SILVER: Yes.
17	MR. FITZGERALD: Dr. Berenji?
18	MEMBER BERENJI: Yes.
19	MR. FITZGERALD: Dr. Markowitz?
20	CHAIR MARKOWITZ: Yes.
21	MR. FITZGERALD: Dr. Redlich?
22	MEMBER REDLICH: Yes.

1	MR. FITZGERALD: Mr. Domina?
2	MEMBER DOMINA: Yes.
3	MR. FITZGERALD: Mr. Mahs?
4	MEMBER MAHS: Yes.
5	MR. FITZGERALD: Ms. Pope?
6	MEMBER POPE: Yes.
7	MR. FITZGERALD: Mr. Tebay?
8	MEMBER TEBAY: Yes.
9	MR. FITZGERALD: It's unanimous.
10	CHAIR MARKOWITZ: Then let's discuss
11	two things: one is, the claims we already have,
12	with additional time to review those claims, what
13	are we going to do with our observations?
14	I think there's been a suggestion that
15	we aggregate those, sort of categorize and
16	aggregate those observations, looking at
17	commonalities across claims. Not that that work
18	will necessarily lead to specific recommendations
19	to the Department, but at least it organizes our
20	thoughts and prepares us to perform a more
21	systematic evaluation.
22	So should we do that over the next period

1 of time, and then have a telephone Board meeting 2 in two to three months in order to discuss the aggregated observations about the claims we have 3 4 so far? 5 We just need a sense of the group; we 6 don't need to actually vote on that, I think. 7 MEMBER BERENJI: Yes. CHAIR MARKOWITZ: Okay. So what I 8 think I'll do is, I'll propose, after the Board 9 10 meeting, a way in which we do that so that, in terms 11 of who's reviewing what and which claims are already reviewed, et cetera, so that we come up with a common 12 13 I don't think we need to do that right output. 14 now. 15 MEMBER BERENJI: I'm sorry, I just 16 think it would be good to at least have a couple of general things we can already at least kind of 17 put into respective buckets, at least with respect 18 19 to the industrial hygiene, CMC. 20 At least we can kind of put some general categories, because I feel like those were where 21 we found the issues, so at least we can kind of

set up some sort of form at least some sort of way to organize our though process with respect to the most common issues that we came across.

CHAIR MARKOWITZ: So what we could do with reference to that is, within the next week or two, send in our preliminary issues that we've found so far so that we can begin to develop some categories which we then can use to further look at these claims. So that's a good idea; we'll do that.

There is the issue of our request for 80 additional claims. I detect a lot of enthusiasm. But that is an outstanding request, and we need to -- if we're going to modify it, modify it. But right now, that's our outstanding request to the Board.

Those are in, as I said before, chronic beryllium disease claims, sarcoidosis, interstitial lung disease, and asthma. So do we want to take a look at that again? Do we -- internally, do we have the capacity to review 80 claims?

1	PARTICIPANT: Depends on the time frame
2	
3	MEMBER REDLICH: I think it would be
4	helpful because we've already reviewed those
5	claims, and I think to see if there's been any
6	changes.
7	CHAIR MARKOWITZ: Dr.
8	Friedman-Jimenez?
9	MEMBER FRIEDMAN-JIMENEZ: Why don't we
10	write up what we have now, the ones we've already
11	reviewed, and then when we see what we have, then
12	we can decide what additional diagnoses or
13	additional information we would want to request.
14	But I think we already know a lot of
15	what we want to say, based on reviews we've already
16	done. So why don't we just write them up now and
17	then revisit this in our conference pool and design
18	what we want to request?
19	Do we want a random sample of all claims?
20	Do we want specific diagnoses? Do we want some
21	information on the frequency of each diagnosis?
22	What exactly do we want to ask for?

1	Because we're generating a lot of work for
2	the DOL, and also, we want the most useful
3	information that we can get. We've already
4	reviewed a lot of these claims; we know a lot about
5	what we're going to say in our synopses.
6	CHAIR MARKOWITZ: That's entirely
7	sensible. Here's my problem with that: if we
8	suspend our request and wait to reformulate that
9	for two to three months, there's just a time delay.
10	Obviously, it takes time to identify and prepare
11	those claims.
12	MEMBER REDLICH: So you were on a
13	different subcommittee. We have reviewed prior
14	beryllium and sarcoid claims, and I think that was
15	now two years ago. And I think what we'd like to
16	see is, we have a sense of what has been done, and
17	see of that process has changed at all over this
18	period of time.
19	So I agree potentially for other
20	categories, but there was a subcommittee that did
21	review the respiratory claims.

CHAIR MARKOWITZ: So then the question

is, do we really want 20 claims of each of those 1 2 four different categories? If we shrank the number things of claims, then miqht happen 3 more 4 expeditiously. 5 MEMBER BERENJI: We could to a random 6 sampling of 20 of the CBD, sarcoid, ILD, and asthma, so five from each cohort. 7 MEMBER REDLICH: We had also seen the 8 data on the numbers of new claims in those 9 10 categories, and they were not that huge, is my recollection. You had looked at that, 11 annually, so I think, at least for the -- I think 12 13 we need more than five. Well, there 14 CHAIR MARKOWITZ: Okay. are some numbers between five and 20. 15 16 MEMBER FRIEDMAN-JIMENEZ: We are already over-sampling diagnoses of interest like 17 beryllium, sarcoid, so I would propose that we 18 19 request a random sample of, say, 100 claims that will give us rough, small numbers on the relative 20 frequencies of different diagnoses; that we request 21

the categories you listed; and also, I would like

1 to add cancers: lung cancer, mesothelioma, and leukemias; maybe bladder cancer. 2 MEMBER REDLICH: We do have a piece of 3 4 it. We know from the prior look at the data that John -- that was focused on respiratory, so we do 5 6 have that sense of -- as far as the respiratory 7 But I think that all the other conditions, I think we do want to see what the most common ones 8 are in terms of where to focus our efforts. 9 10 So I think, in terms of the cancers and the like, I'd first like to see where the big buckets 11 are; where the --12 13 MEMBER FRIEDMAN-JIMENEZ: A random 14 sample will answer that question, the most common. It won't answer the question of how many leukemias 15 16 and the rare ones. John had 17 MEMBER REDLICH: I mean, nicely organized five respiratory diseases -- we 18 19 can probably even pull that up -- asthma, COPD, the number of claims, the percentage accepted. 20 And that did identify areas to target. 21 if we had that for other conditions, then it would

1 help focus. 2 MEMBER FRIEDMAN-JIMENEZ: So you think random sample by strata of respiratory, 3 4 neurological --5 MEMBER REDLICH: I personally No. 6 would stick with our current request of the 20 For those of us who are familiar with 7 claims. looking at the respiratory ones, we could go through 8 9 those rather quickly. 10 It would favor for other conditions that the request we've already put in, which is to get 11 the sort of basic data on what those claims are: 12 13 cancer, neurologic. And then when we see that, decide which claims outside of the respiratory 14 15 arena --16 MEMBER FRIEDMAN-JIMENEZ: Could request a data run that would just give us the 17 diagnoses of everybody, of all claims, so that we 18 19 could see the relative frequencies of them, rather than giving us all the information on each case? 20 MEMBER REDLICH: That's what 21 was

requested, basically, already.

1	MEMBER FRIEDMAN-JIMENEZ: Okay, but
2	that would be a separate request than the actual
3	medical records.
4	MEMBER REDLICH: Can I I can quickly
5	pull up John's
6	CHAIR MARKOWITZ: But the question is,
7	for outstanding requests, four pulmonary
8	conditions, 20 claims each; do we really need 20
9	denied claims each?
LO	MEMBER REDLICH: We probably don't need
L1	the 20 accepted for each.
L2	CHAIR MARKOWITZ: No, our request was
L3	only denied claims, actually, for the five
L4	pulmonary conditions. So do we need 10 denied
L5	sarcoidosis?
L6	I'm questioning the Board's ability to
L7	thoroughly evaluate a large number of claims, and
L8	I hesitate to ask the Department for products that
L9	represent considerable work if it's
20	MEMBER REDLICH: So what if we just did
21	15?
22	MEMBER FRIEDMAN-JIMENEZ: I would

propose that we request, say, 10 denied, oversample the denied, and five accepted. I think we should look at some of the accepted. I think there's some useful information there. But we don't need as many, I don't think, and we could concentrate our efforts on the denied claims but still look at accepted, smaller numbers. We probably don't need 20. CHATR MARKOWITZ: For these

conditions, given the fact that we've looked at some of these things before, is it really useful to look at accepted claims?

MEMBER BERENJI: I think it's good to look at accepted claims because you can look at what was done right, and I feel that it's good to provide -- at least inform the DOL that there are things that are working in the process. I think it's good to have that.

And then you can also look at the denied claims, and then you're able to kind of bring up the themes and the issues that were seen on a repeated basis. So I think it's good to have both.

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1 CHAIR MARKOWITZ: Dr. Dement? 2 MEMBER DEMENT: I'd like to weigh in If we're looking at consistency, there's on that. 3 4 no way to do that unless you look at accepted and 5 denied claims. There's no way to assess that. 6 I mean, we can concentrate on more denied than 7 accepted, but I think you have to look at both. CHAIR MARKOWITZ: So is it the sense 8 that 15 claims of each of those conditions, 10 9 10 denied and five accepted, is a better formula? I think it should be 11 MEMBER BERENJI: five and five; at least a total of 10. I just feel 12 13 that we don't have the manpower in the field to reveal those cases unless we create a separate 14 working group to focus on those only. And I really 15 like what Dr. Friedman-Jimenez was mentioning, 16 doing a random sample. That way we're able to 17 capture more cases and be able to identify more 18 19 diagnoses. 20 I mean, at least we could create two working groups in parallel, and Dr. Redlich could 21 kind of focus on the respiratory. I'm happy to 22

1 kind of focus on the general sampling, because I 2 do feel that will capture a lot more information if we're able to cast a wider net as opposed to 3 4 just homing in on respiratory, even though I do 5 feel that it's important. 6 I feel there's already been work done. 7 I think we really need to focus on capturing other diagnoses: cancers, other types of diseases that 8 we don't even have a clue about. 9 10 MEMBER FRIEDMAN-JIMENEZ: Well, the 11 reason I proposed the random sample was mainly to get the information on relative frequencies of 12 13 different diagnoses, especially the common ones. But that's already been requested in a different 14 form that's much less labor intensive. 15 16 So maybe we could go through that information and then make a second request on the 17 ones we feel we want to oversample and look at in 18 19 detail and actually go through claims. 20 But to go through 100 random-sample cases is a lot of work, and we'd probably be better 21 22 spending our time on five and five or 10 and five,

1	of the specific diagnoses of interest.
2	So I would withdraw the random sample
3	idea, give what Carrie's saying that we've already
4	requested, the information that would answer that
5	main question.
6	Is that true, John, that we can get that
7	information from what we've already requested, for
8	all diagnoses, including injuries? Including
9	everything? Has that already been requested?
10	MEMBER DEMENT: I thought we had
11	requested it.
12	CHAIR MARKOWITZ: Are you talking about
13	the Power Point that
14	MEMBER REDLICH: I have it here. I
15	just couldn't
16	CHAIR MARKOWITZ: I sent that Power
17	Point to Carrie. Maybe in the briefing book. I
18	think it might have Dement Power Point or something
19	like that, slides.
20	MEMBER REDLICH: I'm sending it.
21	CHAIR MARKOWITZ: In any event, let me
22	continue the conversation. Dr. Dement, you were

1	
2	MEMBER DEMENT: I sort of lost my train
3	of thought.
4	MEMBER FRIEDMAN-JIMENEZ: What has
5	been requested in terms of the overall view of the
6	database and the diagnoses? The frequency of
7	diagnoses?
8	MEMBER DEMENT: I think we concentrated
9	on the respiratory conditions in that one. We were
10	given a I think it was in an Excel file, a data
11	dump, and all I did with that was to pull it into
12	some programs I can use to summarize the data.
13	But I don't think we got everything, and that was
14	old, anyway, and we're a couple of years out.
15	So we've got two issues: I think the
16	program can provide that summary, similar to what
17	we have; or alternatively, I guess we could do
18	another overall data dump and do it ourselves maybe.
19	That's part two.
20	Now, I'm willing to take it on if that's
21	something the Board wants to do, but it's in and

of itself, a little bit labor-intensive.

1	CHAIR MARKOWITZ: Steve Markowitz.
2	I'm looking at that request from December 10th,
3	and it included the 20 most common conditions in
4	descending order for which claims had been filed
5	since 2013. Then we provided a sample table of
6	what we wanted it to look like, which reflected
7	John's work previously.
8	We then requested 10 most common
9	neurologic conditions, 10 most common cancers, and
10	the 10 most common renal or kidney disorders. So
11	that's the information that would be very useful,
12	yes.
13	MEMBER FRIEDMAN-JIMENEZ: That sounds
14	excellent, and what is the status of that? Are
15	we going to receive that any time soon, that we
16	can use it in our decision-making of what we want
17	to look at in detail?
18	CHAIR MARKOWITZ: Mr. Vance, are you
19	still on the phone?
20	MR. VANCE: Yes, I am.
21	CHAIR MARKOWITZ: Do you have any
22	I don't know if this is within your area, but do

1	you have any sense on the progress on that part
2	of the request?
3	MR. VANCE: Not at this time. I know
4	that we've got a lot of different requests floating
5	around.
6	CHAIR MARKOWITZ: Okay. So we need to
7	make our decisions today without knowing when that
8	information will come.
9	MEMBER REDLICH: I think this is what
10	John had done before and, John, you're welcome to
11	speak. I think part of our new request for data
12	was based on this data, such as when we happened
13	reviewed the sensitization claims, we thought those
14	were very reasonably adjudicated, and we did not
15	request more of them.
16	So we were and also taking into
17	account their trends, and there were trends over
18	the past 10 years in terms of more asthma, more
19	COPD. So we had this in mind.
20	John might want to quickly run through,
21	or I can, either one, but please speak up. I think
22	let's see, the first slide is that the first

1	of the data slides? I think if you go to the first
2	of that section.
3	So this, I believe, was the total number
4	of claims over it was a 10-year period of time
5	under each category. I think it will be more
6	helpful to go through the next slide.
7	Also, this is not given individuals,
8	because a given individual could have more than
9	one claim. So an individual could have a claim
10	of beryllium and COPD. But the next slide this
11	one just gives you some idea of the trends.
12	It shows the third column down is the
13	total number of claims under, let's say, CBD
14	beryllium sensitization. CS is chronic silicosis,
15	approved and denied under each of those.
16	And I believe this was the because
17	a given condition could recirculate, so
18	CHAIR MARKOWITZ: These are just
19	counter-claims where people sent
20	MEMBER REDLICH: That's right. We had
21	highlighted certain trends so that there were more
22	of the CBDs, and also as a percentage approved,

1 that the CBD claims, that percentage had gone down, 2 that may have been because -- or reasonable claims had been approved, but we were just looking at the 3 4 trends. I think the next slide --5 (Simultaneous speaking.) 6 7 MEMBER REDLICH: Yes, because it was the year, I believe, is referring to the year that 8 9 the claim was processed. That could be processed 10 for a pre. I think over time, more of them are in post 1996 simply because of the timing. 11 The next slide, this was additional --12 13 it's the same organization. These were the additional conditions we had gotten, the data John 14 had analyzed for: asthma, COPD, ILD, and sarcoid. 15 16 So I think you can see that from 2005 the number of asthma and COPD claims had gone up. 17 The ILD was really a total of only 21 total claims. 18 19 Most of those were denied, so that was the reason we had an interest in looking at more of those, 20 and sarcoid was a relatively small number of claims, 21

but they were denied. So we had targeted our recent

request with this in mind.

CHAIR MARKOWITZ: One thing that this shows -- Steve Markowitz -- is that we request five or 10 approved sarcoidosis claims, we're not going to get any. Similarly with ILD, we're going to get a large percentage of those that had been approved. So we might consider modifying that. But I didn't mean to interrupt you.

MEMBER REDLICH: I thought that that was the upper limit, that there might be not as many claims in the category that we requested.

I think the next slide -- please, John, speak up, because you did this. These were the denial reasons, and I believe this was not -- these were the reasons that John -- there was a reason given in the database.

And the reasons varied somewhat for the different conditions, so for CBD and BS, beryllium sensitization, the most common reason was medical information insufficient. For chronic silicosis, it was some additional reasons too, in terms of whether the employee was covered.

1	For some of these, we saw examples of
2	them: where there was a claim for beryllium
3	sensitization and there wasn't a result of the test,
4	and the claim that someone submitted that, and there
5	was a final adjudication, all of which was
6	appropriate.
7	I think there's one more slide. These
8	were for, then, the additional COPD, asthma, and
9	these were a different negative-positive result
10	for COPD, and I think we saw we've seen some
11	examples of that. So it is a different reason than
12	insufficient medical information, and same with
13	asthma and interstitial lung disease. So we had
14	an interesting focus on these conditions.
15	CHAIR MARKOWITZ: Is there anything
16	else
17	MEMBER REDLICH: I think some of these
18	negative-positive results, based on the records
19	that we had reviewed, relate to the interpretation
20	of the exposure information, and the issues that
21	we discussed yesterday at length.

And I think -- is there one more?

That's it, I think. So our feeling was that it would be useful to have this data updated for the past two years to see if there's been a change in these trends, since we have been working on this and meeting, and then to use that information to best target where we focus our efforts.

But, George, I totally agree, this is focused on the respiratory component.

CHAIR MARKOWITZ: Right.

MEMBER BERENJI: But I just want to comment; Dr. Dement and Dr. Redlich, you guys did a great job at least kind of developing the methodology. It would be great to able to kind of apply this basic methodology to other organ systems, perhaps Parkinsonism, manganism. We could definitely apply the same methodology, because it looks like we've already got it down to a science, pretty much.

MEMBER DEMENT: I wouldn't necessarily call it a science at this point. The data came in an Excel file, and it took a bit of going back and forth of the program staff to interpret some

of the fields. 1 2 I guess we got it down to a possibility. If it's the Board's desire that we expand that 3 4 look at it across the board for 5 conditions, then I guess I could take that on if 6 needed. John did this; 7 MEMBER REDLICH: simply took his tables and formatted them to put 8 them into the Power Point. 9 10 CHAIR MARKOWITZ: Well, we're going to take a break, and I'll try to look at -- in our 11 request, we did request data for a variety of 12 13 conditions in a certain form which reflected what we've been looking at. So let me just take a look 14 at that detail, and then we can discuss that 15 16 further. But I propose that we take a break and 17 reconvene in 10 minutes, at 20 of 11, if that's 18 19 all right. 20 (Whereupon, the above-entitled matter went off the record at 10:29 a.m. and resumed at 21 10:46 a.m.) 22

engaged public here. We're going to get started.

Okay. So we have a pending data request on a number of important conditions, including pulmonary disease, neurologic, cancer, and kidney disease, and the most common conditions overall, organized in a way that should be similar for all the outcomes.

We're not sure when that data request will be fulfilled, but it's pending, and it will be useful.

I would just suggest that we, on the claims request, we modify our current request. Pending is four outcomes, 20 each, so that's a total of 80 claims. And so the question is, how many of those claims do we actually want, believe that we can review if it's less than 20? And then, what's the breakdown, accepted versus denied?

I would suggest that requesting claims for any other conditions such as cancer and the like, that we wait until we get the data so that we can make an intelligent choice about --

1	meanwhile, we get additional claims, we'll have
2	our hands full taking a look at those claims. Dr.
3	Redlich?
4	MEMBER REDLICH: I think the point
5	about wanting to look at some accepted claims is
6	appropriate.
7	MEMBER FRIEDMAN-JIMENEZ: Move closer.
8	MEMBER REDLICH: I'm sorry. So I think
9	that the point made that we should need to look
10	at both accepted and denied claims, so we could
11	do something like five accepted and 15 denied for
12	the different conditions.
13	I'm open to others. I found that the
14	pulmonary ones, once you are familiar with them,
15	can be reviewed rather quickly.
16	CHAIR MARKOWITZ: It is true. Dr.
17	Redlich reminded me at the break that for these
18	outcomes, as opposed to COPD, and as opposed to
19	Parkinson's disease, the path for decision-making
20	is much clearer because it's set out in part by
21	the regulation in the statute, for instance,

chronic beryllium disease, sarcoidosis and, to some

1	extent, asthma. So reviewing those claims is more
2	straightforward, takes less time, than the claims
3	that we've looked at to date.
4	But then you're proposing the same
5	number, 20 divided by five accepted and 15 denied.
6	MEMBER REDLICH: Yes.
7	CHAIR MARKOWITZ: For the four
8	conditions. And my question is, do we need that
9	many?
10	MEMBER REDLICH: Based on our past
11	experience, there were a number of claims that were
12	very appropriately adjudicated. So we want to
13	identify any issues, I think, that the team denied
14	would be appropriate.
15	CHAIR MARKOWITZ: We need some
16	consensus on this, because we're
17	MEMBER REDLICH: John and Kirk also
18	reviewed them. I'm open to other suggestions.
19	CHAIR MARKOWITZ: We need some
20	consensus on this, because this is work that we're
21	going to do ourselves. So this is synonymous with
22	a commitment by the Board to get this claim review

done. Board is smaller this iteration than it was 1 2 previously, so if we're going to stick with a request for 20, we should just be assured that we're 3 4 going to be able to do that work. 5 Would you remind us MEMBER DEMENT: 6 again of what we've asked for, because I'm a little 7 I know we have 80, but how do they distribute themselves? 8 9 CHAIR MARKOWITZ: Sure. So t.he 10 original request was for a total of 120. of those were for Parkinson's disease, and we have 11 The remaining 100 of those were for lung 12 diseases, and it was each of five different 13 conditions. So five times 20 is 100. 14 We have the ones for COPD, so there are 15 16 four pending pulmonary disease requests, and those four conditions are chronic beryllium disease, 17 sarcoidosis, interstitial lung 18 disease, and 19 asthma. The request was that only denied claims 20 should be included, and the most recent claims 21 available should be selected. We should exclude 22

1 claims that were previously reviewed by the Board. 2 Let me just say, the Department of Labor has already done work. The pool claims, 3 4 compliance with this request. We should not 5 reverse or modify our request. That, in any way, 6 subverts any work they've done to-date on these 7 claims. We're not sure where they are with this request, but just, the request is not to undo work 8 that they've done so far. 9 10 MEMBER MAHS: Ron Mahs. Was there a 11 chance you could continue with those 20 and just ask for an additional five accepted? Because those 12 13 were all denied that you asked for correct? 14 you asked for? CHAIR MARKOWITZ: You mean increase the 15 16 number to 25? Well, whatever we're 17 MEMBER MAHS: allotted to do each, if we can get to them all, 18 19 that's fantastic. If we can't get to them all, 20 at least we've got the opportunity there. Well, I mean, the CHAIR MARKOWITZ: 21 22 problem is that it appears to be considerable work

1	on the part of the Department to secure and provide
2	these claims, so I don't think we should make a
3	request unless we believe we can do our work on
4	those claims.
5	MEMBER REDLICH: So I had proposed that
6	we reduce the number from 20 to 15 of the denied
7	claims, but also include five accepted.
8	CHAIR MARKOWITZ: Okay.
9	MEMBER REDLICH: It is still 20, but
10	if people want to reduce that further, I other
11	people's thoughts.
12	CHAIR MARKOWITZ: Right.
13	MEMBER REDLICH: My guess is also that
14	in that number what happened last time was, there
15	was overlap. So it would be like COPD, the same
16	claim could end up in both buckets, because the
17	person could have a claim for COPD and
18	pneumoconiosis, so the total number of people was
19	less than the number of requests.
20	CHAIR MARKOWITZ: Okay. So the
21	proposal is that we stay with 20 claims request
22	for each of the four conditions, but modify the

1 request to include 15 denied and five accepted. 2 However, if DOL has already done the work to provide the 20 denied claims, that would be fine. 3 4 So does anybody want any departure from I don't think we need to formulate an 5 that? 6 official recommendation about that. 7 MR. FITZGERALD: No. In fact, that's one of the reasons we created the form that we 8 9 discussed yesterday, to try to avert this sort of 10 issue where there is maybe some uncertainty or 11 changes going on in terms of the thinking. We want the Board to able to really think 12 13 about what the requests are, formulate the data requests, be very specific about what the use of 14 that data is going to be so that we're not grappling 15 16 with trying to address those issues. I think the form will actually help us, and that process will 17 get better. 18 19 CHAIR MARKOWITZ: Okay. So let's move 20 on. Let's get back to the agenda. Mr. Tebay? MEMBER TEBAY: Calin Tebay. Before we 21 22 leave today, I know we're running out of time, but

1 I'd like some help in this matter where the Board will make a formal recommendation regarding the 2 3 IH response. 4 Basically, what they're saying is the 5 lack of data is lack of risk, or the fact that an 6 OEL or a PEL determines a diagnosis. 7 I think we should come up with some kind if recommendation here to modify that response or 8 that language so it doesn't almost set the claim 9 10 -- I mean, the IH is driving the diagnosis at this 11 It seems to be; maybe I'm not communicating that correctly, but I'd like to -- maybe we can 12 13 make a formal recommendation to change how that's being worded. 14 CHAIR MARKOWITZ: So just a point of 15 16 clarification. You're talking about the boilerplate language about post-'95 exposures. 17 That the lack of data means that it doesn't exceed 18 19 20 MEMBER TEBAY: Not meeting regulatory limits is seeming to drive the direction of the 21 claim. 22

1 CHAIR MARKOWITZ: Okay. Discussion? 2 And I would say that this relates to Task 4 of the in duty to assess the quality of the Board: 3 4 industrial hygiene evaluation. 5 if we're going to elaborate So 6 recommendation, then we need to actually put that 7 text on the board and see if we can come to This relates to the recision of 8 agreement. Circular 15-06. So how do we want to phrase this? 9 10 MEMBER DEMENT: This is John. I think 11 it ought to be phrased first in stating what we've observed based on the case review. 12 13 So the observation is that the ΙH 14 assessments continue to use the phrase and the 15 determination that exposures in the past, 16 mid-1990s, would not exceed regulatory limits, but without supporting information, both with regard 17 to levels and what regulations are actually being 18 19 referred to. So that's an observation. 20 I think the second part of it is, the Board recommends that this language be omitted from 21 22 the IH report, and the basis for determination

1	exposures in mid-1990s and be stated by the IH that
2	is period. That's a first draft.
3	And basis for exposure determination
4	be provided by the IH in the report. Be provided
5	by the IH in the report.
6	CHAIR MARKOWITZ: Well, their
7	statement that no monitoring data exist as evidence
8	of exceeding regulatory limits, to play devil's
9	advocate, would be their basis for their exposure
10	determination. We don't have any data to suggest
11	it's above the limits. That's the basis for our
12	exposure determination.
13	MEMBER DEMENT: The alternative would
14	be to the rescinding the circular, and the
15	observation is it continues to use the language
16	contained in the circular, basically.
17	I don't know how to phrase this
18	perfectly, but just say get it out of there.
19	CHAIR MARKOWITZ: How about that the
20	absence of monitoring data post-1995 should not
21	be automatically interpreted as representing an
22	absence of risk?

1	MEMBER DEMENT: I would say, in absence
2	of exposure or risk.
3	CHAIR MARKOWITZ: Right.
4	MEMBER DEMENT: Because there's two
5	things that are going to be addressed by that
6	statement: one is the absence of exposure, and
7	the second is, the assumption is, if you were within
8	the regulatory limits, there is no risk. And we
9	know that not to be the case for many materials.
10	CHAIR MARKOWITZ: Well, that
11	observation will be in the rationale for this.
12	It will be addressed, unless you want to put it
13	here, right in the front.
14	MEMBER DEMENT: And I think we should
15	modify this second part of the recommendation:
16	the basis for a negative exposure determination
17	be provided by the IH.
18	CHAIR MARKOWITZ: So to fill out this
19	first line, The Board has observed that industrial
20	hygiene assessments or rather, recent industrial
21	hygiene assessments appear to frequently use
22	stereotypic language, indicating that the absence

1	of monitoring data above the established regulatory
2	levels.
3	So I think indicating has to be changed
4	to citing. They use stereotypic language that cite
5	the absence. After cite, you can just take out
6	that.
7	So I wonder there on the last line where
8	we talked about the basis for negative exposure
9	determination be provided, whether we should add,
10	if available?
11	MEMBER DEMENT: I guess, back to our
12	discussion yesterday, just trying to get to the
13	rationale behind that determination. Sometimes
14	it's based on monitoring data; sometimes it may
15	be based on professional judgment. If so, that's
16	what it is; it's professional judgment based on
17	IH. I'd just like to see that in there.
18	CHAIR MARKOWITZ: Right. And that can
19	be perfectly acceptable.
20	MEMBER DEMENT: Sometimes it's just
21	common sense.
22	CHAIR MARKOWITZ: Right. Dr.

## Friedman-Jimenez?

MEMBER FRIEDMAN-JIMENEZ: In the spirit of providing, not just an explanation of the problem but a proposed solution, I think that the data largely don't exist at all. What we're seeing is a paucity of monitoring data in general, not just monitoring data above some limit.

And the reality is that many of these jobs do have some information on exposure that's reflected the SEM and in the industrial hygienist's knowledge of those kinds of jobs. These are people that have specific knowledge of many of these jobs.

And so the exposure assessment will need to reflect the industrial hygienist's expertise and judgment. I think that word is useful: judgment, as to what level of exposure actually existed in the absence of individual or area monitoring data for that particular site and person.

It's going to have to be job-specific, not individual-specific, and that's the reality.

So I think we should suggest that they use

industrial hygiene literature as well as their specific knowledge of these work sites to generate these exposure assessments.

Because exposure assessments have to be generated, and we don't have this Holy Grail of individual or area monitoring data. So we have to suggest something, I think.

CHAIR MARKOWITZ: Well, I disagree,

CHAIR MARKOWITZ: Well, I disagree, actually. They could talk to the claimants. They could find out what actually happened post-'95 in the workplace.

They could find out what they did and whether that disruption likely produced results, because they're not going to find them in the text, and their professional judgment is great, but DOE is a very big complex, and they've been everywhere and assessed all those jobs.

So there are multiple sources, but frankly I think upgrading their interaction and understanding of what actually happened in the workplace to that claimant post-'95 would be a good place to start. So I don't want to be specific

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FRIEDMAN-JIMENEZ: So this MEMBER prohibition against subjective information, think we have to address that. What people imagine is objective measurement really isn't objective at all; it reflects a lot of variables. So I agree with you completely, that starting with occupational health questionnaire, the individual's perception of exposure, important information that has to be factored in by the industrial hygienist.

CHAIR MARKOWITZ: Well, and DOL recognizes that, because part of their assessment is the occupational health questionnaire and whatever affidavits are submitted and the like.

So they recognize the legitimacy of that information. We're talking about amplifying it; we're talking about emphasizing it for post-'95 where it's not clear what was going on.

MEMBER TEBAY: What -- Calin Tebay.

But we have hundreds of IHs on these sites, and each working group has a -- what do they call them

-- they're a project IH for each individual working group.

If they're using professional judgment, the contracted IHs, why aren't they reaching out to these project IHs or these work groups, along with the workers and saying, What were the exposures that exist?

I can tell you from working at a couple of sites in different places, that I think a lot of professional judgment is used. I don't think there's a bunch of IH data that exists for each one of those work groups, but the professional judgment is there to say, These are the possible potentials. We don't know what levels, because we don't monitor for them, but there's definitely these potentials for these exposures.

But once again, the IHs are not being forced to reach out to the worker or the other resources at all. They're just making a boilerplate response and walking away. Why aren't they being forced to reach out to the people with the information?

1 CHAIR MARKOWITZ: Dr. 2 Friedman-Jimenez? MEMBER FRIEDMAN-JIMENEZ: 3 George Friedman-Jimenez. 4 One potential problem with that 5 is that it puts the individual IHs in the workplace 6 in an awkward position. They are responsible for 7 the health and safety of the people on their site, and for them to say, Well, the exposures were really 8 pretty substantial and could have led to health 9 10 defects, would maybe be difficult for them. 11 So it puts them in a difficult position and almost a conflict of interest. I'm not sure 12 13 that that's going to lead to objectively better estimates of the exposures. So we'd have to think 14 I don't know the specifics, but --15 about that. Sure, I understand. 16 MEMBER TEBAY: MEMBER POPE: It seems like the cases 17 I've looked at, the IHs that were making these 18 19 blanket statements, saying that the exposure was 20 low, and I totally agree with Dr. Dement's comment about, if you don't know, then say no, the exposure 21 did not exist. 22

1 But for you to blanket and say that the 2 exposure was low just to forward that claim through, I think we've seen a lot of that, where the IHs 3 4 making that statement. So the CMC is 5 concurring with that assessment from the IH. 6 CHAIR MARKOWITZ: Mr. Vance, are you still on? 7 MR. VANCE: Yes, I am. 8 9 CHAIR MARKOWITZ: So an interesting 10 question has been raised about the industrial hygienists that you have that are working on these 11 cases, claims whether any consideration of reaching 12 13 out. I know that there's a request 14 whatever records DOE has, but has it ever been 15 16 discussed, any sort of communication with the currently-employed industrial hygienists at the 17 sites? 18 19 MR. VANCE: Not that I'm aware of. would always be getting information from the 20 Department of Energy with regard to any individual 21 22 monitoring data that we have on an employee, but

1 as the Board noted, oftentimes that information 2 might not be very comprehensive or cover the entire working history of an employee. 3 The context of our discussions up to 4 5 this point have been mostly focused on what 6 information would be best obtained through the 7 occupational health questionnaire modifications to that process, but I don't recall 8 discussing specifically or engaging in any kind 9 10 formal interaction between our industrial 11 hygienists and site industrial hygienists. Okay, thank you. 12 CHAIR MARKOWITZ: So 13 we're looking at -- we have language of -- we're looking at language of a recommendation. Before 14 we receive an official motion to accept this 15 16 recommendation, is there any change in the language that we're looking at that anybody proposes? 17 I would like to make a MEMBER MAHS: 18 19 statement if I can, Ron Mahs. 20 CHAIR MARKOWITZ: Sure. MEMBER MAHS: In the last 15 years 21 before I retired, I was general foreman at Y-12 22

and the other two plants. We toured many buildings every day, because I had people working all over the place. And in all of those years, about the only thing I saw IH ever monitor was for asbestos or for radiation.

If you're on the job looking for toxin or something, the safety person assigned to that job did it. IH had no contact or no idea what the exposures would be.

CHAIR MARKOWITZ: Thank you.

MEMBER REDLICH: Carol Redlich. I'11 just add that I think the cases that we've reviewed to date, the prior ones, the major reason we disagreed, in cases where we did disagree with the final adjudication was where the CMC interpreted the IH report differently, given our expertise in occupational lung disease, occupational medicine, industrial hygiene exposure, based the information we had from the occupational health questionnaire, the type of work the person did, and the time period, that we felt that the SEM was not accurately representing the exposure, and that

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1 the CMC, then, didn't have our expertise to also, 2 let's say, look at the questionnaire and put a lot of weight on the SEM that we felt was not accurately 3 4 reflecting the exposure. One example, just being a miner of 15 5 6 years where the SEM said that the only relevant 7 exposure as far as COPD was aluminum or -- and knowing the nature of what mining work is like, 8 we felt that that was not correct. 9 So we knew how 10 to interpret the SEM. So I think that's why it's so important that that wording be modified. 11 CHAIR MARKOWITZ: When you say that 12 13 wording, you're talking about what we're looking 14 at now? 15 MEMBER REDLICH: Yes, exactly. 16 Because I think that, given the nature of the physicians who are the CMCs, given that they're 17 more limited expertise in pulmonary occupational 18 19 medicine conditions are weighing the SEM very 20 heavily. CHAIR MARKOWITZ: So Ι 21 have some 22 suggestions for the language: the Board has

1 observed, comma, based on review of a limited number 2 of recent claims, comma, that -- leave that second recent in, just to be duplicative, but you can take 3 4 out that the before the recent -- appear frequently 5 So we don't have a split infinitive. 6 And then above the established 7 regulatory levels in the mid-1990s, so we're clear what time frame we're talking about. Any other 8 9 suggestions on the wording? Yes, Dr. Silver. 10 MEMBER SILVER: Should we put in a phrase that draws attention to Circular 15-06, 11 rescinded by Circular 17-04? 12 That is what we're 13 talking about, despite the official rescinding of Circular 15-06, the Board has observed --14 CHAIR MARKOWITZ: You know, we could 15 16 put in something to the effect of it. I wouldn't write this, that this appears to contradict the 17 rescision of -- but we actually raised this 18 19 yesterday with Ms. Leiton, and she had a response 20 to that, how this language did not contradict the rescision. 21

So I'm a little concerned that we --

1 it doesn't further our recommendation. Dr. 2 Redlich? My reading, and I MEMBER REDLICH: 3 4 think others can correct me, was that it's not that they cite the absence of monitoring data; they don't 5 state the basis of their conclusion that there is 6 7 no low exposure or low risk. Did you just check what the wording was? 8 I think we should just check what the wording is. 9 10 MEMBER DEMENT: The actual phrase that 11 consistently appears is that no available evidence, 12 personal industrial or area 13 monitoring data, paren close, to support after the mid-1990s, as exposure would have exceeded existing 14 regulatory standards. 15 16 you said, there's no available evidence. So they're sort of saying there's no 17 sampling data. Then I look at what I got from a 18 19 DAR, the request for information, and for the most 20 part there's nothing there except for some radiological monitoring data; very little IH data 21

that I've seen in what I've reviewed so far.

1	So the question is, so nobody sampled,
2	so therefore you assume that because some programs
3	were beginning to be implemented to have industrial
4	hygiene at these sites, that exposure didn't
5	exceed; that's not an appropriate conclusion to
6	draw.
7	MEMBER REDLICH: Yes, okay. So I'm
8	just wondering what the most clearest way to state
9	what our concern is. It might be first that they
10	should, number one, clarify the source of the data
11	that their decision is based on; and number two,
12	that lack of data should not be interpreted as low
13	or no risk. We may have worded it that way.
14	MEMBER DEMENT: I think the source and
15	basis support the negative exposure.
16	MEMBER REDLICH: Yes. But I think we
17	just put in active of what we want.
18	MEMBER DEMENT: So down at the bottom
19	of the recommendation, I guess.
20	MEMBER REDLICH: Okay. So I'm sorry.
21	MEMBER DEMENT: I support the data
22	sources in basis for negative IH reporting.

MEMBER REDLICH: So why don't we just make it more general for whether it's negative or positive, we would like clarification on the source of the exposure data? And then that statement that lack of data should not be interpreted as --

MEMBER TEBAY: So real quick, because I've read the version that you read, what I've seen is the lack of data. But then there's the version that says that there's significant exposure, and there's a lack of data showing that you've been exposed over an OEL or a PEL; therefore, having or contracting some kind of disease or condition would be not -- so I want to make sure we're still covering that portion, right? Because that's what happens.

When they say that there's a lack of information, we know you've been exposed. But it's going to be below an OEL or a PEL, and then when that moves on to the CMC, the CMC then interprets that as, there's no way this person was exposed at enough of a level to create some kind of condition or disease.

1 So as long as this is covering that, 2 I'm good, and I'll rely on you folks to determine that. 3 This is Kirk Domina. 4 MEMBER DOMINA: 5 One of the other issues I have when Mr. Vance was 6 on the phone about the information he gets from 7 DOE; well, DOE didn't have a moratorium destroying records until way after '95. So you're 8 always going to get that answer. 9 10 So even if there was data, they didn't have to keep it. And so to me in my thinking, we 11 need to move that '95 date out because of their 12 13 moratorium of not having records. It's biased against the claimant. 14 MEMBER REDLICH: I think we're all 15 16 saying the same thing; it's just a matter of how we word this recommendation. 17 Well, and what we CHAIR MARKOWITZ: 18 19 could do actually, in the last sentence: the absence of monitoring data post-1995, or evidence 20 of data showing exposure levels of below regulatory 21 22 limits. Does that capture --

1 MEMBER DOMINA: That's better. 2 MEMBER DEMENT: I would say, absence of -- You could say significant exposure. 3 I mean, 4 the issue is -- sometimes they'll say significant 5 exposure, but they did say it's a low regulatory 6 limit so therefore, de minimis. They don't say 7 de minimis, but that's they really interpret it 8 as. So, Kevin, the third 9 CHAIR MARKOWITZ: 10 word from the last, exposure? If you could just put in in, significant, before that. Yes, that's 11 it. 12 13 And omitting MEMBER REDLICH: so 14 language; there's variations on the language, so I think what's most important is that we want 15 16 clarification of the basis of the exposure data, because that's usually not stated, and the absence 17 of monitoring. 18 19 We also concerned about the are 20 lanquage. I just don't think that that is the number one piece, because there's lots of variance 21 22 of language.

1 CHAIR MARKOWITZ: Yes, but the question 2 is whether what we're looking at now captures what 3 we mean. 4 MEMBER REDLICH: So industrial hygiene 5 folks, what do you think would be appropriate 6 information to include as justification for the 7 conclusions that the IH has come up with? There can be lots of MEMBER DEMENT: 8 There can actually be some experience with 9 things. 10 the industrial hygienist's experience with that particular job, that work site, that task, and 11 that's all legitimate. 12 13 There can be published literature that 14 supports in that time frame that exposures were significantly reduced. So we all come to this with 15 16 experiences, knowledge, our own and determinations, if you will. I just think they 17 need to put it there. 18 19 If it's in IH's -- based on my own 20 personal experience and the published literature, likely not have exceeded 21 exposure were to regulatory limits, then that's our basis. 22

1	It does three things: it gives you the
2	basis for the decision; it also sets some parameters
3	about your certainty about that decision.
4	MEMBER REDLICH: I understand. Are
5	they providing the basis of their conclusions?
6	What IH data do they use to determine?
7	MEMBER DEMENT: Yes, I mean
8	MEMBER MAHS: No, is that partly
9	happening?
10	MEMBER DEMENT: We saw yesterday, I
11	think the standard reference list, most of which
12	don't provide a basis for determination of
13	exposures for that job. I mean, it's a standard
14	IH set of references, some of which are actually
15	on some medical texts.
16	CHAIR MARKOWITZ: Very good, very good
17	medical texts, I would add.
18	MEMBER DEMENT: Yes, they're old.
19	Some of them are quite old. They really don't
20	provide a basis for that decision. Now, if you
21	were to go on diesel exhaust, for example, you can
22	go to the literature, and you can find exposure

determinations based on objective measurements that would actually support, in the mid-1990s and early 2000s, that exposures to diesel exhaust in a general way, were reduced.

MEMBER REDLICH: So all I'm -- is it appropriate for us to request as a recommendation

MEMBER REDLICH: So all I'm -- is it appropriate for us to request as a recommendation that they provide a better basis for their assessment?

CHAIR MARKOWITZ: But no. The language currently in there on the second sentence says that we recommend that language be omitted from the industrial hygiene report and that the basis for a negative exposure determination be provided by the industrial hygienist.

MEMBER DEMENT: Yes, I don't know how to get more specific than that. For example, if the document request came back and there were some industrial hygiene monitoring data, not even for that person, but at least for a similar job or a similar location, that could be used. That's legitimate information, so I don't know think we want to box ourselves in to specify exactly what's

1	needed, because it can vary a lot.
2	MEMBER REDLICH: Okay. I agree. So
3	I just think we want the wording to be broad enough
4	so it's both a negative or a low. So I just think
5	that we should start with the request that we want
6	the basis for the determination of the exposure
7	assessment.
8	And we want to get rid of certain
9	language. But I think that the more active thing
10	is, we need the basis for their determining low
11	or no risk.
12	MEMBER DEMENT: Right. You know, if
13	you went on the second sentence after IH report,
14	put a period and then start a new sentence
15	MEMBER REDLICH: Yes, that's fine.
16	MEMBER DEMENT: so now, one line up,
17	IH report, period. Right, okay. Then start a new
18	sentence: The basis is that?
19	MEMBER REDLICH: Yes. I think
20	sometimes they are mentioned as being low, and I
21	think that's
22	MEMBER DEMENT: We're asking for the

1	basis of their determination. So if they determine
2	that it's low, fine. What's the basis?
3	MEMBER REDLICH: Yes, but you have
4	negatives. So I think whether it's low, whether
5	it's negative, we want the basis for the exposure
6	determination provided.
7	MEMBER DEMENT: Right.
8	MEMBER REDLICH: We want it broader.
9	MEMBER DEMENT: So what language do you
10	want, where?
11	MEMBER REDLICH: The basis for the
12	negative or low exposure determination. We want
13	the basis for all exposure determinations. I defer
14	to John and
15	CHAIR MARKOWITZ: My concern I
16	understand that, but it dilutes the impact, because
17	we're really zeroing in on use of specific language.
18	MEMBER REDLICH: Okay.
19	CHAIR MARKOWITZ: I'm afraid our main
20	point may get a little lost or diluted.
21	MEMBER REDLICH: Okay.
22	MEMBER DEMENT: I think it will be,

1	because in some cases IH is making a determination
2	that, prior to this time frame, in the mid-1990s.
3	In some cases they are making a determination that
4	exposures were significant and sometimes not higher
5	than low anyway. So I that's they're using a
6	time frame reference to make that determination.
7	I think that's fine.
8	MEMBER TEBAY: Can I ask a quick
9	question of Mr. Vance? Is he still on the phone?
10	MR. VANCE: Yes, I'm still here.
11	MEMBER TEBAY: This recommendation,
12	how does that get distributed? Because really,
13	there's part of this that apply to different people
14	in the process, right? I mean, you've got the IH
15	that's going to read it; the CMC is going to utilize
16	it, and the CE. The last sentence of it is really
17	important for the CE. How does this get
18	distributed?
19	MR. VANCE: What specifically are we
20	talking about?
21	MEMBER TEBAY: For instance, the last
22	part of this recommendation says, the absence of

1 monitoring data post-1995 -- you follow me there? 2 That piece? Right. You have to keep MR. VANCE: 3 4 in mind that what the Department of Labor is 5 utilizing is the opinion of subject matter experts. 6 So what the Board is always going to struggle with, 7 what the Department of Labor is struggling with is the absence of information. 8 direct, 9 We do not have personal 10 information about many workers, so we leave it to the judgment of the industrial hygienist's team, 11 toxicology team, the medical folks, and other 12 13 experts to give us information. So a lot of the information that you're 14 discussing is directly attributable 15 to 16 industrial hygienist looking at it and saying, This is my best understanding of the information that 17 I have in case, in my knowledge, my education, and 18 19 background in being able to respond to this. So what we did was, we took away the 20 ability of claims examiners who 21 make those

generalizations, and now it's the

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industrial

1 hygienist that's incorporating that information 2 and their best understanding of the information that they're being asked to respond to from an 3 4 extent and nature or duration of exposure. So that's sort of where that information 5 6 comes from. So that's what I think the Director 7 was talking about the other day, is that what we did was, we said, Okay, but this is not a claims 8 examiner generalization any longer. This is an 9 10 industrial hygienist looking at it and applying their best understanding of exposure information. 11 I think Dr. Dement was talking a little 12 13 They operate in sort of these bit about that. generalizations and their own understanding of 14 their expertise as industrial hygienists. 15 16 there can always be a lot of discussion about that interpretation of whether that's accurate or not. 17 But that's the general source of that information. 18 19 Does that make sense? 20 MEMBER REDLICH: Yes. MEMBER SILVER: 21 A comment. 22 CHAIR MARKOWITZ: Go ahead, I'm sorry.

MEMBER SILVER: Yes. Mr. Vance, you said there can always be a lot of discussion, but it can't be an informed discussion unless we crack open the black box of their judgment.

An analogy to radiation for a long time, the health physics profession said, We're the experts; we've got it figured out. The NIOSH Advisory Board has cracked open that black box in various perspectives have been brought to bear on the large number of assumptions that are made in the absence of hard data very often, and we're looking to do the same thing here.

This dovetails nicely with our long-standing recommendation that the industrial hygienist be able to talk to the claimant.

When that additional data from the claimant is part of the determination, it can be laid out with various sources of information; documents and literature; there are models; there's a rational process that starts with a volume of material; the energy that's applied to it; the volume of the work space in which it's diluted;

1 the presence or absence of ventilation systems; 2 the position of the worker in relation to the sources of contaminant. 3 4 You can get that from talking to the 5 worker and spell out as much of that as is available 6 in the hygienist's brain. 7 CHAIR MARKOWITZ: So we do need to move on, so I just want to make sure we don't -- and 8 Silver did this, but we 9 I'm not suggesting Dr. 10 shouldn't repeat comments that have previously been 11 But I'm not suggesting you did that. made. Friedman-Jimenez? 12 13 MEMBER FRIEDMAN-JIMENEZ: quick 14 point of information; question: Does this site exposure matrix have a time dimension? 15 16 differentiate between pre-1995 and post-1995? CHAIR MARKOWITZ: The answer is --17 MEMBER FRIEDMAN-JIMENEZ: 18 That's an 19 area in which individual workers can have perceived 20 a change or no change. Assuming that they have a good memory, and they're not memory loss people 21 with Parkinson's disease, but they may not know 22

1 exactly the identity of the dust that they're 2 exposed to, but they would know if it's decreased a lot recently. 3 4 So I think, again, we should really push 5 for the industrial hygienist having access to ask 6 the claimants what their perceptions were and take 7 that for what it is. I mean, it's a subjective piece of information, but it is a data point where 8 we're really -- there's a real absence of hard data. 9 10 CHAIR MARKOWITZ: Right. So take a 11 look at this language. We need a proposal, a motion on this. 12 13 MEMBER REDLICH: I just would suggest one or two minor modifications of the wording. 14 That recent IH assessments -- it's based on review 15 16 of a limited number, so I think we could take out the appear. 17 That's fine. CHAIR MARKOWITZ: 18 19 MEMBER REDLICH: And there was one 20 other -- just to make it more -- and then the absence, the last sentence? Automatically -- I 21 22 think the word automatically could be removed.

1	CHAIR MARKOWITZ: That's fine also.
2	Friendly amendments accepted. We could also spell
3	out IH; that would help. Dr. Friedman-Jimenez?
4	MEMBER FRIEDMAN-JIMENEZ: The last
5	sentence: The absence of monitoring data
6	post-1995 or evidence of data showing exposure
7	levels below that is ambiguous as to whether
8	you mean the absence of evidence of data showing
9	exposure levels or I think we should clarify
10	that language.
11	CHAIR MARKOWITZ: I think we should
12	have a comma after post-1995. And if you want to
13	put either at the beginning of the sentence, would
14	that help, George? Put either the absence or
15	evidence, to make it clear that there are two
16	distinct conditions.
17	MEMBER FRIEDMAN-JIMENEZ: Yes. I
18	think that clarifies it well.
19	CHAIR MARKOWITZ: Aren't some of you
20	glad you don't work for a university?
21	MEMBER REDLICH: One other friendly
22	suggestion: the first sentence, they use language

1	that cites the absence of monitoring data above
2	the regulatory levels as they're using that data
3	as indication of no risk, or
4	CHAIR MARKOWITZ: They don't say that,
5	actually. They just make the statement; they don't
6	actually draw that conclusion. You know what I
7	mean?
8	MEMBER REDLICH: Corrected?
9	CHAIR MARKOWITZ: Yes. Okay. Is
10	there a motion?
11	MEMBER TEBAY: Yes.
12	CHAIR MARKOWITZ: Okay. That's a
13	motion to accept this recommendation. Is there
14	as second?
15	MEMBER MAHS: Second.
16	MEMBER FRIEDMAN-JIMENEZ: Do you want
17	to take out that and where the cursor is now? So
18	it would be provided by the industrial hygienist
19	in the report?
20	CHAIR MARKOWITZ: So right now it says,
21	The Board has observed, based on a review of a
22	limited number of recent claims, that recent

1 industrial hygienist assessments frequently use 2 stereotypic language to cite the absence monitoring data above the established regulatory 3 levels in the mid-1990s. 4 The Board recommends that this language 5 6 be omitted from the industrial hygienist's report. 7 The basis for a negative exposure determination should be provided by the industrial hygienist in 8 9 the report. 10 Either the absence of monitoring data, post-1995 or evidence of data showing exposure 11 levels below regulatory limits should not 12 13 representing interpreted as an absence of 14 significant exposure or risk. Evidence of data 15 MEMBER REDLICH: 16 versus just data. Do we need the evidence of? We can take that CHAIR MARKOWITZ: 17 evidence off; that's fine, in the interest of 18 19 shortening the recommendation. Okay. Open for discussion, final discussion. 20 MEMBER FRIEDMAN-JIMENEZ: 21 Do we want 22 to only ask for negative exposure determination

1	basis be provided, or also low or all exposure
2	determination? The question that
3	MEMBER REDLICH: I we
4	CHAIR MARKOWITZ: No, no. That
5	dilutes the impact of this is targeted to
6	specific language, and I'm afraid if we expand the
7	domain of this, the impact will be diluted.
8	MEMBER REDLICH: I already took back
9	that suggestion.
10	CHAIR MARKOWITZ: He's reneging it.
11	MEMBER REDLICH: I agree with Steve.
12	CHAIR MARKOWITZ: Okay. Final
13	comments; otherwise, we're going to take a vote.
14	Okay.
15	MR. FITZGERALD: All right, I'll call
16	the role here. Dr. Dement?
17	MEMBER DEMENT: Yes.
18	MR. FITZGERALD: Dr. Friedman-Jimenez?
19	MEMBER FRIEDMAN-JIMENEZ: Yes.
20	MR. FITZGERALD: Dr. Mikulski?
21	MEMBER MIKULSKI: Yes.
22	MR. FITZGERALD: Dr. Silver?

1	MEMBER SILVER: Yes.
2	MR. FITZGERALD: Dr. Berenji?
3	MEMBER BERENJI: Yes.
4	MR. FITZGERALD: Dr. Markowitz?
5	CHAIR MARKOWITZ: Yes.
6	MR. FITZGERALD: Dr. Redlich?
7	MEMBER REDLICH: Yes.
8	MR. FITZGERALD: Mr. Domina?
9	MEMBER DOMINA: Yes.
10	MR. FITZGERALD: Mr. Mahs?
11	MEMBER MAHS: Yes.
12	MR. FITZGERALD: Ms. Pope?
13	MEMBER POPE: Yes.
14	MR. FITZGERALD: Mr. Tebay?
15	MEMBER TEBAY: Yes.
16	MR. FITZGERALD: All right. Vote
17	passes unanimously.
18	CHAIR MARKOWITZ: Okay. We're going
19	to resume with the agenda. We will not finish by
20	11:45 today. Could you bring up the language on
21	the non-cancer outcomes?
22	So this is a brief item. Dr. Silver,

2 to add some, I think, specificity or helpful language to DOL's request to us. They requested 3 4 that we help them with looking at non-cancer outcomes of radiological materials. 5 6 So this is just a reformulation of that 7 language, which actually I think we reviewed at the February 28th meeting and pretty much approved. 8 I just want to put it out there. 9 10 We have no plan to actually work on this as 11 of yet, but I want to make sure it remains on the radar. 12 13 And let me say that I don't think we necessarily need 14 to engage in an extended discussion about the specific words of this. 15 16 long as it get the gist of what we think they might be after, we can submit it, because this is going 17 to go to DOL. They're going to tell us whether 18 19 this is what they had in mind. So, Dr. Silver. 20 MEMBER SILVER: That's the key point: we want DOL to give us feedback on whether this 21

just to remind you, fashioned some language trying

is what they were after.

22

1 CHAIR MARKOWITZ: You want to read 2 this, Ken? MEMBER SILVER: Some of this language 3 4 was adopted in whole cloth from what DOL originally 5 gave us, and then other language here was our 6 reformulation. In reviewing some of the radioactive 7 substances found in DOE sites, the SEM only linked 8 uranium with non-cancerous condition of acute 9 10 tubular necrosis. DEOIC asked the 11 Board to conduct 12 research, peer review, human studies, а ascertain whether there are additional non-cancer 13 14 diagnoses that literature link to exposure to radioactive substances such as uranium, plutonium, 15 16 polonium, thorium, and americium. While all are technically heavy metals, 17 plutonium, polonium, and americium have no stable 18 19 isotopes. Health effects may be based upon non-cancer effects of radiation, high LET alpha 20 radiation in particular, chemical toxicity, or a 21

22

combination thereof.

1 A related set of issues pertains to 2 effects, especially circulatory non-cancer diseases observed in the life span study of atomic 3 bomb survivors with an association with low LET 4 5 radiation exposure. 6 Evidence of such non-cancer effects in 7 nuclear worker cohorts or other occupational groups would be of interest. The Board could offer advice 8 9 the results of its analysis, including 10 recommendations, additional links produced by the division as part of an update to the SEM or its 11 policy quidance. 12 13 CHAIR MARKOWITZ: So were there any comments on this? It's a different matter as to 14 the extent to which Board members are currently 15 16 willing to volunteer to work on this issue, but in any event, this is the reformulation. 17 Anybody have any discussion on this? 18 19 Okay, that's fine. Actually, I think I submitted it, but only within the past week to DOL, 20 obviously there wouldn't be a response. 21

Then we can postpone the issue of who

1 wants to work on this particular subject, so we 2 needn't have that discussion. Thank you. Ken, 3 yes. 4 MEMBER SILVER: Thinking one or two 5 steps ahead, if DOL wants us to take this on, I 6 would do a little outreach to the NIOSH Advisory 7 Board, because they may have thought about this, and dose matters, whether you're talking about 8 cancer or non-cancer effects, and they have some 9 10 top-shelf expertise. Okay. So let's wait 11 CHAIR MARKOWITZ: until they get back to us, and then we can consider 12 13 whether we have -- we're also one person short on the Board, whether we have time and resources to 14 address this issue. 15 16 The next item on the agenda is review of public comments. I thought that we should --17 we received a number of comments that were posted 18 19 on our website, the written submissions, and there 20 were a number of comments yesterday. I thought we should just take a few 21

minutes to mention some of those and also address,

1 I think, the status of some of the issues raised 2 in those public comments. I've done some of that. If others want 3 4 to chime in, that's fine. There was one comment 5 we received about raising the issue of the quality 6 of an industrial hygiene assessment on a peripheral 7 neuropathy case, that certain relevant exposures were not adequately considered. 8 The submitter also provided the example 9 10 with personal information, identifiable information, deleted. 11 Frankly, I think this issue falls within 12 13 discussions about the claims and about our assessing the adequacy of the industrial hygienist 14 So it's an argument, actually -- in 15 assessment. 16 this case it was peripheral neuropathy, which is probably a fairly common claim, but I think we're 17 going to address this in our recommended claims 18 19 evaluation. 20 There was one comment citing the ombudsman 2017 report, that it contained items 21

relevant to our mission, and thankfully, Mr. Nelson

1 and Ms. Felin have attended this meeting and 2 yesterday gave us an update on the 2017 report, which we should review as a board. 3 4 There is a comment on -- a couple of comments actually, on this issue that we just made 5 6 a recommendation about. So we've addressed that. 7 There's a comment on impairment. came in a couple of months ago, actually, having 8 to do with the question of what the policy of the 9 10 program is. Let me just read this short section 11 here, because it encapsulates what the issue is. This a is January 28th, 2019 letter from 12 13 It has come to our attentions that DOL's ANWAG: Division of Energy Employees, Occupational Illness 14 and Compensation changed their policy regarding 15 16 assigning impairment ratings for pulmonary disease. 17 This policy is not published on the 18 DEOIC's website. It is our understanding that this 19 policy was issued only to DOEIC's contract medical 20 consultants and not to private practice impairment 21

specialists. And then it gives an example of use

of impairment ratings.

So I think what we need to do is simply ask for clarification from DOL about what this involves so that we can understand the issue. Not right now, Mr. Vance, but we just should request some clarification.

And this pertains to our Task number 4. I'm already assimilating that data form that you used. This relates to Task number 4, which is our obligation to look at the consistency, objectivity, and quality of the medical input into the claims process.

A couple of comments that came up yesterday: I did ask the Department of Labor, Ms. Leiton if, as a standing request, if we could have their participation in all of the board meetings, which I had not asked prior to the February 28th meeting, and they have agreed.

They will either attend in person or attend by phone our meetings to be available for clarification. I want to thank you, Mr. Vance, for being available today.

1	A question arose yesterday about
2	whether this board needed to re-approve or confirm
3	the prior board's recommendations. So this is a
4	question I've always I saw no need in that.
5	I assume those recommendations still to have full
6	weight or standing. Isn't that right, Doug?
7	MR. FITZGERALD: Yes, I mean, the Board's
8	made determinations of which and the agency
9	yes. I would say that is correct.
10	CHAIR MARKOWITZ: Okay. There's also
11	a question raised, which we've talked about briefly
12	before, that the industrial hygiene assessment
13	seems to focus on a limit of seven toxins, despite
14	the fact that the SEM frequently has a much larger
15	number of toxins in association with any given job
16	title.
17	I think that's something we should
18	discuss in the future.
19	MEMBER DOMINA: Steve.
20	CHAIR MARKOWITZ: Yes.
21	MEMBER DOMINA: On the comment just
22	before there that Doug was talking about, I saw

1	a letter from Ms. Hearthway where we had asked in
2	a prior board about an advisory committee or someone
3	to help us, and she said that this board, if I
4	remember correctly, did not ask for it, and so it
5	was null and void, something to that effect.
6	I think that letter was like, maybe last
7	fall, early
8	CHAIR MARKOWITZ: Well, yes. I think
9	that refers to the previous board's request for
10	resources, which we've taken care of today.
11	MEMBER DOMINA: All right. I just want
12	to be clear.
13	MR. FITZGERALD: I think there was
14	discussion within the past boards about resources,
15	but I don't think there was any formal request that
16	came forward from the Board.
17	CHAIR MARKOWITZ: There was a comment
18	yesterday about a preponderance of evidence. The
19	Board needs to look at that issue, a: whether it's
20	relevant to our assigned tasks, and then b: if
21	so, what it means. But we're not going to do that
22	now. We actually need to read that comment, I

1 think, which will be in the transcript. 2 I think actually that same comment has been made before, so if it's relevant to our 3 4 assigned tasks, then we need to look at that and 5 discuss that. 6 Any else on the public comments that I didn't -- I didn't review all of them, but many 7 of them. 8 I can give a brief update on the 9 Okav. 10 Presumption for Solvent-Induced Hearing Loss. haven't forgotten about it, but we also have not 11 prepared a response to DOL's response, pretty much 12 13 rejection of recommendation regarding our solvent-induced hearing loss. 14 I think they did, in the most recent 15 16 version of the Procedure Manual, 3.0, I believe they've added a couple more solvents to the list. 17 believe that may have come out 18 of 19 recommendation; I don't quite remember. 20 that's great. But we will continue to look at that issue and see whether there's evidence that we can 21

assemble that would be persuasive to Department

of Labor.

Any comments on that? Okay. Maybe the new Board member will be an expert in solvent-induced hearing loss, with any luck.

I want to, in a few minutes, go over what we expect to do in the next couple of months, the things that we've assigned ourselves, but I think we should spend a couple of minutes. And I'm thinking we don't have that much more, so that we can just continue to work and then adjourn, instead of taking a lunch break.

But I'd like to put on our agenda some reflection, a little bit of time to reflect on the workings of the Board and whether there are some alternative structure, alternative means, or additional communication that would improve the functioning of the Board. So this is a moment when, if you have suggestions on how we work and how we can be more effective, then let's discuss them.

So while you're thinking, I would note that this Board has not developed committees per se. The previous board did, and those committees

had meetings, telephone meetings that were open to public.

We have a single working group with this board, which is on the Parkinson-related disorder, which we intend to continue. There's no particular reason why those discussions wouldn't be open to the public as a committee, except for the fact that it may just prolong the process. But we should discuss that, I guess. Dr. Dement?

MEMBER DEMENT: I guess we had a working group on the OHQ as well. I think the process of having that working group report back to the Board anything that's at least for the discussion development, I think that works reasonably well, other than having standing committees, per se.

CHAIR MARKOWITZ: In the past, working groups -- we have not scheduled them. The Federal Register had the six weeks' advanced notice. It gave us a lot of flexibility. On the downside, it meant that the public didn't have as much access to the discussions. What do we want to do about that?

MEMBER DEMENT: This is John again. I guess, unless there are some objections from the public, we could change where we are. I think it gives us more flexibility to have a meeting to exchange information and work on issues; no decisions are made, obviously. They're just working groups to assemble data and facts to present to the Board for discussion.

And those discussions, whatever the working groups bring forward, need to be open to the public. I think it works reasonably well unless there are some objections to it.

CHAIR MARKOWITZ: The committees had broader domains. Working groups are really very targeted, task-specific, and it frankly is helpful to be able to have the flexibility of having those discussions on a more frequent basis without scheduling them two months ahead of time.

I'm thinking about Parkinson's disease, for instance. That group could easily make excellent progress over the next period of time, and then have a discussion.

1 So unless there's a big objection, I 2 would agree that for these two working groups, we continue them as they are, but be thinking, first 3 4 of all, the logic -- the discussion that is had 5 in those working groups be brought both to the full 6 Board, which has the benefit of the full Board being 7 involved, but also has the benefit of the public having access to the thinking that comes out of 8 9 those working groups. 10 I think that would address the issue 11 public access adequately. But that's Silver? 12 opinion. Dr. 13 When a meeting is MEMBER SILVER: announced in the Federal Register, our DFO and 14 staff make all 15 assorted sure that relevant 16 documents get posted on the web. I'm happy with the working group arrangement, but to strike a 17 balance with public transparency, let's just try 18 19 to be scrupulous about posting any and all documents that don't contain PII. 20 Our review sheet, for example, for the 21

claims and any other things that the working groups

develop before the next full Board meeting. 1 2 CHAIR MARKOWITZ: That's reasonable. I mean, in the Parkinson's disease, some of those 3 4 documents were published articles, and actually 5 we discussed that. We published the names of the 6 articles, but we couldn't publish the articles 7 because there are restrictions on those. But if we were, for instance, to come 8 up with a draft of an OHQ, yes, we could put that 9 10 online. As it stands now, it's not up to us to 11 come up with a new draft; it's actually up to the Department to look at the advice that we've given 12 13 them. Okay. So we'll go, then, as we've been 14 proceeding, and if we hear a lot of objections, 15 16 or if we, in our self-monitoring, think that we're engaging in discussions that the public ought to 17 have access to, then we'll change our way of 18 19 working, if that's all right. Any other aspects of the way the Board 20 works that you think need attention, could be 21

improved, aside from me getting the agenda out with

a little bit more notice? Dr. Friedman-Jimenez? 1 2 MEMBER FRIEDMAN-JIMENEZ: I want to make a request that I've made before, which is that 3 4 the medical records that are provided in PDF files 5 be provided as searchable PDFs, rather 6 bitmapped PDFs. It would save us a lot of time 7 in reviewing; it would probably save the CEs and CMCs a lot of time too. 8 It's an extra step, running it through 9 an optical character recognition program, but I 10 think it would really add to the efficiency of the 11 record review. 12 This is John. 13 MEMBER DEMENT: Some of these documents are barely readable, and some of 14 them are not readable at all. 15 16 MEMBER FRIEDMAN-JIMENEZ: Sure. MEMBER DEMENT: And so there's going 17 to be a lot of garbage that comes out of the optical 18 19 character reading. The other thing that I've experienced in optical character reading is that 20 sometimes you can't rely on the words that come 21 22 out of that thing.

1	So indexing of where things are in this
2	big file, what page number the OHQs or what page
3	number the CMC report is on IH report you
4	know, key pieces of information of where you can
5	find them, that would probably be of much greater
6	benefit to me. I can go in and read them and know
7	I'm reading the exact words, rather than maybe
8	something I have to verify.
9	MEMBER FRIEDMAN-JIMENEZ: So you're
10	saying having someone actually go through the file
11	and index where the different key sections are?
12	MEMBER DEMENT: Well, I
13	MEMBER FRIEDMAN-JIMENEZ: That would
14	work too, yes.
15	MEMBER DEMENT: Yes, maybe that's sort
16	of this you know, we've asked for some help to
17	do this. It would greatly cut down on our time
18	if that were done. You know, we could review the
19	cases much more quickly if that were done. That
20	wouldn't be a great burden, I don't think. It would
21	take some time on the individual's part.
22	But now, I would say that half the time

1 I spend on these cases was on my thumb, going through 2 them, back and forth, back and forth, trying to find something. That's a waste of time. 3 4 CHAIR MARKOWITZ: Okay. Let me ask, 5 actually, the Department: Communications from the 6 public come addressed to me. I'm assuming that 7 those are disseminated to the Board without my asking, assuming that the Department decides that 8 it's relevant to the Board's mission. 9 That's true? 10 MR. FITZGERALD: Yes, that is true. 11 CHAIR MARKOWITZ: Okay. 12 MR. FITZGERALD: I mean, what is 13 relevant to the Board's work is posted unless it involves individuals and things that are more 14 case-specific things. We make a determination and 15 16 make sure the information with the Board is not public because it's not necessarily prudent to 17 share some of the things. 18 19 But if it involves the Board's work or 20 -- I will also say, we also get correspondence and communication from to the Board that is not in their 21 22 It may go on to the program; those things area.

1	get referred to the program for a response.
2	MEMBER REDLICH: We can't hear you.
3	MR. FITZGERALD: I'm sorry; I don't
4	know why this microphone is giving me so much
5	trouble.
6	So basically, yes, we do post everything
7	that's relevant to the Board's work on the website;
8	however, sometimes we get information that's very
9	case-specific, and it's not prudent to share that
LO	information with the public.
L1	We may alert the Board to the issue.
L2	Whether or not it's in their scope, it could be
L3	contextual: here's some information that you
L4	should be aware of, but it doesn't necessarily
L5	reflect on what the Board's work is.
L6	Other times information and questions
L7	and correspondence comes in to the Board and to
L8	the Chair that's really not in the Board's purview,
L9	and we refer that on to the program usually, where
20	it's best responded to.
21	CHAIR MARKOWITZ: Okay. Dr.
22	Redlich, you wanted to

1	MEMBER REDLICH: Just a request. In
2	terms of our requests of information, do we know
3	when we will be getting the additional materials
4	that we requested such as the data and the other
5	cases? I think it would I understand if it takes
6	time or if certain of our requests may not be
7	reasonable in the form they were in, but it would
8	just be very helpful, and I think enable us to plan
9	our time better if we knew this will take us
10	approximately this much time, or could we get part
11	of the request, so we just know what the plan is?
12	MR. FITZGERALD: Yes, that was actually
13	one of the reasons we created the form, the request
14	process we created, so that there can be some
15	limited back and forth between the Board and the
16	program, refining the request to make sure that
17	we're doing the most efficient
18	MEMBER REDLICH: Okay.
19	MR. FITZGERALD: as possible. Or
20	if the data you were asking for may not be available
21	in the form you're requesting it in, but maybe

there's proxy data that we could provide or

1 something of that nature, and then determine what 2 the time frame would be for making that deliverable. MEMBER REDLICH: Okay. 3 4 CHAIR MARKOWITZ: Okay. So we expect 5 we'll probably have a telephone meeting in a few 6 months; probably three months, and then another 7 in-person meeting in about six months. I'll talk about that in a moment. 8 I just want to talk about what we're 9 10 going to do in the next three months, so there's some understanding. I have a list; I think maybe 11 it's a full list, not quite sure. 12 13 But we're going to continue to look at the claims we have, and we're going to develop, 14 each and every one of us, a list of concerns about 15 16 the claims, and at the telephone meeting, we're going to try to identify a set of common concerns 17 that appear across claims. 18 19 If, in the interim, we aggregate, we can discuss them as to what the most sensible 20 organization of them is going to be. 21 So at the next telephone meeting, then, 22

we will probably not spend a lot of time reviewing claims, you'll be happy to know, but we will be shooting for a list of, however long it is, of issues that are of concern.

And that's going to help us when we have the opportunity to conduct a larger, systematic evaluation of the claims.

Two working groups: I think the occupational health questionnaire, I think it's a question of just waiting for DOL to respond to the recommendation that we made at the February 28th meeting. And there were two other recommendations we made, one on asthma, one on asbestos, and we're waiting on response to those as well.

On the Parkinson's disease working group, great work so far. Clearly, we can make additional progress on both -- there are four aspects to DOL's request, but they really focus on the diagnosis, advice on the diagnosis, and advice on causation. So I think we can continue to make progress.

1	Now, that working group right now is
2	Dr. Mikulski, Ms. Pope, and Mr. Mahs, right?
3	Okay. So I'll join that what's that? And I'll
4	join that working group.
5	But if anyone else wants to participate,
6	you're welcome. If you want to think about it,
7	you can join next week too; it will remain open.
8	We're going to get, I think, an update
9	on the status of the approved recommendations from
10	the Department. I think this was a dashboard
11	issue. Our hope is to get an update and then a
12	periodic update automatically, not by request when
13	we think about it.
14	And finally from this meeting,
15	hopefully Carrie has developed a list of some action
16	items which I will review and circulate to see if
17	we've gotten it right, see if we've left anything
18	off and the like.
19	Are there any other issues that we have
20	promised to look at over the next period of time
21	that have come out of this meeting? Yes.
22	MEMBER POPE: The replacement of Dr.

1 Cassano. 2 CHAIR MARKOWITZ: Yes. Well, we've been told that pretty soon, the announcement will 3 4 come out in the Federal Register, and then we tell 5 all our friends and get them to apply, those who 6 are still talking to us. And I think, Doug, you said yesterday 7 that we expected appointment by someone during the 8 Hopefully, it will be by the time we have 9 summer. 10 the telephone meeting so they can participate, but if not, hopefully several months before the next 11 in-person meeting so they can get oriented. 12 13 Any additional comments, questions? 14 Any you want to say? 15 MR. FITZGERALD: No. I just want to 16 thank the Board for all their work. We've covered a lot of territory over the last day and a half, 17 and I appreciate all your efforts. 18 19 I thank the public for participating and coming here to listen to deliberations, and 20 I think the SIDEM; the contract staff here is really 21

with

the

job

great

doing

22

and

logistics

1 coordinating all the travel, meeting set-up and everything else, and Carrie Rhoads, my alternate 2 DFO. 3 4 CHAIR MARKOWITZ: And I want to echo 5 those thanks. Kevin becomes, next week, the most 6 popular person, because that's the person we submit 7 our reimbursement form to, and if you don't submit it, he'll remind you. So don't worry about that; 8 he's very good about that. 9 10 The next in-person meeting: the way we select the location has generally been where 11 -- who's next on the list in terms of the number 12 13 of claims or claimants? We try to hit that 14 qeographic area. I haven't looked at the list lately, 15 16 so I don't know. NTS? All right. Okay. But we're going to base that decision on objective data, 17 not lack of data. 18 19 And the telephone meeting at three months arrives towards the end of July. 20 have a bit of a challenge, because of people's 21 different schedules, scheduling that, so we'll

1	trying to schedule that sooner rather than later.
2	But be responsive to the request to schedule that,
3	because we may have to go back and forth a bit.
4	We'll try to be as inclusive as we can. Mr. Tebay?
5	MEMBER TEBAY: Could we add to the
6	agenda to revisit the borderline test results for
7	beryllium sensitization or the diagnostic
8	criteria? Before, when we did that, it was based
9	on two borderline tests. They're utilizing three
10	borderline tests. I think we could just add it
11	to the agenda as a reminder to discuss, rather than
12	but I would appreciate it if we could do that.
13	CHAIR MARKOWITZ: Yes, that's fine,
14	particularly if DOL has anything new to add, because
15	we've made that recommendation, and they haven't
16	accepted it, based on the statute. But we should
17	keep it on the agenda when we can.
18	MEMBER TEBAY: Thank you.
19	CHAIR MARKOWITZ: So I guess the
20	meeting's adjourned. Thank you.
21	(Whereupon, the above-entitled matter
22	went off the record at 12:13 p.m.)