## **Official Transcript of Proceedings**

## NUCLEAR REGULATORY COMMISSION

Title: Advisory Committee on Reactor Safeguards

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1	UNITED STATES OF AMERICA
2	NUCLEAR REGULATORY COMMISSION
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4	ADVISORY COMMITTEE ON REACTOR SAFEGUARDS (ACRS)
5	$514^{\text{TH}}$ MEETING
6	+ + + +
7	FRIDAY,
8	July 9, 2004
9	+ + + +
10	The meeting was convened in Room T-2B3
11	of Two White Flint North, 11545 Rockville Pike,
12	Rockville, Maryland, at 8:30 a.m., Dr. Mario V.
13	Bonaca, Chairman, presiding.
14	MEMBERS PRESENT:
15	MARIO V. BONACA ACRS Chairman
16	GRAHAM B. WALLIS Vice-Chairman
17	F. PETER FORD Member
18	THOMAS S. KRESS Member
19	GRAHAM M. LEITCH Member
20	DANA A. POWERS Member
21	VICTOR H. RANSOM Member
22	STEPHEN L. ROSEN Member-at-Large
23	WILLIAM J. SHACK Member
24	JOHN D. SIEBER Member
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1	ACRS STAFF PRESENT:
2	JOHN T. LARKINS Executive Director, ACRS/ACNW
3	SAM DURAISWAMY Technical Assistant,
4	ACRS/ACNW, Designated Federal Official
5	Ralph Caruso ACRS Staff
6	David Cullison NRR Staff
7	John Hannon NRR Staff
8	Robert Elliot NRR Staff
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1	P-R-O-C-E-E-D-I-N-G-S
2	8:29 a.m.
3	CHAIRMAN BONACA: Good morning. We are
4	here for the purpose of writing our reports. And
5	yesterday we heard that on the final generic letter
6	potentially to the pre-blockage of emergency
7	circulation building's design basis accidents of
8	PWR. There have been additional changes to the
9	generic letter. And so this is not an official
10	meeting in the sense of we simply, we want to
11	have some information regarding these changes so
12	that we can make a decision whether or not we're
13	going to write the report ourselves or not, at this
14	time.
15	So I would like to turn to the staff and
16	see if you can give us some insights.
17	MR. HANNON: Thank you. My name's John
18	Hannon, I'm plant systems branch chief. And I have
19	Dave Cullison and Rob Elliot from the staff with me
20	this morning.
21	Let me start out by reminding that the
22	outcome we're seeking with this generic letter is to
23	assure a long-term core cooling capability to PWRs,
24	to make sure their performance capability is
25	adequate for that. And what you've been witnessing,

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1	both in the subcommittee meeting and the full
2	committee meeting, is the staff is being exercised
3	over a process issue: what we can or cannot put into
4	a generic letter. You've seen two different
5	versions of the letter, and I'm sure it appears
6	frustrating to you that we've been whipsawed over
7	this. And it's doubly frustrating to the staff.
8	But just to recap. The public comment
9	version was geared for establishing compliance with
10	50.46, and it was an information request. Much of
11	the public comment we got was suggesting that, look,
12	let's call a spade a spade. This is a back-fit.
13	Just tell us what you want us to do. So it was
14	based on much of that comment that we revised the
15	generic letter and came to the subcommittee with
16	that version that was written towards based on a
17	back-fit, and asking for action to be taken. And
18	then during subsequent review by our OGC staff, they
19	concluded it was too much like an order. And we
20	wound up modifying it, taking it back closer to the
21	original version which was an information request
22	back to a compliance orientation. And so that's
23	what we briefed the full committee, on that version.
24	Now, we have I have Rob Elliot here.
25	We've uncovered a policy paper that was written back

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1	in 1999 that's just been passed out which tends to
2	provide us some flexibility. It seems to us that it
3	provides us some flexibility that would enable us to
4	do either one of these two approaches. And we have
5	that under active consideration right now.
6	DR. SHACK: I would point out the
7	website on generic communications, generic letters,
8	says that the first purpose it gives to a generic
9	letter is to, you know, you can request
10	calculations.
11	MR. HANNON: Understood. If I could,
12	I'd like to let Rob try to explain the origin of
13	that and how we came to where we are.
14	MR. ELLIOT: This is Rob Elliot.
15	Basically, back in 1999 we decided to put out some
16	changes to our generic communication process in
17	order to clarify them. There had been a number of
18	stakeholder comments. There was confusion about
19	differences between a bulletin and generic letter.
20	I'm sure many committee members probably remember
21	that because we probably came to you all with this
22	paper when we wrote it. And at the time we sent a
23	paper to the Commission where we indicated what the
24	purposes of each generic communication were, and the
25	process we would use for putting out generic

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1	communications.
2	And for some reason there seems to be
3	some confusion about whether or not requesting
4	action in a generic letter is something we can do.
5	But if you read the Commission paper that we handed
6	out to all the members, you'll see that it's clearly
7	stated on Page 3, where it talks about bulletins and
8	generic letters that both of those can request
9	either action or information.
10	And what we said is we recognize that
11	even though we're requesting action, the industry
12	perceives requested action as the actual
13	implementation of a regulatory burden. And so we
14	committed at that time to performing a limited cost-
15	impact analysis as part of doing the back-fit
16	analysis for the generic letter or bulletin. So we
17	revised our process a little bit, but we reserved
18	the right to request action, if need be, through
19	generic letters.
20	We took this to OGC yesterday, and we
21	were not able to get a decision from them as to
22	whether they agreed that we could use the generic
23	letter that we brought to the subcommittee. We are
24	tempted to go that way. They asked for a little bit
25	more time to be able to review both the information

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1	we gave them on the policy paper and what they had
2	commented on before, before they would come up with
3	a decision. And so we felt it necessary to come to
4	you, explain to you what we are trying to do so that
5	you wouldn't be surprised.
б	John?
7	MR. HANNON: So I'd like Dave now, if he
8	would, to try to explain our next steps and where we
9	would like to go with this.
10	MR. CULLISON: Yes, I'm Dave Cullison
11	from the staff. What we're trying to I guess get
12	from you all today is since we don't have a
13	definitive OGC decision on whether we can ask for
14	action or not is that the committee make a
15	recommendation on which path we should go, whether
16	the information-only path, or the required action
17	path, if possible. Requested action path, that's
18	correct. We can't require an action.
19	DR. WALLIS: These are procedural
20	matters that we don't usually advise about.
21	Process. We don't usually give advice about this
22	kind of thing.
23	CHAIRMAN BONACA: You are going to
24	lawyers to get advice on how in fact you are doing.
25	It is for us to comment to that to make sure you get

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1	it.
2	MR. HANNON: As an alternative to
3	actually commenting on which our preferred path
4	would be, what we would be seeking is a letter from
5	you to tell us to go ahead and issue one form in
6	this generic letter. Because we need to do that in
7	order to move forward towards a resolution.
8	CHAIRMAN BONACA: I think something has
9	to be issued. Something has to move. I understand
10	the concern is the issue of compliance. Compliance
11	with what? Compliance with the current
12	requirements. They are required to be complied
13	with. Or compliance with the intent, which is the
14	way of providing cooling. And now I certainly think
15	that the committee wants to see the units complying
16	with intent. And so there has to be some movement.
17	But I'm puzzled because you may remember during the
18	presentation, at the end we had a statement from NEI
19	that took exception with the generic letter. They
20	seemed to prefer the regional approach. I don't
21	think they were attempting to simply get compliance
22	with current requirements.
23	Do you have some insights on why they
24	would think that way?
25	MR. HANNON: Well, it's a process issue.

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1 It turns on whether we're going to be asking a 2 licensee to put themselves on report or not. And 3 that's objectionable. And we can deal with that as 4 a process issue. There are ways we can enable a 5 licensee to make changes to their plant while they are still are obliged to comply with their current 6 7 licensing basis. And once those changes have been implemented, they can adopt a new licensing basis, 8 and wouldn't have to address that compliance 9 question. And that was the way the subcommittee 10 11 version was geared, to not have to have. And that's 12 why, you may recall, that NEI didn't object to that version of the letter. 13 14 CHAIRMAN BONACA: Okav. The other 15 question I have, the second question I have is what's the benefit of issuing a letter now when 16 17 there is no NEI guidance behind at the same time. Ι mean, the letter is going to sit there. 18 19 MR. HANNON: That's a valid question. 20 And our approach has been to get this information 21 out as soon as we had it available, as soon as it 22 was ready. We've always had it on the schedule to 23 be published in August, late August, I think it's 24 the 23rd of August. And recognizing that the 25 methodology review would be completed at a later

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1	time. It's now scheduled for the end of September.
2	And we took the comments that we got
3	during the public comment period, which suggested
4	that we should key the response date on the actual
5	publication of the SE methodology. And we're going
б	to do that, and that has been built in to both
7	versions of the letter.
8	MR. ROSEN: I guess you really didn't
9	answer that question, except that's your answer
10	was `That's the way we want to do it. We want to
11	get it out as soon as we can,' even though you've
12	acknowledged that there's nothing licensees can do
13	with it till they get the guidance. So it seems to
14	me impractical to ask licensees to respond through a
15	question and compliance when they don't have the
16	guidance to determine whether they're in compliance
17	or not.
18	MR. HANNON: Agree, and that's the
19	benefit of using the subcommittee version, because
20	it doesn't ask for the compliance question to be
21	addressed.
22	CHAIRMAN BONACA: And so in fact you're
23	requesting some information at a later date.
24	DR. WALLIS: So you want a kind of carte
25	blanche letter that says issue any kind of generic

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1	letter that you eventually are satisfied with?
2	MR. HANNON: We have two versions.
3	We're going to go with one of them.
4	DR. WALLIS: But it seems to me there
5	might well be a third one. The way this thing is
6	bouncing around, there might be some kind of
7	compromise which might not appeal to us at all. I
8	don't know. I can't tell. We don't know what it's
9	going to be. You're looking for approval of a
10	letter that we haven't really seen.
11	MR. ROSEN: Which we haven't seen any
12	public reaction to, obviously, because they can't
13	react to something they haven't seen either. You
14	know, I always take into account what the public and
15	the other stakeholders say before I would venture a
16	response. So here I'm going to be, as part of the
17	committee, asked to do that without any input from
18	stakeholders, people affected.
19	MR. HANNON: You have had input from the
20	stakeholders on the one version of the letter, the
21	one that had the compliance orientation with an
22	information request. And that's why we came to the
23	subcommittee with a different version, the one that
24	addressed those two issues. So that's our preferred
25	pathway. And there may be some modifications, but I

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1	would not consider them to be significant. That
2	comes from CRGR. Because we are going to have to
3	take that version to CRGR for approval. So they may
4	have some wordsmithing, but I wouldn't expect it to
5	be significant.
6	DR. WALLIS: Well, the original draft
7	that I wrote for the letter that the committee might
8	have approved, I did say that the subcommittee
9	thought that the version that we saw was an
10	improvement, because it actually specified actions,
11	and it asked for specific calculations. And that
12	all seems to have disappeared from the next version.
13	We rather liked the idea of saying thou shalt do
14	these specific things, and use some guidance, and
15	come up with some mechanistic predictions and so on.
16	That seemed to disappear.
17	MR. HANNON: That's our preferred path
18	now.
19	DR. WALLIS: That's our preferred path
20	too, if we got the chance to influence the outcome.
21	MR. ELLIOT: This is Rob Elliot again.
22	But I guess that gets back to what we originally
23	asked, is whether you could endorse a specific path
24	or both paths. And it's my understanding the
25	committee doesn't normally comment on that. We

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13 1 would welcome you taking an exception to the rule 2 though. 3 DR. WALLIS: Well, from the technical 4 point of view, it's nice to say some technical 5 requests rather than this vague thing about go out and show compliance in some way or other. And I 6 7 think NEI liked the idea of using this -- I mean, this whole thing has driven their effort, which 8 9 seems to be a good faith one, to produce a guidance aimed at responding to the subcommittee version of 10 11 the letter. 12 We don't have anything MR. HANNON: further, unless you have any questions or comments. 13 14 DR. WALLIS: You haven't been to CRGR 15 yet? 16 MR. HANNON: That's correct. We're scheduled. 17 They may do something else 18 DR. WALLIS: with this letter. 19 20 MR. HANNON: We want to take the 21 subcommittee version to CRGR. We're scheduled to do 22 that on the tenth of August. CHAIRMAN BONACA: We don't know what's 23 24 going to be in this letter. 25 DR. KRESS: I think we can assume it's

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1	pretty much like the subcommittee letter.
2	MR. ROSEN: How about the letter you
3	just gave us? It starts with
4	DR. KRESS: It's pretty much like the
5	subcommittee letter.
6	DR. SHACK: If OGC will agree to it.
7	DR. KRESS: Well, that's
8	MR. ELLIOT: One thing I forgot to
9	mention is this policy paper that we sent forward in
10	1999 was concurred in by Karen Cyr of OGC. So we
11	believe we're conforming with it, it's just a matter
12	of convincing the working level who want to consult
13	with their management that this is appropriate. I
14	can't say absolutely we think we're going to end up
15	with that product, but we think we have a strong
16	basis for saying that the subcommittee product is
17	appropriate and consistent with policy.
18	DR. KRESS: What we could do if we like
19	that letter is say they should go forth with a
20	letter that calls for and spell out the things
21	that we think is going that letter will have.
22	DR. SHACK: We can't ask for them to do
23	something that's illegal.
24	DR. KRESS: It does look like we can
25	ask. If it turns out to be illegal, why it's just -

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1	- can do anything they want to.
2	MR. SIEBER: When are they going to
3	issue it?
4	CHAIRMAN BONACA: Earlier in the year we
5	wrote a letter that said we have to move on, we have
6	to issue a document.
7	DR. WALLIS: Last year.
8	CHAIRMAN BONACA: Provide guidance, and
9	you have to do this, because it has to be fixed.
10	And consider all these elements. So already we came
11	up and said.
12	DR. KRESS: I think our letter should be
13	consistent with that.
14	CHAIRMAN BONACA: Why should we repeat
15	that? I mean, I'm afraid that in repeating that we
16	get caught into the discussion that is taking place
17	right now, which is totally not to do with the
18	elements of technical issues. Really it is to do
19	with compliance, what we should be doing. My only
20	fear is that we step in a minefield with some
21	opinions.
22	DR. KRESS: Well, I take a little
23	different view. This issue has significant safety
24	implications. We're allowed to weigh in on
25	significant safety implications, and I think we

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<pre>1 ought to tell them what we think ought to be done. 2 In spite of the fact there are process questions an 3 legal questions. 4 DR. SHACK: They're going to do the sam 5 thing in either case. The question is whether it's 6 a matter of compliance, or it's essentially a case 7 of verbalizing the design basis. Now, NEI has a 8 very strong reason for not making it a compliance 9 issue. 10 DR. KRESS: Well, I think we ought to 11 skirt around that. You know, that's for these guys</pre>	e
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11 skirt around that. You know, that's for these guys	
12 to decide, for them to iron out. But in our letter	
13 we can say we think we ought to ask for whatever we	
14 think we ought to ask for. In my opinion it would	
15 be pretty much like our letter that we had, the	
16 first letter we had.	
17 CHAIRMAN BONACA: Before we changed	
18 anything. Which said this letter is a better	
19 version than the earlier version.	
20 DR. KRESS: Yes.	
21 MR. ROSEN: What bothers me about this	
22 is it starts off by saying this is the request that	
23 addressees submit information to confirm compliance	•
24 It's not bashful about that. It says.	
25 DR. WALLIS: They're going to change	

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1that. They're going to go back to the old version.2MR. ROSEN: They are going to change3that? So what are we I don't know what I'm4supposed to be sending you a letter approving then.5MR. SIEBER: My question is6MR. ROSEN: I thought this was what we7were talking about.8MR. HANNON: That's the version we9shared with the full committee, but the previous10version we shared with the subcommittee.11MR. ROSEN: Now I don't know what I'm12supposed to be13MR. SIEBER: My question is just one of14form. If you don't require compliance, doesn't that15de facto make it a back-fit? Where a licensee could16then say I don't want to do that unless you go17through the cost-benefit business.18MR. HANNON: Well, that is correct. And19the version that we shared with the subcommittee did20treat it as a back-fit.21MR. ROSEN: Well, I guess I wasn't at22the subcommittee meeting, so I'm not sure whether23MR. SIEBER: I wasn't either. On the24other hand, if you get into that, I'm sure that if25it is a \$3 or \$4 million change for each licensee		17
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	18
1	that they'll say it's a back-fit. You know, do the
2	cost-benefit. And then you'll be stuck.
3	DR. KRESS: No, it's a compliance back-
4	fit. You don't have to do a cost-benefit on that.
5	MR. SIEBER: If it's a compliance issue,
б	you ought to say it.
7	MR. ELLIOT: In 1999, we committed to
8	even if we were using the compliance back-fit, to do
9	a limited cost-benefit analysis. So we are doing
10	that analysis.
11	DR. KRESS: But you just do it for
12	information.
13	MR. ELLIOT: Right.
14	DR. KRESS: But, you know, clearly this
15	is a question of compliance with the spirit of that
16	law. And I would call it a compliance back-fit.
17	The spirit of the law is clear. You've got to
18	provide long-term cooling. And I don't care if the
19	guidance they had before is wrong or what. That's
20	the spirit of the law.
21	MR. SIEBER: Well, the original guidance
22	
23	DR. SHACK: The way they ask the
24	question is very different in the two generic
25	letters. Any action that they require the licensee

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	19
1	to take will be a compliance back-fit, but the way
2	they ask the question is quite different. Which is
3	why NEI
4	DR. KRESS: They're asking whether or
5	not if they're in compliance.
6	MR. ROSEN: Does OGC agree that this is
7	a compliance issue?
8	MR. CULLISON: This is Dave Cullison.
9	OGC does agree that it is a compliance issue, at
10	least on the high level, like we're discussing.
11	There is a requirement for long-term cooling. The
12	question here is that whether or not when they
13	perform an analysis, if they perform an analysis
14	under the information letter, that they have to
15	compare their current configuration against this new
16	information, and determine if they're in compliance
17	today. Or can they do this analysis, determine what
18	their configuration should look like, implement
19	those, update the licensing basis, and then
20	determine compliance. And it's a matter of I guess
21	you could say timing, when they determine when we
22	ask the compliance question.
23	The letter that was sent out for
24	comment, the draft comment, which the letter we
25	presented to the full committee, they're very

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	21
1	called a back-fit? Why doesn't the staff just
2	follow the back-fit process?
3	MR. CULLISON: This is a compliance
4	exception in that, as with discussions on OGC on
5	this issue, that this information it's one of
6	these issues where if we'd known this information
7	when we issued the original guidance, we would have
8	included it.
9	DR. KRESS: We shouldn't constrain the
10	Commission and the Agency to fix things that are
11	basically safety problems that don't meet the spirit
12	of the law just because they gave some bad guidance
13	at one time. If they know better now, well they
14	ought to be able to go back and fix it without going
15	through the cost-benefit issue of a back-fit. If
16	it's a problem, fix it.
17	MR. ROSEN: As long as the licensees
18	know how to fix it, that's fine.
19	DR. KRESS: Well, right now I would say
20	the letter ought to say
21	DR. POWERS: Why is it the NRC's
22	responsibility to tell them how to fix it?
23	MR. ROSEN: Because they took on the
24	burden originally to tell them how to design.
25	MR. SIEBER: No.

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	22
1	DR. KRESS: No. And of course what
2	they'll do is use the NEI guidance, and that'll
3	become the ad hoc way to fix this. That's all right
4	with me. I just want to get going with something.
5	MR. SIEBER: I'm still struggling with -
6	- I even struggle with the 60 days after the
7	guidance appears clause in the letter. Why is it
8	the NRC's responsibility?
9	MR. LARKINS: Licensees are responsible
10	for showing compliance with 50.46.
11	MR. SIEBER: That's exactly right.
12	MR. LARKINS: The staff gave them one
13	way to do it, okay, and they generally followed
14	that. Some of them did not, not necessarily, but
15	most of them did. And now you're giving them
16	another way to do it. If you think that the way
17	that they have done it is incorrect, then I would
18	think you'd be in an enforcements phase. What do
19	the enforcement people say about this?
20	MR. SIEBER: I think that's raising a
21	red herring here. I think you have discovered
22	something. It says what everybody was thinking in
23	the past is probably not correct. Now we know
24	something more. Not as much as we would like to
25	know, but we know something more. Does it seriously

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	23
1	affect your plant or not. Okay? Now, Question
2	Number 1. If it does affect your plant, what can
3	you do about it? Question, Number 2. Question
4	Number 3 is what will you do about it.
5	DR. KRESS: I think the letter ought to
6	ask for that
7	MR. SIEBER: And I'd say do it right
8	now. And if any
9	MR. LARKINS: 50.46 says what you have
10	to do.
11	MR. SIEBER: And what it says is you've
12	got to assure long-term cooling. Okay, now the
13	question is by asking them can you, in light of the
14	additional information we have.
15	MR. LARKINS: Right.
16	DR. WALLIS: 50.46 says you have to take
17	immediately take the appropriate action. It's
18	very clear.
19	DR. KRESS: I don't know about the
20	immediate. I don't think this is an urgent issue.
21	MR. LARKINS: If a vendor doing a LOCA
22	calculation discovers that the peak cladding
23	temperature is 2202 degrees, and somebody signs off
24	and says, yes, this is the number, a licensee is
25	obliged to take immediate action to bring that plant

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1	in compliance with the 2200 criteria.
2	DR. POWERS: Okay, now tell me what
3	immediate is. Does that mean within one nanosecond?
4	No, it does not. Does that mean within one day?
5	MR. LARKINS: Listen to the plant
6	person. What does he say?
7	MR. SIEBER: It's one hour.
8	MR. LARKINS: One hour.
9	MR. ROSEN: That's a tech spec word
10	"immediate".
11	MR. SIEBER: You have to reduce power.
12	MR. LARKINS: What they usually do, the
13	vendors, is they have something else that they can
14	either Well, what I'm telling you is, I'm telling
15	you what they do when they go over 2200 degrees.
16	DR. POWERS: There's no question of 2200
17	degrees right now. It's a question of long-term
18	cooling right now. Stay on the topic, please.
19	Please stay on the topic.
20	MR. LARKINS: Long-term cooling is the
21	last of the five criteria in 50.46. The criteria in
22	50.46 are 2200, 17 percent, hydrogen generation
23	DR. POWERS: Would you please
24	MR. LARKINS: and long-term cooling.
25	DR. POWERS: I'm not going to talk with

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	25
1	you anymore if you're going to be off the topic.
2	DR. WALLIS: Well, he's using them as
3	examples of other regulations.
4	DR. POWERS: I think it's an
5	inappropriate example, okay? Now, you've discovered
6	that you don't have the capability of long-term
7	cooling. Okay. The request is that you immediately
8	fix it. We have a risk-informed regulatory machine.
9	That means based on risk considerations, we can
10	define what immediate is. And we've done so,
11	effectively.
12	DR. KRESS: I think that's the way to
13	view this thing.
14	DR. WALLIS: But there's I mean you
15	haven't used much risk information in resolving
16	this.
17	DR. KRESS: Well, all we know is it's
18	not that big of a deal from the standpoint
19	DR. WALLIS: Well, it's all hearsay. I
20	mean, the bulletin said either show that you're in
21	compliance or take other actions. And only one
22	plant fixed the sump. The others took a lot of
23	actions, which were quite varied I understand.
24	MR. SIEBER: How do you know one plant
25	took?

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26 1 DR. WALLIS: Now, what was the measure 2 of safety achieved by all these other actions? Did 3 it make the problem go away or not? We asked for 4 specific measures of success of these other actions 5 at the subcommittee meeting, and it seems to be a qualitative thing, rather than saying, yes, they 6 7 reduced the CDF to the point where it doesn't 8 matter, or something like that. DR. SHACK: No, they reduced it to the 9 10 point that they think they can wait until 2007. 11 DR. WALLIS: Well, it doesn't matter 12 It doesn't matter that we don't take today, then. immediate action today. They all did that? 13 14 DR. SHACK: All licensees have responded 15 with their actions. We have them under review. They're still under review. 16 DR. WALLIS: 17 So we don't know what --MR. HANNON: We don't know the full 18 19 But we do know that of the plants that we extent. 20 have reviewed, they have taken action. 21 DR. KRESS: A lot of those actions are 22 going to be difficult in a PRA space. 23 The original imperative DR. WALLIS: came from the suggestion that the CDF was really 24 25 quite hot. And it was these other actions that made

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1	the problem seem less immediate.
2	DR. KRESS: If we still thought it was
3	that high, we would classify this as urgent.
4	DR. WALLIS: There would be an order or
5	something.
б	DR. KRESS: I think that we've decided
7	is not correct information. And if the CDF effect
8	is such that we don't have to call it urgent, we can
9	do it on a measured basis. But I agree with Dana's
10	line of thinking on this.
11	DR. WALLIS: It's the line of thinking
12	that a member of the public might have, essentially.
13	A sensible member of the public.
14	DR. KRESS: It might be the view of a
15	member of the public, but it seems to me like the
16	appropriate view.
17	DR. WALLIS: I had a question about
18	50.54(f), which is cited in the second version here.
19	In this 1999 document you've just given us, it says
20	that generic letters will typically not invoke
21	50.54(f) unless the NRC has been unable to obtain
22	needed information through other means. So if you
23	invoke it, licensees are going to come back and say
24	why are you doing this.
25	MR. CULLISON: We did have some public

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1	comments on the 50.54(f) from the NUBAR, the
2	industry's tactic group. And when we went through
3	the OGC this time with the letter, we originally
4	went up to them without a reference to 50.54(f), and
5	their recommendation was to put it back in to make
6	our request stronger.
7	And all these legal issues, we are
8	deferring to OGC.
9	MR. ROSEN: Well, that's not a legal
10	issue, that's a judgment issue. I mean, to make it
11	stronger is not a legal issue. Just their judgment
12	is it should be stronger. I don't agree that's a
13	legal.
14	DR. KRESS: Well, you're not always
15	you're not constrained to actually do what the
16	public comments ask for. You just take them into
17	consideration. Then you do what you think is the
18	right regulatory approach. You know, you can take
19	these into consideration and see if it's the right
20	thing to do, but I don't think you're constrained to
21	do what the public comments say you have to.
22	DR. WALLIS: I guess we could write you
23	a carte blanche letter and say we're in favor of
24	issuing a generic letter. We've seen various
25	versions. Any one of these would be acceptable to

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1	us in order to get things rolling. Thank you very
2	much.
3	DR. POWERS: You're thinking very highly
4	of yourself today.
5	DR. WALLIS: A one-page carte blanche.
6	CHAIRMAN BONACA: I would be willing to
7	consider a letter if we make the point that there
8	are different versions being evaluated right now of
9	the communications, and that we are not making a
10	judgment on the way it's going to happen. Simply
11	that we believe there should be a communication,
12	follow the appropriate guidance, so that the
13	licensees can move on and fix this problem.
14	DR. KRESS: I think I would be more
15	specific on what the letter ought to ask for. It
16	ought to ask for the things Dana mentioned. It
17	ought to ask for them to make an evaluation whether
18	they're in compliance with the spirit of the law.
19	And if not, tell us what they're going to do about
20	it, and when. And I'm sure they'll use the NEI.
21	MR. ROSEN: The bulletin's already done
22	that.
23	DR. WALLIS: That's why we said the
24	second version was good.
25	DR. KRESS: Well, why do we need another

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1	letter?
2	DR. WALLIS: It's appropriate for the
3	purpose of gathering information to confirm
4	compliance.
5	DR. SHACK: Because you don't want to
6	rely on those compensatory actions forever. You
7	know, you've reduced your risk to some sort of
8	manageable state that doesn't require an immediate
9	shutdown or downgrading of all PWRs, but you want to
10	take some the further action.
11	DR. KRESS: I don't see that that says
12	are you in compliance with the spirit of the law.
13	They're only going to do that if they do the NEI
14	calculation.
15	MR. SIEBER: Which are undefined.
16	DR. KRESS: That's the part I'm saying
17	they have to decide whether they're in compliance
18	with the spirit of the law according to the NEI
19	methodology. And that's what I would ask for. If
20	they are, okay. If they're not, fix it, and tell us
21	how you're going to fix it. That's different than
22	the bulletin, I think.
23	DR. SHACK: Well, that's different than
24	the bulletin, yes.
25	DR. KRESS: I think that's what Dana was

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1	asking.
2	DR. SHACK: He's happy with the second
3	version of the letter, the original version.
4	DR. KRESS: I think that's the one I'm
5	happy with.
6	DR. WALLIS: We could take our "yes"
7	letter, and we could simply rewrite the paragraph
8	which says that the latest version we've seen is
9	okay, and we could also say that the other version
10	was okay, and in fact we prefer the way in which it
11	specifically mentions the calculations to be
12	performed. Then the "yes" letter would go simply as
13	it is, essentially, with a few changes.
14	DR. KRESS: Yes, I think that would be
15	good.
16	DR. FORD: In this 1999 thing, it says
17	here under generic letters, "Generic letters will
18	not be issued without prior staff interaction with
19	industry and public." Do I assume, therefore, that
20	the version that we saw yesterday, will it be in
21	fact discussed with NEI?
22	DR. SHACK: It looks like the one that
23	went out for public comment. It's certainly been
24	discussed.
25	DR. FORD: No, the one that we saw at

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1	the Thermal Hydraulics Subcommittee meeting had been
2	discussed.
3	DR. SHACK: No.
4	DR. FORD: Yes?
5	MR. ELLIOT: That was a version that was
6	changed in response to public comments.
7	DR. FORD: We received a copy of a
8	we've seen two before the one yesterday.
9	DR. SHACK: Right. Let's review
10	DR. FORD: Let me tell you what I
11	understand, Bill. The one that we saw and we
12	essentially we agreed upon technically, and which
13	the industry said "no problem" is the one that we
14	looked at in Thermal Hydraulics.
15	DR. WALLIS: It's this one that has the
16	blue lines and the red additions and things. That's
17	the one.
18	DR. SHACK: No. Prior to that.
19	DR. WALLIS: The one before. That was
20	like that.
21	DR. SHACK: Take all the blue lines and
22	additions out of that one.
23	DR. FORD: And that's the one
24	DR. WALLIS: That's right.
25	DR. FORD: that the public agreed

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1	with?
2	DR. SHACK: No, because that wasn't the
3	one that went out for public comment. That was
4	created in response to the public comment.
5	DR. FORD: Well, let me ask the
б	question. The one that we saw yesterday that we're
7	supposed to be pronouncing on has not there's
8	another factor. The one that we saw yesterday has
9	not been discussed with the public?
10	DR. SHACK: That's correct.
11	MR. ELLIOT: Not specifically.
12	DR. SHACK: Not specifically.
13	MR. ELLIOT: Though it's very similar to
14	the first version.
15	DR. FORD: But that's the one that Tony
16	Patrianni, they all went crackers over.
17	DR. SHACK: That's what Tom says. You
18	have to listen to the public comment.
19	DR. FORD: Was that meant to be the
20	public comment? He hadn't even seen it.
21	DR. SHACK: He had seen the original
22	version, which is the one they had the public
23	comment on, which was very much like that final one.
24	MR. ROSEN: That's what you say, but the
25	public hasn't had a chance to agree with that.

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1	MR. ELLIOT: I don't think the
2	Commission paper meant to imply that we would, every
3	time we changed the generic letter, try to go out
4	and get public comment again. I don't think that
5	was the intent.
6	DR. SHACK: It's an absolutely, invert
7	iterative process.
8	DR. WALLIS: It would go on forever.
9	MR. ELLIOT: It was meant to say, you
10	know, that we would solicit public comment on first
11	draft, and then we would revise it.
12	DR. WALLIS: Well, let me try something.
13	Suppose we wrote you a letter which said if you have
14	all these uncertainties, you really shouldn't send
15	this letter right now. You should wait, and you
16	should sort things out, and you should also wait
17	until the guidance has been approved, and then you
18	could have a nice package which can go, and it's all
19	clear what they have to do.
20	MR. ROSEN: Then we can see it, and so
21	can the public. And then we can
22	DR. WALLIS: Would that throw a wrench
23	into your work somehow?
24	MR. HANNON: It would just mean a delay
25	in the issuance of the

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	35
1	DR. WALLIS: Why is that a critical
2	thing?
3	MR. HANNON: Again, it
4	DR. WALLIS: You get whipped if you
5	don't meet a schedule or something?
6	MR. HANNON: We're attempting to get
7	this out of our off our plate so we can focus on
8	the methodology. As long as this is continuing to
9	bounce back and forth it's a diversion, and keeping
10	some of our staff occupied that could otherwise be
11	working on the methodology.
12	MR. ROSEN: So we're doing it to make
13	the staff's processes work better. And the balance
14	of that is it will create quite a bit of difficulty
15	in the industry because the question of compliance
16	comes into play.
17	MR. HANNON: Not if we issue the version
18	that was discussed with the subcommittee, which is
19	our preferred path.
20	MR. ROSEN: But that's the version I
21	haven't seen, I guess.
22	DR. WALLIS: You haven't seen that one?
23	MR. ROSEN: No.
24	DR. KRESS: Somebody get him a copy of
25	that.

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1	DR. SHACK: It was sent to you. Just
2	look in your packet.
3	DR. WALLIS: It's this one that has all
4	the lined out because they changed it.
5	DR. SHACK: No, it's not that one,
б	because I think
7	DR. WALLIS: No, it's the one before
8	that.
9	MR. ELLIOT: We can bring extra copies
10	of the second version.
11	DR. WALLIS: Well, you see the problem
12	my colleagues have is those who weren't at all the
13	meetings don't know what it is they might be
14	approving.
15	DR. FORD: I don't know. In terms of
16	There was a version issued which we discussed at the
17	Thermal Hydraulics meeting where the guys from NEI
18	said, yes, we are happy with this, and looked at all
19	the legal aspects, the technical aspects. Everybody
20	said fine. The thing we saw yesterday was not that
21	document. And that's what I'm concerned about.
22	DR. KRESS: That's because OGC didn't
23	say fine.
24	DR. FORD: Exactly. Exactly.
25	CHAIRMAN BONACA: My understanding was

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1	that this was the original letter.
2	DR. WALLIS: No, no, no. That is about
3	Version 5 that was lined out. We have seen several
4	things. It's evolved through this whole period.
5	The one in there is not the same.
6	DR. SHACK: I believe if you take all
7	the line-in/line-outs out of that one, you have the
8	subcommittee version.
9	DR. WALLIS: Do you?
10	MR. LARKINS: I'm not sure. Let me ask.
11	The comparative tech version is compared against the
12	subcommittee version or comparative against the
13	original version that went out for public comment?
14	MR. HANNON: The subcommittee version.
15	MR. LARKINS: Okay, that's good then.
16	Okay, yes.
17	MR. CULLISON: And I have a copy of the
18	subcommittee version that we can make copies for and
19	give you immediately.
20	MR. LARKINS: Then, if you take the
21	line-in and line-out version
22	MR. ROSEN: And read only the line-outs,
23	though.
24	MR. LARKINS: Read the line-outs, and
25	then you've got that version.

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1	MR. ROSEN: Read the line-outs, but not
2	the red inserts.
3	DR. FORD: Have you got a sample of that
4	version?
5	MR. ELLIOT: Well, we can get you a
6	clean version to make it easy on your eyes.
7	DR. WALLIS: I'm not sure we have
8	exactly that version, but it's very similar to the
9	subcommittee version.
10	DR. SHACK: I never verify exactness.
11	DR. WALLIS: There may be a semicolon.
12	MR. SIEBER: Yes, the more we discuss
13	it, the more I think
14	DR. SHACK: Does it have requested
15	action?
16	DR. WALLIS: Oh, that's the first
17	original. Request to perform evaluation is the
18	first. It should be in here. We didn't have this
19	at the time the package was put together.
20	DR. SHACK: Now, this is not the this
21	is the subcommittee.
22	MR. ELLIOT: Theron's making copies for
23	the committee right now, the subcommittee version.
24	No, we're going to bring it in to you in just a few
25	seconds here.

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1	MR. ROSEN: I get lots of questions
2	wrong even when I know what they are. It would help
3	a lot if I knew what the question was. Then I have
4	a chance of getting
5	DR. WALLIS: Well, let me make a
6	proposal here that we take and send the
7	CHAIRMAN BONACA: I think we should
8	first of all
9	DR. WALLIS: Discuss this?
10	CHAIRMAN BONACA: Yes, get off at
11	this point get off the record and thank the staff
12	for their input. I think we understand the
13	situation now. And then we'll make a decision. We
14	may have to take this off.
15	DR. WALLIS: John, may I ask you if
16	anything changes between now and eleven o'clock or
17	something that you come back?
18	(Laughter)
19	MR. HANNON: We'll be sure and do that.
20	Thank you, sir.
21	MR. SIEBER: Why wait so long, it could
22	flip four or five times.
23	MR. LARKINS: John, when do you expect
24	to hear from OGC?
25	MR. HANNON: They're working on it right

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	40
1	now. We got their attention last night, so
2	hopefully very shortly.
3	DR. WALLIS: So there might be new
4	information.
5	MR. ROSEN: There might be something
6	before eleven o'clock.
7	MR. SIEBER: have OGC come down here?
8	MR. HANNON: There was some interoffice
9	communication that had to take place, so I can't
10	predict on how long that's going to take. It may be
11	that they weren't able to get to Karen Cyr. And if
12	that's the case today, then we have to go into next
13	week.
14	MR. ROSEN: Did everybody find a copy of
15	the subcommittee version? They say they're making
16	copies of it.
17	MR. SIEBER: Well, you can get it off
18	the computer.
19	CHAIRMAN BONACA: Okay.
20	DR. WALLIS: Can also we hear from the
21	members who haven't expressed an opinion yet?
22	MR. SIEBER: He's handing out a
23	different letter.
24	CHAIRMAN BONACA: We can go off the
25	record. So we're going to go off the record. Thank

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		41
1	you.	
2	(Whereupon, the foregoing matter went	
3	off the record at 9:14 a.m.)	
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