PROPOSED TUBERCULOSIS AND BRUCELLOSIS REGULATORY FRAMEWORK

PUBLIC MEETING

PUBLIC MEETING, held on May 24, 2011, at the Renaissance Concourse Atlanta Airport Hotel, One Hartsfield Center Parkway, Atlanta, Georgia, commencing at 8:30 a.m., before Natalie Gail Sheckton, Court Reporter and Notary Public in and for the State of Georgia.

1	2 PUBLIC MEETING				
2	APPEARANCES:				
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4	DR. B	ILLY HA	RTMAN		
5	DR. S	COTT MA	RSHALL		
6	DR. L	EANN TH	OMAS		
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1	3 PUBLIC MEETING
2	PUBLIC MEETING
3	MAY 24, 2011
4	ANNOUNCER: Good morning. Since we
5	are such a small group, we thought we would
6	be a little more informal than our past
7	meetings. We thought we would have more of
8	a public conversation-style meeting today
9	since we have so little today.
0	So, if you will, move forward.
1	Well, thanks again for coming. I'm sorry
2	we're such a small crowd. but that is all

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rry we're such a small crowd, but that is all right. We'll get into a more fruitful discussion today.

My name is Lynn, and I work in the policy division for the USDA. I'm just here to moderate the meeting. We're going to take a little more of an informal structure today.

We do have two presentations we still want to go through because everybody came here to hear what our new proposed tuberculosis and brucellosis regulatory framework will be.

> We have two presentations. The

1	4 PUBLIC MEETING
2	first one will be from our working group
3	members, and the second will be from Dr.
4	Leann Thomas.
5	Just logistics very quickly. You
6	have a packet that is a packet of all the
7	presentations. If you would like to make a
8	formal statement for the record, that is also
9	contained inside the packet. Name tags are
10	inside your packet.
11	Restrooms are just outside across the
12	hall. We won't rotate today for breakout
13	stations, which you will see in the agenda;
14	we will stay here.
15	I'm trying to think if there is
16	anything else we need to cover. I think
17	that is it.
18	Our first presentation will be from
19	the working group perspective. Dr. Bill
20	Hartman and Dr. Scott Marshall will provide
21	that followed by Dr. Leann Thomas.
22	Let me go ahead and ask Dr. Hartman
23	and Dr. Marshall to come forward.

DR. HARTMAN: Well, good morning,

everybody. Good morning to a group we have

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today here to talk about this. We're here as a public meeting to locate the regulatory framework for brucellosis and tuberculosis.

The first thing I want to start out by saying is I really support this framework. We have had TB -- and I have talked to a couple of you already -- in Minnesota now after being free of it for 30 years. We have had a five-year effort toward it, and one of things that I recognize in the midst of this is the framework or the regulations we had prior to this were worse than the disease.

It impacted our cattle industry in Minnesota more so than the disease itself. I recognized pretty quickly we had to do something to change that. We're not here to get in the way of the cattle industry, but we're in the business of helping the cattle industry, and we want to get rid of these diseases too while we're doing it.

I want to give you some background in a minute on these diseases and why we think these changes are necessary, and then

1	6 PUBLIC MEETING
2	Dr. Marshall is going to provide you with
3	some information on the working group, how it
4	functioned, and how it was put together.
5	So the purpose of this new framework
6	is to have an adaptable program so that the
7	USDA can work with tribes and eliminate these
8	diseases. The key word there is
9	adaptability. We need to be able to change
10	on the fly to adapt to some of the
11	challenges that I'm going to talk to you
12	about in a minute that we have had with the
13	disease, and those challenges are changing.
14	I'm going to talk a little bit about
15	the background of these diseases. I think
16	most of you are pretty familiar with TB and
17	brucellosis and the eradication programs, but
18	it is just to give you a little bit of

And you're going to change slides for me.

ANNOUNCER: Yes.

background.

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DR. HARTMAN: Thank you.

So these are diseases that we have been trying to eliminate from cattle herd in

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the United States from the early 1900s, especially with TB. Brucellosis started a little bit later. These are diseases we're trying to eliminate not only because they affect cattle and economics, but also they're both diseases that can affect people, and I know that was a lot more important in the past, prior to antibiotics and some of the challenges that we have with people but are still significant issues. We don't want diseases in our livestock that can affect people.

So we made tremendous progress over that period of time. We have gone from a pretty high prevalence of brucellosis and about a five percent prevalence of TB in cattle to where the prevalence is unmeasurable, but we're having some challenges in getting that final step where we totally eliminate these diseases. It is a bigger challenge than maybe we thought it would be.

And I think the significant thing with that is states like Minnesota, Indiana, South Dakota, California, Texas, and I think

there is some infection in Kentucky with TB that have been free of this disease for 20 to 30 years, and Minnesota it was 30 years we were free. We're suddenly finding individual cases of the disease. We're having that challenge of getting those last few infected herds taken care of.

And similar with brucellosis in the greater Yellowstone area, there is infection in the elk and venison, and infection keeps going back in the cattle herds there. So, we have that challenge, and it is a big challenge. It's a political battle as well as a disease battle, and we want to try and make the final steps with that as well.

Next slide.

So what are those challenges that are new to these programs? The first one is the one I just mentioned, the disease in wildlife. In Minnesota we found an infected cattle herd, and then we found three more rapid fire, and they were all beef cattle herds and then DNR, because of the situation in Minnesota, said we better look at the

2 deer as well.

We located the deer. The first year they found one positive animal, and then a land owner got a permit and shot a deer, and that one was positive as well.

That creates some really interesting dynamics because not only are you dealing with the Board of Animal Health and the Board of Agriculture, but now you're dealing with DNR, and you're also dealing with a disease you can't contain.

When you have got -- I always told the people from DNR that when you are dealing with cattle, my job is a lot easier because cattle can be rounded up and tested. Deer can't be rounded up and tested. They have to be dealt with in a completely different manner.

It also creates some dynamics where you have to have an area that is quarantined off from the rest of the area so to speak, and it becomes a bigger challenge to do that. How do you monitor the movement of animals?

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You really need the support of the entire state, and in Minnesota that is when it started being effective is when the governor flew up there. He provided the resources and the authorities to the Board of Animal Health and to the other agencies to be able to deal with the disease.

And he also appointed a TB coordinator who then oversaw all the activities of all the agencies within the state.

So if I needed the highway patrol to stop cars up there and check them to make sure they had the right paperwork, I had the support to do that. So this is a very necessary thing.

So we're dealing with wildlife for TB, but also brucellosis as I mentioned in the elk and deer in the greater Yellowstone area, and that is going to be a tremendous challenge, but we need to not punish those states surrounding the greater Yellowstone area involved while we're trying to get rid of the disease.

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The other challenge we're having is livestock don't stay on little farms anymore their entire lives and never go anywhere. Livestock in the United States move on a regular basis.

I was at a meeting with the National Pork Board, and we were talking about planning for foot and mouth disease and how we would do that, and we started talking about the swine production systems that are in place now, and they no longer respect state borders. So pigs are moving from North Carolina to Iowa to Minnesota on a regular basis and back and forth.

They're also -- they can't be stopped. In other words, if you have got some baby pigs that are ready to move, they have to move within three days because there is nowhere for them to be. There are pigs coming behind them.

So the way we raise cattle and hogs is different than we did before. In any one of our sale barns in Minnesota, we have or the larger ones all have cattle from 20

different states on any one day that go through that, so animals move a lot more than they did in the past, and our operations are bigger.

In Minnesota when I started, the average dairy herd was about 30 head, and now the average dairy group is over 100 animals, and we have bigger herds, and I realize Minnesota doesn't have some of the bigger herds in the country, but it is moving in that direction for sure.

The other thing is, the challenge has been incomplete traceability. So when we found this infected herd in Minnesota, we went to the owner. It so happened he had a very good identification system on his cattle, and he had excellent records so we were able to do good tracing in and out of his herd, and some of the other records we found were not so good. They were on the back of a shopping bag and that sort of thing, so we were not as able to trace, so we need to improve that system, and we're working on doing that.

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The diagnostics for these two diseases are not perfect. When you do TB testing in cattle, it's your -- an example that we had is we had one herd in Minnesota that we tested three times before we finally found the infection.

So we had two whole negative tests on that herd, and on the third test it tested positive. It is the same with brucellosis, they're not perfect, and that is a limitation we have to work within.

The other challenge that we have with TB at least is Mexico is at a different level of TB eradication than in the United States.

I'm on a binational committee that deals with Mexico in the United States and tries to harmonize what we're doing, and they are making a lot of progress I will tell you that, but they have a lot more infection in Mexico than they do in the United States where we're routinely importing cattle from Mexico, and so we have to figure out a way to manage that, too, so we're not bringing

the disease in and allowing it to spread once it gets in Minnesota or in the United States.

I'm from Minnesota, so I often refer back to that, but next slide, please.

So why these changes? And I will tell you that not only are these rule changes necessary at a federal level, but they're also necessary at the state level, and we are making changes in Minnesota as we speak to try and adapt our program, too, and it is very similar to what the USDA is doing.

So one of the things that I found in the 20 years that I have been doing this kind of work is that our regulations are difficult to change, and that is both from the state side and the federal side.

For us in Minnesota to get a rule change through takes two years on the average. Two years. So if you were in a business and it took you two years to adapt to changes surrounding you, you would be out of business pretty quickly. We have to

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change that so that we're more adaptable both from the federal and the state side.

So what we're proposing to do in Minnesota and federally is to put some of the key things in regulation, but then the things that we need to be able to adapt more quickly put into policies or memos so they can be changed quickly. Now, they have to be changed with public input of course, but they can change more quickly and we can adapt.

The other thing that I think is a challenge now is money. The funding for these programs has not increased. It has decreased over time, and there are limits, and the way that we have been trying to deal with depopulating big herds is using emergency funding provided by the federal government. That funding is no longer available, and there is a limit on these budgets, and the budgets may actually become less. So we're going to have to find unique ways of dealing with these diseases that are not where the funding is limited.

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Next slide, please.

So veterinary services started off this process by listening to state coroners, and that is an important thing. By the way, I'm glad you all could make it this morning because it is so important that we get your feedback on "are we headed in the right direction or not," and we think we are, but we need to know that the cattle industry all over the United States is behind them and is supportive.

One of the things in the feedback that the USDA got on this was from an organization called the United States Animal Health Association and that group --

I'm member of that group, and Carter is a member of that group, and the livestock industry is part of that group, and they have the TB and Brucellosis Committee, and they passed resolutions urging the USDA to make these changes. I think there is a lot of input from all over the United States for that.

The other thing is this working

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group has tried to align with international standards, so how do we deal with these diseases not only in the United States but how do other countries deal with it? So we're on a par with them so that when we're training with other countries they understand what we're doing, and we understand what they're doing, and we trust each other.

There have been several public forums to talk about this. The first one was -the series was in December of 2008 that the
USDA put on. After that, the USDA did
some internal listening sessions talking to
their employees about what they thought would
work best, and then one that I was part of
putting together was the future of the
National Tuberculosis Program, and that was
in July of 2009 in Denver, and it brought in
about 150 people and really a lot of input
from the cattle industry on how we should
move ahead.

And there was very much in line with the results of that, and very much in line with what the USDA is proposing here.

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The other one was for brucellosis.

Veterinary services met with representatives from Idaho, Montana, Wyoming and the National Park Service in June of 2009, and that was about the endemic brucellosis in the elk and venison in the greater Yellowstone area and how to manage that.

That meeting resulted in a development of four core principals which include prevention and surveillance, disease response, and disease management, and the roles of state and federal agencies.

Slide six, please.

So from that input of VS produced two concept papers, and those concept papers for each disease were the basis for what this working group started with, and the objectives of both programs were the same, and I will just list those for you.

To mitigate the introduction of these diseases, to enhance surveillance, to increase options for managing infected herds, modernize the regulatory framework and transition both programs from a state classification system

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2 to a risk-basis only approach.

So that last one is really important. We have been dealing with these diseases by classifying states free, modified, accredited, having a five-tier system for states, and when Minnesota found TB we dropped two levels in that status. Well, I tell you what, you don't want to drop. is not a good place to be. We got out of there as soon as we could. We have a long state, and in some states like California it is even worse because they're even longer. Up where we had TB in Minnesota was a seven-hour drive from St. Paul, and it was 300 miles away from where other cattle herds were, and there was really no reason for those people to be punished, but there were herds in North Dakota and Canada that were way closer than the herds in Minnesota. this is a better way to do it. It makes more sense to me.

With that, I would like to introduce Dr. Marshall from Rhode Island, and he is going to finish this up.

1	20 PUBLIC MEETING
2	DR. MARSHALL. Thank you, Dr.
3	Hartman.
4	Once this group was formed, I was
5	asked to be on it, and I was kind of took
6	back a little bit, why would I have been
	-
7	there from Rhode Island, he asked me, and
8	I'm a working group dealing with tuberculosis
9	and brucellosis.
10	A couple of reasons, TB is not the
11	diagnosis. We have been a free state since
12	1972 with TB and brucellosis since 1982. My
13	counterpart in the state of Connecticut pokes
14	a little fun at me saying that he thinks
15	that dairy cattle ought to be listed as an
16	endangered species in Rhode Island.
17	I think the only reason I was asked,
18	I'm hoping well, it makes sense that I
19	could bring the perspective of a small
20	win/win state where a zoning could have
21	impacts and bring the perspective of the
22	state that has not had to deal with these

I will say that in my background, TB probably has had an impact on me. My

problems in a while.

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grandfather was a dairyman, and I still live on the same farm that he had, and he lost his herd in 1958 because it was condemned with TB. On that horse then, he decided to get out of the dairy business and take up his hobby and passion, which is race horses, which is what I grew up with, and if he had not lost his herd to TB.

So anyway, the perspective that I will bring and Dr. Hartman can bring a lot of experience dealing with these things, I'm going to bring the perspective of what it was like working with the working group formation of it and the challenges we had going forward.

The purpose of the working group was that both concept papers state that APHIS will work closely with stakeholders to obtain input on the proposed strategy, program standards, surveillance plans, and other policy concepts before publishing any proposed regulation and throughout the regulatory process.

Given this commitment in similarities

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in the proposed directions for both TB and brucellosis programs, VS decided to convene a single working group to discuss the regulatory concepts of both programs.

One of the first things that was introduced to us in the room is the concept of one proposed rule that would cover both TB and brucellosis. Immediate knee jerk reaction, I think, to most members of the group is these are two very different diseases, why would we want to go down that road, but I think it became clear that we could basically come up with one overarching rule. If we had left the rule kind of generic, understanding there are going to be different program standards for both diseases, and it took a little bit of time to get our head around that, but we finally did, think that group eventually reached reconsensus probably within three or four weeks into the process before we decided that this is the way we wanted to go, but that we could write very generic rules that would not have to be changed everytime that

something changed in a program, put all the details in program standards, which were much more changes in the program standards.

Next slide please.

So the Charge and Deliverables, the charge of this working group was to develop a comprehensive regulatory framework for both TB and brucellosis program.

We're now holding serious bold meetings to request your input and comment on proposed regulatory framework. I think it is something we're also very uneasy with the way the group was composed, and we'll have to get into the conversation a little bit later, is that it was basically imposed VS personnel, state, tribal leaders. We didn't have any industry input.

We were very much concerned that we didn't have industry input at that point, and we were reassured there would be a time and a place for that, and this is the time and place, so we really encourage you to speak up and say what you like and what you don't like about it.

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Based on the products of the working group comments VS receives during the public meetings about this framework, APHIS will develop regulatory text to be published as a proposed rule in 2011.

After a public comment period and necessary revisions, APHIS's goal is to publish a final rule by 2012. Okay. So it has been a pretty ambitious schedule. We started this in September of last year.

Next slide please.

The working group membership was not charged a Federal advisory committee. Its membership was limited to federal employees, representatives of state, tribal, or local governments. And again, conspicuously asking from that was there any kind of industry through -- we're approximately 20 members.

And we had state and animal health Officials; six of us, and there were five of us.

Originally -- I actually ended up with seven.

It was myself from Rhode Island, and we had Dr. Hallstead from Michigan, Dr. Susan Keller from North Dakota, and Dr. Bill Brydon

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from Idaho and Dr. Dee Ellis from Texas as the state veterinarians. Dr. Hartman came in a little bit later in the process and referred. He showed up after all the hard work was done. I'm just kidding. I truly appreciate his perspective.

Dr. Ellis, because of some issues they were having in Texas -- I swear Texas must have, it is so ambitious. The Texas State Vet Office has probably got a huge wall with every animal disease imaginable on it. One side of the wall are diseases Texas doesn't have today and the other side is diseases Texas does out there today and is constantly shifting the diseases over. It can't be an easy job.

We had two state wildlife officials, one from Minnesota and we had another one from Montana, so these people are very in tune with the challenges that wildlife infection brings to these disease programs, four tribal representatives, seven VS regional or area offices, two legislative and Public Affairs and one regulatory analysis and

26 1 PUBLIC MEETING 2 development. 3 Next slide, please. So Principles of the New Framework 4 5 Change, through the new framework, APHIS will 6 implement a flexible yet coordinated approach 7 to TB and brucellosis disease control and 8 management that embraces the strengths and 9 expertise of states, tribes, and producers. The overarching objectives of the TB 10 11 and brucellosis programs is to detect the 12 disease rapidly. That is not going to 13 change. That has been the goal since that 14 action. Again, these are all concepts, 15 objectives that have been since the inception 16 of TB and brucellosis programs, eradicate the 17 disease when possible and document disease 18 for domestic and international trading 19 partners. 20

Next slide.

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So here is where my naivety comes in, when we saw these program elements, I'm looking at going down the list, and while this one is easy, this one difficult, this one is easy, this one is difficult, and I

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think I have got them all about wrong.

Programs state requirements, you know,
I thought that that may be difficult for
some states because we need resources to deal
with, Rhode Island needed one of them, and
that actually turned up that we're all
pulling in the same direction on this, and
that is not a big deal.

Zoning in a state where every other state is right on top of one another, and at least conceptually see that TB detected could be spilling into Massachusetts or Connecticut. I thought there would be some issues with that. Again, it didn't seem like that was a program.

Indemnity, now again, my being naive about it that we protect the herd with the TB, and the federal government is just going to come in and buy it and depopulate and it kind of opens my eyes quite a bit when I hear that the federal government has about a million dollars to deal with infected herds and \$500,000 to deal with brucellosis infected herds. So the reality of it is,

1	28 PUBLIC MEETING
2	that ain't the way it is going to happen.
3	Indemnity, it happened to be the
4	most contentious issue. It was one that we
5	could not reach a consensus agreement on with
6	the members.
7	Interstate movement controls and
8	importation requirements, those are things we
9	have to take a long hard look at. Right
10	now I live in a very restrictive state. We
11	have a total staff of myself and a
12	technician. So we have always had the idea
13	that we're going to build up this gigantic
14	wall and moat our areas and keep TB out of
15	our state, and if that is the most effective
16	resource, I think that is debatable on that,
17	and approval of procedures related to
18	official tests and laboratories.
19	With that, I would turn it over to
20	Dr. Leann Thomas.
21	ANNOUNCER: Just very quickly from a

ANNOUNCER: Just very quickly from a process standpoint, originally we had anticipated going through Dr. Thomas's presentation and then have a series of break out sessions, but since we're such a small

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group, we thought what we would do because there are eight elements, as Dr. Marshall just indicated, as Dr. Thomas finishes one, we thought we would open it up to question and discussion.

If you will, I will allow her to get through the first element, and we will proceed on to the second elements, and so forth.

I will go ahead and leave the microphone at the table. If you would like to identify yourself feel free. Again, the meeting is being transcribed, so your name would be identified, so feel free not to identify yourself if you prefer not to as well.

Dr. Thomas.

DR. THOMAS: While Lynn is pulling up my presentation, I just wanted to express my thanks to those of you who are here today as well as to the working group members and the technical reps who came to assist with the suggestions we're going to be having.

And you will hear this said several

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times, is that we want public input. We need your input, and while we will be recording your comments here today, I would also encourage those of you to submit written comments as well as we move forward with this process.

Next slide.

So as Scott mentioned, the vision for this new regulation and this process is that we have developed draft regulatory framework. This is not the regulatory text. What you're going to see in the slides is first to the framework. The regulatory, the text, is still to be developed and actually we will rely on individuals and that is their job; they write regulations. But we wanted to make sure the framework captured the framework here so it is the concepts we're requesting your feedback.

We are looking to develop a single rule but as Scott indicated, we don't mean to imply or suggest that the two diseases are the same. Actually, they're very different. But if you will, look at the

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framework that has been generated. If you're trying to eradicate a new disease, you would still be looking at what would likely be those same eight elements. If you want to do surveillance you want to be able to manage effective herds.

You want to be able to manage effective herds at the investigations. You want to have importation requirements in place for that disease. So again, we looked at the background or the elements that would be needed for any disease to help provide the framework for these two diseases, and ultimately what that will give us is if we develop and put these regs into place, it will provide us greater flexibility in the long run if we do have to incorporate additional diseases into the regulation so single rule ensures consistency, improves our flexibility and also provides a great relief to administrative burden. Bill mentioned the problems, but the challenges in Minnesota with changing their regulations and I would ask Bill a question. I suspect that if you

32 1 PUBLIC MEETING 2 can do one rule change as opposed to two 3 rules, there is significant decrease in that 4 administrative burden at the state side, and 5 certainly that is how we feel on the federal 6 side as well. 7 DR. HARTMAN: (Nods head.) 8 DR. THOMAS: It was also mentioned 9 that the regulations will only include those 10 comments that are absolutely necessary but 11 otherwise, we will be placing guidelines, 12 policies into programs standards. 13 Next slide. 14 The working group had several 15 discussions on, well, what are the scope of 16 these regulations, and are you going to 17 include Brucella melitenis or Brucella suis. 18 And based on funding issues, we're 19 going to maintain the current agents that we 20 currently regulate, that being Mycobacterium 21 bovis and Brucella abortus. 22 That doesn't mean that we're going 23 to know our situations, for instance, where

we might have Brucella suis in a dairy in

We want to ensure that our

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Florida.

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regulations as we currently do have the flexibility that in the event of an epi investigation on that situation, that we would be able to indemnify and move that animal and do further diagnostics and actually document that it wasn't the abortus, it was suis. So I don't want to suggest that the flexibility that we currently have with such agents as Brucella suis, we will maintain that in the new program.

Likewise, there was discussion about extending the Brucella program and as I recall, TB into sheep and goats, and again because of funding issues, we don't have surveillance streams set up in sheep and goats, so based on those factors we determined that new regulations would include, as they include now, cattle and bison, captive cervids. We currently don't have a captive cervid regulation for Brucella, but with these new regulations we do plan on including captive cervids to both TB and Brucella suis.

The program or state requirements,

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and early on we had a lot of discussion, and I think I should point out at this point that the intent here is not to back away from state, federal and tribal programs.

That is not the intent here. We will work cooperatively in those states that have TB or brucellosis to help eradicate. So we're not backing away from that.

We are, however, as was mentioned, proposing that we move from a current five-tier system for both diseases to a three-tier system, with the critical component being the state's comprehensive health plan, and what that animal health plan does is it defines the activities that the State would put into action if the disease, these diseases were to be found in our state, and I have another slide that provides more details.

Under program requirements, we have such general categories as must have the rules, regulations, or infrastructure in place to implement a Brucella or TB program.

One of the questions that came up

was -- I will pick on Scott being from a small state, and Rhode Island is not the only state that this situation applies -- is that he mentioned that I think he has two or three personnel in the whole state, and that could be problematic if you had a number of herds that were affected with TB or brucellosis.

And the essence or what I think is really the beauty of what we're talking about are there are ways in addition to federal assistance, federal personnel that might be available is that there are neighboring states that he could have an arrangement, MOU if you will, so that in the situation where they might need his assistance that he can provide assistance and vice versa.

So we're not saying that the states have to take on these whole programs, but we're looking at ways, which there is more than one way to skin this cat.

And I think appropriately is that in this day and time with the prevalence is so low that it would make sense for Rhode

Island to have six or seven people chasing down diseases that for the last 40, almost 40-something years they have not seen within Rhode Island.

A critical component is reporting requirements, and then finally compliance and accountability. And I think I see this particular element as being the legal linchpin for these new regulations because while we recognize that our current status system is punitive as well alluded to, we also have to ensure that we don't make it so flexible that it doesn't have any backbone. So I think ensuring that the regulations require compliance and accountability for this tier system is really important.

And one of the areas that we really need your feedback on is are there certain things that require draconian action. We certainly want to move away from what Bill mentioned, the two status level decrease, which you can comment with that decrease in status level, there was a significant increase in testing requirements, and we're

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trying to get way from that, but are there certain circumstance where that needs to be written into the regulations or should it be in the program standards?

More about the state plan, and this is a written health plan, one for TB, one for brucellosis, and again, some of the activities would be that you want to make sure the State educates their legal authority on resources, surveillance, how they would handle an affected herd. Do they have a high risk subpopulation? I think the folks in Michigan recognize they have probably one of the highest high risk of population in its wildlife, but also is this a state that receives a large number of imported cattle, be it from Mexico, be it from Canada, be it from Australia, are there a high number of animals being feed-lotted in their state? What mitigation activities are going on in this state?

Again, is this a "One Size Fits All" for animal health plan? No, it would be different. I suspect the animal health plan

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for Rhode Island would be different from Minnesota if they were to write those documents today. Does that mean that -- so the animal health plans are not a "One Size Fits All" but there probably will be components that the State must include in their plan.

So how does the state plan tie into this concept of a status system, and the status as we were currently defining it, the statuses are, it's a consistent state, or provisionally consistent or it is inconsistent.

With an inconsistent state, what we want to do is define what the consequences are if a state is deemed to be inconsistent.

Next slide.

So I have spoken a little bit about the general program requirements. A state must develop infrastructure laws and regulations to implement and enforce these programs. It must implement a reportable disease process for both TB and brucellosis, and as I just mentioned, must develop and

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implement a comprehensive animal health plan.

I didn't mention it on the previous slide, but because this will be a new activity, although most states can't have components of an animal health plan, and one of our commitments is we will provide some script in play for states to complete their animal health plan.

The other thing that is under discussion is that what we would love to develop is an IT system mechanism where this could all be done electronically as opposed to hard copy, and so that is in our future vision for these programs.

But as much as possible, we want to be able to automate this, this aspect of the program, and make it easier for, number one, everybody to submit the information, as well as for everyone to have access to it.

A component very critical to the program is we want these animal health plans to be transparent. So the reporting requirements, again, to highlight the need for transparency in the success of this

program, and as a sidebar, the reporting aspect, I can't stress how critical it is because states -- at least it is my hope that if we develop the appropriate reporting systems on affected herds on epi investigations that the inclination or the perceived need to implement interstate moment restrictions at the state level will be less, but it is a definitely a challenge that we face.

Lastly, the animal health plan should include a description of how the states and tribes will coordinate their reporting. When the working group was meeting, we had numerous discussions about whether or not a tribe would be required to have its own animal health plan, and actually that will be something that the regulations would allow some degree of flexibility, and some areas with some tribes it was discussed that it might be a state or tribal animal health plan, whereas in other situations it may be the tribes that actually have their own plan, and it would be entirely dependent on the,

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as I see it, somewhat the antonymy or the infrastructure that the tribe has established and is used to working with them.

So, lastly, as I mentioned, that compliance and accountability is going to be built into the system, and a compliant state would not be subject to federal interstate moving restrictions or testing requirements.

How would we handle noncompliant states? Well, I think our current approach is that when their disease prevalence reaches a certain point, we increase testing requirements, while we recognize that there are other mechanisms and those include reduction to inconsistent status, or imposition of other consequences such as increased testing requirements, loss of funding, or increased surveillance requirements, and there may be other noncompliant actions or consequences that we develop based on your input, so certainly that is not an inconclusive list there.

So with that and actually this being a small group is we have some questions or

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perhaps before we go to the questions, based on the presentation that you have heard, do you have any questions about how the state status would work or do you have concerns because we're going away from a system that has been in place?

The TB program is roughly coming pretty close to a 100 years. It is quite a pick-on-the-female gender because it is quite an old lady and brucellosis is not far behind. So do you have any concerns about this approach?

MR. WHITE: Johnny White with

Georgia Cattlemen's Association. Dr. Hartman

brought up the penalty currently imposed on

the state, for one corner of the state

having issue, and I guess my concern would

be that a state could be penalized for not

having a state vet or Animal Health Board

that is appropriately functioning to the

program requirements that you're establishing

and not having a TB or brucellosis program,

so I'm flipping the whole scenario on its

head. I guess that would be from a producer

43 PUBLIC MEETING 1 2 perspective. I would not want my producers 3 being "penalized" in their ability to market 4 their animals or move them freely when there 5 is, in fact, no real animal health problem 6 in the state. 7 Did that make sense? 8 DR. THOMAS: Uh-huh. 9 Since we're having somewhat of an 10 informal discussion -- and maybe, Bill and 11 Scott, you can help me out. I think every 12 state has a state vet? 13 DR. HARTMAN: Either an animal 14 health official or a state veterinarian, and 15

I think right now they're all veterinarians, but there was a time when one was not a veterinarian.

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And I think the infrastructures in all states and obviously the states with livestock have more infrastructure. I thought that was an excellent idea you had, Lynn, that if it was a small state with limited resources that they would have a memoranda of understanding with another state to be able to utilize those resources.

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We have learned a tremendous amount about TB, and we have some very talented people working with TB, and we'd be happy to share those with other states. We would probably charge them for that, though.

DR. THOMAS: One of the things that I didn't mention in my presentation and I am amiss in doing so, is we have talked about an advisory board as a concept we want to use for these regulations, and an advisory board is a group of individuals that provide recommendations, evaluations. For instance, it could be a state animal health plan. It could be compliance with reporting requirements, analysis of surveillance plans, and we would like, as I said, we would like the regulations to include and incorporate this concept.

One of the difficulties that we have is probably from the aspect of our Federal Advisory Committee Act, which is our FACA Regulations as Scott alluded to in his presentation. FACA Regulations were the reasons we didn't include industry in this

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group because the federal government has -I'm using the term flexibility in getting
input, recommendations, feedback from other
federal and/or state and/or tribal
representatives.

We have -- it is the reason we were actually able to put the working group together, but it was, and I recognize, that some consider it to be a clear end flaw with the working group that industry group was not represented at the table. That is because of FACA.

FACA issues -- we have to make sure we comply with FACA when we stand up an advisory board, so we're still in the process of getting legal feedback from our office of general counsel regarding how we could set up an advisory board and who might be able to participate in a board.

But are there, is there any type of similar structure that exists within Georgia or is anybody aware of other states that have groups of individuals that are to provide input, analysis and recommendations

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into their animal health programs?

UNIDENTIFIED SPEAKER (1): We have had some advisory boards over the years on various things and y'all need some diseases like that we have had working groups or advisory committees on.

Of course, we have a new commissioner in Georgia, and he divided up to what the department does about 21 or 22 disciplines and has from three to seven people serving as a working group for those disciplines, and they will come up and make a report in June to a steering committee made up of about 40 or so people that will kind of get put in place. It is a strategic plan for the Department of Agriculture. So I think we have got some things in place that we could put together an advisory board together for brucellosis and TB and, of course, we have had a pretty strong program over the years to deal with any infection and deal with it, you know, years ago. The USDA published their White Paper on brucellosis, that being the final

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brucellosis meeting, and of course that was the final brucellosis meeting I have been to.

But Dr. Dees punched me and he said that they even asked before they stole your program because we had been doing a lot of the things, and we had been aware over the years that we was the state that started 365 day tests, four negative tests over a year, restricting all the infected cattle within a mile and a half, vaccination mandatory in a herd specific and depopulate in 30 days, so all of those kinds of things.

And I think our industry -- you know, I said one time that I was the one that was somewhat dumb enough to try to get things approved through the system, but our industry in Georgia has supported the Department of Agriculture and our rules, and we have a good -- we never had anybody buck us real hard about testing or doing what we needed to do.

I think we could establish an advisory committee for us if we needed to.

MR. COLLINS: I'm Jim Collins and I

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work with the Southeastern Livestock Network, and I would like to comply with what Dr. Black and Josh said. Like I said, I work

with cattle organizations from West Virginia to Louisiana, and depending on the state they vary in terms of how far this process is fleshed out. We try to serve as a clearing house when necessary to work with the industry, to work with our state vets across this region, and having just come back from Southern Animal Health, Dr. Black, Dr. Cobb and several others, are on the forefront of industry's concerns, and, I guess, I'll commit from organization in that region wherever, however we can work an advisory

role or facilitate more of what is going on. We want to do that because we want to

20 DR. THOMAS: A question back.

recognize how critical this

Based, Carter, based on what you indicated about Georgia is if we were to stand up an advisory board, I think probably the way the working group was thinking about it is that it would be national in scope, but perhaps

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that is just my own interpretation of the discussions is that do you think that having, if you will, 50 -- or how would the advisory board, such an advisory board at the state level, how would we ensure that it covered national situations and that it reflected a national viewpoint of perspective as opposed to a situation where it might be somewhat local?

UNIDENTIFIED SPEAKER 1: You know, I think that, you know, I was speaking from an advisory board just for the state to get that done. You know, we've had a situation in this country where a disease program was somewhat controlled by an advisory board, so it is not an undoable-type thing. I think that it worked very well in that.

That control board met twice a year, once at NIAA and the other time at USAHA, and we kind of were divided up, and we had a fall reporting time or a spring reporting time, but I think that worked very well, and I can vision the same type thing would work well with brucellosis and TB.

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DR. HARTMAN: We would be very
supportive of the idea in Minnesota, and with
all due respect, when we were dealing with
TB in Minnesota, I had the the first herd
owner that we dealt with had a purebred,
current case, black Angus herd, and had been
developing that herd for 30 years and a lot
invested in it and not only financially but
emotionally. The day we depopulated his
herd, he rode out on his horse and cried.
It was a very emotional experience.

But the reason I bring all that up is he started referring to decisions that were made by the USDA in our program as the people in Washington, D.C. as the TB gods. He didn't feel like there was any state or industry input into these decisions that were made, and that is what an advisory committee can do. It allows for the producers to have some input on how their program is managed.

It is not a decision from up high.

It's a decision made by all. So I think it is an excellent concept, and I think it could work both at state and federal levels.

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In every program we have had in Minnesota, we have had an advisory board because we need that input from the people that were affected.

As a matter of fact, Minnesota, we are run by a board, so I answer to a five-member citizen board whom are veterinarians and three livestock persons, and it works incredibly well. So I think the concepts are very good.

UNIDENTIFIED SPEAKER 2: I think the one question to answer is the rule. Where does the rule fit in the decision-making process? Is there somebody that the board advises or is the decision of the board final? So I think that is something we need to sort out. Also, it inspires the competition of the board, obviously industry representatives, federal, state regulators, I would argue that these are floating members of the board for particular states, so there would be natural concrete membership that is therefore virtually everything, and the state is affected, so Minnesota and Rhode Island

1	PUBLIC MEETING	
2	state veterinarians are part of that board	
3	for those decisions. That is just a	
4	thought.	
5	DR. THOMAS: With this discussion,	
6	we're all saying the concept with an advisory	
7	board be one of the elements or one of the	
8	components for animal health documented in	
9	your health plan, so I don't think we	
10	captured that during the working group.	
11	Should there be a different advisory	
12	board for brucellosis and one for TB? We	
13	talked about one specific pursuit of rabies,	
14	and could you have the same group of	
15	individuals at the state and/or national	
16	level that would serve that rule or would	
17	you have two?	
18	DR. HARTMAN: My opinion would be in	
19	the direction we're moving in Minnesota	
20	and I think you are federally, too, instead	
21	of programs by disease, have them by species.	
22	So I think you could have a cattle	
23	health advisory group that could function for	
24	either disease.	
25	UNIDENTIFIED SPEAKER 1: I think	

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probably that is right that we would be better off if we had it by the species rather than disease because, you know, you don't have, most folks that raise hogs don't have a lot of cattle industry on their place, but I think that would be a wise way to do it would be if you go by species groups.

UNIDENTIFIED SPEAKER: I would just say based on you see how many producers want to be at meetings all the time, so if you could knock it out with one board to be a whole lot further down the road.

DR. THOMAS: It may be that this question is more for after we presented the rest of the elements, but I will go ahead and ask it, and you can be thinking about it.

We have talked about the role of the advisory board as wanting to provide recommendations and veterinary services regarding certain actions, whether they should or should not be taken, but are there areas of noncompliance with the framework or the

1 PUBLIC MEETING 2 concepts that we're presenting that there 3 should be a defined consequence? 4 And maybe we'll go ahead and move on 5 to the next element. 6 DR. HARTMAN: Well, if you don't get 7 tired of hearing me, I will give you my 8 opinion when I want. 9 I think the whole purpose of the 10 program from the federal side is to monitor 11 what the states are doing, and I think there 12 are two things that are important. 13 One is in that monitoring there 14 should be other people monitoring as well as 15 the federal government. I think that was 16 important in Minnesota. We need these 17 advisory groups to be monitoring what is 18 going on, but clearly, if we're not doing 19 enough in Minnesota to contain this disease, 20 we're creating risks for other states. 21 has got to be consequences. 22 So should they be strictly defined?

Probably, so that it is clear when you don't do what you need to do to contain the disease, there is going to be some pretty

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55 1 PUBLIC MEETING 2 serious consequences because you're putting 3 other states' cattle industry at risk. 4 DR. THOMAS: Any other comments 5 about state or program requirements? And 6 there will actually be time at the end if 7 any questions come to mind that we will 8 address before we wrap up. 9 Next element is zoning. We brought 10 zoning up into two components, that being 11 short-term containment and long-term 12 containment. 13 And as was referenced earlier, we're 14 not talking about draconian changes. We're 15 still going to go after TB or 16 brucellosis-affected herd, and in really the 17 same manner and fashion that we are today, 18 but under short-term containment those are actions that are necessary when you find an 19 20 affected herd or you find that you have got

Bill described a scenario of finding
TB in deer in Minnesota; likewise, what do
you do when you find TB or brucellosis in a

the presence of disease in wildlife without

livestock involvement.

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56 1 PUBLIC MEETING 2 wildlife population in your state? 3 And the last or the other aspect to 4 zoning is a long-term containment plan, and 5 we, the working group, talked about 6 generically greater than one year. Is there 7 anything magical about greater than one year? 8 No. We pulled it out of the air. 9 And one of the questions that has 10 already come up regarding the one year was, 11 well, does that mean that if you have a TB affected herd that is under a test and a 12 13 remove, do you have to move into a long-term 14 containment plan? No, not necessarily. 15 We looked at long-term containment in a the situation that if over a year's time 16 17 frame you were seeing increases in the 18 number, your number of affected herds were 19 increasing, that would be the time that you 20 would need to kick in a long-term 21 containment. 22 So next slide. 23 As you indicated, short-term 24 containment is what we're doing now.

There is a standard

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are quarantined.

2 investigation according to the protocols that 3 have been described.

The plan is outlined in the state or tribes animal health plan. The goal of the containment is eradication, and it would end with a release of a quarantine.

Now, I think that it is important to stress at this point in time when we call this "short-term containment" under the category of zoning is are we going -- we the federal government -- are we going to go in and specifically designate an area that is under a quarantine? No.

What we're talking about doing is, in this situation is, relying on the states to implement quarantines on affected herds to implement movement restrictions from those herds so that we can assure the other states that surrounded or the rest of the U.S., as the case may be, is that that disease is contained within that zone.

So it is not a formal zone as we currently have in place for the split state status that Minnesota or Michigan or New

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Mexico is currently under, so it won't be a defined zone that you would see in the regulations.

Next slide.

So I mentioned that long-term containment is if eradication of the disease cannot be achieved within a year, then a long-term containment plan would be developed by the state or tribe.

Here is a reference to the advisory board where this would be an area where we would want the advisory board, either state, local and/or both to look at the containment plan and to see if it should be approved or if there were areas of concerns, how should the plan be modified so that it meets the federal government and the advisory board's approval?

We also recognize that in some situations there needs to be a risk assessment conducted. What is the actual level of risks associated with finding both wildlife and domestic livestock affected with TB or brucellosis? There may be situations

59 PUBLIC MEETING 1 2 that we do have to do a risk assessment. 3 Ultimately, VS would approve or 4 disapprove of the containment plan, and then 5 lastly, it would end with the eradication of 6 the disease. 7 In some situations, we recognize, for 8 instance, in the GYA, long-term containment 9 plan may go on for a long time, and it is 10 a very difficult thorny issue, and certainly 11 questions have been raised regarding the 12 wildlife component to disease control. 13 And Bill, I think you described it 14 eloquently when you said that you can't round 15 up a herd of deer or your deer to test them 16 to find out what their TB status is. 17 So it is a different -- totally 18 different, and I'm not sure it is appropriate 19 to say that you can manage TB or brucellosis 20 in venison. It is a very difficult concept. 21

I will say, although we don't have any wildlife representatives in the audience, that we do not have the authority to manage wildlife, which makes this situation very difficult.

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The veterinary service have been thinking about the need to have less punitive regulations as well as those that are risk-based, and we actually have already implemented a couple.

The first one being the TB Federal Order, and what the TB Federal Order does is that there is no automatic downgrade for an entire state. If a state was to have two or more herds for TB within a specified time frame under the current regulations, that would be an automated downbreak; however, the TB Federal Order supercedes that, and as long as the herd is under a quarantine, there is an epi investigation, if there is any additional surveillance, we're not downgrading those states, and we have employed this, and it has been successful to date.

The brucellosis interim rule that was published this past December 27 recognized that we needed to have a regulation in place that addressed the situation where we had nine brucellosis-affected wildlife.

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considered to be high risk, and currently we

2	And so in the interim rule, similar
3	to what we're talking about, moving to
4	generically for the U.S. is in the
5	brucellosis interim rule, a state that is

7 have four states that are considered to be

8 high risk. The three GYA states and Texas.

And if you're wondering why Texas has been class free for less than five years. So due to concerns regarding the fact that it's a fairly new class free state, it is still under, has that high risk monitor for brucellosis. So the management plan of the GYA, the focus there is those GYA states, number one, will set up their zone, and this is a situation where the federal government is not determining what the zone should be, is the GYA states are putting forth a zone, and they have to justify that zone size based on surveillance and other criteria, but we're not going to dictate to them that this is the area that we want to see where you have mitigation activities in place. We do have the option

62 1 PUBLIC MEETING 2 of reviewing that zone and making comment and 3 feedback. 4 But I think getting to the comment 5 you, Bill, made about there may be situations 6 that really the states know their own 7 infrastructure and the risks, and this is 8 more of it is more appropriate for the 9 states to be defining their zones in the 10 mitigation. 11 And certainly this is not having a state define their zone, but under the 12 13 existing split state status where you have 14 several tiers, for instance, for TB, it is 15 the states who provide the zone to veterinary 16 services, and so we would continue to do 17 that under this element. 18 Next slide. 19 So any questions or any comments 20 about zoning? 21 UNIDENTIFIED SPEAKER: Thank you, Dr. 22 Thomas. One of the interesting concepts that

UNIDENTIFIED SPEAKER: Thank you, Dr. Thomas. One of the interesting concepts that came up in the zoning discussion, which I think is new to at least most of us in here is considering different compartments of an

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industry as such being a zone.

Think about tuberculosis, and is tuberculosis eradication the same in the dairy industry and the beef industry or are there a certain sets of risks in one industry that are not present in the other? And I don't think we have covered that in the discussion, but there is particular mitigations that would be more effective possibly in tackling our national TB program if the dairy was considered as a zoning.

DR. THOMAS: Thanks, Dave, for mentioning that because the concept of a high risk population and another question that comes up is that, for instance, dairy heifers, should all dairy heifers be subject to interstate movement requirements?

DR. HARTMAN: I just have a comment about how it is functioning now with these zones that are being created around the greater Yellowstone area. What our state has done is develop each of those zones in each of those states and how they're doing it, and we have based our import requirements on

2 that.

I would rather see this advisory board evaluate it so that all 50 states don't have to do the same thing over and over again, so if we had an advisory board that we trusted that could evaluate what these states were doing, then we could all act in harmony, and we would not have to repeat the work, so I think that is another thing that an advisory board could do.

DR. THOMAS: We also have the advantage, if you will, of Katie Pertache with us here today, and so I should have said this early on, for those of you who are in the working group or technical reps, feel free such as they just did, if there are questions or comments that you want to add to the discussion, please feel free to do so.

So Katie, I don't know if you have any questions or clarifications that you would like to provide to the folks here, but please feel free or anybody feel free who was involved in discussions.

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2	MS. PERTACHE: We do have the
3	question to pose to everyone along the lines
4	with what Dr. Hartman was talking about.
5	Every state has the option to add additional
6	restrictions, animals coming into their state,
7	and so is the advisory board concept going
8	to be enough to provide satisfaction of other
9	states that these animals are free to be put
10	into place or are there additional things we
11	should be doing? Should the risk assessment,
12	for example, done in Minnesota, was it
13	adequate to help provide security to other
14	states that animals outside the zone are, in
15	fact, free? Is there something we could be
16	doing to help states along with that process?
17	DR. MARSHALL: I think risk
18	assessment as you said, timely reporting

rting is probably the most important thing we as state veterinarians have a real-time feel about what is going on with other states in the country, and that is the most important as far as I'm concerned.

DR. HARTMAN: My comment would be that it was a real learning process for us

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to get a disease that nobody else wanted.

We have had some real challenges with

surrounding states in how they treat us, and

it also speaks to the ability to change on

the fly.

So, for instance, Minnesota has not found an affected TB herd in over two and a half years. Last year when we tested and we did the surveillance on our deer in that area, none were positive.

And I'll give you a story. The president of the Minnesota State Cattlemen's Association had a wholesale, and he sold a bull to somebody in Illinois, and when they called to find out what they had to do to get that bull to Illinois, they found out they had to have a test -- and by the way he lives about 200 miles away from where we did have TB -- and he had to test his entire herd within the last year.

Well, he is an accredited herd, and in a free zone when you're accredited you have to test every two years, and it had been about a year and a half since he did a

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herd test, so they were going to make him repeat the herd tests. Well, he has got 900 cows. So it is not a small matter you can test.

Then in addition to that, they wanted him to test that bull within 90 days of going to their state and, then within 30 days of going to their state. It simply could not be done and does that make sense?

Should states have that ability to create unusual regulations that really are not based on risks but are just, it's the law?

And when I talked to the state veterinarian in Illinois, he acknowledged that, "No, it doesn't make sense, but it's the law."

So we need to have things that are more adaptable as time changes. We can't have these strict laws that don't allow us to do commerce between states for no reason.

UNIDENTIFIED SPEAKER: I just had a question as we're moving to this zone consent, and you mentioned earlier also about

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1 PUBLIC MEETING 2 the fact that you were a lot closer to some 3 other states being affected than the southern 4 part of your own state, and I guess just for 5 somebody that is not a state vet, how is 6 that going to work? Is there a potential, 7 in other words, if we had a case in northwest Georgia, it could be northeast 8 9 Alabama, southern southeast Tennessee, you 10 know, that could be your zone, and how is 11 that going to work with three state vets 12 potentially trying to figure this out? 13 DR. THOMAS: I think how we would 14 deal with it now is that -- Bill, I'll pick 15 on Minnesota because you're here -- in a 16 situation where there was an affected herd, 17 and let's just use a scenario that is close 18 to it. What does Minnesota border or what 19 is a good state? 20 DR. HARTMAN: Up there, there is 21 Canada and North Dakota. 22 DR. THOMAS: Thank you. My 23 geography is not so good. 24 So if it is close to the North

Dakota border, is that certainly the North

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Dakota state that would be advised, but specifically, if there are any traceouts or any sort of epidemiological association with herds in that state, there would be -- that herd put under quarantine and tested. So we would use the existing framework, and when that, so that herd would be quarantined, so it would be part of the existing process that we currently use.

We have talked about the scenario when you go into a long-term containment plan, and that would be a scenario we would expect that state to have a long-term containment plan. Again, but it would be based on risks just because it is a GOP political border in itself is not reason for or against that animal health plan. You have to look at what the risks are and certainly any of the other states.

Does anyone want to comment?

DR. HARTMAN: How it actually worked in Minnesota was since it was about 20 miles from the most northern infected cattle herd to Canada -- it was only 20 miles.

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Actually, no, it was less than that, so when we drew a ten-mile circle around the affected herds to test all the herds in that area, the circle went into Canada.

Now, we don't have any authority to make Canada testing there, probably nor does the USDA, but they did, and they tested all the herds that were in their part of that ten-mile circle.

North Dakota is about 60 miles away, and so they were never really impacted by it, but they had a great concern about deer. They were nervous about the deer early on, especially that we happened to find where infected deer were, and so they did surveillance on the eastern part of North Dakota, too.

So I think it is already working.

We're already cooperatively working together,

and I don't think there is any state that

wants to let the disease sit and not do

anything about it. So I don't think that is

going to be a huge issue, and I think people

are going to cooperate regionally and all the

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UNIDENTIFIED SPEAKER: I know in my other life I was in private practice and I happened to live in northwest Georgia and crossed the line nearly every day, and we did have brucellosis in that area over there at one time, and I had one herd of cows, and I had one herd of cows, a farm, that depended on which catch pen they penned them in, they were Alabama cows, and the other pen was Georgia cows, but we dealt with that, and I would call the federal office over there in Alabama and the state vet's office and let them know that we had problems there, and it was very close to the line and, of course, most of the time -- and we did the same thing with them -- if they notified they had one, a lot of times we would tell them to test the Georgia herds, and let one person be responsible for all testing.

But it worked out well for us. We never had that kind of problem, but it does exist.

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MS. PERTACHE: I guess I will ask another question. With all these potential for diseases, there is a potential for wildlife reservoir, so when you find a case, and infected herd, should it be a requirement that wildlife are tested, and should it also be a requirement in an adjacent state when there is close proximity?

UNIDENTIFIED SPEAKER: I think it probably should if you have susceptible wildlife in the area. Now, we have never in Georgia proven that white tail deer get Brucella.

And I did a good bit of work because one time when I was in practice and I had some herds that were bordering the wildlife management area, and everybody was blaming the deer on that, and I even got the DNR to collect some samples for me, and submitted them to the brucellosis lab, and we never found any problems with that, but white tail deer will get TB, and I think it would be a very good idea that we require surveillance in those areas, especially if

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And if you're in the western part, the northern part where you have got elk load, you're going to have to do that, too, for brucellosis, but we have never found white tailed deer to have brucellosis.

DR. THOMAS: Did you have a comment, Bill?

DR. HARTMAN: I hate to dominate, but we have been through all these issues, so I think it is important. I think absolutely. We have shown that particularly with TB, it is not unusual for it to be sitting in an area for a while with high deer density but there is a strong chance that it is not going to spread into the deer, so not to do that surveillance would be irresponsible, not only to the rest of the country but to your own state because you need to know where the disease is and if it is in the deer, and if you don't deal with it, then instead of having Minnesota's situation you have Michigan's, and it is not -- I don't envy them, the challenge they

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have, because the disease was so prevalent and so widespread by the time they found it. It's a huge job to manage.

So I think there needs to be a requirement for wildlife surveillance and in certain circumstances where there is a risk that it is spreading to wildlife, and I think that would be part of the advisory board's decision is, do they need to do it or not and the consequences for not doing it would be similar to not dealing with it in the cattle herd. You lose your consistent status or whatever that status may be.

UNIDENTIFIED SPEAKER: I just want to ask a question, and I know this is not the wildlife management crew, but anecdotally, what kind of testing goes on in wildlife? I mean, is there any program out there now to test?

DR. THOMAS: The way we work is we work with our wildlife agencies as well as through the states. Typically, what we have done is provide funding for those

surveillance activities to some degree.

Bill, you may want to elaborate on how the wildlife has been surveyed in Minnesota, but in the GYA states we have provided funding that supports the surveillance.

We also have through, if you will, initiatives or programs that have been put in place for the GYA. Those animals, they're coming outside of the GYA during the winter months. Those animals are captured and subject to surveillance testing, so we may have somewhat of a better idea of the instance of brucellosis at least with the advice from the GYA from an ongoing program that is associated with activities when those animals move of to greater Yellowstone Park.

DR. HARTMAN: In Minnesota, our experience was DNR is just as concerned about these diseases in their herd and the deer as they are in the cattle herd, so we had no problem with them doing a surveillance.

And on a larger scale, they will do surveillance statewide if necessary to

determine if there is any prevalence in this disease.

We require early on to prove that there was no TB here or anywhere else in the state. I don't think that is widespread in the country, that there is a lot of statewide testing going on, but if there is no reason to believe it is there, I don't think we need to be doing that routine surveillance.

As far as dealing with the disease and wildlife, our DNR, their plan was to reduce the densities of the deer in the area, stop and baiting and feeding.

Actually, we don't allow baiting in the state. They banned feeding in that area, and there was a lot of feeding going on probably not only by recreational uses but by the cattlemen that liked seeing deer around, so they were feeding them, and lastly to do surveillance.

Our way of reducing deer density
when we're lucky enough to get together and
push back with this is we had helicopters in

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the area shooting deer. We had sharp shooters shooting deer. We had liberalized hunting regulations so that if you were a land owner you could shoot deer whenever you saw them.

You could get permits to hunt as many as five deer in a season, and so we did a lot of work to try and reduce the density of the deer and then stop the aggravation of the deer in the area.

And finally for those cattle herds we did a cattle herd buyout, so in that area when we were finding infected deer, we brought out 46 of the 68 producers that were raising cattle in the area, and we continued to make payments to them until we cleaned it up, and then for the 22 herds that were left in that area, they had to have wildlife risk assessment every year, and they had to develop ten-foot deer-proof fences around the winter feeding areas when they started feeding, and we're hoping that those really strong regulations have helped us eliminate disease.

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UNIDENTIFIED SPEAKER: One of the reasons I ask is because we have some deer feeding and overbaiting and feral hogs, so there is a lot of that here in south Georgia.

UNIDENTIFIED SPEAKER: I think in Georgia, DNR would be jumping on it if we find TB, and they would really do the surveillance for us there.

DR. THOMAS: I think we'll take a ten-minute break before we start the next element.

(Whereupon, there was a Break taken.)

DR. THOMAS: I will go ahead and get started with the third element. The first component of surveillance is national surveillance. So for both TB and brucellosis we will continue to have national surveillance. As it currently exists, that is now slaughter surveillance via blood samples that are collected at slaughter or TB or suspicious granulomas that are collected for TB at slaughter. Any other surveillance that would be appropriate to include in the

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national surveillance, we will do so.

And one of the questions that came up in the earlier session was that, well, what happens -- you know, does this say is this a state or federal requirement, and this is when we talk about national surveillance, we're talking about the federal role in ensuring we're able to collect samples that are federally approved slaughter house facilities.

Targeted surveillance would be one of our components, and that is where you have a situation you're monitoring an atlas called the population and as described in your animal health plan.

And then the third component is any other surveillance that is being used to support state or tribes of zoning efforts, and this would include testing associated with movement controls, testing for other, if you will, zoning activities that go on, and any other methods of disease detection that you're using for that area zone.

This next one, the last element in

regards to animal ID is we will maintain consistency with the traceability proposed regulations. We're not going to be proposing anything new and above anything new or different than the proposed traceability rule when it is published; however, we will be indicating for certain program activities such as vaccination testing that is conducted as a result of an epi investigation, that those animals must be officially ID'd.

And although I suspect this group, given location may feel a bit differently, I want to stress this is not a forum for the discussion of the traceability of proposed regs, but we do recognize to have effective surveillance you have to have your animals ID'd, and I will just leave it at that.

And rather than stop here and have questions, I'm actually going to go to the next element. Effective herd management and epidemiological investigation. The regulations will provide a definition of terms, how do you fine an affected herd? Are there other groups of animals whereby we should provide a

list or a definition, such as an affected feedlot, et cetera? We will provide for the process and identification of the people involved in decision making. Who makes the call for an affected herd? How is that call made? Who is involved? And it also similarly provides a process and identification of the individuals that are involved with the herd plan. Who makes those? Who implements them? Who oversees them, et cetera?

Next slide.

The regulations will provide for the development of investigation reporting requirements and time frames for those epi investigations. It will allow consequences that if epi investigations are not properly conducted or within specified time frames, again, the thing Bill alluded to it earlier is what are the consequences if a state is not conducting the epi investigation to time frames maybe or conducting an epi investigation at all.

Having said that we want to have

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consequences, we also recognize that there may be valid circumstances that require forbearances and time frames, so as our existing regulations do.

And then recognizing that under affected herd management, we will continue to evaluate a policy of the testing or removal procedures for both TB and brucellosis, and in doing so we recognize that there may be some need at the state level for facilities to receive high risk or restricted movement animals.

So, again, I think this goes back to the concept for quarantine feed lots.

Definitely we want these to be terminal feed lots, but what mitigations can be put in place to move affected animals -- not affected animals, excuse me -- exposed animals, not an infected herd, so that they provide, or I should say, that they lessen the penalty for the producer?

That is it. So a very brief overview of surveillance and affected herd management and epi investigations. The

83 1 PUBLIC MEETING 2 latter element is many of the concepts we have in place now we will continue to have 3 4 in place. Where we can, we will put as 5 much as of the guidance as we can into 6 program standards as opposed to the 7 regulation. 8 And as Bill alluded to in his 9 presentation, the advantage of that is we can 10 change them more quickly. It will be 11 through a public notification and comment 12 period, but that procedure takes much less 13 time than doing a proposed and then a final 14 regulation, which is ultimately if you do it 15 in two years on the federal side, you're 16 doing good. 17 So any comments or questions about 18 those two elements? 19 UNIDENTIFIED SPEAKER: Well, thinking 20 along the lines of epi investigations and 21

timeliness, how long do you think an investigation should take?

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UNIDENTIFIED SPEAKER: Well, I always felt like if when I had an infected, brucellosis infected herd, I wanted an epi

document in my office within 15 days of the
time that that BMO was notified. That does
not mean that this epi investigation is over
with because an epi investigation is an
ongoing process that oftentimes you find
additional herds as you begin to work through
an area that may or may not and, of course
with the TB epi I'm sure which I have not
been involved. Well, I was involved in one
when I was in practice, but that was the
last break of TB that we had in Georgia in
1975. So I'm sure that that those epis are
going to take a great deal longer because
you have got to go back further and do all
the traces and traceouts in which brucellosis
is somewhat a shorter-term situation in most
cases, but I have got no answer, but Bill,
you may have a better idea.

DR. HARTMAN: I think this is an area where flexibility is important because in Minnesota with our TB investigation, we eventually found all infected herds, and they were all beef cattle herds, but they led to over 1,200 traces in and out of those herds

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into nine different states, and if you have
got that situation, you have got the time
frames have to be based on, again, I think
an evaluation of this advisory board and
brucellosis epidemiologists. Are they moving
quickly enough on it? But I don't think you
could say it has got to be done the next
number of days. I just don't think it fits
all of the situations that you would find.

UNIDENTIFIED SPEAKER: Would it fit
better to have the time frame associated with
the stages of the investigation as in so
many days to an initial report and then
every 30 days or every 60 days pick a number
that status updates are flowing appropriately
as monitored more than a cutoff point, and
it is also supposed to be finished?

DR. HARTMAN: Yeah, I would agree with that. That makes sense.

UNIDENTIFIED SPEAKER: I realize this is for industry input, but certainly from a market figure standpoint, that is preferable, where if there is going to be any fallout on a market standpoint from the unknown, and if

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you have it in certain stages and time frames within those stages, that is reassuring to most of the domestic market as well as outside of that.

MS. PERTACHE: I have a question from the State's perspective. What about transparency? What level of reporting would you suggest at each of these stages, and to whom will that reporting be transparent to?

DR. MARSHALL: Well, just covering with the eased yield frame, I think that, you know, that whole transparency seems to work fine. You know, we're getting timely reports, and we're getting what appears to be very transparent reports, and I think we feel very comfortable in the state level knowing what is going on, so that could serve as a good model.

DR. THOMAS: And Scott, I have been out of the office a lot, but it looks like the mechanism of reporting, of EHD reporting are the state vets are submitting a report to VS and then they're disseminating it after the state vet has cleared the info as

2 opposed to VS documenting or providing the 3 reporting.

DR. HARTMAN: I would have to agree that we should be doing for brucellosis and TB what we're doing for these equine diseases.

Right now, we have no mechanism for North Dakota to find out what is going on in Minnesota, other than Minnesota having to report to them and to every other state in the country.

And if we had a national report that was provided every month on progress within states handling these diseases, I think that everybody would be comfortable with getting that report, and it would be up to the states that are having issues with the disease to provide the information to veterinary services.

But we don't have that right now, and part of the response to not having that right now is uncertainty, and uncertainty leads to higher important requirements frequently when it is not necessary.

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it is not a dairy herd which we frequently hear, particularly with TB, that a dairy producer can survive under a test and remove but a beef producer cannot.

So if he has animals that test negative that need to be moved, for instance, into a feed lot, do you agree there should be mitigations put into place to recognize the need to, I will say, send those animals to slaughter -- well, we currently have the mechanism to send them to slaughter -- but to feed those animals out?

UNIDENTIFIED SPEAKER: I think if they were going to do that they would have

89 1 PUBLIC MEETING 2 to go to a quarantine feed lot if it still 3 exists. 4 DR. THOMAS: Is that something we 5 should put into our regulations that it is 6 up to the State to determine whether or not 7 they want to have quarantine feed lots? 8 UNIDENTIFIED SPEAKER: I think it 9 has always been the State's privilege to do 10 that. We did have one at one time in 11 Georgia, but I said what happened there we would never have another one if I had 12 13 anything to do with it. 14 UNIDENTIFIED SPEAKER: Well, from industry perspective, obviously, our producers 15 16 want as much flexibility to survive the 17 situation, but at the same time, you have to 18 deal with the health issues, and we don't 19 want to affect the rest. That is the only 20 thing that makes any sense. 21 UNIDENTIFIED SPEAKER: Just to follow 22 up on that, also. I sure appreciate Dr. 23 Black's comment. As I look at what he 24 mentioned earlier, the funding for

depopulation, and I see indemnity as "X", and

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I think in my mind that is the only guide in which discussing going to the quarantine lot comes in play. Is that a fair statement that if you know funding is going to be decreased then that puts it in more perspective?

DR. HARTMAN: And I think a couple of the issues with that when we found in Minnesota even when we depopulated herds, and we frequently find a plant that would be willing to slaughter these animals but they would pay a dime on a dollar for them, so they were heavily discounted.

So if you were going to do that, you would have to have the mechanism to compensate the owner for that loss in value, and my guess is when you have had quarantine feed lots, they didn't pay as much as the nonquarantine feed lots, so again, the quarantine feed lot you would have to have a mechanism whatever they were discounted for their cattle because they were going to a quarantine feed lot; otherwise, it wouldn't work.

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2	DR. THOMAS: So perhaps that is a
3	good way to talk about indemnity. And
4	Scott, I will repeat something that you
5	indicated in your presentation that this was
6	the most contentious element that we
7	discussed, and I think it is certainly when
8	you talk about the removal of animals from a
9	producer's farm where it's their lives, and
10	for some of them it is not just their life
11	but it is the family, what the family has
12	been doing for generations.

So we talked about indemnity in great detail, and what I'm going to present after this slide is the VS on position and certainly does not represent a consensus by the working group.

Before I go into that, I want to provide you some background, and actually it's a couple of things that have already been mentioned today is that we do have flat and actually for this fiscal year it is a declined federal budget for both of these diseases.

TB decreased by \$800,000, and

brucellosis decreased by a half million, and it is correct that we have -- well, I wish we had \$100 million for TB indemnity -- \$1 million in TB indemnity and \$500,000 in brucellosis.

The budgets are put for TB. The way that we have paid indemnity in the past was that we have gone for emergency funding or we have gone through, specifically CCC, which is the Commodity Credit Corporation, which is federal funding, that we apply for, but that request has to be approved through the department and because of fiscal deficits as well as we have used the argument that with TB is that those are the last herds. If we can take these we'll eradicate these diseases, and you can only use that argument so many times before people will become skeptical about your argument.

And so the reliance on CCC funding is we're not going to be able to rely on that in the future. If you look at the average funding from 2007 to 2010, we have spent roughly \$5 million in TB indemnity

annually. So roughly four million dollars and again, this is average for the purposes of discussion. We have spent \$4 million in indemnity. So you can see as we move past our reliance on indemnity funds, it is going to be harder and harder to indemnify herds for TB.

Particularly, and it does appear that in the near future we will be seeing increases in our line item. For those of you who are familiar with our proposal for funding for 21012, which is as opposed to going for specific funding for TB and brucellosis, we're looking a line item entitled "Cattle Health."

And whether it is cattle health or

TB or brucellosis in 2012, we don't know

what the funding is going to be called, but

the funding will remain the same. It is not

going to change. So in 2012 we are looking

for a decrease.

The final bit of background that I would like to provide is that it roughly takes 60 days after an appraisal is done for

a herd to be depopulated. I'm not referring necessarily to what you may know as diagnostic purchases. I don't know the average time frame, but roughly when a herd is depopulated, it takes about 60 days to get that herd off of the ground, and that is primarily I'm referring to TB information, but I suspect it is similar for a brucellosis affected herd.

Next slide.

So the regulatory components for indemnity we will provide definitions for those terms that are specific to indemnity, and we will indicate that our indemnity payments are subject to the availability of federal funds, and that is not new. Our existing regulations indicate that.

And then we'll describe the approach to indemnify because what we want to do is -- what we want to do with other components of this framework is we want to keep only what is necessary from the regulations and have program standards that will provide the details to the indemnity process.

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So what we're proposing in this framework is that we will provide value for the individual animals using a calculator, and a calculator would be based on such criteria as the age, the weight. For dairy animals, milk production, and it would reflect regional values.

There would be a defining process for updating the calculator. The calculator would be developed with input from others, and the indemnity paid would be 100 percent of the fair market value minus the salvaged value when the animal was slaughtered. There is no appeal process.

One of the factors around an appeal process is we would not be able to remove those animals quickly, and then you might have a 60-day time frame where you have got affected animals on the farm.

The veterinary services have contracted without side of the government to develop a calculator, and the beef calculator was developed by Dr. Peel, P-E-E-L, and it did have outside review, and it covers bred

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cows, heifers and calf pairs and herds, and the base price is based on the cow's value with consideration given for the cow's age, the cow's weight, the calf, age, quality and considering the pricing in five different regions.

But the purpose of this is just provide you an example of what has been developed. There is also been a dairy calculator developed, but for the purposes of this example it is much more complicated, and I do want to go into it and certainly I'm not a calculator development specialist, so this was just an example to give you an idea of how we might develop a calculator. So I think at the last meeting, we had quite a bit of discussion about indemnity, and we will go ahead and open it up for comments, and we also have some questions. comments?

I apologize for that. I was anxious to get to the comments or the questions.

Then the last thing we'll do is the regulations will describe those eligible

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indemnity payments, other than the actual payments for the animals, which includes transportation and disposal. We do not plan on paying for any cleaning and disinfection that might be required, although we would consider under certain circumstances to fight for the disinfectant.

So the questions that we had for indemnity, what criteria should be considered to develop a calculator? I will ask another question and maybe it will stimulate some conversation. Several groups in Michigan indicated that they saw a difference in the circumstances wherein a calculator might appropriately be used, and they saw a distinction between an animal that you were wanting to purchase as to what we would refer to as a diagnostic purchase versus a depopulation of a herd, and there was some discussion around if you have reactors or suspects that you're wanting to purchase for diagnostic purposes, they indicated that if you could rapidly remove that animal, i.e., use the calculator, get the animal off of

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the farm, get the producer paid based on that, there was value in the use of the calculator.

Having said that, there was significant discussion about high value stock, registered animals, genetic material, MMD, regardless of whether it was for a diagnostic purchase or whether it was for an indemnification of a depopulated herd that such factors needed to be considered.

But going back to my real deal, do you see any value in the use of a calculator for diagnostic purposes?

UNIDENTIFIED SPEAKER: I guess there is, you know, multiple ways to look at that as a starting point, and I have a lot of respect for Dr. Peel and his work throughout his career, but it maybe gives you a starting point, and again just in this room and in the meetings last week, just across the southeast, the varying degrees, if you will, of staffing or appointed careers of different members within those states so that representation in terms of negotiating with

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producers are trying to quickly get to that value, and I guess in my mind is that the calculator may be a good starting point.

I would ask back to the point in terms of flexibility. I don't know if it is the single answer. I guess I would also say in today's environment where threat is a bigger issue, too, and I have not looked at his breakout for what the regions are, there would be a lot of difference in terms of value, if you will, a better grade feeder calf that is in southeast Georgia versus one that may be in western Mississippi, and it is not taken into account here, and we could figure the rest of the day discussing genetic material, high value animals, and I'm going to tell you there are some of us from the producer side of it looking at where there is opportunity for a public private partnership to come up with solutions to that, and that is a whole other day's discussion to your original point as you look at the feeder calf end of it, that needs to, particularly in today's environment, be looked

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at as closely as the bulls and cows, and also keep in mind the transportation variation as well.

DR. THOMAS: You said there has been discussion around a public/private and do you care to elaborate any on that because we had similar discussions in Michigan?

UNIDENTIFIED SPEAKER: I guess at this point there are some sessions going on, again, at the federal level looking at where there is a fit, and I stress "a fit" because there is nothing that takes the place of the role that I listened to, but where there is the opportunity for, you know, if Josh has cows worth \$4,000 to him, and I'm seeing Dr. Blackshoes, and it's worth real market value, how is there a means to somehow allow Josh, if you will, to buy up seven if he does have to be depopulated through a public/private partnership, he can get the real value of that animal, him, but for the rest of us that are out here, Dr. Blackshoes and his neighbors, that animal is depopulated quickly.

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And it would be all of the warranty-type scenario is what it would be, again, for back to work there has got to be, and I use the term a public/private partnership. Before we get there all these questions you're asking from industry needs to have some feedback to look realistically.

And again, I operate within the political environment, too, and we all know over the years there are certain producers and certain districts that may have had one value and may not be the same in the other. I don't think anyone argues that nor wants to discuss that, but we're about getting to the right answers, and that is where a calculator, as I said, is a baseline.

One doctor used the example the other day where he had gotten three animals in North Carolina that had been delivered, and his local guy put a value within a matter of hours, and I'm not sure how many days it was until he officially got a value and got a nice report of \$100 for each animal.

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And, again, I'll give him credit, there is nobody as good as Bruce in a lot of states, but that is some of the variations we have in states, but from an industry perspective, we want to work with you to get the answers.

UNIDENTIFIED SPEAKER: Well, you already brought up the feed stock registered business, and I guess one of the concerns with the, you know, null of bill process and a formula is, you know, your leaving those guys very vulnerable in that process.

And the other point in that is the discussion we had earlier where "Bull," which is a huge feedstock producer in Minnesota, is sending bulls from Florida to Washington state every year, and that is over a million doctors out of their production sale every year.

Those feedstock guys -- and there is a good chance, and I understand there is a lot of females, too -- those are getting disbursed as breeding animals a long way away, so the scenario is right for that to

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be a problem potentially in the breeding herd. So there needs to be a realistic scenario, I guess, developed for that eventuality.

I've seen the same thing in dairy over there, and you said your formula was a little more complex, and maybe that is good I'm hoping for the dairy folks but, I mean, on any given week in Georgia there is an animal sold from, you know, a seven-month old heifer sold for \$400 at a sale barn and \$10,000 at a purebred sale, and that is the same color, same weight, so I don't know with your shrinking budget has to do with that.

DR. THOMAS: And another question with the shrinking budget and the limited indemnity funds is -- just a couple of comments -- our current Authority Animal Protection Act does not cover replacement value. It's full market value. So I think it is important that I state that just because of what we have the ability to do.

But, secondarily, given the limited

104 PUBLIC MEETING 1 2 funding we have for indemnity, if we're 3 talking about full market value minus salvage value, if we have a situation where we even 4 5 get into a couple of large herds or where we 6 have many smaller herds but on a higher end 7 is that our current regulations have a cap 8 of \$3,000, and so is it -- what is our 9 responsibility overall toward indemnity? Do we try and spread it out as much as possible 10 11 or do we allow the payment of full market 12 value minus salvage to the first producer who happens to have the lucky, so to speak, or 13 14 the luck to have the first TB affected herd 15 at the first of the year? 16 UNIDENTIFIED SPEAKER: First of the 17 fiscal year. 18 DR. THOMAS: Thank you, yes. 19 UNIDENTIFIED SPEAKER: If you have 20 it in the fall you get funded or in the 21 spring I guess you don't or whatever. 22

I guess from -- as we're moving forward with the reduced budget, and you know, Jim brought up the public/private partnership, and it will take a lot of heavy

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lifting to get it done, but in time -- and Jim has already been working on it and it's a big educational component -- but the market, you know, where there's a market for a gap or warranty insurance or, whatever you want to call it, I don't know, but maybe that is something we need to discuss with RMA like Jim is already doing, but maybe --DR. THOMAS: What is RMA? I'm

sorry.

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UNIDENTIFIED SPEAKER: Risk management.

UNIDENTIFIED SPEAKER: Just for a point for the record, what I'm discussing will not be under RMA, and that is the reason I carefully used the term "warranty," but I guess for a moment I would like to pass to Dr. Hartman in the scenario used right now. Is that lucky herd -- should all those dollars, you know, be paid out? Can you shed any light? And I have got a comment after that.

DR. HARTMAN: I probably could go on for an hour, but we depopulated 12 herds,

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and the biggest herd took that entire million dollars, so in that one herd, the first herd we found would have been done.

eradication program if you don't have money for indemnity? I don't know a single one of those producers under our current system if there had not been emergency funding identified that we would have had a solution for it. We had nowhere -- if we couldn't have depopulated them, we then couldn't have done test and remove because they're all beef cattle producers, and they all produce calves, and every year those calves have to go somewhere, and if they have to go somewhere where they're to get half the value they normally get, they're out of business.

So we have got to, as we're discussing this, be realistic about what we can accomplish with a million dollars in indemnity, and I sat across the kitchen table from these people, and they would have not liked a calculator -- I'll tell you that right now -- because they knew how much

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their cattle were worth. There are appraisers in that area that knew what their cattle were worth, and they could have told us what they were worth in one day.

Instead we had the system that was in effect when we depopulated these herds. The USDA had to contract with somebody to do the indemnity, and the person who won the contract frequently was not from Minnesota. In fact, they were never from Minnesota, they were from California and Michigan, and the one person who came didn't understand the --well, he understood the dairy industry but did not understand the beef category industry and did not understand in that part of Minnesota.

So after a very long complicated process of him developing a process of him developing what he thought it was appraised for, the owner said, "That is not going to work."

So there was the appeal process.

You said, "60 days for removing the animals."

We only wish. It was three to six months

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before those animals left the farm in the northwestern part of Minnesota.

So I'm not defending the old way.

It is not working very well or it was not working at the time we depopulated, but if there was a mechanism to offer the owner what the calculator says, if that is right on target, they'll accept, and if not, there should be some local appraisal that is allowed and can be done quickly, and the owner is offered what that appraisal is.

If they're still are not satisfied with that, there should be an appeal process. Anything else less, I just can't see it working unless there is some sort of insurance or private or public partnership.

UNIDENTIFIED SPEAKER: Now, you see why I passed to him first. I thought back to my comments earlier on the calculator that it may be a good starting point, but as to your point, and I guess I can't imagine. I guess that is what happens in the bidding process is why you would have someone removed from that local environment.

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And back to Dr. Marshall's point he made last week about this local guy putting a value on it in a matter of hours. So I guess my question would be or point would be as we look there and move down the road is I think all of us realize, if you will, the dollars are not likely to increase dramatically.

As we think about solutions, you know, I wonder how any of these systems have got to be built to where we certify the producers to do all things in terms of protection, and all things they can do in terms of working with their -- whether local vets or state vets or industry, and as you said, those guys may not have had an alternative. Likewise, every one of them wanted the disease to go away.

And so I think that is -- part of this indemnity discussion is how you tie these objectives together. I don't have the answer for you today. I will say there has been an awful lot of work done and thought done in trying to recognize these things and

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come to an answer, and I guess the high valued animals at some point in terms of getting to -- and my family is in the commercial cattle business -- and we would be one of those if something like this hit and if we had a disease, and in a very, very small community you could hit \$30,000 cows that fast, just because of the way cattle moves, and we all have lots of 60 herds next to each other, and we all cross over each other.

In that case, any one of the three of us right there in our immediate area would max out your dollars probably, and that is disturbing as we're in the commercial state selling business, and also we're one of the hundreds across the country.

So, again, there is not an easy answer to this. I think there has to be some combination to that is where we go today.

UNIDENTIFIED SPEAKER: I guess my concern is the appeal process. Why are you putting that in there?

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DR. THOMAS: Part of it is it's a legal issue associated with the transparent -- going forward with the transparent process that everybody agrees on the calculator.

The second is to the issue of the appeal process that Bill alluded to. The appeal process, as it currently exists is extremely onerous, and it is time consuming, so if you have that appeal process, you're not going to be able to get the animals quickly off that farm. And maybe that is an inherent problem with the appeal process and the length of time.

UNIDENTIFIED SPEAKER: Like Jim said, some combination with the calculator with an appeal. You pay on the calculator, and you can get an appraisal done in a week from a local person. You can move, and you can establish an initial value and have the appeal process behind.

DR. HARTMAN: And our experience was the delays were not necessarily the appeal's process, it was the initial appraisal process, and in my opinion it was not

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necessary, but it was federal regulations on how it had to be done and how it was implemented in our state that led to that, so I think the appeal process could be done very quickly if it was managed quickly.

UNIDENTIFIED SPEAKER: One last comment. In every one of these cases we have discussed -- and I'm going to reflect back to something Dr. Black told me early on in my career, and in every one of these cases they can move very quickly, depending upon the relationship and the partnership between industry and animal health officials, and he could keep us here the rest of the day with telling us stories for having a relationship and knowing how to address those producers is the answer, I guess.

And I will wrap up all my other comments by saying that from an industry perspective, you are absolutely committed to continue to move aggressively down that road because frankly there is fewer dollars, and you have got to be able to tie that into the total equation as we move forward.

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board.

DR. THOMAS: There was one suggestion that was discussed in the first meeting, and that was to look at a situation -- and I think that this is where you're going -- to bring industry into the process. The folks in Michigan felt it was critical to have industry as part of the advisory

As well, there's a suggestion that industry would help with funding, and I mentioned the checkoff fund, and I recognize there are issues associated with that fund, and it is very closely regulated as to what it can be used for, but I think the folks in Michigan are echoing what you are saying or what you said here. If you want industry, you can't expect industry to help with this issue if you're not going to have them at the table officially, so I just want to recognize that we have to figure out how we're going to work around that, and I think there may be options that we can, but just to recognize it is going to be a work around for us.

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UNIDENTIFIED SPEAKER: Well, I know that -- I don't think there is any way the federal government can pick up the tab on all of this, and the producers are going to lose some, a great deal in some instances whenever they get a disease but, you know, you have got to look at the situation. The government didn't give them the disease and, you know, some of these producers have done some very risky things over the years.

But, two, you have got to -- that producer needs to realize if he is a feedstock producer that stock is not worth nothing once you get TB.

And regardless if you clear it up, if you test and remove, who is going to touch those cows? A lot of people won't never come back and buy again.

So you have got to look at it basically that all of a sudden those cattle become whatever they're worth at slaughter. So that is something.

And I know the producers don't want to look at it that way, but in reality that

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is what it is. I don't think the USDA has got enough money to buy all these cattle and pay producers, what they're worth on the fair market value.

DR. HARTMAN: And we had those discussions too early on, "Your herd is not worth anything." After going through 12 herds, what I figured out is none of these people did anything wrong, and none of them deserved to have this disease or a herd worth nothing.

eradication program, you can't allow individuals to be damaged like that. So even though they all got fair market value, they were all damaged. They were set back in years in developing their herds, and during these periods of time that indemnity process was going on, they were feeding cattle with no benefit for them feeding the cattle.

And I got some pretty high numbers on what it cost them for the two months or three months to feed their cattle while we

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are doing that and all, and it is a very complicated issue.

And an example of what we did in Minnesota that may be helpful is we used federal dollars, but we used a lot of state dollars as well, and for the year 2009 there was an assessment on the sale of cattle in Minnesota of \$1 over the checkoff dollar, and that money was used -- well, it was a million dollars riding on it, and it was not used necessarily for indemnity, but it was used for cattle herd buyout, so in that buyout the producers recognized the impact this disease was having on the state, and they were willing to contribute. They did that long-term.

UNIDENTIFIED SPEAKER: Emergency checkoff.

DR. HARTMAN: He said that it was an emergency checkoff. It was developed in response to the TB issue.

DR. THOMAS: Do you still have the capability to collect or are you still collecting those checkoff funds?

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DR. HARTMAN: No, that was a one-time assessment in 2009, and that is the only time it was authorized. It would have to be authorized again as far as that.

DR. THOMAS: Any other comments or questions about indemnity?

Okay. I think we'll move on. And the next element is "interstate movement controls."

So with these regulations, we want to be sure that we have the ability for movement controls in our state or tribal area and movement control for animals in which a disease risk -- there is risk of disease transmission.

We want to have the authority to define the types and classifications of the animals and herds that might be subject to movement control, for instance, breeder animals out of a high risk area. The consequences may be applied for the lack of implementation, maintenance, or compliance with risk mitigation measures or noncompliance for other restrictions, and we recognize that in

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certain situations that the mitigation that a state puts in place may diminish or even preclude a need for controls.

A couple of examples for that would be feed lots, quarantine feed lots as another example. In addition, the use of approved disease management plans, and that latter one is a fairly specific reference to what is ongoing in GYA in their high risk zones.

In fact, they have a number of mitigation activities that include herd risk assessment, vaccination plan, movement requirements when animals are moving out of the zone, so it was just a recognition that depending on the strength of the animals, if you will, animal health plan, their disease management plan, that there is a recognition that we being the federal government may not need to institute specific movement requirements but we want to be able to have the ability to do so in the event we need to.

And then lastly, we like the administrator clause in our regulation, and

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this is actually a good thing. The administrator clause allows us to make a variance from the regulation as long as it doesn't put other animals at risk or increase the risk of disease transmission.

So we want to have the administrator clause and, again, this is another area where we feel the advisory board could play a role in providing recommendations, if you will, to veterinary services on whether or not they felt a certain state or stage should have a variance from any sort of movement controls.

So I think with that I will go ahead and open it up for questions or comments.

We want these movement controls based on the risks, and it is not necessarily associated, per se, with a certain status level although if we get comments back, but it is going to be -- we can't be flexible. We're going to have to make a stand on some of these issues.

Again, as I indicated earlier, recognizing that this is a lot of information

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to provide to you and for you to digest just within really what is a couple of hours, I encourage everybody to please provide those written comments.

DR. HARTMAN: I think this interstate movement is a big deal, and there is a thing called "federal preemption." And what that means and they're going to use it, they're proposing to use it in the animal disease traceability program, and I won't describe that to you, but there may be a purpose for it because one of the things I hear frequently from veterinarians and producers is why are the regulations different from every single state in the United States? Can't we come to some agreement on what regulations should be in place for the movement of these animals to be consistent in all given states? We have an opportunity here for all states to participate developing these regulations. Can't we come to some consensus of how we manage it so we're managing it the same, and you can't build a barrier beyond that state? It is not reasonable. We all agreed to

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this, and we're all open to this. I can
sense that there are states that are going
to say, "Wait a minute, we will decide how
we're going to protect our own cattle," but
if we do that, the harm that happens is what
happened in Minnesota and has happened to
other states is their regulations that they
put in place are not reasonable. They're
not based on risks, but they have the
authority to do that, so I think it's a real
question, "Can't we come to a consensus?"
As scientists, should we not be able to say
these are what the risks are, and these are
the interstate movement requirements we will
all agree on, and we will stick with that?
And that would be my opinion.

UNIDENTIFIED SPEAKER: Are there risks of groups that should be subject to interstate moment controls with regard to the status, and if so which one would you think?

DR. THOMAS: Can you repeat that?

UNIDENTIFIED SPEAKER: Are there

obvious groups of cattle, production-type or some other unified thing which should be

122 1 PUBLIC MEETING 2 subject to movement controls, testing 3 requirements in that regard to the state 4 status, and if so, what type are you 5 thinking for one example go to the owner 6 event stock? 7 UNIDENTIFIED SPEAKER: I quess 8 would say from some of the regional 9 discussions have come about in that class 10 specifically, if you will, animals that are 11 for exhibition or whatever it may be, and I 12 know particularly related to TB, there has 13 been some issues there with that, you know, 14 and that is when we have got to have 15 discussion, regardless of the state status, 16 of how we look at how we handle that and 17 then break them down by classic category. 18 UNIDENTIFIED: Well, I think your 19 rodeo stock is one thing. Another issue 20 that I feel very strongly about is the dairy

replacement heifers with TB, and thank God I'm one that was bull-headed not to drop my regulations whenever one went free.

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A lot of folks are scrambling trying to get the regulations back in place, but

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the problem is so many of these dairy
heifers, you look at the ear tags and you
say, well, they come from Indiana, Illinois,
that end of the country, but in reality they
could have come from anywhere, and they get
to those grow out facilities out there, and
that is where they get to them vaccinated or
get a better ear tag stuck in.

So I think dairy heifers, regardless of where they come from, I want them tested coming to Georgia if they're six months age or older.

DR. MARSHALL: I think that idea makes a lot more sense than just to say we're going to have restrictions from all animals coming from Minnesota. We can look at high-risk populations and target them. It think that is spending resources a lot more responsibly than just by the GOP political zone versus industry risk.

DR. HARTMAN: I would argue that I don't think with limited resources and who pays for that testing for TB when it goes for interstate movement, the producer. So

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instead of spending the money testing cattle
that are moving in your state, dairy heifers
or whatever, we invest in the program and
the parts that are likely to find the
disease, and there are probably other experts
that could tell you more about how many
times that has paid off.

So right now I think there is some 20-year or so states that require all dairy cattle from states to be tested that come into their state.

I would ask the questions, "Can we evaluate overtime? Can we say start at such and such date? How much did that cost the cattle industry, and how much infection did they find?" And if that is not a good way of finding infection should we do that and invest money to things in finding the disease? I don't know the answer to it, but I'm nervous that that isn't a very effective way of detecting the disease.

In Minnesota, we detected it by slaughter surveillance. In Michigan, they protected it because somebody shot a deer and

1 PUBLIC MEETING 2 found infection in it. Now they're finding 3 it by doing herd testing, but I think it is 4 rare we found the disease by testing animals 5 for interstate movement. 6 I'm not sure what the answer is, but 7 I think it needs to be evaluated, and we

need to do what is cost effective.

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DR. THOMAS: Bill mentioned the preemption issue, and I don't want to belabor that point, but it is something as we go through the development is what we want to accomplish by the claiming of preemption is that we create a level playing field for all states.

That's the intent so we just want to go on the record that we recognize that it We don't want to have 50 is an issue. states with 50 different requirements in place for the interstate movement of animals.

So the next element --

DR. HARTMAN: Before we leave that, could I ask the people that are here from Georgia what they think about that, the preemption. Is that a bad thing, a good

PUBLIC MEETING thing? And I'm not sure I know the answer, but I would be interested in your perspective. Do you think that is something we should be shooting for or should we leave it alone? DR. THOMAS: If I could just clarify, we currently in our regulations do not outright say that claim preemption, i.e., when we claim preemption is that the federal

when we claim preemption is that the feder government in essence is saying you cannot put more stringent requirements in place above what the federal standard is.

As a result of the executive order by this current administration, we actually have been directed that we have come -- we have to indicate what our position is. Are we going to claim preemption or are we not going to claim preemption? If we provide, we have to have regulatory tests as well as an explanation within what we refer to as the preamble of the regulation.

So it is not something. It's easy to stay silent on the issue, but we are being -- our current directions from this

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current administration are that we have to develop a position and state that position.

UNIDENTIFIED SPEAKER: Well, you know I look back at the days whenever we had brucellosis, and Georgia was one of the dirty dozen. It was one of the last 12 states that still had brucellosis.

If you look, if the states that wound up being the last, they're states that did just exactly the minimum that the UMR requires, and I feel like that, you know, had I been told that I shouldn't have done no more than what the UMR requires, I would not have been free in 1999. It would have been 2003, '04, '05 because we did some things that was, you know, unheard of in those days, and for many years I never admitted where a lot of it came from, but there was four of us sitting in a motel room with a fifth of Jack Daniels one night.

UNIDENTIFIED SPEAKER: This is being recorded.

UNIDENTIFIED SPEAKER: It made good sense to do some things that we had to do,

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and we had regs in sale barns that nobody -everybody said that they don't happen in our
states, but 22 known reactors through the
market one day and five months later I was
tracing infected herds, and one thing was
common, at least one of the reactors in
every herd was in that market that day.

Our commissioners, our state

veterinarian at the time said that it was

time we done something and that we stopped

known reactors going through the sale barn,

and we never had another wreck, but those

are the kind of things that if I had had

preemption, I would have still been here

sitting here worrying about brucellosis for a

lot of years back.

UNIDENTIFIED SPEAKER: I would just say that from an industry perspective, Dr.

Hartman's idea of having a sit-down and working through it from the state vet's perspective would seem to be a better solution in my mind that from a top down, this is it, and getting consensus ability, but from our perspective I don't have

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heartburn. We want the markets to work, and we went the cattle to be able to move, and we want them to be healthy, you know, and that is whatever it takes.

DR. HARTMAN: I think what would be important and you brought it up -- if there was a federal preemption, the federal government would have to be amenable to making changes in their program like you did quickly so that if there was, if there were things going on that it could be adjusted, but if there was availability it could address some of those issues, and I think that is also where an advisory board would come in and they would be able to look at those situations and say, "Yes, we need to make some changes to this quickly because this is not working."

UNIDENTIFIED SPEAKER: Well, even whenever the USDA developed their White Paper on brucellosis and made the recommendations that basically was Georgia's program, it was still a recommendation, and nobody had -- it was recommended that everybody follow that,

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but it was not mandatory that everybody would follow that.

So, you know, if they're going to make changes and it is going to have preemption, then they have to be amenable to making those changes in a timely fashion because, like I say, I went over a period of five or six years, and I went to three meetings, one in Memphis, Tennessee, one up here in north Georgia, Unicoi State Park, and then the final meeting, the third meeting was down at Southern Animal Health in Pointe Clear, Alabama, and they have all billed as the "Big Brucellosisment."

DR. THOMAS: Oh, I can't resist, but actually at USAHA we had had discussion, and we had a -- we were claiming the farewell to bovine brucellosis, and then we had a heart, so we're delaying that because luckily our last herds have been what I have referred to as the high risk areas, but you do go out on a limb when you say "this is it," and you find a herd the next day.

So with that, I would like to move

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on to importational requirements. We have broken these up really based on the activities and where those activities occur into three different areas, pre-import, import, and post-import.

Under pre-import, we will review a country or zone based on the 11 factors and 9 CFR, part 92.

And those regionalization regs that exist today. We're not proposing any changes there. We want to use that existing framework that allows us based on a country's request to evaluate their ability to import animals and products in some situations into the U.S., and based on that evaluation we would determine what import requirements would be required to safely import animals into the U.S.

We would monitor the country's zone and changes for changes that would trigger potentially increase and import requirements. This is a rough high level description of how we're handling Mexico now. So there are a lot of similarities for what we're

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proposing to the new regulations that exists; however, we're not proposing that we would use the existing status system.

What we're looking at is when we go in and we evaluate, we want to see what the risks are from a state or zone and then have based on those risks, if you will, have some standard requirements in place for import.

If you'll notice -- and this is by intent -- we're talking about Mexican import requirement. We want to look at our regs specifically for the import of all animals.

For those countries or zones that have not been evaluated, we want to determine if there are mitigations that could be put in place, that is, it allow some level of imports. We currently have that in place for our TB regs, our importation related, requirements related to TB. If a state in the situation has not been evaluated, those animals can come in, but it is only for immediate slaughter only, and in that specific incident related to what we had in place for Mexico.

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So moving from pre-import, we have importation, and we want to ensure that the first point of concentration or mingling after entry is identified on that import paperwork, and that records are maintained to facilitate the tracing of animals.

And then further if those animals are moved in interstate, after that first point of concentration, then the state or destination must be notified.

Next slide.

For post-import, the general work requirements include the continuity of official indication, be that with ear tag and/or the paperwork that is following the animal.

We would hope that official ID that comes in with the animals is not removed; although, we have some concerns that actually ID is being removed.

For those animals that move in interstate commerce, after that first point of concentration, we want an interstate certificate or brand inspection for that

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movement, and for post-entry restrictions and long-term testing requirements, we want to ensure that imported steers and spayed heifers that enter the U.S. as part of the food production chain, that they're maintained separately from domestic breeding cattle during pasturing. We are considering some component of pasturing in this new regulatory framework based on the feedback we're getting back. That was somewhat redundant, sorry about that.

Then here is the concept about the periodic testing of event and rodeo cattle. It would be helpful to get your comments about the testing of this particular commodity group.

It was actually a difference of opinion on dairy heifers for interstate movement, but what about periodic testing of event rodeo cattle? The majority of these animals are imported from Mexico. We do occasionally find disease in these animals. Is this a group that some sort of periodic testing or interstate movement requirements

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should be imposed?

NIDENTIFIED SPEAKER: I don't know how we're going to accomplish it but we get -- I'm sure, Ken, y'all do, too -- we get steers that are Mexican steers, and they come in from, to a point when they cross the border, and then they may get moved somewhere else in Texas, Louisiana or wherever, and they're broken up from that, and we don't even get a health certificate coming in, so it is a nightmare.

DR. THOMAS: Do you want that health certificate? Or would you like it?

UNIDENTIFIED SPEAKER: I would like it. I know we had a trace back on a steer that turned up positive, and they traced the cattle from Georgia, and I don't know where else, but they went through a dealer in Kentucky, and we had to go look, and they did finally tell us that it looked like it was Mexican origin TB on that steer, and we went and looked and found two places that there had been Mexican steers, and one of them by the time we got trace back was not

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the same set of steers that was there at the time that this animal would have been exposed, and those animals had gone on and probably been slaughtered by then, but that is the kind of thing that happens, but we've no way of knowing when or where those animals were tested or what was tested after they crossed the border.

DR. HARTMAN: I think this may be one of the most important questions we ask. The bacteria we found in Minnesota was most similar to what is found in the southwestern United States and Mexico, and Minnesota might have been TB free and not had this episode if we had better controls on what happens with cattle coming in for Mexico.

And, again, I acknowledge everything that Mexico is doing, and they're doing a lot to try and eradicate the disease, but it is more advanced in certain parts of Mexico than it is in others.

I think you have hit a lot of the key points on here what to do. I mean, we do allow TB in from Mexico every single

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year. The number of cases has gone down dramatically, but in my mind it is not where it should be until it is zero. We need to make sure that we're not importing this disease from another country. We have got enough to deal with already in the United States with developing penicillin, and a very important question, and I think we really need to examine all new, regulating animals that come in that could be potentially dangerous to others.

DR. THOMAS: Any comments about the concept of having imported cattle, the concept of pasturing? Do you feel you can safely pasture our domestic breeding stock, not pulsing volt, but fence line or ten-feet separation?

UNIDENTIFIED SPEAKER: Well, I think we all know that the majority of the Mexican stock that comes in is not ready to go to the feed lot, and they're grazed for a period of time.

And I don't know what the distance should be, but they have got to be kept away

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from the domestic stock and variably if somebody has got 30 herd of cow out there and then they get a bunch of Mexican steer and they're going to graze them for a period of time while they're held up in the same pasture, and that is where we're getting exposure, but I don't know what the distance should be, but there is going to have to be some separation in there to keep them away from breeding herds.

DR. HARTMAN: I forgot one thing.

When Minnesota dropped its status from accredited free and modified to accredited, and we had a lower status in some states and in Mexico, and our restrictions to sell cattle in the state were higher than they were from Mexico.

So we have to level the playing field, and I don't know how it can be done or if it can be done, but I think it is an important issue.

DR. THOMAS: Any other comments or questions about import? If not, we'll move on to the last element.

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"a lack of interest," but I think there is one area that because of the diagnosis, particularly of TB, is for the most part coming from our National Veterinary Service's Lab, so we tend to feel that it is of critical importance really for any disease program to define what is the official test, what are the official tests, and where does the testing take place and under what circumstances?

So for the last element, we will provide definitions for appropriate terms or pertinent terms, and we will include them.

We're trying to be forward thinking in those definitions.

There are diagnostic tests that are purported to be of use, of possible use as a Penside test, and so we want to ensure we're forward thinking in our definitions to include terms, such terms, for instance, as a "Penside test." They will provide for a process of initial approval and recertification of continued approval, official

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diagnostic tests, official la	boratories, and
when appropriate, official tes	sters, and that
official tester signs back to	inside
technology as well as the cu	rrent use of a
test for brucellosis, so we'r	re recognizing
what we're currently doing un	nder certain
circumstances what we may be	doing in the
future for improvements on di	agnostic tests.

And the changes to the process for approval for laboratory or diagnostic tests, we would propose to indicate the regs, how that change would take place, but we don't want to necessarily include that specific change in the regulations. This is an area where we use our program standards, and we'll notify the public of a change and allow them the opportunity to provide comment, but using such short -- that notification and comment is much quicker than a rule change, a proposed rule change with a final rule.

One of the differences that I can almost guarantee you you will see is currently under the brucellosis regs, and we have a significant amount of information

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about the diagnostic tests, how the tests are run, what are the cutoffs, and all that type of information would be included in the program standards as opposed to the regs again for the flexibility that it offers us, and the regs will only contain those terms they have to.

So any questions or comments?

Actually, I have another slide.

There will be a mechanism to withdraw

laboratory or suspend its approval, and the regs will reference the need for quality assurance and quality control for testing laboratories as well as proficiency testing of authorized personnel to conduct the tests.

So any questions about laboratory approval or official tests? Did any questions come up during the break regarding the elements that we discussed during the break? So any general thoughts about this framework?

Well, recognizing it is a lot of information to digest, and in the information packet that you received, you received the

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public notice and it has the sites and the mechanisms by which you can submit your comments.

So I would just encourage everybody to submit your written comments and read the framework that was provided actually in your packet as well as it is available online, and just provide us your feedback.

The timeline for this process is that we plan on getting a proposed rule out in the fiscal year -- not fiscal year, but the timeline is to get a proposed rule out in 2011 and a final rule out in the calendar year of 2012, so it is a very ambitious timeline, and that is why your written comments are so important because we don't want to put a rule out there that you can't live with or you send us comments that suggest what are you doing. So we don't want any surprises.

Another question that came up that I have mentioned is the program standards and how would you know what those program standards are? The program standards will be

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published along with the actual proposal, so you will have the opportunity to see the program standards.

And in case you're wondering what I'm referring to, the program standards, those program standards are what we commonly refer to now as the uniform method and rules, the UMRs, and there are other policy documents, VS notices. We're looking at incorporating and providing general directions that support the implementation of this program in the form of those program standards, so they will be available for your comments as well when the rule comes out, and many of them will be similar to those that exist today, particularly in the areas of infected herd management, epi investigation, interstate movement, the diagnostic tests, official tests, and so we want to make sure that we keep those items that are working for the program and improve those that need improving.

So with that, if there are not any additional comments or questions I just want

1	144 PUBLIC MEETING
2	to thank everybody for their participation
3	today. I really appreciate it.
4	(Whereupon, this Public Meeting was
5	concluded at 12:30 p.m.)
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145 1 PUBLIC MEETING 2 CERTIFICATE 3 4 STATE OF GEORGIA: 5 COUNTY OF FULTON: 6 7 I hereby certify that the foregoing 8 transcript was reported, as stated in the 9 caption, and the questions and answers 10 thereto were reduced to typewriting under my 11 direction; that the foregoing pages represent a true, complete and correct transcript of 12 13 the evidence given upon said hearing, and I 14 further certify that I am not of kin or 15 counsel to the parties in the case; am not 16 in the employ of counsel for any of said 17 parties; nor am I in any way interested in 18 the result of said case. 19 Matalo Hal Shecton 20 21 22 NATALIE GAIL SHECKTON 23 24 DATED: JUNE 4, 2011

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