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Suwalsky, Connie

From: Porretta, Mary
Sent: Wednesday, May 19, 2004 3:42 PM
To: Suwalsky, Connie
Subject: FW: Comments for FSIS BSE regulations

Connie—There are some comments on the AMR rule in this set of comments.

-----Original Message-----

From: Smith, Kevin [mailto:KSmith@usmef.org]
Sent: Friday, May 14, 2004 10:43 AM
To: 'Mary.porretta@fsis.usda.gov'
Subject: Comments for FSIS BSE regulations

Thank you!

May 7, 2004



FSIS Docket Clerk, Docket #01-033DF,
Room 102, Cotton Annex, 300 12th and C Street, SW
Washington, DC 20250-3700

*Putting U.S. Meat
On the World's
Table
Since 1976*

The U.S. Meat Export Federation (USMEF) appreciates the opportunity to provide comments to FSIS in regards to the Emergency Final Rules (Docket numbers: 03-025IF, 03-048N, 01-033IF, and 03-038IF) which have been enacted in response to the discovery of one Canadian origin dairy cow in Washington State, USA which was diagnosed and confirmed to have Bovine Spongiform Encephalopathy. The USMEF is a non-profit trade association working to create new opportunities and develop existing international markets for U.S. beef, pork, lamb and veal. The USMEF has eight distinct sectors, representing the entire U.S. red meat production, processing and distribution system. Allied industries, which provide critical inputs to the red meat industry, are also active on the USMEF Board of Directors. Over 90 percent of U.S. red meat exports are from red meat companies represented by USMEF. USMEF provides market intelligence for international markets and addresses concerns or problems faced by its membership in these international markets. USMEF commends the Food Safety Inspection Service along with other agencies of the United States Department of Agriculture (USDA) for their rapid response to this issue and for the means by which the investigation was conducted along with the efficient methods by which information and results of the investigation were conveyed. While we commend FSIS for the speed at which the final rules were implemented under emergency conditions, we have several questions and comments surrounding several of the details of the rules.

Denver

Hong Kong

London

Mexico City

Monterrey

Moscow

Osaka

Seoul

Shanghai

Singapore

Our comments and questions are located in the following pages and in the additional attachment accompanying this document. We apologize for sending an attachment to these comments; however, due to the detailed photographs which are included in the comments and the sheer size of the file, we have no choice but to send the

July 1, 2003

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St. Petersburg attachment in Adobe Acrobat format. Thank you for your attention to our concerns.

Taipei Sincerely

Tokyo

Philip M. Seng
President/CEO
U.S. Meat Export Federation

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9 CFR Parts 309, 310, 311, 318, and 319
Prohibition of the Use of Specified Risk Materials for Human Food and Requirements for the
Disposition of Non-Ambulatory Disabled Cattle

The definition for Specified Risk Materials (SRM) is consistent with the Canadian definition and it is stated on this page that this was a purposeful decision. We question the reasons behind patterning the United States definition of Specified Risk Materials following the conclusive results that the origin of the index animal was in fact Alberta, Canada. This would indicate that the United States is still without an indigenous case of BSE while Canada could now claim two indigenous cases. It would appear that the response and the definition of the SRM material within the United States should be based on the most current peer reviewed science. It is USMEF's understanding that countries which wish to export beef and beef products to the United States must also comply with this new rule. How will FSIS provide assurances that products imported by the United States were produced in compliance with the new regulations? Will there be surveillance, auditing, or monitoring to ensure compliance?

Has a date been determined when FSIS will publish compliance guidelines for use by small and very small establishments?

As written, there will be two ways which be allowed for use in determination of animal age; one through the use of documentation and the other through the use of dentition of each individual animal. It is also stated that because the National Animal Identification System for the U.S. is not yet complete there is not a uniform standard of documentation which FSIS can rely on to verify the age of cattle slaughtered. There are certain establishments within the U.S. which have company-specific animal identification systems in place. Will FSIS accept private systems of animal identification and private certification of animal age at the time of slaughter or must certification be supplied by an entity within USDA such as the Agriculture Marketing Service?

What is the basis for the FSIS calculation for the estimate of burden to be placed on establishments? We also ask if estimates for burden have been calculated for small and very small establishments.

We request that FSIS review pertinent research regarding the distal ileum of the small intestine and we request that FSIS allow for the separation of the distal ileum from the remaining portion of the small intestine (jejunum and duodenum) and that the remaining portion of the small intestine be allowed for processing and human consumption.

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Comments regarding the removal of the distal ileum of the small intestine are attached in a separate document. Due to the detailed photographs which accompany the attached file, the sheer size of the file makes it necessary to condense the information into an Adobe Acrobat format.

9 CFR Parts 301, 318, and 320

Meat Produced by Advanced Meat/Bone Separation Machinery and meat Recover (AMR) Systems

This rule states that AMR product may not contain any brain, trigeminal ganglia, spinal cord, or dorsal root ganglia. Is the detection limit or "allowable" level of SRM material the same as published in previous regulations? Will the measurement of these substances be strictly under the jurisdiction of USDA or will individual entities have the ability to test and certify that products meet the requirements of this final rule?

It is our understanding that the brain, spinal cord, trigeminal ganglia, and dorsal root ganglia are not Specified Risk Materials if they originate from animals under the age of 30 months. If they are not SRM products, it is our understanding that their presence in AMR product would represent a "misbranding" or a labeling issue and not a food safety concern and thus they would be handling accordingly. Why does AMR product produced from materials derived from animals less than 30 months of age need to be part of a HACCP plan if this represents a misbranding issue and not a food safety concern?

As vertebral column is defined as a Specified Risk Material only from animals over the age of 30 months, USMEF feels that beef stock, beef extracts, and other beef products derived from bones, should be able to include the vertebral column from animals under the age of 30 months. This product should not be required to be identified as having the potential to contain CNS tissues. Vertebral column bones derived from animals under the age of 30 months is not considered a Specified Risk Material and thus poses extremely minimal risk to humans.

We also request clarification as to the meaning of the statement made which reads: "FSIS will ensure that materials that could present a significant risk to human health, but whose infectivity status cannot be readily ascertained are excluded from the human food supply".

What is the basis for the FSIS calculation of the estimate of burden? Was a calculation of the estimate of burden conducted for small and very small establishments?

9 CFR Part 313

Prohibition of the Use of Certain Stunning Devices Used to Immobilize Cattle During Slaughter

As it is stated in this document, FSIS is not aware of any United States slaughter establishment utilizing air injection stunning equipment. As this is the case, what is the purpose of implementing a rule which prohibits the use of such equipment? It would appear that this rule would be unnecessary and that it would be perhaps detrimental to consumer confidence in the safety of US beef production systems.

FSIS indicated having knowledge that air-injection stunning devices are available and allowable in countries that may export beef products to the United States. How will FSIS assure compliance to this regulation in foreign countries?

FSIS indicates that Australian law does not ban the use of air-injection stunning methods. Will FSIS restrict the importation of Australian beef products if the Australian law continues to be inconsistent with U.S. regulations regarding air-injection stunning?

USMEF concurs with FSIS in their decision to reject the option of a performance standard. A performance standard and testing for CNS emboli would be costly and unwieldy to both industry and government enforcement officials.

Bovine Spongiform Encephalopathy Surveillance Program

As stated in the document, FSIS will not apply the mark of inspection to products until the results of the APHIS BSE testing was known by FSIS and the establishment. As it is now known that the carcasses of tested animals will be "held" until results of BSE testing are known, does this apply to all products from the tested animals? Must all offal, viscera, hide, etc. be held along with the carcass of the tested animal until results of the APHIS test are known? May items which are destined for inedible rendering be allowed to pass through to inedible rendering prior to the receiving of the results of the BSE test as these products will be excluded from the human and ruminant food chain?

Preliminary Analysis of Interim Final Rules and An Interpretive rule to Prevent the BSE Agent From Entering the U.S. Food Supply

Page 21: Table 1. Annual estimates of average SRM amounts affected by the SRM Interim final rule.

USMEF requests the origin or basis for the USDA estimate of 11 pounds yielded per animal of small intestine (including the distal ileum). USMEF estimates of this weight have been greater than 12 pounds. A one pound increase would increase the volume by 14.5 million pounds.

Page 23: SRMs excluded from the human food supply.

USMEF is not in favor of any of the alternatives presented by FSIS. USMEF feels that Alternative 3 would be the most favorable, however, USMEF feels that a procedure should be available which would allow for the removal of only the distal ileum. Please see our comments on page three along with the accompanying attached document. This would allow the salvage of the remainder of the small intestine.

Page 25 - Use of 'Trepas' and price

In *Table 4. Average net revenue loses due to exclusion of SRMs under the Interim Final Rule*, small intestine is classified as Casings and Trepas. Trepas are an export product for Mexico, and since edible intestines are also exported to Japan and some other markets, we request clarification on the quantity and analysis for the Trepas classification. Does this only include product destined for Mexico, or does the analysis for Trepas include small intestine destined for other markets also?

Also, the price per pound for Trepas is listed at \$0.37 cents. USMEF's analysis shows Trepas priced at approximately \$0.55 cents per pound in December 2003. We request verification of Trepas values prior to the BSE discovery.

Page 26: Table 5. Comparison of Average Change in Potential Human Exposure and cost of Regulatory Alternatives.

USMEF would like to make the point that the removal of the small intestine does not make a significant impact in the reduction in human exposure; however, the removal of the small intestine has a substantial impact on the cost. USMEF feels that a procedure should be available which would allow for the removal of only the distal ileum. Please see our comments on page three along with the accompanying attached document. This would allow the salvage of the remainder of the small intestine.

Page 31 Salvage Value of Non-ambulatory Disabled Cattle

In *Table 6 - Cost of prohibiting use of non-ambulatory disabled cattle from human food use*, Salvage value is shown to be zero. Under the proposed rule, these animals would be eligible for inedible rendering. USMEF requests clarification on why the proposed analysis does not consider the rendering value of these animals.

Page 33: HACCP plan development, record keeping and certification.

While USDA has estimated the costs associated with development of the programs to implement the interim final rules, USMEF feels that there are many additional costs incurred by U.S. companies which have not been included. USMEF feels that additional labor, employees, and training costs should be included in this analysis and these costs impact the U.S. industry.

Page 36 Cost impacts of SRM interim final rule

In *Table 8 - Summary: Cost impacts of the SRM interim final rule*, the estimated costs of SRM ban and regulations averages \$37.1 million. A recent study conducted by *Cattle-Fax* estimates costs related to SRM removal to be approximately \$100 million. There is a considerable disparity between these numbers. USMEF requests a more comprehensive analysis comparing the two estimates which would include input from industry resources.

Page 43: Domestic Economic Impacts

The impact of BSE on changing U.S. beef production levels may be insignificant due to the supply situation prior to the discovery. The U.S. cowherd is currently in the eighth year of contraction which drove prices to record levels prior to the BSE discovery. Although the discovery of BSE has impacted prices negatively, they remain well above previous year prices at levels typically needed to spur heifer retention and eventually, production increases.

U.S. Meat Export Federation estimates the negative price impact on live cattle due to the BSE discovery at \$10.25 per hundredweight (cwt). This scenario is calculated by estimating impact of additional weekly pounds of beef supplied domestically rather than exported and adding to it the lost premiums paid for variety meats internationally. The calculations are shown below:

Loss of premiums paid for variety meats:

	Pro BSE (December) Price *	March Price *	Value loss per pound	Million lbs produced **	Lost Value
Tongues	3.80	0.90	\$ (0.90)	100	\$ (291,400.500)
Intestine (Tripe)	0.55	0.03	\$ (0.54)	215	\$ (116,275.500)
Skin	3.11	2.00	\$ (1.11)	201	\$ (223,070.700)
Liver	0.35	0.10	\$ (0.25)	318	\$ (78,952.500)
Tail	0.90	0.27	\$ (0.72)	215	\$ (155,031.000)
					\$ (664,745.200)
			U.S. commercial slaughter (excluding crossbreds)	28,710,000	

* Prices from USDA reports in December 2002 and January 2003
 ** Based on U.S. Heifer/Steer Slaughter

Value / Hd
\$ (0.32)
{0}
\$ (2.11)
\$/cwt live

Combined Price Impact Including Supply Effects:

x

Although USMEF is optimistic regarding the current BSE bans, the 2004 estimated annual flow used in the previous table assumes that current bans will continue through 2004. This is done to estimate annual losses from the bans currently in place.

The combined calculations in the previous tables show a decrease in live cattle prices by approximately \$10 per hundredweight due to the discovery of BSE.

Page 58 Restoration of Beef Export Markets

The text states, "About 40 countries have banned beef from the United States." We have determined that approximately 64 countries initially banned U.S. beef products and that currently 15 of these markets have reopened to some degree. We request an update of this statistic in the text.