



United States
Department of
Agriculture

Food Safety
and Inspection
Service

Washington, D.C.
20250

MAY 28 2002

Dr. Halldor Runolfsson
Chief Veterinary Officer
Ministry of Agriculture
Solvholgata 7
150 Reykjavik
Iceland

Dear Dr. Runolfsson:

The Food Safety and Inspection Service (FSIS) has completed an on-site audit of Iceland's meat inspection system. The audit was conducted from October 8-18, 2001. Enclosed is a copy of the final audit report. Iceland's comments have been included as Attachment G.

During this audit, the FSIS auditor noted several deficiencies. These deficiencies were discussed in our February 4, 2002, letter to you, which transmitted the draft final audit report. During our next audit of Iceland, which is scheduled for August 2002, FSIS expects that lamb heads will not be removed prior to veterinary disposition, that findings of inadequate HACCP implementation have been corrected, and that Iceland is using an AOAC-approved method for testing samples for generic *Escherichia coli*.

If you have any questions regarding the audit or need additional information, please contact me at 202-720-3781. My fax number is 202-690-4040 and you may also reach me by email at sally.stratmoen@fsis.usda.gov.

Sincerely,

/s/ Sally Stratmoen
Chief, Equivalence Section
International Policy Staff
Office of Policy, Program Development
and Evaluation

Enclosure

cc:

Philip Letarte, Agriculture Counselor, FAS, American Embassy, Denmark
Gundi Bragason, Minister Counselor, Agriculture Affairs, Embassy of Iceland
Borghildur Magnúsdóttir, FAS, US Embassy, Reykjavik
Ed Brown, Economic/Commercial Counselor, US Embassy, Reykjavik
Linda Swacina, Acting AA, FSIS
Martiza Colon-Pullano, SAIIS, OPPDE
John Wilson, Area Officer, FAS
John Prucha, ADA, OPPDE
Karen Stuck, IEPS, IPS, OPPDE
Donald Smart, Director, Review Staff, OFO
Amy Winton, State Department
Nancy Goodwin, ES, IPS, OPPDE
Country File (Iceland—FY 2002 draft final audit report to CVO)

FSIS:OPPDE:IPS:ES:N Goodwin:bw:5/23/02:720-9187:5/20/02:Iceland-FY 2002 final audit report to CVO



United States
Department of
Agriculture

Food Safety
And Inspection
Service

Technical
Service
Center

Suite 300, Landmark Center
1299 Farnam Street
Omaha, NE 68102

AUDIT REPORT FOR ICELAND OCTOBER 9 THROUGH OCTOBER 18, 2001

INTRODUCTION

Background

This report reflects information that was obtained during an audit of Iceland's meat inspection system from October 8 through October 18, 2001. Five establishments are certified to export meat to the United States and all were audited. All of these were slaughter establishments and conduct some further processing.

The last audit of the Icelandic meat inspection system was conducted in October 2000. Five establishments were audited and all were acceptable. These establishments were 22, 23, 31, 40 and 81. The following concerns were reported at that time: HACCP was poorly understood in all plants, especially the measurement of critical control limits, and pre-shipment review was not being done in any plants. The residue control program had an excessive turn-around time for results being in excess of four months. Species testing on finished product was not being done.

Iceland is eligible to export beef, pork and sheep meat to the United States at this time.

From January to September 30, 2001, Iceland establishments exported nearly 5,000 pounds of lamb to the U.S. Port-of-entry rejections were for missing shipping marks (0.6%).

PROTOCOL

This on-site audit was conducted in four parts. One part involved visits with Icelandic national meat inspection officials to discuss oversight programs and practices, including enforcement activities. The second entailed an audit of a selection of records in the meat inspection headquarters facilities preceding the on-site visits. The third was conducted by on-site visits to establishments. The fourth was a visit to two laboratories, one performing analytical testing of field samples for the national residue testing program, and the other conducting species testing on finished product and antibiotic screening tests. A farm was also visited and questions were asked about the use of antibiotics, vaccines and other chemical compounds and the procedures in place to prevent these chemicals from entering the food chain.

Iceland's program effectiveness was assessed by evaluating five areas of risk: (1) sanitation controls, including the implementation and operation of Sanitation Standard Operating Procedures (SSOPs), (2) animal disease controls, (3) residue controls, (4) slaughter/processing controls, including the implementation and operation of Hazard Analysis and Critical Control Point (HACCP) systems and the *E. coli* testing program, and (5) enforcement controls.

During all on-site establishment visits, the auditor evaluated the nature, extent, and degree to which findings impacted on food safety and public health, as well as overall program delivery. The auditor also determined if establishment and inspection system controls were in place. Establishments that do not have effective controls in place to prevent, detect and eliminate product contamination/adulteration are considered unacceptable and therefore ineligible to export products to the U.S., and are delisted accordingly by the country's meat inspection officials. (This was the case with one establishment—see below.)

RESULTS AND DISCUSSION

Summary

Effective inspection system controls were found to be in place in four of the five establishments audited. One establishment (40) was found to be unacceptable. Details of audit findings, including compliance with HACCP, SSOPs, and testing program for generic *E. coli*, are discussed later in this report.

As stated above, some concerns had been identified during the last audit of the Icelandic meat inspection system, conducted in October 2000. During this new audit, the auditor determined that these concerns had been addressed and corrected with the exception of pre-shipment review, which was still not being done.

Entrance Meeting

On October 9, 2001, an entrance meeting was held in the Reykjavik offices of the Icelandic Meat Inspection Division of the Ministry of Agriculture, and was attended by Dr. Sigurður Örn Hansson, Chief of Iceland Meat Inspection; Mr. Edwin Brown, Economic/Commercial Officer, U. S. Embassy; Ms. Borghildur Magsúsdóttir, Assistant Economic/Commercial Officer, U.S. Embassy and Dr. M. Douglas Parks, International Audit Staff Officer, USDA.

Topics of discussion included the following:

1. Audit itinerary.
2. Species testing of finished product.
3. Animal disease status.
4. Compliance and enforcement of meat regulations.
5. Subjects to be covered on the audit (SSOP, HACCP and generic *E. coli* testing).

6. Farm visit, the national residue program and the turn around time for residue sample results.

Headquarters Audit

There had been no changes in the organizational structure or upper levels of inspection staffing since the last U.S. audit of Iceland's inspection system in October 2000.

To gain an accurate overview of the effectiveness of inspection controls, FSIS requested that the audits of the individual establishments be led by the inspection officials who normally conduct the periodic reviews for compliance with U.S. specifications. The FSIS auditor (hereinafter called "the auditor") observed and evaluated the process.

The auditor conducted a review of inspection system documents at the time of the on-site audit. The records review focused primarily on food safety hazards and included the following:

- Internal review reports.
- Supervisory visits to establishments that were certified to export to the U.S.
- Training records for inspectors and laboratory personnel.
- Label approval records such as generic labels, and animal raising claims.
- New laws and implementation documents such as regulations, notices, directives and guidelines.
- Sampling and laboratory analyses for residues.
- Pathogen reduction and other food safety initiatives such as SSOPs, HACCP programs and generic *E. coli* testing.
- Sanitation, slaughter and processing inspection procedures and standards.
- Control of products from livestock with conditions such as tuberculosis, cysticercosis, etc., and of inedible and condemned materials.
- Export product inspection and control including export certificates.
- Enforcement records, including examples of criminal prosecution, consumer complaints, recalls, seizure and control of noncompliant product, and withholding, suspending, withdrawing inspection services from or delisting an establishment that is certified to export product to the United States.

No concerns arose as a result the examination of these documents.

Government Oversight

All inspection veterinarians and inspectors in establishments certified by Iceland as eligible to export meat products to the United States were full-time Inspection Service employees, receiving no remuneration from either industry or establishment personnel.

Establishment Audits

Five establishments were certified to export meat products to the United States at the time this audit was conducted. All five establishments were visited for on-site audits. In four of the five establishments visited, both Iceland Inspection Service inspection system controls and establishment system controls were in place to prevent, detect and control contamination and adulteration of products.

Laboratory Audits

During the laboratory audits, emphasis was placed on the application of procedures and standards that were equivalent to U.S. requirements. Information was also collected about the risk areas of government oversight of accredited, approved, and private laboratories, and *intra*-laboratory quality assurance procedures, including sample handling and methodology.

The Icelandic Fisheries Laboratory in Reykjavik was audited on October 9, 2001. Except as noted below, effective controls were in place for sample handling and frequency, timely analysis, data reporting, tissue matrices for analysis, equipment operation and printouts, minimum detection levels, recovery frequency, percent recoveries, and corrective actions. The methods used for the analyses were acceptable. No compositing of samples was done.

The Institute for Experimental Pathology Laboratory in Reykjavik was audited on October 17, 2001. This laboratory does species testing of finished product samples sent from the exporting establishments each month that product for export to the U.S. is produced. It also does microbiological screening tests for antibiotic residues on samples sent from the exporting establishments as directed by the national residue program. Any positive screen test is sent to Denmark for identification of the antibiotic residue. Results are returned in about two weeks.

The check sample program for both laboratories did meet FSIS requirements. Some substances used for standard solutions preparation were outdated and daily working solutions were not dated in the Fisheries Laboratory and the *Bacillus* spores used for the screen tests were outdated at the Pathology Laboratory.

Microbiological testing for *Salmonella* is not applicable in Iceland, as sheep products are the only meat products exported to the United States. Generic *E. coli* testing for minor species is done in private laboratories in the slaughter establishments.

Establishment Operations by Establishment Number

The following operations were being conducted in the five establishments:

Beef, sheep, swine and horse slaughter and boning – two establishments (23 and 81)

Beef, sheep and horse slaughter and boning – one establishments (22)

Beef, sheep and swine slaughter and boning—one establishment (40)

Sheep slaughter and boning—one establishment (31)

SANITATION CONTROLS

Based on the on-site audits of establishments, Iceland's inspection system had controls in place for water potability, chlorination procedures, back siphonage prevention, hand washing facilities, sanitizers, pest control program and monitoring, temperature control, lighting, operations work space, inspector work space, and ventilation.

Sanitation Standard Operating Procedures (SSOPs)

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOPs were met, according to the criteria employed in the U.S. domestic inspection program. The data collection instrument used accompanies this report (Attachment A).

The SSOPs were found to meet the basic FSIS regulatory requirements, with only occasional minor variations, except in Establishment 40. In Establishment 40, there was a major problem with condensation falling on exposed carcasses in the carcass cooler. Condensate, from overhead structures not cleaned and sanitized daily was falling on approximately 25% of 450 carcasses in the cooler. These carcasses were removed to another area and trimmed before going to the boning room.

Cross-Contamination

The bung dropping procedure on the lamb slaughter lines of four of the five establishments visited was resulting in fecal contamination. However, contamination was being trimmed prior to the CCP for zero tolerance. These were Establishments 22, 31, 40 and 81. A different procedure will be investigated as soon as possible, was the commitment by the inspection officials.

Product Handling and Storage

Meat products in plastic bags were found to be stored on the floor of the freezer in Establishment 40. This will be remedied very soon.

Personnel Hygiene and Practices

The procedures for personnel hygiene were in place in all establishments and were effective.

The sanitation control findings that are of major concern, and the proposed actions are as follows:

1. Bung drop procedure—A new system will be devised as soon as possible in all plants.
2. Condensate falling on exposed product—The carcasses were removed to another area and trimmed. The cause in Establishment 40 was a switch that was turned off by unauthorized personnel. The switch will be secured.
3. Over-spray at the carcass wash in the slaughter department was falling from overhead structures onto exposed carcasses in two establishments (40 and 81). The procedure was changed to prevent this from happening.
4. The final trim station was not properly manned due to an accident (Est. 22) or the trimming was not completely effective (Ests. 81 and 23). These problems were given attention and solved at once and will have closer supervision in the future.

ANIMAL DISEASE CONTROLS

Iceland's inspection system had controls in place to ensure adequate animal identification, ante-mortem and post-mortem inspection procedures and dispositions, condemned and restricted product control, and procedures for sanitary handling of returned and rework product.

There is one area of concern in the slaughter area pertaining to post mortem procedures. The heads of all lambs slaughtered are removed after bleeding and are not available for diagnosis and veterinary disposition with the carcass.

There were reported to have been no outbreaks of animal diseases with public-health significance since the previous U.S. audit.

All animals in Iceland have individual identification for trace back of disease and residue.

RESIDUE CONTROLS

Iceland's National Residue Testing Plan for 2001 was being followed, and was on schedule. The Icelandic inspection system had adequate controls in place to ensure compliance with sampling and reporting procedures and storage and use of chemicals.

A sheep farm near Hvammstangi was visited and the owner and wife were interviewed along with their attending veterinarian. The Chief of Meat Inspection, Dr Sigurd Hansson was also present. There was mandatory individual animal identification soon after birth. Tags and ear cuts or notches insure trace back of any animal to its origin. There is almost no movement of livestock from farm to farm and it is closely regulated by Icelandic law. All antibiotics,

vaccines and other chemicals are only sold on the veterinarian's prescription and each carries a withdrawal notice from the veterinarian. Each treatment is recorded and the books were complete and up-to-date. There are no central markets for livestock in Iceland so all are sold directly to the slaughtering establishment.

SLAUGHTER/PROCESSING CONTROLS

The Icelandic inspection system had controls in place to ensure adequate ante-and post-mortem inspection procedures and dispositions, control and disposition of dead, dying, diseased or disabled animals, humane handling and slaughter and disposition of inedible materials generated in the establishments.

HACCP Implementation

All establishments approved to export meat products to the U.S. are required to have developed and implemented a Hazard Analysis – Critical Control Point (HACCP) system. Each of these systems was evaluated according to the criteria employed in the U.S. domestic inspection program. The data collection instrument used accompanies this report (Attachment B).

The HACCP programs were found to meet the basic FSIS regulatory requirements with the following exceptions:

1. Critical control limits were not measurable for one CCP (dirt on wool) in all establishments, they were judgmental. The critical control points that were involved were changed to control points (CP) or good manufacturing procedures (GMP).
2. Preventative actions were not recorded in Establishments 40, 81 and 23. These were initiated immediately.
3. Verification procedures were not written into the program in Establishments 31, 23 and 22. This was corrected immediately.
4. Pre-shipment review was not done in all establishments audited. The procedure is now being done in all establishments as a result of this audit.

Testing for Generic *E. coli*

Iceland has adopted the FSIS regulatory requirements for generic *E. coli* testing.

All five of the establishments audited were required to meet the basic FSIS regulatory requirements for generic *E. coli* testing, and were audited and evaluated according to the criteria employed in the U.S. domestic inspection program. The data collection instrument used accompanies this report (Attachment C).

The *E. coli* testing programs were found to meet the basic FSIS regulatory requirements with one exception:

1. Two establishments were using testing methods that were neither AOAC approved nor were the procedures submitted for an equivalence determination. The testing methods will be submitted as soon as possible for that determination.

Additionally, establishments had adequate controls in place to prevent meat products intended for Icelandic domestic consumption from being commingled with products eligible for export in the U.S.

ENFORCEMENT CONTROLS

Inspection System Controls

With the exception of the unacceptable establishment (Est. 40), the Icelandic inspection system controls [control of restricted product and inspection samples, boneless meat reinspection, shipment security, including shipment between establishments, prevention of commingling of product intended for export to the United States with domestic product, monitoring and verification of establishment programs and controls (including the taking and documentation of corrective actions under HACCP plans), inspection supervision and documentation, the importation of only eligible livestock or poultry from other countries (i.e., only from eligible countries and certified establishments within those countries), and the importation of only eligible meat or poultry products from other countries for further processing] were in place and effective in ensuring that products produced by the establishment were wholesome, unadulterated, and properly labeled. In addition, adequate controls were found to be in place for security items, shipment security, and products entering the establishments from outside sources

Testing for *Salmonella* Species

Salmonella testing is not required in Iceland's establishments that are certified to export meat products to the United States because Iceland only exports meat from sheep and FSIS has not established *Salmonella* performance standards for sheep.

Species Verification Testing

At the time of this audit, Iceland was not exempt from the species verification-testing requirement. The auditor verified that species verification testing was being conducted in accordance with FSIS requirements.

Monthly Reviews

These monthly reviews were being performed by the Icelandic equivalent of a District Supervisor. He is a veterinarian with many years of experience. Dr. Hansson was in charge of the exporting establishments.

The internal review program was applied equally to both export and non-export establishments. Internal review visits were not always announced in advance but were sometimes unannounced by the reviewer. These reviews are preceded by a letter sent to each establishment once per year. This letter address facilities, equipment and operating conditions in the establishment. These are directed to the 17 District Veterinarians. This letter is followed by an on-site review by the Chief of Meat Inspection one or two times per year. The records of audited establishments were kept in the inspection offices of the individual establishments, and copies were also kept in the central offices in Reykjavik, and were routinely maintained on file for a minimum of 3 years.

In the event that an establishment is found, during one of these internal reviews, to be out of compliance with U.S. requirements, and is delisted for U.S. export, before it may again qualify for eligibility to be reinstated, an in-depth review is conducted, and the results are reported to Dr. Halldor Runolfsson, Chief Veterinary Officer for evaluation. A plan is formulated for corrective actions and preventive measures.

Enforcement Activities

There have been no formal investigations regarding violations of the legislation of slaughtering, meat processing and meat handling under the jurisdiction of the Veterinary Services during the past year because no violations were revealed. There are no provisions in Icelandic legislation on meat and meat processing which prohibit persons, that have been prosecuted and found guilty of an offense to the legislation, to start working in the meat industry after having served their sentence.

Exit Meeting

An exit meeting was conducted in Reykjavik on October 18, 2001. The participants included Mr. Hakon Sigurgrimsson, Iceland Ministry of Agriculture; Dr. Gisli Sverrir Halldorsson, Iceland Veterinary Officer for Import and Export, Mr. Edwin Brown, Economic/Commercial Officer, U.S. Embassy, Ms. Borghildur Magnusdottir, Assistant Economic/Commercial Officer, U.S. Embassy, and Dr. M. Douglas Parks, International Audit Staff Officer, USDA. The following topics were discussed:

1. Audit results and the delistment of Establishment 40. The required procedures for relistment were discussed and explained to the Icelandic officials.

2. The dropping of lamb heads before veterinary evaluation was discussed. The Iceland officials noted that they would apply for an equivalence determination in the near future to the International Policy Staff.
3. Critical control limits for critical control points were discussed and the officials said that these standards were now better understood and would be changed to proper limits in the future in the HACCP plans of the exporting plants.
4. The requirements for pre-shipment review and the records of the same were discussed and a commitment to assure they are promptly implemented was given by the Icelandic officials.
5. The turn-around time for residue test results was reported to have been reduced from four months to three months.

CONCLUSION

The inspection system of Iceland was found to have effective controls to ensure that product destined for export to the United States was produced under conditions equivalent to those which FSIS requires in domestic establishments, except for Establishment 40 which was delisted because of deficiencies revealed during the on-site audit. The concerns noted in the HACCP programs were corrected, as were the deviations in the SSOP program. Commitments were forthcoming from Icelandic officials to correct all of these deviations as soon as possible. A testing method for generic *E. coli* testing will be submitted very soon for an equivalence determination by the International Policy Staff. Five establishments were audited: four were acceptable, and one was unacceptable. The deficiencies encountered during the on-site establishment audits, in those establishments which were found to be acceptable, were adequately addressed to the auditor's satisfaction.

Dr. M. Douglas Parks
International Audit Staff Officer

(signed)Dr. M. Douglas Parks

ATTACHMENTS

- A. Data collection instrument for SSOPs
- B. Data collection instrument for HACCP programs
- C. Data collection instrument for *E. coli* testing
- D. Data collection instrument for *Salmonella* testing (*not applicable*)
- E. Laboratory Audit Forms
- F. Individual Foreign Establishment Audit Forms
- G. Written Foreign Country's Response to the Draft Final Audit Report

Data Collection Instrument for SSOPs

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOPs were met, according to the criteria employed in the U.S. domestic inspection program. The data collection instrument contained the following statements:

1. The establishment has a written SSOP program.
2. The procedure addresses pre-operational sanitation.
3. The procedure addresses operational sanitation.
4. The pre-operational procedures address (at a minimum) the cleaning of food-contact surfaces of facilities, equipment, and utensils.
5. The procedure indicates the frequency of the tasks.
6. The procedure identifies the individuals responsible for implementing and maintaining the activities.
7. The records of these procedures and any corrective action taken are being maintained on a daily basis.
8. The procedure is dated and signed by the person with overall on-site authority.

The results of these evaluations were as follows:

| Est. # | 1. Written program addressed | 2. Pre-op sanitation addressed | 3. Oper. sanitation addressed | 4. Contact surfaces addressed | 5. Frequency addressed | 6. Responsible indiv. identified | 7. Documentation done daily | 8. Dated and signed |
|--------|------------------------------|--------------------------------|-------------------------------|-------------------------------|------------------------|----------------------------------|-----------------------------|---------------------|
| 40 | √ | √ | √ | √ | √ | no | no | √ |
| 31 | √ | √ | √ | √ | √ | no | √ | √ |
| 81 | √ | √ | no | √ | √ | √ | √ | √ |
| 23 | √ | √ | √ | √ | √ | √ | √ | √ |
| 22 | √ | √ | √ | √ | √ | √ | √ | √ |

Data Collection Instrument for HACCP Programs

Each of the establishments approved to export meat products to the U.S. was required to have developed and implemented a Hazard Analysis – Critical Control Point (HACCP) system. Each of these systems was evaluated according to the criteria employed in the U.S. domestic inspection program. The data collection instrument included the following statements:

1. The establishment has a flow chart that describes the process steps and product flow.
2. The establishment has conducted a hazard analysis that includes food safety hazards likely to occur.
3. The analysis includes the intended use of or the consumers of the finished product(s).
4. There is a written HACCP plan for each product where the hazard analysis revealed one or more food safety hazard(s) reasonably likely to occur.
5. All hazards identified in the analysis are included in the HACCP plan; the plan lists a CCP for each food safety hazard identified.
6. The HACCP plan specifies critical limits, monitoring procedures, and the monitoring frequency performed for each CCP.
7. The plan describes corrective actions taken when a critical limit is exceeded.
8. The HACCP plan was validated using multiple monitoring results.
9. The HACCP plan lists the establishment's procedures to verify that the plan is being effectively implemented and functioning and the frequency for these procedures.
10. The HACCP plan's record-keeping system documents the monitoring of CCPs and/or includes records with actual values and observations.
11. The HACCP plan is dated and signed by a responsible establishment official.
12. The establishment is performing routine pre-shipment document reviews.

The results of these evaluations were as follows:

| Est. # | 1. Flow diagram | 2. Hazard analysis conducted | 3. Use & users included | 4. Plan for each hazard | 5. CCPs for all hazards | 6. Monitoring is specified | 7. Corr. actions are described | 8. Plan validated | 9. Adequate verific. procedures | 10. Adequate documentation | 11. Dated and signed | 12. Pre-shipment doc. review |
|--------|-----------------|------------------------------|-------------------------|-------------------------|-------------------------|----------------------------|--------------------------------|-------------------|---------------------------------|----------------------------|----------------------|------------------------------|
| 40 | √ | √ | √ | √ | √ | no | √ | √ | √ | √ | √ | no |
| 31 | √ | √ | √ | √ | √ | √ | √ | √ | no | √ | no | no |
| 81 | √ | √ | √ | √ | √ | no | √ | √ | √ | √ | √ | no |
| 23 | √ | √ | √ | √ | √ | no | √ | √ | no | no | √ | no |
| 22 | √ | √ | √ | √ | √ | √ | √ | √ | no | √ | √ | no |

Data Collection Instrument for Generic *E. coli* Testing

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for generic *E. coli* testing were met, according to the criteria employed in the U.S. domestic inspection program. The data collection instrument contained the following statements:

1. The establishment has a written procedure for testing for generic *E. coli*.
2. The procedure designates the employee(s) responsible to collect the samples.
3. The procedure designates the establishment location for sample collecting.
4. The sample collection is done on the predominant species being slaughtered.
5. The sampling is done at the frequency specified in the procedure.
6. The proper carcass site(s) and/or collection methodology (sponge or excision) is/are being used for sampling.
7. The carcass selection is following the random method specified in the procedure or is being taken randomly.
8. The laboratory is analyzing the sample using an AOAC Official Method or an equivalent method.
9. The results of the tests are being recorded on a process control chart showing the most recent test results.
10. The test results are being maintained for at least 12 months.

| Est. # | 1. Written procedure | 2. Sampler designated | 3. Sampling location given | 4. Predominant species sampled | 5. Sampling at the req'd freq. | 6. Proper site or method | 7. Sampling is random | 8. Using AOAC method | 9. Chart or graph of results | 10. Results are kept at least 1 yr |
|--------|----------------------|-----------------------|----------------------------|--------------------------------|--------------------------------|--------------------------|-----------------------|----------------------|------------------------------|------------------------------------|
| 40 | √ | no | no | √ | √ | √ | no | √ | √ | √ |
| 31 | √ | √ | no | √ | √ | √ | no | √ | √ | √ |
| 81 | √ | √ | √ | √ | √ | √ | no | √ | √ | √ |
| 23 | √ | √ | √ | √ | √ | √ | no | no | √ | √ |
| 22 | √ | no | √ | √ | √ | √ | no | √ | no | √ |

U.S. DEPARTMENT OF AGRICULTURE
 FOOD SAFETY AND INSPECTION SERVICE
 INTERNATIONAL PROGRAMS

REVIEW DATE

NAME OF FOREIGN LABORATORY

October 9,
2001

Icelandic Fisheries Laboratory
 Trace Analytical Laboratory

FOREIGN COUNTRY LABORATORY REVIEW

FOREIGN GOV'T AGENCY
 Ministry of Fishes'

CITY & COUNTRY
 Reykjavik, Iceland

ADDRESS OF LABORATORY
 P O Box 1405
 121 Reykjavik

NAME OF REVIEWER
 Dr. M. Douglas Parks

NAME OF FOREIGN OFFICIAL
 Dr. Sigudur Hansson

| Residue Code/Name | | | Hg | Pb | Cd | As | | | | | | | | | | |
|------------------------------|------------------------------|--------|-----------------|----|----|----|---|--|--|--|--|--|--|--|--|--|
| SAMPLING PROCEDURES | REVIEW ITEMS | ITEM # | EVALUATION CODE | | | | | | | | | | | | | |
| | Sample Handling | 01 | | A | A | A | A | | | | | | | | | |
| | Sampling Frequency | 02 | | A | A | A | A | | | | | | | | | |
| | Timely Analyses | 03 | | A | A | A | A | | | | | | | | | |
| | Compositing Procedure | 04 | | O | O | O | O | | | | | | | | | |
| | Interpret Comp Data | 05 | | O | O | O | O | | | | | | | | | |
| Data Reporting | 06 | | A | A | A | A | | | | | | | | | | |
| ANALYTICAL PROCEDURES | Acceptable Method | 07 | EVALUATION CODE | A | A | A | A | | | | | | | | | |
| | Correct Tissue(s) | 08 | | A | A | A | A | | | | | | | | | |
| | Equipment Operation | 09 | | A | A | A | A | | | | | | | | | |
| | Instrument Printouts | 10 | | A | A | A | A | | | | | | | | | |
| QUALITY ASSURANCE PROCEDURES | Minimum Detection Levels | 11 | EVALUATION CODE | A | A | A | A | | | | | | | | | |
| | Recovery Frequency | 12 | | A | A | A | A | | | | | | | | | |
| | Percent Recovery | 13 | | A | A | A | A | | | | | | | | | |
| | Check Sample Frequency | 14 | | A | A | A | A | | | | | | | | | |
| | All analyst w/Check Samples | 15 | | A | A | A | A | | | | | | | | | |
| | Corrective Actions | 16 | | A | A | A | A | | | | | | | | | |
| International Check Samples | 17 | | A | A | A | A | | | | | | | | | | |
| REVIEW PROCEDURES | Corrected Prior Deficiencies | 18 | EVAL. CODE | A | A | A | A | | | | | | | | | |
| OTHER REVIEW | | 19 | EVAL. CODE | | | | | | | | | | | | | |
| | | 20 | | | | | | | | | | | | | | |

SIGNATURE OF REVIEWER

M Douglas Parks DM

DATE

9 Oct 2001

| | | | |
|--|---|--|---|
| FOREIGN COUNTRY LABORATORY REVIEW <i>(Comment Sheet)</i> | | REVIEW DATE October 9, 2001 | NAME OF FOREIGN LABORATORY Icelandic Fisheries Laboratory Trace Analytical Laboratory |
| FOREIGN GOV'T AGENCY Ministry of Fishes | CITY & COUNTRY Reykjavik, Iceland | ADDRESS OF LABORATORY P O Box 1405 121 Reykjavik | |
| NAME OF REVIEWER Dr. M. Douglas Parks | NAME OF FOREIGN OFFICIAL Dr. Sigudur Hansson | | |

| RESIDUE | ITEM | COMMENTS |
|---------|------|--|
| | | Some of the standard compounds as received from the manufacturer were outdated. Daily working solutions were not dated. |

FOREIGN COUNTRY LABORATORY REVIEW

October 17,
2001

Institute for Experimental Pathology, Keldur

FOREIGN GOV'T AGENCY
 Ministry of Education

CITY & COUNTRY
 Reykjavik, Iceland

ADDRESS OF LABORATORY
 1S-112 Reykjavik

NAME OF REVIEWER
 Dr. M. Douglas Parks

NAME OF FOREIGN OFFICIAL
 Dr. Sigurdur Hansson

| Residue Code/Name | | 100 | 200 | 300 | 500 | | | | | | | | | | | |
|------------------------------|------------------------------|--------|-----------------|-----|-----|---|--|--|--|--|--|--|--|--|--|--|
| SAMPLING PROCEDURES | REVIEW ITEMS | ITEM # | EVALUATION CODE | | | | | | | | | | | | | |
| | Sample Handling | 01 | A | A | A | A | | | | | | | | | | |
| | Sampling Frequency | 02 | A | A | A | A | | | | | | | | | | |
| | Timely Analyses | 03 | A | A | A | A | | | | | | | | | | |
| | Compositing Procedure | 04 | O | O | O | O | | | | | | | | | | |
| | Interpret Comp Data | 05 | O | O | O | O | | | | | | | | | | |
| Data Reporting | 06 | A | A | A | A | | | | | | | | | | | |
| ANALYTICAL PROCEDURES | Acceptable Method | 07 | A | A | A | A | | | | | | | | | | |
| | Correct Tissue(s) | 08 | A | A | A | A | | | | | | | | | | |
| | Equipment Operation | 09 | A | A | A | A | | | | | | | | | | |
| | Instrument Printouts | 10 | A | A | A | A | | | | | | | | | | |
| QUALITY ASSURANCE PROCEDURES | Minimum Detection Levels | 11 | A | A | A | A | | | | | | | | | | |
| | Recovery Frequency | 12 | A | A | A | A | | | | | | | | | | |
| | Percent Recovery | 13 | A | A | A | A | | | | | | | | | | |
| | Check Sample Frequency | 14 | A | A | A | A | | | | | | | | | | |
| | All analyst w/Check Samples | 15 | A | A | A | A | | | | | | | | | | |
| | Corrective Actions | 16 | A | A | A | A | | | | | | | | | | |
| International Check Samples | 17 | A | A | A | A | | | | | | | | | | | |
| REVIEW PROCEDURES | Corrected Prior Deficiencies | 18 | A | A | A | A | | | | | | | | | | |
| OTHER REVIEW | | 19 | | | | | | | | | | | | | | |
| | | 20 | | | | | | | | | | | | | | |

SIGNATURE OF REVIEWER

M Douglas Parks Dr

DATE

17 Oct 2001

| | | | |
|--|--|--|--|
| FOREIGN COUNTRY LABORATORY REVIEW <i>(Comment Sheet)</i> | | REVIEW DATE October 17, 2001 | NAME OF FOREIGN LABORATORY Institute for Experimental Pathology, Keldur |
| FOREIGN GOV'T AGENCY Ministry of Education | | CITY & COUNTRY Reykjavik, Iceland | ADDRESS OF LABORATORY 1S-112 Reykjavik |
| NAME OF REVIEWER Dr. M. Douglas Parks | | NAME OF FOREIGN OFFICIAL Dr. Sigurdur Hansson | |

| RESIDUE | ITEM | COMMENTS |
|---------|------|--|
| | | <p>There was "white out" used to change entries in the records.</p> <p>Bacillus spores used in the antibiotic tests were outdated.</p> |

| | | | |
|--|-----------------------------|---|---|
| U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS FOREIGN PLANT REVIEW FORM | REVIEW DATE Oct 15, 2001 | ESTABLISHMENT NO. AND NAME Kaupfelag V-Hunvetninga est. 22 | CITY Hvammstangi COUNTRY Iceland |
|--|-----------------------------|---|---|

| | | |
|--|--|---|
| NAME OF REVIEWER Dr. M. Douglas Parks | NAME OF FOREIGN OFFICIAL Dr. Sigurdur Orn Hansson | EVALUATION <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable |
|--|--|---|

CODES (Give an appropriate code for each review item listed below)
 A = Acceptable M = Marginally Acceptable U = Unacceptable N = Not Reviewed O = Does not apply

| | | | | |
|--|---|---------|--|---------|
| 1. CONTAMINATION CONTROL | Cross contamination prevention | 28 A | Formulations | 55 A |
| (a) BASIC ESTABLISHMENT FACILITIES | Equipment Sanitizing | 29 A | Packaging materials | 56 A |
| Water potability records | Product handling and storage | 30 A | Laboratory confirmation | 57 A |
| Chlorination procedures | Product reconditioning | 31 A | Label approvals | 58 O |
| Back siphonage prevention | Product transportation | 32 A | Special label claims | 59 O |
| Hand washing facilities | (d) ESTABLISHMENT SANITATION PROGRAM | | Inspector monitoring | 60 A |
| Sanitizers | Effective maintenance program | 33 A | Processing schedules | 61 O |
| Establishments separation | Preoperational sanitation | 34 A | Processing equipment | 62 O |
| Pest --no evidence | Operational sanitation | 35 A | Processing records | 63 O |
| Pest control program | Waste disposal | 36 A | Empty can inspection | 64 O |
| Pest control monitoring | 2. DISEASE CONTROL | | Filling procedures | 65 O |
| Temperature control | Animal identification | 37 A | Container closure exam | 66 O |
| Lighting | Antemortem inspec. procedures | 38 A | Interim container handling | 67 O |
| Operations work space | Antemortem dispositions | 39 A | Post-processing handling | 68 O |
| Inspector work space | Humane Slaughter | 40 A | Incubation procedures | 69 O |
| Ventilation | Postmortem inspec. procedures | 41 A | Process. defect actions -- plant | 70 O |
| Facilities approval | Postmortem dispositions | 42 A | Processing control -- inspection | 71 O |
| Equipment approval | Condemned product control | 43 A | 5. COMPLIANCE/ECON. FRAUD CONTROL | |
| (b) CONDITION OF FACILITIES EQUIPMENT | Restricted product control | 44 A | Export product identification | 72 A |
| Over-product ceilings | Returned and rework product | 45 A | Inspector verification | 73 A |
| Over-product equipment | 3. RESIDUE CONTROL | | Export certificates | 74 A |
| Product contact equipment | Residue program compliance | 46 A | Single standard | 75 A |
| Other product areas (inside) | Sampling procedures | 47 A | Inspection supervision | 76 A |
| Dry storage areas | Residue reporting procedures | 48 A | Control of security items | 77 A |
| Antemortem facilities | Approval of chemicals, etc. | 49 A | Shipment security | 78 A |
| Welfare facilities | Storage and use of chemicals | 50 A | Species verification | 79 A |
| Outside premises | 4. PROCESSED PRODUCT CONTROL | | "Equal to" status | 80 A |
| (c) PRODUCT PROTECTION & HANDLING | Pre-boning trim | 51 A | Imports | 81 A |
| Personal dress and habits | Boneless meat reinspection | 52 A | | |
| Personal hygiene practices | Ingredients identification | 53 A | | |
| Sanitary dressing procedures | Control of restricted ingredients | 54 A | | |

| | | | |
|--|--------------------------|--|-------------|
| FOREIGN PLANT REVIEW FORM (reverse) | REVIEW DATE | ESTABLISHMENT NO. AND NAME | CITY |
| | Oct 15,2001 | Kaupfelag V-Hunvetninga est. 22 | Hvammstangi |
| | | | COUNTRY |
| | | | Iceland |
| NAME OF REVIEWER | NAME OF FOREIGN OFFICIAL | EVALUATION | |
| Dr. M. Douglas Parks | Dr. Sigurdur Orn Hansson | <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable | |

COMMENTS:

- 17--Condensate was dripping from overhead surfaces not cleaned and sanitized daily onto a product packaging table ready for use.
- 27--An employee was cutting thru the rectum and continued the cut thru the tail without sanitizing the knife.
- 27--A product trimmer at the final trim station was removed due to an accident but was not replaced and the product flow continued not trimmed adequately.
- SSOP--Preventative action not being recorded.
- HACCP--No written verification procedures were included in the plan.
- HACCP--Pre-shipment review was not being done.
- E.coli testing--The procedure did not designate the employee responsible to collect the samples.
- E.coli testing--The carcass for sampling was not selected randomly.
- E.coli testing--No process control chart has been produced.

| | | | |
|--|-----------------------------|---|--------------------|
| U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS FOREIGN PLANT REVIEW FORM | REVIEW DATE Oct 16, 2001 | ESTABLISHMENT NO. AND NAME Solufelag A-Hunvetninga est. 23 | CITY Blonduos |
| | | | COUNTRY Iceland |

| | | |
|--|--|--|
| NAME OF REVIEWER Dr. M. Douglas Parks | NAME OF FOREIGN OFFICIAL Dr. Sigurdur Orn Hansson | EVALUATION <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable |
|--|--|--|

CODES (Give an appropriate code for each review item listed below)
 A = Acceptable M = Marginally Acceptable U = Unacceptable N = Not Reviewed O = Does not apply

| | | | | | |
|---------------------------------------|---------|--------------------------------------|---------|-----------------------------------|---------|
| 1. CONTAMINATION CONTROL | | Cross contamination prevention | 28 A | Formulations | 55 A |
| (a) BASIC ESTABLISHMENT FACILITIES | | Equipment Sanitizing | 29 A | Packaging materials | 56 A |
| Water potability records | 01 A | Product handling and storage | 30 A | Laboratory confirmation | 57 A |
| Chlorination procedures | 02 A | Product reconditioning | 31 A | Label approvals | 58 O |
| Back siphonage prevention | 03 A | Product transportation | 32 A | Special label claims | 59 O |
| Hand washing facilities | 04 A | (d) ESTABLISHMENT SANITATION PROGRAM | | Inspector monitoring | 60 A |
| Sanitizers | 05 A | Effective maintenance program | 33 A | Processing schedules | 61 O |
| Establishments separation | 06 A | Preoperational sanitation | 34 A | Processing equipment | 62 O |
| Pest --no evidence | 07 A | Operational sanitation | 35 A | Processing records | 63 O |
| Pest control program | 08 A | Waste disposal | 36 A | Empty can inspection | 64 O |
| Pest control monitoring | 09 A | 2. DISEASE CONTROL | | Filling procedures | 65 O |
| Temperature control | 10 A | Animal identification | 37 A | Container closure exam | 66 O |
| Lighting | 11 A | Antemortem inspec. procedures | 38 A | Interim container handling | 67 O |
| Operations work space | 12 A | Antemortem dispositions | 39 A | Post-processing handling | 68 O |
| Inspector work space | 13 A | Humane Slaughter | 40 A | Incubation procedures | 69 O |
| Ventilation | 14 A | Postmortem inspec. procedures | 41 A | Process. defect actions -- plant | 70 O |
| Facilities approval | 15 A | Postmortem dispositions | 42 A | Processing control -- inspection | 71 O |
| Equipment approval | 16 A | Condemned product control | 43 A | 5. COMPLIANCE/ECON. FRAUD CONTROL | |
| (b) CONDITION OF FACILITIES EQUIPMENT | | Restricted product control | 44 A | Export product identification | 72 A |
| Over-product ceilings | 17 A | Returned and rework product | 45 A | Inspector verification | 73 A |
| Over-product equipment | 18 A | 3. RESIDUE CONTROL | | Export certificates | 74 A |
| Product contact equipment | 19 U | Residue program compliance | 46 A | Single standard | 75 A |
| Other product areas (inside) | 20 A | Sampling procedures | 47 A | Inspection supervision | 76 A |
| Dry storage areas | 21 A | Residue reporting procedures | 48 A | Control of security items | 77 A |
| Antemortem facilities | 22 A | Approval of chemicals, etc. | 49 A | Shipment security | 78 A |
| Welfare facilities | 23 A | Storage and use of chemicals | 50 A | Species verification | 79 A |
| Outside premises | 24 A | 4. PROCESSED PRODUCT CONTROL | | "Equal to" status | 80 A |
| (c) PRODUCT PROTECTION & HANDLING | | Pre-boning trim | 51 A | Imports | 81 A |
| Personal dress and habits | 25 A | Boneless meat reinspection | 52 A | | |
| Personal hygiene practices | 26 A | Ingredients identification | 53 A | | |
| Sanitary dressing procedures | 27 U | Control of restricted ingredients | 54 A | | |

| | | | |
|--|--------------------------|---------------------------------|--|
| FOREIGN PLANT REVIEW FORM (reverse) | REVIEW DATE | ESTABLISHMENT NO. AND NAME | CITY |
| | Oct 16, 2001 | Solufelag A-Hunvetninga est. 23 | Blonduos |
| | | | COUNTRY |
| | | | Iceland |
| NAME OF REVIEWER | NAME OF FOREIGN OFFICIAL | | EVALUATION |
| Dr. M. Douglas Parks | Dr. Sigurdur Orn Hansson | | <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable |

COMMENTS:

- 19--The compartments of the visera conveyor were not properly cleaned between uses.
- 27--The exposed end of the carcass tail of many carcasses had fecal contamination due to an error in the dressing procedure.
- 27--Three of twenty carcasses examined in the carcass chiller were observed with fecal material on them.
- SSOP--Preventative action is not being recorded.
- HACCP--The limits of one CCP were not measurable, they were judgemental.
- HACCP--There were no written verifications in the plan.
- HACCP--Corrective action was not documented when there was a failure.
- HACCP--No pre-shipment review was being done.
- E.coli testing--The carcasses selected for sampling were not selected randomly.
- E.coli testing--Laboratory testing procedures are not AOAC and the equivalence has not been determined.

FOREIGN PLANT REVIEW FORM

REVIEW DATE: Oct 12, 2001
ESTABLISHMENT NO. AND NAME: Nordlenska Ehf Husavik est. 31

CITY: Husavik
COUNTRY: Iceland

NAME OF REVIEWER: Dr. M. Douglas Parks
NAME OF FOREIGN OFFICIAL: Dr. Sigurdur Orn Hansson
EVALUATION: Acceptable Acceptable/ Re-review Unacceptable

CODES (Give an appropriate code for each review item listed below)
A = Acceptable M = Marginally Acceptable U = Unacceptable N = Not Reviewed O = Does not apply

| | | | | | |
|---------------------------------------|---------|--------------------------------------|---------|-----------------------------------|---------|
| 1. CONTAMINATION CONTROL | | Cross contamination prevention | 28 A | Formulations | 55 A |
| (a) BASIC ESTABLISHMENT FACILITIES | | Equipment Sanitizing | 29 A | Packaging materials | 56 A |
| Water potability records | 01 A | Product handling and storage | 30 A | Laboratory confirmation | 57 A |
| Chlorination procedures | 02 A | Product reconditioning | 31 A | Label approvals | 58 A |
| Back siphonage prevention | 03 A | Product transportation | 32 A | Special label claims | 59 O |
| Hand washing facilities | 04 A | (d) ESTABLISHMENT SANITATION PROGRAM | | Inspector monitoring | 60 A |
| Sanitizers | 05 A | Effective maintenance program | 33 A | Processing schedules | 61 O |
| Establishments separation | 06 A | Preoperational sanitation | 34 A | Processing equipment | 62 O |
| Pest --no evidence | 07 A | Operational sanitation | 35 A | Processing records | 63 O |
| Pest control program | 08 A | Waste disposal | 36 A | Empty can inspection | 64 O |
| Pest control monitoring | 09 A | 2. DISEASE CONTROL | | Filling procedures | 65 O |
| Temperature control | 10 A | Animal identification | 37 A | Container closure exam | 66 O |
| Lighting | 11 A | Antemortem inspec. procedures | 38 A | Interim container handling | 67 O |
| Operations work space | 12 A | Antemortem dispositions | 39 A | Post-processing handling | 68 O |
| Inspector work space | 13 A | Humane Slaughter | 40 A | Incubation procedures | 69 O |
| Ventilation | 14 A | Postmortem inspec. procedures | 41 A | Process. defect actions -- plant | 70 O |
| Facilities approval | 15 A | Postmortem dispositions | 42 A | Processing control -- inspection | 71 O |
| Equipment approval | 16 A | Condemned product control | 43 A | 5. COMPLIANCE/ECON. FRAUD CONTROL | |
| (b) CONDITION OF FACILITIES EQUIPMENT | | Restricted product control | 44 A | Export product identification | 72 A |
| Over-product ceilings | 17 A | Returned and rework product | 45 A | Inspector verification | 73 A |
| Over-product equipment | 18 A | 3. RESIDUE CONTROL | | Export certificates | 74 A |
| Product contact equipment | 19 M | Residue program compliance | 46 A | Single standard | 75 A |
| Other product areas (inside) | 20 M | Sampling procedures | 47 A | Inspection supervision | 76 A |
| Dry storage areas | 21 A | Residue reporting procedures | 48 A | Control of security items | 77 A |
| Antemortem facilities | 22 A | Approval of chemicals, etc. | 49 A | Shipment security | 78 A |
| Welfare facilities | 23 A | Storage and use of chemicals | 50 A | Species verification | 79 A |
| Outside premises | 24 A | 4. PROCESSED PRODUCT CONTROL | | "Equal to" status | 80 A |
| (c) PRODUCT PROTECTION & HANDLING | | Pre-boning trim | 51 A | Imports | 81 A |
| Personal dress and habits | 25 A | Boneless meat reinspection | 52 A | | |
| Personal hygiene practices | 26 A | Ingredients identification | 53 A | | |
| Sanitary dressing procedures | 27 A | Control of restricted ingredients | 54 A | | |

| | | | |
|--|--------------------------|--|---------|
| FOREIGN PLANT REVIEW FORM (reverse) | REVIEW DATE | ESTABLISHMENT NO. AND NAME | CITY |
| | Oct 12,2001 | Nordlenska Ehf Husavik est. 31 | Husavik |
| | | | COUNTRY |
| | | | Iceland |
| NAME OF REVIEWER | NAME OF FOREIGN OFFICIAL | EVALUATION | |
| Dr. M. Douglas Parks | Dr. Sigurdur Orn Hansson | <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable | |

COMMENTS:

- 20--The edge of a packing table had residues from previous day's uses in the cutting department.
- 19--The bung hook was not sanitized between uses(animals) in the slaughter department.
- SSOP--No preventative action recorded for operational sanitation procedures.
- SSOP--The individual responsibe for implementing and maintaining the activities was not named in the procedure.
- HACCP--There were no written verification procedures in the plan.
- HACCP--The plan is not signed by the overall on-site authority.
- HACCP--Pre-shipment review was not done.
- E.coli--The procedure does not designate the plant location for sample collecting.
- E.coli--The carcass selection is not done randomly.

FOREIGN PLANT REVIEW FORM

REVIEW DATE: Oct 11, 2001
ESTABLISHMENT NO. AND NAME: Slaturfelagid Bubot est. 40

CITY: Hornafjordur
COUNTRY: Iceland

NAME OF REVIEWER
Dr. M. Douglas Parks

NAME OF FOREIGN OFFICIAL
Dr. Sigurdur Orn Hansson

EVALUATION
 Acceptable Acceptable/ Re-review Unacceptable

CODES (Give an appropriate code for each review item listed below)

A = Acceptable M = Marginally Acceptable U = Unacceptable N = Not Reviewed O = Does not apply

| 1. CONTAMINATION CONTROL | | Cross contamination prevention | 28 A | Formulations | 55 O |
|---------------------------------------|---------|--------------------------------------|----------|-----------------------------------|---------|
| (a) BASIC ESTABLISHMENT FACILITIES | | Equipment Sanitizing | 29 A | Packaging materials | 56 O |
| Water potability records | 01 A | Product handling and storage | 30 30 | Laboratory confirmation | 57 O |
| Chlorination procedures | 02 A | Product reconditioning | 31 A | Label approvals | 58 O |
| Back siphonage prevention | 03 A | Product transportation | 32 A | Special label claims | 59 O |
| Hand washing facilities | 04 A | (d) ESTABLISHMENT SANITATION PROGRAM | | Inspector monitoring | 60 O |
| Sanitizers | 05 A | Effective maintenance program | 33 A | Processing schedules | 61 O |
| Establishments separation | 06 A | Preoperational sanitation | 34 A | Processing equipment | 62 O |
| Pest --no evidence | 07 A | Operational sanitation | 35 A | Processing records | 63 O |
| Pest control program | 08 A | Waste disposal | 36 A | Empty can inspection | 64 O |
| Pest control monitoring | 09 A | 2. DISEASE CONTROL | | Filling procedures | 65 O |
| Temperature control | 10 A | Animal identification | 37 A | Container closure exam | 66 O |
| Lighting | 11 A | Antemortem inspec. procedures | 38 A | Interim container handling | 67 O |
| Operations work space | 12 A | Antemortem dispositions | 39 A | Post-processing handling | 68 O |
| Inspector work space | 13 A | Humane Slaughter | 40 A | Incubation procedures | 69 O |
| Ventilation | 14 A | Postmortem inspec. procedures | 41 A | Process. defect actions -- plant | 70 O |
| Facilities approval | 15 A | Postmortem dispositions | 42 A | Processing control -- inspection | 71 O |
| Equipment approval | 16 A | Condemned product control | 43 A | 5. COMPLIANCE/ECON. FRAUD CONTROL | |
| (b) CONDITION OF FACILITIES EQUIPMENT | | Restricted product control | 44 A | Export product identification | 72 A |
| Over-product ceilings | 17 U | Returned and rework product | 45 A | Inspector verification | 73 A |
| Over-product equipment | 18 U | 3. RESIDUE CONTROL | | Export certificates | 74 A |
| Product contact equipment | 19 A | Residue program compliance | 46 A | Single standard | 75 A |
| Other product areas (inside) | 20 A | Sampling procedures | 47 A | Inspection supervision | 76 A |
| Dry storage areas | 21 A | Residue reporting procedures | 48 A | Control of security items | 77 A |
| Antemortem facilities | 22 A | Approval of chemicals, etc. | 49 A | Shipment security | 78 A |
| Welfare facilities | 23 A | Storage and use of chemicals | 50 A | Species verification | 79 A |
| Outside premises | 24 A | 4. PROCESSED PRODUCT CONTROL | | "Equal to" status | 80 A |
| (c) PRODUCT PROTECTION & HANDLING | | Pre-boning trim | 51 A | Imports | 81 A |
| Personal dress and habits | 25 A | Boneless meat reinspection | 52 A | | |
| Personal hygiene practices | 26 A | Ingredients identification | 53 A | | |
| Sanitary dressing procedures | 27 U | Control of restricted ingredients | 54 A | | |

| | | | |
|---|--------------------------|-----------------------------|--|
| FOREIGN PLANT REVIEW FORM (reverse) | REVIEW DATE | ESTABLISHMENT NO. AND NAME | CITY |
| | Oct 11,2001 | Slaturfelagid Bubot est. 40 | Hornafjordur |
| | | | COUNTRY |
| | | | Iceland |
| NAME OF REVIEWER | NAME OF FOREIGN OFFICIAL | | EVALUATION |
| Dr. M. Douglas Parks | Dr. Sigurdur Orn Hansson | | <input type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input checked="" type="checkbox"/> Unacceptable |

COMMENTS:

17--Extensive condensate was falling onto exposed carcasses in two carcass chillers from overhead structures that were not cleaned and sanitized daily. Condensate was falling on approximately 25% of 450 carcasses in these chillers.

30--Frozen carcasses in plastic bags were stored directly on the floor in the freezer.

18--Overspray in the carcass wash area was dripping onto exposed carcasses.

27--The bung-dropping procedure resulted in fecal contamination in about 20% of the carcasses.

SSOP--The individual responsible for implementing and maintaining the activities was not named in the plan.

SSOP--Corrective and preventative actions are not recorded.

HACCP--In one critical control point, the limits were not measurable but were judgemental.

E.coli testing--The procedure does not designate the employee responsible to collect the samples.

E.coli testing--The procedure does not designate the plant location for sample collecting.

E.coli testing--The carcass selection for testing was not random.

| U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS | | REVIEW DATE | ESTABLISHMENT NO. AND NAME | | CITY |
|---|---------|--|-------------------------------|--|---------|
| FOREIGN PLANT REVIEW FORM | | Oct. 10, 2001 | Slaturfelag Sudurlands est 81 | | Selfoss |
| | | | | | |
| NAME OF REVIEWER Dr. M. Douglas Parks | | NAME OF FOREIGN OFFICIAL Dr. Sigurdur Orn Hansson | | EVALUATION <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable | |
| CODES (Give an appropriate code for each review item listed below) A = Acceptable M = Marginally Acceptable U = Unacceptable N = Not Reviewed O = Does not apply | | | | | |
| 1. CONTAMINATION CONTROL | | Cross contamination prevention | 28 U | Formulations | 55 A |
| (a) BASIC ESTABLISHMENT FACILITIES | | Equipment Sanitizing | 29 A | Packaging materials | 56 A |
| Water potability records | 01 A | Product handling and storage | 30 A | Laboratory confirmation | 57 A |
| Chlorination procedures | 02 A | Product reconditioning | 31 A | Label approvals | 58 A |
| Back siphonage prevention | 03 A | Product transportation | 32 A | Special label claims | 59 O |
| Hand washing facilities | 04 A | (d) ESTABLISHMENT SANITATION PROGRAM | | Inspector monitoring | 60 A |
| Sanitizers | 05 A | Effective maintenance program | 33 A | Processing schedules | 61 O |
| Establishments separation | 06 A | Preoperational sanitation | 34 A | Processing equipment | 62 O |
| Pest --no evidence | 07 A | Operational sanitation | 35 A | Processing records | 63 O |
| Pest control program | 08 A | Waste disposal | 36 A | Empty can inspection | 64 O |
| Pest control monitoring | 09 A | 2. DISEASE CONTROL | | Filling procedures | 65 O |
| Temperature control | 10 A | Animal identification | 37 A | Container closure exam | 66 O |
| Lighting | 11 A | Antemortem inspec. procedures | 38 A | Interim container handling | 67 O |
| Operations work space | 12 A | Antemortem dispositions | 39 A | Post-processing handling | 68 O |
| Inspector work space | 13 A | Humane Slaughter | 40 A | Incubation procedures | 69 O |
| Ventilation | 14 A | Postmortem inspec. procedures | 41 A | Process. defect actions -- plant | 70 O |
| Facilities approval | 15 A | Postmortem dispositions | 42 A | Processing control -- inspection | 71 O |
| Equipment approval | 16 A | Condemned product control | 43 A | 5. COMPLIANCE/ECON. FRAUD CONTROL | |
| (b) CONDITION OF FACILITIES EQUIPMENT | | Restricted product control | 44 A | Export product identification | 72 A |
| Over-product ceilings | 17 A | Returned and rework product | 45 A | Inspector verification | 73 A |
| Over-product equipment | 18 U | 3. RESIDUE CONTROL | | Export certificates | 74 A |
| Product contact equipment | 19 M | Residue program compliance | 46 A | Single standard | 75 A |
| Other product areas (inside) | 20 A | Sampling procedures | 47 A | Inspection supervision | 76 A |
| Dry storage areas | 21 A | Residue reporting procedures | 48 A | Control of security items | 77 A |
| Antemortem facilities | 22 A | Approval of chemicals, etc. | 49 A | Shipment security | 78 A |
| Welfare facilities | 23 A | Storage and use of chemicals | 50 A | Species verification | 79 A |
| Outside premises | 24 A | 4. PROCESSED PRODUCT CONTROL | | "Equal to" status | 80 A |
| (c) PRODUCT PROTECTION & HANDLING | | Pre-boning trim | 51 A | Imports | 81 A |
| Personal dress and habits | 25 A | Boneless meat reinspection | 52 A | | |
| Personal hygiene practices | 26 A | Ingredients identification | 53 A | | |
| Sanitary dressing procedures | 27 U | Control of restricted ingredients | 54 A | | |

| | | | |
|--|--------------------------|--|---------|
| FOREIGN PLANT REVIEW FORM (reverse) | REVIEW DATE | ESTABLISHMENT NO. AND NAME | CITY |
| | Oct. 10, 2001 | Slaturfelag Sudurlands est 81 | Selfoss |
| | | | COUNTRY |
| | | | Iceland |
| NAME OF REVIEWER | NAME OF FOREIGN OFFICIAL | EVALUATION | |
| Dr. M. Douglas Parks | Dr. Sigurdur Orn Hansson | <input checked="" type="checkbox"/> Acceptable <input type="checkbox"/> Acceptable/ Re-review <input type="checkbox"/> Unacceptable | |

COMMENTS:

18--Overspray at the carcass wash area was dripping from overhead facilities onto exposed carcasses.

19--The cutting remnants on the end of plastic pipes located in the corners inside of an exposed product container were loose and likely to fall into the product.

27--The bung dropping procedure resulted in fecal contamination in approximately 20% of the carcasses.

28--Between the hide removal area and the final trim approximately 10 % of the carcasses had fecal contamination and no effort was made to correct the dressing procedure.

SSOP--No preventative action being recorded.

SSOP--No records were kept for operational sanitation.

HACCP--One critical control point had critical limits that were not measurable and were judgemental.

HACCP--Pre shipment review was not being done.

HACCP--No written verification procedures in the plan.

E.coli testing--The carcass selection was not random.

Original Message

From: KristensenH [SMTP:KristensenH@fas.usda.gov]
Sent: Monday, April 29, 2002 1:03 PM
To: Stefan, Gary
Subject: Svör v. skyrslu

Please see the attached from Dr. Hansson, Iceland.

Hasse Kristensen

Forward Header

Subject: Svör v. skyrslu
Author: sigurdur.hansson@lan.stjr.is at FAS
Date: 4/29/2002 4:30 AM

Dear Hasse Kristensen

With reference to your e-mail this morning I forward a letter to Ms Sally Stratmeoen regarding our comments to the draft report on the USDA audit last October. The letter was sent to Washington directly and also through the American Embassy in Reykjavik and the Foreign Ministry in Reykjavik. Please let me know if it does not arrive in Washington.

Please appologize our late response.

Best regards

Sigurdur Örn Hansson

----- Forwarded by Sigurdur Örn Hansson/LAN/NotesSTJR on 29.04.2002 08:24 -----

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Reykjavík,

April 15, 2002

Ref:

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SÖH/--

The Chief Veterinary Officer has received your letter dated FEB 4 2002 where FSIS invited us to provide comments regarding the information in the report on your on - site audit of Iceland's meat inspection system conducted from October 8-18, 2001.

We have studied the report and do not wish to comment on the factual information contained therein.

We look forward to our cooperation and your initiative with respect to a teleconference to discuss outstanding matters.

Sincerely yours

Sigurdur Örn Hansson DVM MSc
Deputy Chief Veterinary Officer