

United States Department of Agriculture

Food Safety and Inspection Service Washington, D.C. 20250

JAN 1 6 2007

Mr. Greg Read
Executive Manager, Exports and Food Policy
Australian Quarantine and Inspection Service (AQIS)
Edmund Barton Building
GPO Box 858
Canberra ACT 2601
Australia

Dear Mr. Read:

Enclosed is the final report of the Food Safety and Inspection Service (FSIS) audit of Australia's meat inspection system conducted August 10 through August 30, 2006.

Comments from the government of Australia have been included as an attachment to the final report. We appreciate clarification regarding AQIS oversight of *Salmonella* testing at certified establishments and recognize FSIS' previous equivalence ruling of allowing random observation by AQIS inspectors. Accordingly, we have corrected the FSIS audit report by removing the finding that daily oversight of *Salmonella* testing was not being performed by AQIS.

If you have questions regarding the audit or audit report, please contact me by telephone at 202-720-3781, by facsimile at 202-690-4040, or by e-mail at sally.white@fsis.usda.gov.

Sincerely,

Sally White

Director

International Equivalence Staff Office of International Affairs

Enclosure

FINAL

JAN 1 6 2007

# FINAL REPORT OF AN AUDIT CARRIED OUT IN AUSTRALIA COVERING AUSTRALIA'S MEAT INSPECTION SYSTEM

August 10 through August 30, 2006

Food Safety and Inspection Service United States Department of Agriculture

#### TABLE OF CONTENTS

- 1. INTRODUCTION
- 2. OBJECTIVE OF THE AUDIT
- 3. PROTOCOL
- 4. LEGAL BASIS FOR THE AUDIT
- 5. SUMMARY OF PREVIOUS AUDITS
- 6. MAIN FINDINGS
  - 6.1 Government Oversight
  - 6.2 Headquarters Audit
  - 6.3 Audit of Regional and Local Inspection Sites
- 7. ESTABLISHMENT AUDITS
- 8. RESIDUE AND MICROBIOLOGY LABORATORY AUDITS
- 9. SANITATION CONTROLS
  - 9.1 SSOP
  - 9.2 Sanitation
- 10. ANIMAL DISEASE CONTROLS
- 11. SLAUGHTER/PROCESSING CONTROLS
  - 11.1 Humane Handling and Slaughter
  - 11.2 HACCP Implementation
  - 11.3 Testing for generic Escherichia coli
  - 11.4 Testing for Listeria monocytogenes
- 12. RESIDUE CONTROLS
- 13. ENFORCEMENT CONTROLS
  - 13.1 Daily Inspection
  - 13.2 Testing for Salmonella
  - 13.3 Species Verification
  - 13.4 Monthly Reviews
  - 13.5 Inspection System Controls
- 14. CLOSING MEETING
- 15. ATTACHMENTS TO THE AUDIT REPORT

#### ABBREVIATIONS AND SPECIAL TERMS USED IN THE REPORT

ATM Area Technical Manager

AQIS Australian Quarantine & Inspection Service

CCA Central Competent Authority – (AQIS for this report)

CCP Critical Control Point

CFR U.S. Code of Federal Regulations

E. coli Escherichia coli

ELMER E-Legislation Manuals and Essential References

FOM Field Operations Manager

FSIS Food Safety and Inspection Service

MOU Memorandum of Understanding

MSQA Meat Safety Quality Assurance

NATA National Association of Testing Authorities

NOID Notice of Intent to Delist

OPV On-Plant Veterinarian

PR/HACCP Pathogen Reduction/Hazard Analysis and Critical Control Point

Systems

RTE Ready-to-Eat

Salmonella species

SSOP Sanitation Standard Operating Procedures

#### 1. INTRODUCTION

The Food Safety and Inspection Service (FSIS) of the United States Department of Agriculture conducted an audit of the Australian meat inspection system August 10 through August 30, 2006.

An opening meeting was held on August 10, 2006, in Canberra with the Central Competent Authority (CCA) – Australia Quarantine Inspection Service (AQIS). At this meeting, the audit team confirmed the objective and scope of the audit, the auditors' itineraries, and requested additional information needed to complete the audit of Australia's meat inspection system.

Representatives from AQIS' headquarters and/or representatives from AQIS' regional and local inspection offices accompanied each auditor during the reviews.

#### 2. OBJECTIVE OF THE AUDIT

The objective was to (1) determine whether the concerns identified during the 2005 audit had been appropriately addressed, and (2) evaluate the performance of AQIS with respect to government oversight and enforcement of the AQIS and FSIS regulatory requirements relative to maintaining an inspection system equivalent to that of the United States. This included special emphasis regarding government oversight of CCA's microbiological laboratory program and cold store facilities used for freezing and storing meat products destined for the United States, and knowledge and application of the FSIS regulatory requirements.

In pursuit of the objective, the following localities were visited:

Competent Authority Visits			Comments
Competent Authority (Interviews with AQIS	Central	1	Canberra
Officials)	Regional Office	3	New South Wales, Queensland and Victoria
	Local Office	10	Establishments/Cold Storage Facility
Laboratories (Microbiology)	4		
Meat Slaughter / Processing Es	9		
Cold Storage Facilities	1		

#### 3. PROTOCOL

This on-site audit was conducted in four parts. One part involved visits with AQIS officials to discuss oversight programs and practices, including enforcement activities. The second part involved an audit of a selection of records and personnel interviews in the country's inspection headquarters and regional offices. The third part involved on-site visits to 10 establishments: nine slaughter and processing establishments and one cold storage facility (ID

Warehouse). The fourth part involved visits to four private laboratories certified by AQIS to conduct microbiological testing of meat products destined for the United States. Program effectiveness determinations of Australia's inspection system focused on five areas of government controls and oversight and five areas of risk: (1) sanitation controls, including the implementation and operation of SSOP, (2) animal disease controls, (3) slaughter/processing controls, including the implementation and operation of HACCP programs and a testing program for generic *E. coli*, (4) residue controls, and (5) enforcement controls, including testing program for *Salmonella*.

During the establishment visits, the auditor evaluated the nature, extent and degree to which findings impacted on food safety and public health. The auditor also assessed how inspection activities are carried out by AQIS and determined if controls were in place to ensure that the production of meat and meat products were safe, unadulterated and properly labeled.

At the opening meeting, the audit team explained that Australia's meat inspection system would be audited against the following standards: (1) FSIS regulatory requirements, as applicable, (2) AQIS requirements specific to exporting meat and meat products to the United States, and (3) FSIS equivalence determinations specific to Australia. FSIS requirements include, among other things, daily inspection in all applicable certified establishments, monthly supervisory visits to certified establishments, humane handling and slaughter of animals, ante-mortem inspection of animals and post-mortem inspection of carcasses and parts thereof, the handling and disposal of inedible and condemned materials, sanitation of facilities and equipment, residue testing, species verification, and requirements for HACCP, SSOP, and testing for generic *E. coli, Salmonella*, and *Listeria monocytogenes* (*Lm*).

The FSIS equivalence determinations were made under the provisions of the World Trade Organization Sanitary/Phytosanitary Agreement, and are:

- 1. Establishment employees collect carcass samples for *Salmonella* testing monitored by AQIS on a random basis, and private laboratories analyze *Salmonella* samples.
- 2. Slaughtering equines for Australia's domestic market in the same establishment (Est. # 3416) where bovines are slaughtered for export to the United States.
- 3. Allow the use of the MPSC rinse and chill technique on bovines slaughtered in establishments certified to export to the United States
- 4. Allow Australia to export meat to the United States from sheep and swine carcasses whereby post-mortem inspection would be conducted without examination of the heads. This is permitted only when tissue from the heads was not saved for human consumption.
- 5. The following laboratory testing methods for the detection of generic *E. coli*: AOAC 998.08, AOAC 991.14, and AS 5013.15-2004
- 6. The following laboratory testing methods for the detection of *E. coli* O157:H7: AOAC 2000.14, FDA BAM Chapter 4A (Sept 2002 protocol), *E. coli* O157:H7 BAX 0157:H7, AOAC 996.09, AOAC 996.10, AOAC 2000.13, and ISO 16654:2001.
- 7. The following laboratory testing methods for the detection of *Salmonella*: AOAC 978.24, AOAC 989.14, AOAC 992.11, AOAC 996.08, AOAC 998.09, AOAC 999.08, AOAC 999.09, AOAC 2000.07, AOAC 2001.07, AOAC 2001.08, AOAC 2001.09, AOAC OM 2003.09, and AOAC 5013.10-2004.
- 8. The following laboratory testing methods for the detection of Lm:

# AOAC 995.22; AOAC 997.03; AOAC 999.06; AOAC 996.14; AOAC 2002.09; AS 1766.2.16.1998; FDA BAM Chapter 10 (January 2003); BAX *Lm*.

#### 4. LEGAL BASIS FOR THE AUDIT

The audit was undertaken under the specific provisions of U.S. laws and regulations, in particular:

- The Federal Meat Inspection Act (21 U.S.C. 601 et seq.).
- The Federal Meat Inspection Regulations (9 CFR Parts 301 to end), which include the PR/HACCP regulations.

#### 5. SUMMARY OF PREVIOUS AUDITS

Final audit reports are available on FSIS' website at: <a href="http://www.fsis.usda.gov/regulations/Foreign">http://www.fsis.usda.gov/regulations/Foreign</a> Audit Reports/index.asp.

Previous audits of Australia's inspection system have indicated repeated non-compliance with FSIS regulatory requirements regarding documentation and implementation of HACCP and SSOP. In addition, the following concerns were identified during the last two audits:

#### May 2005 Audit

- 1 establishment delisted for non-compliance with requirements for effective food safety and sanitation procedures, and history of non-compliance resulting from the 2004 audit. *This establishment was relisted following corrective actions.*
- 1 establishment received a Notice of Intent to Delist (NOID) for non-compliance with requirements for effective food safety and sanitation procedures. *Corrective actions were taken within thirty days of receiving the NOID*.
- Laboratories using non-FSIS approved testing methods for the detection of pathogens and residues.
- Incorrect sample size (25 grams) used for the detection of *Salmonella* in Ready-to-eat (RTE) products.
- Inadequate oversight by AQIS of laboratories conducting microbiological and chemical testing of meat products being exported to the United States.
- Inadequate frequency of inspection by AQIS regarding cold stores.

#### June 2004 Audit:

- 1 establishment delisted for failure to have daily inspection. *This establishment had never exported to the United States*.
- 3 establishments received an NOID for not complying with various FSIS regulatory requirements. *Corrective actions were taken within thirty days of receiving the NOID*.
- 2 establishments were cited for inadequate implementation of post-mortem inspection requirements, i.e., heads of slaughtered cattle were not clearly identified with the carcasses. *Corrective actions were taken immediately*.
- 8 of 14 establishments were cited for inadequate government oversight.

#### 6. MAIN FINDINGS

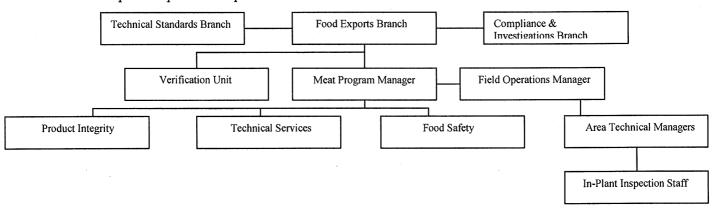
#### 6.1 Government Oversight

All official veterinarians and inspectors assigned to establishments certified by AQIS to export meat and meat products to the United States are official Australian government employees, receiving no remuneration from either industry or establishment personnel.

AQIS utilizes various levels of government oversight as part of its role in managing its export meat program. In addition to the in-plant verification role by the AQIS inspection team, AQIS uses Area Technical Managers (ATM) to conduct routine supervisory audits and Field Operation Managers (FOM) to conduct in-depth audits as means to help assure compliance with importing country requirements. In addition, AQIS has instituted a verification unit, separate from daily inspection activities, to address specific non-compliance or potential non-compliance issues. This verification unit reports directly to the National Manager of AQIS' Food Exports Branch, which is the same governing body that oversees Australia's meat inspection program.

#### 6.1.1 CCA Control Systems

AQIS has the organizational structure and staffing to assure uniform implementation of the U.S. import inspection requirements.



AQIS utilizes an interactive computer program called ELMER 3 as an essential part in assisting inspection personnel including providing a list of the U.S. import inspection requirements. ELMER 3 is accessible by in-plant inspection staff in certified establishments and is managed by the Meat Program Manager and respective staff.

#### 6.1.2 Ultimate Control and Supervision

AQIS has the ultimate legal control over and supervision of the official activities associated with the exports of meat products to the United States. In regard to the 2005 FSIS audit issue concerning frequency of inspection at cold stores, AQIS has instituted the FSIS requirement of quarterly inspection at cold stores. In Australia, certified cold stores do not handle exposed product and, therefore, are required by FSIS to implement only sanitation performance standards and have a minimum inspection frequency of once every three months.

In regard to the 2005 audit issue concerning government oversight of laboratories conducting microbiological or chemical testing of meat products destined for the United States, the following has been instituted as the means of addressing the FSIS 2005 audit concerns.

- Residue Testing Program A memorandum of understanding (MOU) between AQIS and the National Residue Survey, which is part of the Product Integrity Animal and Plant Health Division, was established since the last FSIS audit. Both parties are operating entities within the Australian Department of Agriculture, Fisheries and Forestry. This MOU gives AQIS increased oversight and involvement in the conduct of residue testing programs that support the export of meat products to the United States.
- Microbiology Testing Program A program within Technical Services has been added to AQIS since the last FSIS audit. This program is designed to increase direct oversight and involvement in the conduct of the microbiology testing programs that support the export of meat products to the United States. The functions of this program include independent audits of AQIS certified microbiological laboratories, reviews of sampling and testing methods to assure compliance with the AQIS and FSIS requirements, and reviews of the NATA, which is an Australian company that has an MOU with AQIS to perform 3<sup>rd</sup> party audits of AQIS certified laboratories.

AQIS has satisfactorily addressed the FSIS 2005 audit concerns. However, during this audit, FSIS has identified the following issue and requested AQIS to address:

• For the *Listeria monocytogenes* and *E. coli O157:H7* testing programs, establishment personnel are sampling, packaging, and submitting samples to private laboratories. Since these are regulatory sampling programs, the functions shall be performed by AQIS inspection personnel and tested in government laboratories.

Although these non-compliance issues occurred, FSIS does not believe it lead to questionable sampling and testing results. FSIS requires AQIS to submit the alternative programs for equivalence evaluations.

#### 6.1.3 Assignment of Competent, Qualified Inspectors

#### 6.1.3.1 Employment

Official veterinarians and inspectors are employed by AQIS either as permanent or contract employees. In either case, they are official government employees having the authority to carry out official AQIS inspection requirements.

#### 6.1.3.2 Training

AQIS has implemented various training programs for its inspection personnel, which include induction training for all newly hired veterinarians and meat inspectors and ongoing training for OPVs and senior meat inspectors. Induction training, which includes AQIS inspection requirements, must be successfully completed before trainees become authorized officers.

Ongoing training includes developmental seminars given in a class-room environment and inplant (hands-on) training. AQIS uses ELMER 3 as a tool to provide training modules and competency verification for the OPVs. Such training is required to be conducted within a specified time frame and is monitored by the ATMs.

#### 6.1.4 Authority and Responsibility to Enforce the Laws

The Export Control Act of 1982 and applicable regulations give AQIS the authority and responsibility to enforce Australia's meat inspection laws including meat and meat products produced for export to the United States. From this law, AQIS has implemented regulations to enforce the FSIS inspection requirements.

During this audit, the following concerns were identified:

- One establishment received an NOID due to non-compliance with zero tolerance of foreign materials (i.e., several lamb carcasses, passed for boning, were contaminated with fecal matter, seeds, grease, and/or wool).
- In three other establishments, AQIS was not enforcing all of the U.S. regulatory requirements.

Specific deficiencies are noted in the attached individual foreign establishment audit checklists.

#### 6.1.5 Adequate Administrative and Technical Support

As previously noted in this report, AQIS has implemented a program designed to increase direct oversight and involvement in the conduct of the microbiology testing programs that support the export of meat products to the United States. In addition, AQIS signed an MOU with the National Residue Survey as a means to increase oversight and involvement in the conduct of residue testing programs that support the export of meat products to the United States.

#### 6.2 Headquarters Audit

The auditors conducted a review of inspection system documents at headquarters, regional offices, and inspection offices of the 10 audited establishments. 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 United States
- Label approval records such as generic labels and animal raising claims.
- New laws and implementation documents such as regulations, notices, directives and guidelines.
- Sanitation, slaughter and processing inspection procedures and standards.
- 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 deficiencies arose as a result the examination of these documents.

#### 6.2.1 Audit of Regional and Local Inspection Sites

The FSIS auditors reviewed government oversight and enforcement activities at AQIS' regional offices of New South Wales, Queensland, and Victoria, and the inspection offices of the 10 audited establishments.

#### 7. ESTABLISHMENT AUDITS

Ten establishments certified by the government of Australia were audited. This included nine slaughter and/or processing establishments and one cold storage facility (ID Warehouse). One establishment was issued an NOID by AQIS due to non-compliance with zero tolerance of foreign material (i.e., several lamb carcasses, passed for boning, were contaminated with fecal matter, seeds, grease, and/or wool). This establishment was able to retain its certification for export to the United States as the establishment corrected all deficiencies within 30 days of the date the establishment was reviewed, and AQIS verified the corrective actions.

Specific deficiencies are noted in the attached individual foreign establishment audit checklists.

#### 8. RESIDUE AND MICROBIOLOGY LABORATORY AUDITS

No residue laboratories were audited. The audit of laboratories conducting microbiological testing was limited to verification of testing methods used by laboratories and oversight by the AQIS. The following four microbiology laboratories were visited:

- Est. 555 (EG Green @ Sons PTY LTD) North Dandalup
- Est. 235 (Dinmore Laboratory) Brisbane
- EML Consulting Services Laboratory Melbourne
- Silliker Microtech Laboratory Sydney

#### 9. SANITATION CONTROLS

As stated earlier, the FSIS auditors focused on five areas of risk to assess Australia's meat inspection system. The first of these risk areas that the FSIS auditor reviewed was Sanitation Controls.

Based on the on-site audits of establishments, and except as noted below, Australia's inspection system had controls in place for SSOP programs, all aspects of facility and equipment sanitation, the prevention of actual or potential instances of product cross-contamination, good personal hygiene practices, and good product handling and storage practices.

In addition, and except as noted below, Australia's inspection system had controls in place for water potability records, chlorination procedures, back-siphonage prevention, separation of operations, temperature control, work space, ventilation, ante-mortem facilities, welfare facilities, and outside premises.

#### 9.1 SSOP

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOP were met, according to the criteria employed in the U.S. domestic inspection program. The SSOP in all establishments was found to meet the basic FSIS regulatory requirements with the exception of the following:

- 3 of 10 establishments were cited for SSOP deficiencies. Examples of deficiencies included:
  - o Fecal contamination, seeds, hair and fecal speck were observed on several carcasses ready for boning. The carcasses were immediately trimmed.
  - O The conveyor belt used for handling edible product in a mutton boning room was observed with numerous deep cuts. Establishment was scheduled to replace belt.
  - O Several boxes in freezer were damaged by forklift and product inside boxes was compromised. The establishment officials took immediate corrective action.

Specific deficiencies are noted in the attached individual foreign establishment audit checklists.

#### 9.2 Sanitation

There were no major concerns in this area. Specific deficiencies are noted in the attached individual foreign establishment audit checklists.

#### 10. ANIMAL DISEASE CONTROLS

The second of the five risk areas that the FSIS auditor reviewed was Animal Disease Controls. These controls include ensuring adequate animal identification, humane handling and humane slaughter, control over condemned and restricted product, and procedures for sanitary handling of returned and reconditioned product. The auditor determined that Australia's inspection system had adequate controls in place with exception of the following issue:

• In one establishment, spinal cord removal occurred after final carcass inspection by the inspection official and verification by company employee.

There had been no outbreaks of animal diseases with public health significance since the last FSIS audit.

#### 11. SLAUGHTER/PROCESSING CONTROLS

The third of the five risk areas that the FSIS auditor reviewed was Slaughter/Processing Controls. The controls include the following areas: ante-mortem inspection procedures; ante-mortem disposition; post-mortem inspection procedures; post-mortem disposition; ingredients identification; control of restricted ingredients; formulations; processing schedules; equipment and records; and processing controls of cured, dried, and cooked products.

The controls also included the implementation of HACCP systems in 9 of 10 establishments and implementation of a generic *E. coli* testing program in slaughter establishments.

#### 11.1 Humane Handling and Slaughter

There were no observed deficiencies in this area.

#### 11.2 HACCP Implementation

All slaughter and processing establishments certified to export meat products to the United States are required to have developed and adequately implement a HACCP program. Each of these programs was evaluated according to the criteria employed in the U.S. domestic inspection program.

The HACCP programs were reviewed during the on-site audits of the nine slaughter and processing establishments. The one cold store reviewed was not required to implement a HACCP program.

All nine establishments had adequately implemented the HACCP requirements with the exception of the following issue:

• In two establishments, there was a deviation from CCP 1 for fecal contamination.

Specific deficiencies are noted in the attached individual foreign establishment audit checklists.

#### 11.3 Testing for Generic E. coli

Australia has adopted the generic E. coli testing methods that met the PR/HACCP criteria.

Nine of 10 establishments audited were required to meet the basic FSIS regulatory requirements for generic *E. coli* testing and were evaluated according to the criteria employed in the U.S. domestic inspection program.

There were no observed deficiencies in this area.

#### 11.4 Testing for *Listeria monocytogenes*

RTE meat products sampled for the detection of Lm are being collected by establishment employees and are being tested in private laboratories. Lm testing is a regulatory program and samples should be collected by AQIS and analyzed in government laboratories.

#### 12. RESIDUE CONTROLS

The fourth of the five risk areas that the FSIS auditor reviewed was Residue Control records. Australia's residue program is controlled by the government's National Residue Survey, which is part of the Ministry of Agriculture, Fisheries, and Forestry, and separate from AQIS.

As previously mentioned in this report, AQIS signed an MOU with the National Residue Survey as a means to increase oversight and involvement in the conduct of residue testing programs that support the export of meat products to the United States. This was established in response to a 2005 FSIS audit issue.

#### 13. ENFORCEMENT CONTROLS

The fifth of the five risk areas that the FSIS auditor reviewed was Enforcement Controls. These controls included the enforcement of inspection requirements such as required inspection coverage and the testing program for *Salmonella* and species verification.

#### 13.1 Daily Inspection in Establishments

Inspection was being conducted daily in all nine slaughter and processing establishments audited.

In regard to the one cold store facility audited, AQIS had inspection coverage of once every three months. Cold stores were operating under AQIS' Meat Safety Quality Assurance (MSQA) validation program and only handle non-exposed product. The operation at these cold stores is similar to establishments operating in the United States as ID Warehouses.

#### 13.2 Testing for Salmonella

Australia has adopted the FSIS requirements for testing for *Salmonella* with the exception of the following equivalent measure(s).

- Establishment employees collect samples monitored by AQIS plant management team on a random basis.
- Private laboratories analyze samples.

Nine of the 10 establishments audited were required to meet the basic FSIS regulatory requirements for *Salmonella* testing and were evaluated according to the criteria employed in the U.S. domestic inspection program.

Testing for Salmonella was properly conducted in all nine establishments.

#### 13.3 Species Verification

Species verification was being conducted in those establishments in which it was required.

#### 13.4 Monthly Reviews

During this audit, it was found that in all establishments visited, monthly supervisory reviews of certified establishments were being performed and documented as required. The only exception was inspection at the one cold storage facility. This establishment operates under the Australian MSQA program and inspection personnel were performing quarterly inspection coverage of these facilities.

#### 13.5 Inspection System Controls

The CCA had controls in place for ante-mortem and post-mortem inspection procedures and dispositions; restricted product and inspection samples; disposition of dead, dying, diseased or disabled animals; shipment security, including shipment between establishments; and prevention of commingling of product intended for export to the United States with product intended for the domestic market.

Areas of concern were:

• 4 of 10 establishments audited had deficiencies regarding enforcement of some aspects of FSIS regulatory requirements.

In addition, controls were in place for the importation of only eligible livestock from other countries, i.e., only from eligible third countries and certified establishments within those countries, and the importation of only eligible meat products from other counties for further processing.

Lastly, adequate controls were found to be in place for security items, shipment security, and products entering the establishments from outside sources.

#### 14. CLOSING MEETING

A closing meeting was held on August 30, 2006, in Canberra with the CCA. At this meeting, the primary findings were presented by the lead auditor.

The CCA understood and accepted the findings.

Mr. Steven A. McDermott Lead Auditor

### 15. ATTACHMENTS

Individual Foreign Establishment Audit Forms Foreign Country Response to Draft Final Audit Report

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT DATE		3. ESTABLISHMENT NO. 4. NAME OF COUNTRY			
Teys Bros Pty Ltd,	8-25-06		007 Australia			
Rockhampton,	5. NAME OF AUDITOR(		R(S)	6. TYPE OF AUDIT		
Queensland,	D. 0	4 - TT.1				
Australia		to Urba		X ON-SITE AUDIT DOCUMEN	T AUDIT	
Place an X in the Audit Results block to inc		compl	ance with requireme	ents. Use O if not applicable.		
Part A - Sanitation Standard Operating Procedures (	SSOP)	Audit		rt D - Continued	Audit	
Basic Requirements 7. Written SSOP		Results		nomic Sampling	Results	
Records documenting implementation.			33. Scheduled Sample			
			34. Species Testing		ļ	
9. Signed and dated SSOP, by on-site or overall authority.  Sanitation Standard Operating Procedures (SSOP)			35. Residue			
Ongoing Requirements			Part E -	Other Requirements		
10. Implementation of SSOP's, including monitoring of implement	ntation.		36. Export			
11. Maintenance and evaluation of the effectiveness of SSOP's.			37. Import			
<ol> <li>Corrective action when the SSOP's have falled to prevent di product contamination or adulteration.</li> </ol>	rect		38. Establishment Grounds	and Pest Control		
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	tion/Maintenance	· .	
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light		-	
14. Developed and implemented a written HACCP plan .		;	41. Ventilation	<u> </u>	<del> </del>	
<ol> <li>Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective ac</li> </ol>	ctions.		42. Plumbing and Sewage			
Records documenting implementation and monitoring of the HACCP plan.			43. Water Supply			
The HACCP plan is signed and dated by the responsible establishment individual.			44. Dressing Rooms/Lavatories  45. Equipment and Utensils			
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements			46. Sanitary Operations			
18. Monitoring of HACCP plan.			47. Employee Hygiene		<del> </del>	
19. Verification and validation of HACCP plan.			48. Condemned Product Control			
20. Corrective action written in HACCP plan.						
21. Reassessed adequacy of the HACCP plan.			Part F - Inspection Requirements			
22. Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event occ	of the urrences.		49. Government Staffing			
Part C - Economic / Wholesomeness			50. Daily Inspection Coverage			
23. Labeling - Product Standards			51. Enforcement			
24. Labeling - Net Weights 25. General Labeling			52. Humane Handling			
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Mo	oisture)		53. Animal Identification			
Part D - Sampling Generic <i>E. coli</i> Testing	,		54. Ante Mortem Inspection			
27. Written Procedures						
28. Sample Collection/Analysis			55. Post Mortem Inspection	•		
29. Records			Part G - Other Regu	latory Oversight Requirements		
Salmonella Performance Standards - Basic Requ	irements		56. European Community Di	rectives	0	
30. Corrective Actions	( ,		57. Monthly Review			
31. Reassessment			58.			
32. Written Assurance			59.			

Est. # 007, Teys Bros PTY LTD, Rockhampton - Queensland, 8 -25 -06, Slaughter/Bovine

"There were no significant findings to report after consideration of the nature, degree and extent of all observations."

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT D	ATE	3. ESTABLISHMENT NO. 4. NAME OF COUNTRY			
Cargiill Meat Processors,	8-24-06		249	Australia		
Tamwoth, NSW,	5. NAME OF	AUDITO	R(S)	6. TYPE OF AUDIT		
Australia	Dr. C	Dr. Oto Urban		X ON-SITE AUDIT DOCUMENT AUD		
Place an X in the Audit Results block to i	ndicate nor	comp	liance with requirem	ents. Use O if not applicable.		
Part A - Sanitation Standard Operating Procedures  Basic Requirements	s (SSOP)	Audit Results		rt D - Continued onomic Sampling	Audit Results	
7. Written SSOP			33. Scheduled Sample			
8. Records documenting implementation.			34. Species Testing			
9. Signed and dated SSOP, by on-site or overall authority.	***************************************		35. Residue			
Sanitation Standard Operating Procedures (SSO Ongoing Requirements	P)		Part E -	Other Requirements		
10. Implementation of SSOP's, including monitoring of impler	mentation.		36. Export			
11. Maintenance and evaluation of the effectiveness of SSOF	<sup>D'</sup> 8.		37. Import			
<ol> <li>Corrective action when the SSOPs have falled to prevent product contamination or adulteration.</li> </ol>	t direct		38. Establishment Grounds	and Pest Control		
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	ction/Maintenance		
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements	<b>,</b>		40. Light			
14. Developed and implemented a written HACCP plan .			41. Vendiadon		-	
<ol> <li>Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective</li> </ol>	e actions.		42. Plumbing and Sewage		-	
<ol> <li>Records documenting implementation and monitoring of HACCP plan.</li> </ol>	the	·	43. Water Supply			
<ol> <li>The HACCP plan is signed and dated by the responsible establishment individual.</li> </ol>			44. Dressing Rooms/Lavato			
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements			46. Sanitary Operations			
18. Monitoring of HACCP plan.			47. Employee Hygiene			
19. Verification and validation of HACCP plan.			48. Condemned Product Co	ontrol		
20. Corrective action written in HACCP plan.						
21. Reassessed adequacy of the HACCP plan.			Part F - I	nspection Requirements	and the same of th	
22. Records documenting: the written HACCP plan, monitori critical control points, dates and times of specific event of			49. Government Staffing			
Part C - Economic / Wholesomeness			50. Daily Inspection Covers	age		
23. Labeling - Product Standards			51. Enforcement		1	
24. Labeling - Net Weights						
25. General Labeling	· · · · · · · · · · · · · · · · · · ·		52. Humane Handling			
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins	/Moisture)		53. Animal Identification			
Part D - Sampling Generic <i>E. coli</i> Testing		and a Company	54. Ante Mortem Inspection	1		
27. Written Procedures		-	55. Post Mortem Inspection	1		
28. Sample Collection/Analysis						
29. Records			Part G - Other Reg	ulatory Oversight Requirements		
Salmonella Performance Standards - Basic Re	quirements		56. European Community D	Prectives	0	
30. Corrective Actions			57. Monthly Review			
31. Reassessment			58.			
32. Written Assurance			59.			

Est. # 249, Cargill Meat Processors PTY LTD, Tamworth – New South Wales, 8-24-06, Slaughter/Boning/Processing

"There were no significant findings to report after consideration of the nature, degree and extent of all observations."

61. NAME OF AUDITOR Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

Milan 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2 ALIDIT DA		3. ESTABLISHMENT NO. 4. NAME OF COUNTRY			
Ralphs Meat Company PTY LTD,	8-17-06		260	260 Australia		
Seymour,	5. NAME OF AUDITOR		R(S)	6. TYPE OF AUDIT		
Victoria,	Dr. Ota	T Talls ass		X ON SITE AUDIT DOCUMENT		
Australia	Dr. Oto			ON-SITE AODIT DOCUMEN	TAUDIT	
Place an X in the Audit Results block to in		compl	·			
Part A - Sanitation Standard Operating Procedures ( Basic Requirements	(SSOP)	Audit Results		rt D - Continued onomic Sampling	Audit Results	
7. Written SSOP			33. Scheduled Sample			
8. Records documenting implementation.			34. Species Testing			
9. Signed and dated SSOP, by on-site or overall authority.		***************************************	35. Residue			
Sanitation Standard Operating Procedures (SSOP Ongoing Requirements	)		Part E -	Other Requirements		
10. Implementation of SSOP's, including monitoring of implementation	entation.	Х	36. Export			
11. Maintenance and evaluation of the effectiveness of SSOP's	).		37. Import			
<ol> <li>Corrective action when the SSOP's have falled to prevent of product contamination or adulteration.</li> </ol>	lirect		38. Establishment Grounds	and Pest Control	X	
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	etion/Maintenance		
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light		<del> </del>	
14. Developed and implemented a written HACCP plan .		,	41. Ventilation			
<ol> <li>Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective a</li> </ol>	actions.		42. Plumbing and Sewage		<u> </u>	
<ol> <li>Records documenting implementation and monitoring of th HACCP plan.</li> </ol>	е		43. Water Supply		ļ	
The HACCP plan is signed and dated by the responsible establishment individual.			44. Dressing Rooms/Lavato			
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements			46. Sanitary Operations			
18. Monitoring of HACCP plan.		X			<del>                                     </del>	
19. Verification and validation of HACCP plan.		. ^	47. Employee Hygiene  48. Condemned Product Control			
20. Corrective action written in HACCP plan.			40. Condemned roduct of			
21. Reassessed adequacy of the HACCP plan.			Part F - Inspection Requirements			
Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event oc			49. Government Staffing			
Part C - Economic / Wholesomeness			50. Daily Inspection Cover	age		
23. Labeling - Product Standards			51. Enforcement		- X -	
24. Labeling - Net Weights 25. General Labeling			52. Humane Handling			
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/M	Noisture)		53. Animal Identification		<del> </del>	
Part D - Sampling						
Generic E. coli Testing			54. Ante Mortem Inspection	1		
27. Written Procedures			55. Post Mortem Inspection	n ·		
28. Sample Collection/Analysis		ļ	Part G - Other Reg	ulatory Oversight Requirements		
29. Records						
Salmonella Performance Standards - Basic Req	uirements		56. European Community D	Directives	0	
30. Corrective Actions			57. Monthly Review			
31. Reassessment			58.			
32. Written Assurance			59.			

Est. # 260, Ralphs Meat Company PTY LTD, Seymour - Victoria, 8-17-06, Slaugter/Boning

10/18/51 Seeds, hair and fecal speck was observed on 2 carcasses out of approximately 40 in cooler # 5. This was a deviation from CCP 1 for fecal contamination. The two carcasses were immediately trimmed. Additional corrective action included: (1) establishment officials immediately stopping slaughter line and re-inspecting all carcasses and retraining establishment trimmers, (2) all carcasses between cooler and boning room were reexamined by establishment – no findings, and (3) AQIS officials discussed findings with veterinarian-in-charge. FSIS regulation: 9 CFR 416.13c and 417.4(2).

Establishment has procedure whereby carcasses are reexamined for zero tolerance requirements (feces, ingesta, hair, etc) before entering the boning room.

38 Several gaps under two receiving doors in the product loading area were observed creating the possibility of entrance by pests. Gap under one door was probably due to door not being completely closed. Establishment took immediate corrective action including scheduling maintenance repair for one door. FSIS regulation: 9 CFR 416.2(a).

62. AUDITOR SIGNATURE AND DATE

61. NAME OF AUDITOR
Oto Urban. DVM

Welcer 10-6-06

	ENT NAME AND LOCATION	2. AUDIT DATE	:  :	3. ESTABLISHMENT NO. 4. NAME OF COUNTRY		
	ustralian Meat Marketing	8-14-06		572 Australia		
	tive Limited,	5. NAME OF AUDITOR		R(S)	6. TYPE OF AUDIT	
Katanning						
	Western Australia Dr. Oto U				X ON-SITE AUDIT DOCUMEN	T AUDIT
	in the Audit Results block to inc		mpli			
Part A - Sanita	ation Standard Operating Procedures ( Basic Requirements	- ,	udit esults		rt D - Continued onomic Sampling	Audit Results
7. Written SSO	P			33. Scheduled Sample		
8. Records doc	umenting implementation.			34. Species Testing		
9. Signed and o	lated SSOP, by on-site or overall authority.			35. Residue		
Sanitation S	Standard Operating Procedures (SSOP) Ongoing Requirements			Part E -	Other Requirements	
10. Implementa	tion of SSOP's, including monitoring of impleme	ntation.	κ	36. Export		
11. Maintenanc	e and evaluation of the effectiveness of SSOP's	·		37. Import		
	action when the SSOP's have falled to prevent di tamination or adulteration.	irect	·	38. Establishment Grounds	and Pest Control	
13. Daily record	s document item 10, 11 and 12 above.			39. Establishment Construc	tion/Maintenance	
	Hazard Analysis and Critical Control CCP) Systems - Basic Requirements			40. Light		
	and implemented a written HACCP plan .			41. Ventilation		
	the HACCP list the food safety hazards, rol points, critical limits, procedures, corrective a	ctions.		42. Plumbing and Sewage		
16. Records do HACCP pla	cumenting implementation and monitoring of the n.	•		43. Water Supply		-
	P plan is signed and dated by the responsible entindividual.			44. Dressing Rooms/Lavato		-
	Analysis and Critical Control Point ) Systems - Ongoing Requirements			46. Sanitary Operations		1.
18. Monitoring						+
	and validation of HACCP plan.			47. Employee Hygiene		
20 Corrective s	action written in HACCP plan.			48. Condemned Product Co	ontrol	
	d adequacy of the HACCP plan.			Part F - II	nspection Requirements	
22. Records do	ocumenting: the written HACCP plan, monitoring trol points, dates and times of specific event occ	of the		49. Government Staffing		
	t C - Economic / Wholesomeness			50. Daily Inspection Covera	age	
23. Labeling - F	Product Standards					<del></del>
24. Labeling - N	let Weights			51. Enforcement		- X -
25. General La	beling			52. Humane Handling		
26. Fin. Prod. S	Standards/Boneless (Defects/AQL/Pork Skins/M	oisture)		53. Animal Identification		
	Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem Inspection		
27. Written Pro	cedures			55. Post Mortem Inspection	1	
28. Sample Co	llection/Analysis					
29. Records				Part G - Other Regu	ulatory Oversight Requirements	
Salmonella	a Performance Standards - Basic Requ	lirements		56. European Community D	rectives	0
30. Corrective	Actions			57. Monthly Review		
31. Reassessm	nent	·		58.		
32. Written Ass	urance			59.		

Est. # 572, Western Australian Meat Marketing Cooperative LTM, Katanning - Western Australia, Slaughter/Boning

10/51 Conveyor belt used for handling edible product in mutton boning room was observed with numerous deep cuts. There was a potential to contaminate edible product. Establishment was scheduled to replace belt 9 CFR 416.13(c).

10/51 Several boxes in freezer were damaged by forklift and product inside boxes was compromised. Immediate corrective action was taken by the establishment officials 9 CFR 416.13(c).

61. NAME OF AUDITOR Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

### 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT DAT	TE	3. ESTABLISHMENT NO.	4. NAME OF COUNTRY	2
Tasmangroup,	8-18-06		688 Australia		
Brooklyn,	5. NAME OF AUDITOR		R(S)	6. TYPE OF AUDIT	
Victoria,	Dr. Oto Urban		•		
Australia	<u> </u>			ON-SITE AUDIT DOCUMEN	T AUDIT
Place an X in the Audit Results block to inc		ompl			
Part A - Sanitation Standard Operating Procedures ( Basic Requirements		Audit Results		rt D - Continued onomic Sampling	Audit Results
7. Written SSOP			33. Scheduled Sample		
8. Records documenting implementation.			34. Species Testing		
9. Signed and dated SSOP, by on-site or overall authority.			35. Residue		
Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements			Part E -	Other Requirements	
10. Implementation of SSOP's, including monitoring of impleme	entation.		36. Export		
11. Maintenance and evaluation of the effectiveness of SSOP's			37. Import		
<ol> <li>Corrective action when the SSOPs have failed to prevent d product contamination or adulteration.</li> </ol>	irect		38. Establishment Grounds	and Pest Control	
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	ction/Maintenance	
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light		ļ
14. Developed and implemented a written HACCP plan .			41. Ventilation		<u> </u>
15. Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective a	ctions.		42. Plumbing and Sewage		-
16. Records documenting implementation and monitoring of the HACCP plan.	е		43. Water Supply		
The HACCP plan is signed and dated by the responsible establishment individual.			44. Dressing Rooms/Lavato		
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements			46. Sanitary Operations		+
18. Monitoring of HACCP plan.					<del> </del>
19. Verification and validation of HACCP plan.			47. Employee Hygiene 48. Condemned Product Co	ontrol	
20. Corrective action written in HACCP plan.					
21. Reassessed adequacy of the HACCP plan.			Part F - I	nspection Requirements	
22. Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event oc			49. Government Staffing		
Part C - Economic / Wholesomeness			50. Daily Inspection Covera	age	
23. Labeling - Product Standards			51. Enforcement		
24. Labeling - Net Weights			<u> </u>		-
25. General Labeling			52. Humane Handling		-
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/M	loisture)		53. Animal Identification		
Part D - Sampling Generic <i>E. coli</i> Testing	·		54. Ante Mortem Inspection	1	
27. Written Procedures			55. Post Mortem Inspection	n	
28. Sample Collection/Analysis			<u> </u>		
29. Records			Part G - Other Reg	ulatory Oversight Requirements	
Salmonella Performance Standards - Basic Requ	uirements		56. European Community D	Prectives	0
30. Corrective Actions			57. Monthly Review	·	
31. Reassessment			58. BSE		Х
32. Written Assurance			59.	·	

Est. # 688, Tasman Group Services PTY LTD, Brooklyn - Victoria, 8-18-06, Slaughter/Boning

58 Establishment was removing spinal cords after the final carcass inspection station. Establishment officials performed immediate corrective action by relocating spinal cord removal to a place prior to the final carcass inspection station. No carcasses in coolers or in boning room were observed with spinal cord. FSIS regulation: 9 CFR 310.22(d)3.

61. NAME OF AUDITOR Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

Ho Whan 10-6-06

Place an X in the Audit Results block to indicate noncompliance with requirements. Use 0 if not applicable Part A - Sanitation Standard Operating Procedures (SSOP) Bask Requirements  7. Written SSOP 8. Records documenting implementation. 9. Signed and dated SSOP, by on-site or overall authority. Sanitation Standard Operating Procedures (SSOP) Congoing Requirements 10. Implementation of SSOPs, including monitoring of implementation. 11. Maintenance and evaluation of the effectiveness of SSOPs. 12. Corrective action when the SSOPs have falled to prevent direct pedicut contamination or adderation. 13. Daily records documenting in 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) plain. 14. Developed and implemented a written HACCP plan. 15. Contents of the HACCP plain is stend and dated by the responsible establishment individual. 16. Records documenting implementation and monitoring of the HACCP plan. 17. The HACCP plan is stend and dated by the responsible establishment individual. 18. Monitoring of HACCP plan. 19. Verification and validation of HACCP plan. 19. Verification and validation of HACCP plan. 19. Verification and validation of HACCP plan. 20. Corrective action written in HACCP plan. 21. Reassessed adequacy of the HACCP plan. 22. Records documenting in the written HACCP plan, monitoring of the critical control / Molecomeness 23. Labeling - Product Standards 24. Labeling - Product Standards 25. General Labeling 26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. Coll Testing  7. Writen SSOP 8. Aside Septice with requirements and standard with requirements and standard with requirements and standards  8. Aside Septics with requirements and standards  8. Establishment Construction/Adaintenance  9. Establishment C	
South Australia  Dr. Oto Urban  Place an X in the Audit Results block to indicate noncompliance with requirements. Use 0 if not applicable Part A - Sanitation Standard Operating Procedures (SSOP) Basic Requirements  Results  Part D - Continued Economic Sampling  3.3. Schedded Sample  3.4. Species Testing  3.5. Residue  Part E - Other Requirements  Only play Regularements  Dr. Other Requirements  Sanitation Standard Operating Procedures (SSOP) Ongoing Regularements  Dr. Other Requirements  Dr. Other Results  Part E - Other Requirements  Dr. Other Requirements  Dr. Other Results  Part E - Other Requirements  Dr. Other Results  Part E - Other Requirements  Dr. Other Results  Dr. Other Results  Part E - Other Requirements  Dr. Other Results  Dr. Other Results  Part E - Other Requirements  Dr. Other Results  Part E - Other Requirements  Dr. Other Results  Dr. Dr. Other Results  Dr. Dr. Dr. Other Results  Dr. Dr. Dr. Dr. Other Results  Dr.	
Place an X in the Audit Results block to indicate noncompliance with requirements. Use O if not applicable Part A - Sanitation Standard Operating Procedures (SSOP)  Basic Requirements  Result  Resul	
Part A - Sanitation Standard Operating Procedures (SSOP) Basic Requirements  7. Written SSOP 8. Records documenting implementation. 9. Signed and dated SSOP, by on-site or overall authority. Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements 10. Implementation of SSOP's, including monitoring of implementation. 11. Maintenance and evaluation of the effectiveness of SSOP's. 12. Corrective action when the SSOP's have failed to prevent direct product contamination or aduteration. 13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements 14. Developed and implemented a written HACCP plan. 15. Corfective action with the stop and dated by the responsible establishment individual. 16. Records documenting implementation and monitoring of the HACCP plan. 17. The HACCP plan is signed and dated by the responsible establishment individual. 18. Monitoring of HACCP plan. 19. Verification and validation of HACCP plan. 20. Corrective action written in HACCP plan. 21. Results and Critical Control Point (HACCP) Systems - Ongoing Requirements 22. Records documenting: the written HACCP plan. 23. Labeling - Product Standards 24. Labeling - Product Standards 25. General Labeling 26. Fin. Prod. Standards/Soneless (Cefects/AQL/Pork SkinsMolsture)  Part D - Sampling Generic E. coli Testing  13. Scheduled Sample 36. Species Testing 35. Residue  Part E - Other Requirements 36. Export  95. Export  16. Labeling - Product Standards 37. Import 38. Establishment Crounds and Pest Control 40. Light 41. Ventilation 42. Plumbing and Sewage 44. Diressing Rooms/Lavatories 45. Equipment and Utensils 46. Sanitary Operations 47. Employee Hyglene 48. Condemned Product Control 49. Government Staffing 49. Government Staffing 50. Daily Inspection Coverage 51. Enforcement 52. Humane Handling 53. Animal Identification 54. Animal Identification	TIDUA TI
Basic Requirements 7. Written SSOP 8. Records documenting implementation. 9. Signed and dated SSOP, by on-site or overall authority.  Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements 10. Implementation of SSOPs, including monitoring of implementation. 11. Maintenance and evaluation of the effectiveness of SSOP's. 12. Corrective action when the SSOPs have failed to prevent direct product contamination or adulteration. 13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements 14. Developed and implemented a written HACCP plan. 16. Records documenting implementation and monitoring of the HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements 14. Ventilication and availation of HACCP plan. 15. Contents of the HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements 16. Monitoring of HACCP plan. 17. The HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements 16. Monitoring of HACCP plan. 17. Reassessed adequacy of the HACCP plan. 18. Verification and validation of HACCP plan. 19. Verification of HACCP plan. 20. Corrective action written in HACCP plan. 21. Records documenting: the written HACCP plan, ponitoring of the critical control problems, dates and thines of specific event occurrices.  Part C - Economic / Wholesomeness 23. Labeling - Product Standards 24. Labeling - Product Standards 25. General Labeling 26. Fin. Prod. Standards/Boneless (Defects/AQU/Pork, Skins/Moisture)  Part D - Sampling Generic E. coli Testing	
8. Records documenting implementation. 9. Signed and dated SSOP, by on-site or overall authority. 5. Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements 10. Implementation of SSOP's, including monitoring of implementation. 11. Maintenance and evaluation of the effectiveness of SSOP's. 12. Corrective action when the SSOP's have failed to prevent direct product contamination or adulteration. 13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements 14. Developed and implemented a written HACCP plan. 15. Corrective of the HACCP list the foot safety hazards, critical control points, critical	Audit Results
9. Signed and dated SSOP, by cn-site or overall authority.  Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements  10. Implementation of SSOPs, including monitoring of implementation.  11. Maintenance and evaluation of the effectiveness of SSOP's.  12. Corrective action when the SSOPs have faled to prevent direct poduct contamination or adulteration.  13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  15. Corrients of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions of common limits, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan is eigned and dated by the responsible establishment indivitual.  18. Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and three of specific evert occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Product Standards  25. General Labeling  26. Fin. Prod Standards/Boneless (Defeds/ACU/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  37. Import  36. Export  37. Import  38. Establishment Grounds and Pest Control  40. Light  41. Ventilation  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Government Staffing  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortern Inspection	
Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements  10. Implementation of SSOPs, including monitoring of implementation.  11. Maintenance and evaluation of the effectiveness of SSOPs.  12. Corrective action when the SSOPs have failed to prevent direct product contamination or adulteration.  13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan.  15. Cortents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the establishment Individual.  17. The HACCP plan is signed and dated by the responsible establishment Individual.  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Ressessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, desse and times of specific evert occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Product Standards  25. General Labeling  Part D - Sampling  Generic E. coli Testing  Part D - Sampling  Generic E. coli Testing  Part D - Sampling  Generic E. coli Testing	
Ongoing Requirements  10. Implementation of SSOPs, including monitoring of implementation.  11. Maintenance and evaluation of the effectiveness of SSOP's.  12. Corrective action when the SSOPs have failed to prevent direct product contamination or adulteration.  13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan.  15. Corrents of the HACCP list the food safety hazards, critical control points, critical control points, critical control points, critical limits, procedures, corrective actions or product shall have been applied to the HACCP plan is signed and dated by the responsible establishment individual.  17. The HACCP plan is signed and dated by the responsible establishment individual.  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan.  23. Labeling - Product Standards  24. Labding - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  56. Ante Mortem Inspection	
11. Maintenance and evaluation of the effectiveness of SSOP's.  12. Cornective action when the SSOP's have falled to prevent direct product contamination or adulteration.  13. Daily records document item 10, 11 and 12 above.  14. Developed and implemented a written HACCP plan.  15. Cordents of the HACCP list the foot safety hexards, critical control points, critical imits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan.  17. The HACCP plan is signed and dated by the responsible establishment individual.  18. Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, ponitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Product Standards  25. General Labeling  Part D - Sampling Generic E. coll Testing  35. Establishment Grounds and Pest Control  40. Light  40. Light  41. Ventilation  42. Plumbing and Sewage  41. Ventilation  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hyglene  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Requirements  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
12. Corrective action when the SSCP's have faled to prevent direct product contamination or adulteration.  13. Daily records document item 10, 11 and 12 above.  Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan.  15. Cortents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan is signed and dated by the responsible establishment individual.  17. The HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labgling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  27. Animal Identification  38. Establishment Grounds and Pest Control  40. Light  41. Ventilation  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
product contamination or adulteration.  38. Establishment Grounds and Petr Control Pairt B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  40. Light 41. Ventilation  42. Plumbing and Sewage 43. Water Supply 44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. The HACCP plan is signed and dated by the responsible establishment individual.  48. Monitoring of HACCP plan.  49. Verification and validation of HACCP plan.  40. Light 41. Ventilation  42. Plumbing and Sewage 43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  48. Condemned Product Control  49. Government Staffing  49. Government Staffing  49. Government Staffing  50. Daily Inspection Coverage  51. Labeling - Product Standards  52. Labeling - Product Standards  53. Establishment Construction/Maintenance  40. Light  41. Ventilation  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Labeling - Product Standards  53. Animal Identification  Part D - Sampling  Generic E. coli Testing  54. Ante Mortem Inspection	
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan.  15. Contents of the HACCP list the food safety hazards, critical control points, critical control point	
Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan.  15. Cortents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan is signed and dated by the responsible establishment individual.  17. The HACCP plan is signed and dated by the responsible establishment individual.  18. Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control protect control onthe control control onthe points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQU/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  41. Ventilation  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rocms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Morten Inspection	
14. Developed and implemented a written HACCP plan .  15. Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan.  17. The HACCP plan is signed and dated by the responsible establishment individual.  18. Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  41. Veritications.  42. Plumbing and Sewage  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
critical control points, critical limits, procedures, corrective actions.  16. Records documenting implementation and monitoring of the HACCP plan.  17. The HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  43. Water Supply  44. Dressing Rooms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	<del> </del>
HACCP plan.  17. The HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork SkinsMoisture)  Part D - Sampling  Generic E. coli Testing  44. Dressing Rcoms/Lavatories  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	_
17. The HACCP plan is signed and dated by the responsible establishment individual.  Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  18. Monitoring of HACCP plan.  19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  45. Equipment and Utensils  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements  46. Sanitary Operations  47. Employee Hygiene  48. Condemned Product Control  49. Government Staffing  49. Government Staffing  49. Government Staffing  49. Daily Inspection Coverage  49. Daily Inspection Coverage  49. Labeling - Product Standards  40. Daily Inspection Coverage  41. Enforcement  42. Labeling - Net Weights  43. Animal Identification  44. Ante Mortem Inspection	
19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  48. Condemned Product Control  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	×
19. Verification and validation of HACCP plan.  20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  48. Condemned Product Control  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification	_
21. Reassessed adequacy of the HACCP plan.  22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  Part B - Inspection Requirements  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	1
22. Records documenting: the written HACCP plan, monitoring of the critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  49. Government Staffing  50. Daily Inspection Coverage  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
critical control points, dates and times of specific event occurrences.  Part C - Economic / Wholesomeness  50. Daily Inspection Coverage  51. Enforcement  52. Labeling - Net Weights  52. Humane Handling  53. Animal Identification  Part D - Sampling Generic E. coli Testing  54. Ante Mortem Inspection	
23. Labeling - Product Standards  24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  27. Humane Handling  28. Animal Identification  Part D - Sampling Generic E. coli Testing  29. Humane Handling  50. Animal Identification  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
24. Labeling - Net Weights  25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  51. Enforcement  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	
25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  52. Humane Handling  53. Animal Identification  54. Ante Mortem Inspection	X
25. General Labeling  26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)  Part D - Sampling Generic E. coli Testing  53. Animal Identification  54. Ante Mortem Inspection	
Part D - Sampling Generic E. coli Testing  54. Ante Mortem Inspection	+
Generic E. coli Testing 54. Ante Mortem Inspection	
OT Witten Deposition	
27. Written Procedures 55. Post Mortem Inspection	
28. Sample Collection/Analysis	
29. Records Part G - Other Regulatory Oversight Requirements	
Salmonella Performance Standards - Basic Requirements  56. European Community Directives	0
30. Corrective Actions 57. Monthly Review	
31. Reassessment 58.	
32. Written Assurance 59.	

Est. # 866, Lobethal Australia PTY LMT, Lobethal - South Australia, 8-15-06, Slaughter/Boning

46/51 Establishment employee handling inedible product on the floor was observed to be in physical contact with the saw. Immediate corrective action was taken by establishment by re-cleaning product-contact areas of saw. FSIS regulation 9 CFR 416.4(a).

Sto Motor 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT D	ATE	3. ESTABLISHMENT NO. 4. NAME OF COUNTRY			
Tatiara Meat Company PTY LTD,	8-16-06		1614 Australia			
Bordrtown,	5. NAME OF AUDITOR		R(S)	6. TYPE OF AUDIT		
South Australia	Dr. Oto Urban					
	<u> </u>			X ON-SITE AUDIT DOCUMENT	AUDIT	
Place an X in the Audit Results block to inc		compl	_			
Part A - Sanitation Standard Operating Procedures ( Basic Requirements	SSOP)	Audit Results		rt D - Continued onomic Sampling	Audit Results	
7. Written SSOP	•	1100000	33. Scheduled Sample	monic sampling		
8. Records documenting implementation.			· · · · · · · · · · · · · · · · · · ·			
Signed and dated SSOP, by on-site or overall authority.			34. Species Testing			
Sanitation Standard Operating Procedures (SSOP)			35. Residue			
Ongoing Requirements			Part E -	Other Requirements		
10. Implementation of SSOP's, including monitoring of impleme	ntation.		36. Export			
11. Maintenance and evaluation of the effectiveness of SSOP's	•		37. Import	·		
<ol> <li>Corrective action when the SSOPs have faled to prevent di product contamination or adulteration.</li> </ol>	irect		38. Establishment Grounds	and Pest Control		
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	ction/Maintenance		
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light			
14. Developed and implemented a written HACCP plan .			41. Ventilation			
15. Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective a	ctions.		42. Plumbing and Sewage			
16. Records documenting implementation and monitoring of the HACCP plan.			43. Water Supply			
The HACCP plan is signed and dated by the responsible establishment individual.			44. Dressing Rooms/Lavatories  45. Equipment and Utensils			
Hazard Analysis and Critical Control Point				<b>S</b>	-	
(HACCP) Systems - Ongoing Requirements  18. Monitoring of HACCP plan.			46. Sanitary Operations			
19. Verification and validation of HACCP plan.	· · · · · · · · · · · · · · · · · · ·		47. Employee Hygiene			
		<b></b>	48. Condemned Product C	ontrol		
20. Corrective action written in HACCP plan.  21. Reassessed adequacy of the HACCP plan.			Part F - Inspection Requirements			
, <del></del>		<b></b>				
22. Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event occ			49. Government Staffing			
Part C - Economic / Wholesomeness			50. Daily Inspection Coverage			
23. Labeling - Product Standards			51. Enforcement			
24. Labeling - Net Weights 25. General Labeling			52. Humane Handling			
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pcrk Skins/Moisture)			53. Animal Identification			
Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem Inspection			
27. Written Procedures			55. Post Mortem Inspection	n		
28. Sample Collection/Analysis			]			
29. Records			Part G - Other Reg	ulatory Oversight Requirements		
Salmonella Performance Standards - Basic Requ	uirements		56. European Community D	Directives .	0	
30. Corrective Actions			57. Monthly Review		ļ	
31. Reassessment			58.			
32. Written Assurance			59.			

Est. # 1614, Tatiara Meat Company PTY LTD, Bordertown - South Australia, 8-16-06, Slaughter/Boning/Processing

"There were no significant findings to report after consideration of the nature, degree and extent of all observations."

61. NAME OF AUDITOR Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

Stor Mba. 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT D	ATE	3. ESTABLISHMENT NO.	4. NAME OF COUNTRY		
Australian Country Choice Pty Ltd,	8-28-06		1620 Australia			
Brisbane, Murarrie,	5. NAME OF	AUDITO	R(S)	6. TYPE OF AUDIT		
Queensland	<b>D</b>					
	Dr. C	Oto Urba	an	X ON-SITE AUDIT DOCUMENT	AUDIT	
Place an X in the Audit Results block to in	dicate nor	compl	iance with requirem	ents. Use O if not applicable.		
Part A - Sanitation Standard Operating Procedures	(SSOP)	Audit		rt D - Continued	Audit	
Basic Requirements		Results		onomic Sampling	Results	
7. Written SSOP			33. Scheduled Sample		<del></del>	
8. Records documenting implementation.			34. Species Testing			
9. Signed and dated SSOP, by on-site or overall authority.			35. Residue			
Sanitation Standard Operating Procedures (SSOP	)		Part E -	Other Requirements		
Ongoing Requirements						
10. Implementation of SSOP's, including monitoring of implementation of state of the state of th		<del> </del>	36. Export		<del></del>	
11. Maintenance and evaluation of the effectiveness of SSOP's		· · ·	37. Import			
<ol> <li>Corrective action when the SSOPs have falled to prevent opportunity contamination or adulteration.</li> </ol>	direct		38. Establishment Grounds	and Pest Control		
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	ction/Maintenance		
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light			
14. Developed and implemented a written HACCP plan .			41. Ventilation			
<ol> <li>Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective actions.</li> </ol>			42. Plumbing and Sewage			
16. Records documenting implementation and monitoring of the HACCP plan.		-	43. Water Supply			
17. The HACCP plan is signed and dated by the responsible			44. Dressing Rooms/Lavate			
establishment individual.  Hazard Analysis and Critical Control Point			45. Equipment and Utensils	\$		
(HACCP) Systems - Ongoing Requirements			46. Sanitary Operations			
18. Monitoring of HACCP plan.			47. Employee Hygiene			
19. Verification and validation of HACCP plan.	·	<del> </del>	48. Condemned Product C	ontrol		
20. Corrective action written in HACCP plan.			Part F - Inspection Requirements			
21. Reassessed adequacy of the HACCP plan.						
22. Records documenting: the written HACCP plan, monitorin critical control points, dates and times of specific event or	g of the courrences.		49. Government Staffing			
Part C - Economic / Wholesomeness	•		50. Daily Inspection Cover	age		
23. Labeling - Product Standards			51. Enforcement			
24. Labeling - Net Weights			J. Ellorcellien	· · · · · · · · · · · · · · · · · · ·	<del>                                     </del>	
25. General Labeling		<b> </b>	52. Humane Handling			
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Moisture)			53. Animal Identification			
Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem Inspection			
27. Written Procedures			55. Post Mortem Inspectio	55. Post Mortem Inspection		
28. Sample Collection/Analysis			-			
29. Records		1	Part G - Other Reg	ulatory Oversight Requirements		
Salmonella Performance Standards - Basic Req	uirements		56. European Community [	Directives	0	
30. Corrective Actions			57. Monthly Review	•		
31. Reassessment			58.			
32. Written Assurance			59.			

Est. # 1620, Australian Country Choice Production PTY LTD, Cannon Hill - Queensland, 8-28-06, Slaughter/Boning/Processing

"There were no significant findings to report after consideration of the nature, degree and extent of all observation".

61. NAME OF AUDITOR Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

Of When 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT D	ATE	3. ESTABLISHMENT NO.	4. NAME OF COUNTRY	
Fletcher International Exports,	8-23-0	6	2309 Australia		
Dubbo,	5. NAME OF				
New South Wales	D. 04	- T Tule			
	1	o Urban		X ON-SITE AUDIT DOCUMEN	T AUDIT
Place an X in the Audit Results block to inc	dicate non	compl	iance with requiren	nents. Use O if not applicable.	
Part A - Sanitation Standard Operating Procedures (	SSOP)	Audit		art D - Continued	Audit
Basic Requirements		Results	<b>{</b>	onomic Sampling	Results
7. Written SSOP			33. Scheduled Sample		0
8. Records documenting implementation.		ļ	34. Species Testing		
9. Signed and dated SSOP, by on-site or overall authority.			35. Residue		
Sanitation Standard Operating Procedures (SSOP) Ongoing Requirements			Part E	- Other Requirements	
10. Implementation of SSOP's, including monitoring of impleme	entation.	x	36. Export	• :	
11. Maintenance and evaluation of the effectiveness of SSOP's.	•	Ī	37. Import		
Corrective action when the SSOP's have falled to prevent di product contamination or adulteration.	irect	х	38. Establishment Ground	s and Pest Control	T
13. Daily records document item 10, 11 and 12 above.			39. Establishment Constru	uction/Maintenance	
Part B - Hazard Analysis and Critical Control			40. Light		
Point (HACCP) Systems - Basic Requirements  14. Developed and implemented a written HACCP plan .			41. Ventilation		
15. Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective a	ections		42. Plumbing and Sewage		
16. Records documenting implementation and monitoring of the			43. Water Supply		
HACCP plan.  17. The HACCP plan is signed and dated by the responsible			44. Dressing Rooms/Lava	tories	
establishment individual.  Hazard Analysis and Critical Control Point			45. Equipment and Utens	ils	
(HACCP) Systems - Ongoing Requirements			46. Sanitary Operations		
18. Monitoring of HACCP plan.		х	47. Employee Hygiene		
19. Verification and validation of HACCP plan.	·	<u> </u>	48. Condemned Product (	Control	<b>-</b>
20. Corrective action written in HACCP plan.					
21. Reassessed adequacy of the HACCP plan.			Part F -	Inspection Requirements	
Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event occ			49. Government Staffing		
Part C - Economic / Wholesomeness			50. Daily Inspection Cove	rage	
23. Labeling - Product Standards			51. Enforcement		x
24. Labeling - Net Weights			52. Humane Handling		+
25. General Labeling		ļ	JZ. Fruitiane Francing		<u> </u>
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/M	loisture)	0	53. Animal Identification		
Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem Inspection	on	
27. Written Procedures			55. Post Mortem Inspection	on	
28. Sample Collection/Analysis					
29. Records			Part G - Other Re	gulatory Oversight Requirements	
Salmonella Performance Standards - Basic Requ	uirements		56. European Community	Directives	0
30. Corrective Actions			57. Monthly Review		
31. Reassessment			58. NOID		Х
32. Written Assurance			59.		

Est. # 2309, Fletcher International Exports, Dubbo - New South Wales, 8-23-06, Slaughter/Boning/Processing

10/18/51 Fecal contamination was observed on three carcasses in one of the carcass chillers. Two of these deficiencies were initially observed by the AQIS official performing the inspection and the third was observed by the FSIS auditor. This was a deviation from CCP 1 and its critical limit and there was no product check for fecal contamination before going into the boning room. Company documents from the previous night's operation did not indicate any fecal contamination during the verification process. Establishment performed immediate corrective action by trimming and checking carcasses on the AQIS request. FSIS regulation 9 CFR 416.13c and 417.4(2).

12 Direct product contamination of carcasses with seeds, wool, and grease were missed by establishment officials on the kill floor and boning room during the verification process. AQIS official and FSIS auditor pointed out these deficiencies. FSIS regulation 9 CFR 416.15a.

58 AQIS officials issued an Notice of Intent to Delist to establishment.

61. NAME OF AUDITOR
Oto Urban, DVM

62. AUDITOR SIGNATURE AND DATE

Ho Mibon 10-6-06

1. ESTABLISHMENT NAME AND LOCATION	2. AUDIT DA	ATE	3. ESTABLISHMENT NO. 4. NAME OF COUNTRY		
Swire Cold Store PTY LTD,	8-11-06		5253 Australia		
Palmyra,	5. NAME OF AUDITO		R(S)	6. TYPE OF AUDIT	
West Australia	Dr. Oto Urbar				
				X ON-SITE AUDIT DOCUMEN	T AUDIT
Place an X in the Audit Results block to inc		compl			
Part A - Sanitation Standard Operating Procedures ( Basic Requirements	SSOP)	Audit Results	• • • • • • • • • • • • • • • • • • • •	rt D - Continued onomic Sampling	Audit Results
7. Written SSOP			33. Scheduled Sample		<del> </del>
8. Records documenting implementation.			· · · · · · · · · · · · · · · · · · ·		0
Signed and dated SSOP, by on-site or overall authority.			34. Species Testing	The state of the s	0
Sanitation Standard Operating Procedures (SSOP)			35. Residue		,
Ongoing Requirements			Part E -	Other Requirements	
10. Implementation of SSOP's, including monitoring of impleme			36. Export		
11. Maintenance and evaluation of the effectiveness of SSOP's.			37. import		<u></u>
<ol> <li>Corrective action when the SSOPs have falled to prevent di product contamination or adulteration.</li> </ol>	irect		38. Establishment Grounds	and Pest Control	
13. Daily records document item 10, 11 and 12 above.			39. Establishment Construc	tion/Maintenance	
Part B - Hazard Analysis and Critical Control Point (HACCP) Systems - Basic Requirements			40. Light		
14. Developed and implemented a written HACCP plan .		0	41. Ventilation		
<ol> <li>Contents of the HACCP list the food safety hazards, critical control points, critical limits, procedures, corrective ac</li> </ol>	ctions.	0	42. Plumbing and Sewage		
<ol> <li>Records documenting implementation and monitoring of the HACCP plan.</li> </ol>	•	0	43. Water Supply		
<ol> <li>The HACCP plan is signed and dated by the responsible establishment individual.</li> </ol>		0	44. Dressing Rooms/Lavato		X
Hazard Analysis and Critical Control Point (HACCP) Systems - Ongoing Requirements			46. Sanitary Operations		<del>  ^-</del>
18. Monitoring of HACCP plan.		0	47. Employee Hygiene		
19. Verification and validation of HACCP plan.		0	48. Condemned Product Co	ontrol	<del>                                     </del>
20. Corrective action written in HACCP plan.		0			
21. Reassessed adequacy of the HACCP plan.		0 -	Part F - II	nspection Requirements	
22. Records documenting: the written HACCP plan, monitoring critical control points, dates and times of specific event occurred.	of the currences.	0	49. Government Staffing		
Part C - Economic / Wholesomeness			50. Daily Inspection Covera	age	0
23. Labeling - Product Standards			51. Enforcement		
24. Labeling - Net Weights			52. Humane Handling		0
25. General Labeling					+
26. Fin. Prod. Standards/Boneless (Defects/AQL/Pork Skins/Mi	oisture)	0	53. Animal Identification		0
Part D - Sampling Generic <i>E. coli</i> Testing			54. Ante Mortem Inspection		0
27. Written Procedures		0	55. Post Mortem Inspection	• • • • • • • • • • • • • • • • • • •	0
28. Sample Collection/Analysis		0			
29. Records		0	Part G - Other Regu	ulatory Oversight Requirements	
Salmonella Performance Standards - Basic Requ	irements		56. European Community D	rectives	0
30. Corrective Actions		0	57. Monthly Review	·	0
31. Reassessment		0	58.		
32. Written Assurance		0	59.		

Est. # 5253, Swire Cold Storage PTY LTD, Palmyra - Western Australia, 8-11-06, International Distribution Warehouse

45 Falling snow and ice from the freezing units were observed on several boxed product. Immediate corrective action was taken by AQIS and establishment officials by removing snow and ice from effected boxes and assuring product inside boxes was not affected. As an additional corrective action, product will not be held directly under the freezing units. Product affected was not going to the United States. FSIS regulation: 9 CFR 416.3a.

61. NAME OF AUDITOR
Oto Urban. DVM

62. AUDITOR SIGNATURE AND DATE

Oto Muban 10-6-06

Ms Sally White Director International Equivalence Staff Office of International Affairs Food Safety and Inspection Service Washington, D.C. 20250

#### Dear Ms White

Thank you for your letter of 19 October 2006 accompanying the draft final report of the Food Safety and Inspection Service (FSIS) audit of Australia's meat inspection system from 10 August through 30 August 2006. The Australian Quarantine and Inspection Service (AQIS) values the constructive working relationship that exists between our two countries. We are pleased to note the comments of the US audit team at the exit meeting in relation to noted improvements in the Australian meat inspection system.

We note from section 6.1.3 of the audit report that FSIS has concerns regarding AQIS oversight of Salmonella testing. AQIS believes that it has received a favourable equivalence decision on this method of oversight and has enclosed a copy of the equivalence decision and the AQIS Notice to which that decision refers.

Section 6.1.3 also mentions pathogen tesing programs for ready to eat meat and ground meat. AQIS proposes to enhance oversight of Listeria monocytogenes and E. coli 0157:H7 testing by ensuring AQIS inspection staff directly verify *Listeria monocytogenes* and *E. coli 0157:H7* sampling during US destined production runs. This direct verification would include oversight of sampling, packaging and the submission of samples to government approved laboratories. This enhanced level of oversight is consistent with the approach taken with *Salmonella* testing. AQIS proposes to submit an application for equivalence to cover both *L. monocytogenes* and *E. coli 0157:H7* testing. In addition AQIS will ensure that all establishments registered to produce ready to eat products review their environmental control program for *Listeria spp*. AQIS will continue to verify the effectiveness of these programs.

Deficiencies noted by the FSIS auditors in individual establishments have been addressed and the corrective actions verified by AQIS. One establishment received a Notice of Intention to Delist (NOID) and responded immediately to rectify all deficiencies within 30 days. AQIS has verified the corrective actions taken and provided a report to PSIS in a previous submission. Attached for your consideration is a summary of the corrective actions and preventative measures undertaken to address deficiencies identified at individual establishments.

AQIS continues to ensure the performance of establishment based staff is acceptable in relation to regulation and enforcement activities. AQIS is reviewing its verification systems to identify areas requiring improvement particularly at those establishments where the FSIS auditors identified issues.

In summary I would like to take this opportunity to thank you and the FSIS staff involved in the audit process. AQIS is appreciative of the opportunity to respond to this audit report and takes the findings seriously. AQIS is confident that FSIS will find that enhancements to the meat inspection system that are being implemented following the audit will address concerns raised during this audit. AQIS looks forward to the opportunity to demonstrate the results of these changes in subsequent audits.

Yours sincerely

Greg Read

Executive Manager

Exports and Animal Programs Division

2 December 2006

#### Attachments

- 1. FSIS Equivalence Evaluation of Pathogen Reduction/ HACCP Requirements
- 2. AQIS Meat Notice 96/46 referred to in the equivalence decision
- 3. Summary of corrective actions taken at individual establishments