# Cattle protein/mineral medicated supplement

Bulk Ingredients	Micro, Bag, and Hand Add	Medications/Drugs
Corn Soybean Meal Sunflower Meal Calcium carbonate Salt	Vitamin E Copper Sulfate	
Liquids	Packaging Materials	
Fat	Bags & Totes Bag Label Bulk Label Delivery Truck	

# **List of Product Ingredients and Incoming Materials Form**

- Purpose of the List of Product Ingredients and Incoming Materials Form
- Information needed to complete form

Bulk Ingredients	Micro, Bag, and Hand Add	Medications/Drugs
Corn Soybean Meal Sunflower Meal Calcium carbonate Salt	Vitamin E Copper Sulfate	Rumensin 80 Rabon
Liquids	Packaging Materials	
Fat	Bags & Totes Bag Label Bulk Label Delivery Truck	

# List of Product Ingredients and Incoming Materials Form Product Name: Cattle protein/mineral medicated supplement Bulk Ingredients Micro, Bag, and Hand Add Medications/Drugs Com Soybean Meal Sunflower Meal Calcium carbonate Salt Liquids Packaging Materials Fat

#### **Summary**

- Completing the <u>List of Product Ingredients and</u> <u>Incoming Materials Form</u> is one of the preliminary steps in developing a HACCP plan
- The <u>List of Product Ingredients and Incoming</u>
   <u>Materials Form</u> will be used during the hazard analysis
- The <u>List of Product Ingredients and Incoming</u>
   <u>Materials Form</u> will be included in your HACCP plan

# **Product Description Form**

- Purpose of the Product Description Form
- Information needed to complete form

1. Product name(s)	Cattle protein/mineral supplement
2. Product safety properties (Moist., Pro., etc)	High moisture
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label
4. Type of packaging	
5. Shelf life	
6. Where will the product be sold?	
7. Labeling instructions	
8. Special distribution control	

1. Product name(s)	Cattle protein/mineral medicated supplement
2. Product safety properties (Moist., Pro., etc)	
3. How is the product to be used (intended use) and who is the intended consumer?	
4. Type of packaging	
5. Shelf life	
6. Where will the product be sold?	
7. Labeling instructions	
8. Special distribution control	

Product name(s)	Cattle protein/mineral supplement
2. Product safety properties (Moist., Pro., etc)	High moisture
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label
4. Type of packaging	Bulk & Bag
5. Shelf life	
6. Where will the product be sold?	
7. Labeling instructions	
8. Special distribution control	

1. Product name(s)	Cattle protein/mineral supplement
2. Product safety properties (Moist., Pro., etc)	High moisture
3. How is the product to be used (intended use) and who is the intended consumer?	
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6. Where will the product be sold?	
7. Labeling instructions	
8. Special distribution control	

Product De	escription Form		
1. Product name(s)	Cattle protein/mineral supplement		
2. Product safety properties (Moist., Pro., etc)	High moisture		
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label		
4. Type of packaging	Bulk & Bag		
5. Shelf life	Equal to or less than 90 days		
6. Where will the product be sold?			
7. Labeling instructions			
8. Special distribution control			
-			
1998 Approved:	Date:		

Product De	escription Form		
1. Product name(s)	Cattle protein/mineral supplement		
2. Product safety properties (Moist., Pro., etc)	High moisture		
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label		
4. Type of packaging	Bulk & Bag		
5. Shelf life	Equal to or less than 90 days		
6. Where will the product be sold?	Retail or wholesale		
7. Labeling instructions			
8. Special distribution control			
	<u> </u>		
1998 Approved:	Date:		

- Completing the "Product Description Form" is one of the preliminary steps in developing a HACCP plan
- The "Product Description Form" will be used during the hazard analysis
- The "Product Description Form" will be included in your HACCP plan

Product De	escription Form
1. Product name(s)	Cattle protein/mineral supplement
2. Product safety properties (Moist., Pro., etc)	High moisture
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label
4. Type of packaging	Bulk & Bag
5. Shelf life	Equal to or less than 90 days
6. Where will the product be sold?	Retail or wholesale
7. Labeling instructions	In compliance with federal and state regulations
8. Special distribution control	
998 Approved:	Date:

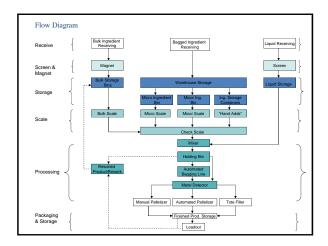
#### **Process Flow**

- Purpose of the Process Flow Diagram
  - Summarize the manufacturing process
  - Assist in hazard analysis
  - Provide immediate reference to critical control points
- Information needed to complete form
  - HACCP team knowledge of the process

1. Product name(s)	Cattle protein/mineral supplement
2. Product safety properties (Moist., Pro., etc)	High moisture
3. How is the product to be used (intended use) and who is the intended consumer?	Feed to animals per instructions on label
4. Type of packaging	Bulk & Bag
5. Shelf life	Equal to or less than 90 days
6. Where will the product be sold?	Retail or wholesale
7. Labeling instructions	In compliance with federal and state regulations
8. Special distribution control	Proper sequencing and flushing

#### **Form Completion Procedure**

- Outline the process flow using a block diagram format.
- Denote ccps on the process flow after performing principle 2, identifying critical control points.

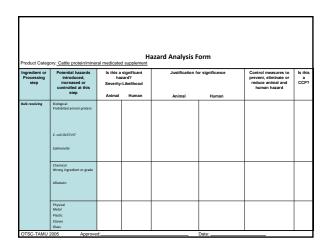


# **Form Completion Procedure**

- The completion of the Hazard Analysis Form involves hazard identification and hazard evaluation
- Each step of the process requires a separate page
- List all the process steps in order of their occurrence and then brain storm to identify hazards throughout the plant
- Perform the hazard evaluation second after the hazard identification has been completed

# Summary

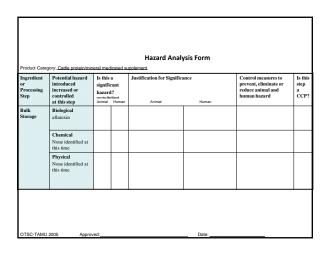
• The "Process Flow Diagram" must be completed prior to hazard analysis and should include the ccp(s) in the HACCP Plan.

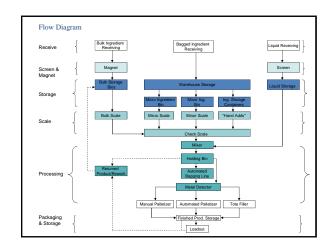


## **Completing the Hazard Analysis Form**

- Purpose of the Hazard Analysis Form
  - Provide some standardization
  - Assist in plan development
- Information needed to complete form
  - Reference material, hazard guide, expertise
  - Forms completed during preliminary steps:
    - Product Description Form,
    - List of Product Ingredients & Incoming Materials Form
    - Flow diagram

Ingredient or Processing Step Magnet	Potential hazard introduced increased or controlled at this step Biological None identified at this time	Is this a significant hazard? sweeky:Melihood Animal Human		Justification for Significance		Control measures to prevent, eliminate or reduce animal and human hazard	Is this step a CCP?
		Animai	Human	Animal	Human		
	Chemical None identified at this time						
	Physical Metal						





Product Cates	gory: Cattle protein/min	eral med	icated su	Hazard Ana	lysis Form		
Ingredient or Processing Step Bulk Scale	Potential hazard introduced increased or controlled at this step	Is this a significant hazard?		Justification for Significance  Animal Human		Control measures to prevent, eliminate or reduce animal and human hazard	Is this step a CCP?
	Biological None identified at this time						
	Chemical None identified at this time						
	Physical None identified at this time						

Ingredient or Processing step	Potential hazards introduced, increased or controlled at this step	ha: Severity	significant zard? :Likelihood	Justification fo		Control measures to prevent, eliminate or reduce animal and human hazard	Is thing
Bulk receiving	Biological Prohibited animal protein	Animal	Human	Animal  Cross contamination by prohibited animal protein (21 CFR:589:2000-1) is a potential source of bovine spongiform	Human  BSE in cattle can cause the human disease variant Creutzfeldt Jakob disease (vCID)	Prohibited animal protein policy, approved supplier, carrier inspection	
	E. coli O157:H7	No	No	encephalopathy (BSE)  Low likelihood in animal feed ingredients	Low likelihood in human food		
	Salmonella	Yes	No	Moderate likelihood in ingredients, a potential source for Salmonellosis	Low likelihood of it causing a human food problem	Approved supplier, cleaning feed manufacturing equipment	
	Chemical Wrong ingredient or grade	Yes	No	Potential source of toxin to animals	Low likelihood of transfer to human food	Approved supplier and testing	
	Affatoxin	Yes	Yes	Toxic to finishing cattle at concentrations above 300 ppb	Transfer to human food when feed to lactating dairy cattle	Sampling and testing incoming ingredients prone to aflatoxin	
	Physical Metal Plastic Stones	Yes Yes Yes Yes	No No No No	Physical hazards can damage animal mouth and digestive system	Low likelihood of transfer to food	Equipment (screens, de- stoning device, metal detectors, and magnets) in place to eliminate hazard	

				Hazard Ana	lysis Form		
Ingredient or Processing Step	Potential hazard introduced increased or controlled at this step	Is this signific hazard sverity:tto Animal	a cant 1?	Justification for Signifi	cance	Control measures to prevent, eliminate or reduce animal and human hazard	Is this step a CCP
Bag Ingredient Receiving	Biological None identified at this time						
	Chemical Mislabeled product Wrong potency of ingredient						
	Physical Metal Other foreign materials						

Product Categ	ory: Cattle protein/min	eral med	icated su	ipplement d			
Ingredient or Processing Step	Potential hazard introduced increased or controlled at this step	Is this a significant hazard? swedy-likelihood Animal Human		Justification fo	r Significance	Control measures to prevent, eliminate or reduce animal and human hazard	Is this step a CCP?
Magnet	Biological None identified at this time	74.1111111	TO THE	~~	1001001		
	Chemical None identified at this time						
	Physical Metal	Y	N	Potential source to control metal	Low likelihood as a human food safety hazard	Routine inspection of magnet	
OTSC-TAMU 2	2005 Approv	ed:			Date:		

Ingredient or Processing Step	Potential hazard introduced increased or controlled at this step	haz	icant	Justification fo	or Significance	Control measures to prevent, eliminate or reduce animal and human hazard	Is thi step a CCP
Bulk Storage	Biological Aflatoxin	N	N	Only accept corn containing 15% moisture or less, short storage time	Only accept corn containing 15% moisture or less, short storage time	Moisture measurement upon corn receipt, storage temperature monitoring and inventory control	
	Chemical None identified at this time						
	Physical None identified at this time						

- Review your Hazard Analysis Form for consistency
- Determine critical control points

Ingredient or Processing Step	Potential hazard introduced increased or controlled at this step	signit haza	ard?		for Significance	Control measures to prevent, eliminate or reduce animal and human hazard	Is this step a CCP?
Bulk Scale	Biological None identified at this time	Animal	Human	Animal	Human		
	Chemical None identified at this time						
	Physical None identified at this time						

#### **CCP Decision Tree Form**

- Purpose of the CCP Decision Tree Form
- Information needed to complete form

Ingredient or Processing Step	Potential hazard introduced increased or controlled at this step	signii hazi	his a ficant ard? nothed Human	Justification fo	er Significance	Control measures to prevent, eliminate or reduce animal and human hazard	Is thi step a CCP
Bag Ingredient Receiving	Biological None identified at this time						
	Chemical Mislabeled product Wrong potency of ingredient	Y Y	N N	Mislabeled products or wrong potency can negatively impact animal performance	Low likelihood of passing through animal into food	Approved supplier program; label inspection at receipt per Receiving Bagged Ingredients SOP; random testing for both hazards	
	Physical Metal Other foreign materials	Y	N	Physical hazards can damage animal mouth and digestive system	Low likelihood of passing through animal into food	Equipment (screens, de-stoning device, metal detectors and magnets) in place to eliminate hazard	

# **Form Completion Procedure**

- Utilize this form for each process step that contains a hazard identified as significant for humans in the "Hazard Analysis Form"
- Complete the CCP Decision Tree Form beginning in the left column
- The HACCP Team should complete the form
- If the process step is a CCP, record this result in the "Hazard Analysis Form"

				cision Tree Fo	rm		
Process / Step	Hazard	Q1a. Do  Q1a. Do  Q1a. Do  preventive measures exist for the identified hazard(s)?  If no-go to Q1b. If yes-go to Q2.	I medicated supplei Q1b. Is control at this step necessary for safety?  If no-not a CCP. If yes— modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely occurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yes-CCP	Q3. Could contamination with Identified hazard(s) occur in excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP, if yes—go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP if yes — not a CPP	CCF No.
Bulk Ing. Receiv -ing Pit							

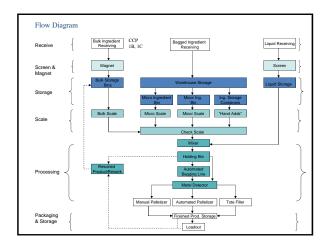
			CCP I	Decision Tree	Form		
Product Ca Process / Step	Hazard	Q1a. Do preventive measures exist for the identified hazard(s)? If nogo to Q1b. If yesgo to Q2.	Otb. Is control at this step necessary for safety?  If no-not a CCP.  If yes— modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely coccurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yes-CCP	Q3. Could contamination with Identified hazard(s) occur In excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP. If yes-go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP if yes — not a CPP	C) Ni
Bulk Ing. Receiv -ing Pit	Prohibited Animal protein	Yes		Yes			

			CCP De	cision Tree Fo	m		
Product Ca	ategory: Cattle	protein/minera	I medicated suppler	ment			
Process / Step	Hazard	Q1a. Do preventive measures exist for the identified hazard(s)?  If no-go to Q1b.  If yes-go to Q2.	Q1b. Is control at this step necessary for safety?  If no-not a CCP. If yes—modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely occurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yes-CCP	Q3. Could contamination with Identified hazard(s) occur in excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP. If yes—go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP If yes — not a CPP	CCP No.
Bulk Ing. Receiv -ing Pit	Prohibited Animal protein						

Product Ca	ategory: Cattle	e protein/minera	CCP I	Decision Tree F	orm		
Process / Step	Hazard	Q1a. Do preventive measures exist for the identified hazard(s)? If no-go to Q1b. If yes-go to Q2.	Q1b. Is control at this step necessary for safety?  If no-not a CCP.  If yes— modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely occurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yesCCP	Q3. Could contamination with Identified hazard(s) occur In excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP, If yes-go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP If yes – not a CPP	CCP No.
Bulk Ing. Receiv -ing Pit	Prohibited Animal protein	Yes		Yes			CCI 1B

			CCP Dec	ision Tree Fo	rm		
Product Ca	tegory: Cattle	protein/minera	I medicated supplen	nent			
Process / Step	Hazard	Q1a. Do preventive measures exist for the identified hazard(s)?  If nogo to Q1b.  If yesgo to Q2.	Q1b. Is control at this step necessary for safety?  If no-not a CCP, ff yes—modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely occurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yes-CCP	Q3. Could contamination with Identified hazard(s) occur in excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP. If yes—go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP if yes — not a CPP	CCF No.
Bulk Ing. Receiv -ing Pit	Prohibited Animal protein	Yes					

Product Ca	tegory: <u>Cattl</u>	e protein/minera	CCP Dec	ision Tree Fo	rm		
Process / Step	Hazard	Q1a. Do preventive measures exist for the identified hazard(s)? If nogo to Q1b. If yesgo to Q2.	Q1b. Is control at this step necessary for safety?  If no-not a CCP, modify step, process or product and return to Q1a.	Q2. Does this step eliminate or reduce the Likely occurrence of the hazard(s) to an acceptable level?  If no-go to Q3.  If yes-CCP	Q3. Could contamination with Identified hazard(s) occur In excess of acceptable Levels or could they increase to Unacceptable levels?  If no-not a CCP, If yes-go to Q4.	Q4. Will a subsequent step eliminate hazard(s) or reduce the likely occurrence to an acceptable level?  If no — CCP If yes — not a CPP	CCP No.
Bulk Ing. Receiv -ing Pit	Aflatoxin	Yes		Yes			CCP 1C

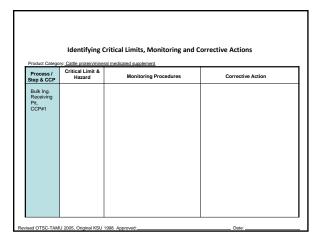


# **Form Completion Procedure**

- Complete the form by listing the process step for the first critical control point (ccp)
- Proceed to establish critical limits (column 2), monitoring procedures (column 3) and corrective action (column 4) for the first ccp
- Repeat this process for all ccps

## Summary

 Completing the "CCP Decision Tree Form" is a tool in deciding whether the process step is a critical control point



# Critical Limits, Monitoring and Corrective Actions Form

- Purpose of the "Critical Limit, Monitoring, and Corrective Actions Form"
  - For each critical control point, establish critical limits, monitoring requirements, and corrective actions necessary if there is a failure to meet a critical limit
- Information needed to complete form
  - Hazard analysis form

	Identifying Critical Limits, Monitoring and Corrective Actions							
Product Catego Process / Step & CCP	ry: Cattle protein/miners Critical Limit & Hazard	al medicated supplement  Monitoring Procedures	Corrective Action					
Bulk Ing. Receiving Pit, CCP#1	Zero Tolerance, Prohibited animal protein							

Identifying Critical Limits, Monitoring and Corrective Actions  Product Category: Cattle protein/mineral medicated supplement						
Process / Step & CCP	Critical Limit & Hazard	Monitoring Procedures	Corrective Action			
Bulk Ing. Receiving Pit. CCP#1	Zero Tolerance, Prohibited animal protein	What will be measured? Cleanout certificate for carriers Bill of Lading from supplier Product labeling Letter of Guarantee (LOG) from approved supplier Presence of prohibited animal protein Where will the CL be measured? Receiving Truck Scale or Dock How will the CL be measured? Visual observation of documentation Purchase only from approved supplier Receiving employee(s) Who will monitor the CL? Receiving employee(s) Every load received into the facility.				

#### **Record Keeping and Verification Form**

- Purpose of the Record Keeping and Verification Form
  - For each critical control point, establish verification and record keeping procedures
- Information needed to complete form
  - "Hazard Analysis Form" and "Critical Limits, Monitoring and Corrective Actions Form"

Process / Step & CCP	Critical Limit & Hazard	Monitoring Procedures	Corrective Action
Bulk Ing. Receiving Pit, CCP#1	Zero Tolerance. Prohibited animal protein	What will be measured? Cleanout certificate for carriers Bill of Lading from supplier Product labeling Letter of Guarantee (LOG) from supplier Presence of prohibited animal protein Where will the CL be measured? Receiving Truck Scale or Dock How will the CL be measured? Visual observation of documentation Purchase only from approved supplier Use of Neogen test strips Who will monitor the CL? Receiving employee(s) How often will the CL be measured? Every load received into the facility.	What caused the deviation?  No documentation or test failure Purchase from non-approved supplier How with the process be corrected? Reject Loads will be implemented to prevent recurrence?  Notify supplier that documentation must be received at delivery Training of purchasing personnel if product purchased from non-approved supplier and appropriate disciplinary action Removal of supplier from approved supplier list what will be the product disposition? Hold product until documentation is received or reject load

Record Keeping and Verification Form  Product Category: Cattle protein/mineral medicated supplement									
Process/Step CCP	Hazard	Records	Responsibility	CCP Verification					
Bulk Ing. Receiving Pit CCP #1									
OTSC-TAMU 2005,	Original KSU 1998	3 Approved:	Date	E					

# Summary

- The "Critical Limits, Monitoring and Corrective Actions Form" is utilized during plan development.
- The team should complete this form as they work through HACCP principles 3, 4, & 5.

Product Category: C	Record Keeping and Verification Form							
Process/Step CCP	Hazard	Records	Responsibility	CCP Verification				
Bulk Ing. Receiving Pit CCP #1	Prohibited animal protein							
OTSC-TAMU 2005,	Original KSU 1998	Approved:	Dar	e:				

Product Category: C	Record Keeping and Verification Form  Product Category: Cattle protein/mineral medicated supplement							
Process/Step CCP	Hazard	Records	Responsibility	CCP Verification				
Bulk Ing. Receiving Pit CCP #1	Prohibited animal protein	Receiving Bulk Ingredients SOP Cleanout certificate from carrier Bill of lading from supplier Product labeling Letter of guarantee (LOG) from supplier Receiving log Approved supplier list Record of testing (test strips) Training log (for purchasing personnel if product came from a non-approved supplier)						
OTSC-TAMU 2005,	Original KSLI 1998	Approved:	Date	p.				

- The "Record Keeping and Verification Form" is utilized during plan development.
- The team should complete this form as they work through HACCP principles 6 and 7.

Product Category: C	Record Keeping and Verification Form  Product Category: Cattle protein/mineral medicated supplement								
Process/Step CCP	Hazard	Records	Responsibility	CCP Verification					
Bulk Ing. Receiving Pit CCP #1	Prohibited animal protein	Receiving Bulk Ingredients SOP Cleanout certificate from carrier Bill of lading from supplier Product labeling Letter of guarantee (LOG) from supplier Receiving log Approved supplier list Record of testing (test strips) Training log (for purchasing personnel if product came from a non-approved supplier)	QA Supervisor Receiving Receiving Receiving Purchasing Receiving Purch. Manager Receiving QA Supervisor						
OTSC-TAMU 2005,	Original KSU 1998	Approved:	Dat	te:					

# **HACCP Plan Summary Form**

- Purpose of the HACCP Plan Summary Form
- Information needed to complete form

Product Category: C	Record Keeping and Verification Form  Product Category: Cattle protein/mineral medicated supplement								
Process/Step CCP	Hazard	Records	Responsibility	CCP Verification					
Bulk Ing. Receiving Pit CCP#1	Prohibited animal protein	Receiving Bulk Ingredients SOP Cleanout certificate from carrier Bill of lading from supplier Product labeling Letter of guarantee (LOG) from supplier Receiving log Approved supplier list Record of testing (test strips) Training log (for purchasing personnel if product came from a non-approved supplier)	QA Supervisor Receiving Receiving Receiving Purchasing Receiving Purch. Manager Receiving QA Supervisor	Short Term Daily review of receiving log and paperwork by QA/QC department  Long Term Operational audit performed by designated management personnel to make sure Receiving Bulk Ingredients SOP is followed					
OTSC-TAMU 2005,	Original KSU 1998	Approved:	Da	te:					

Product C	ategory <u>Cattl</u>	e protein/min	eral medicated	supplement_					
Process		Critical Limits for each CCP	Monitoring			Corrective Verification	Record-		
Step and CCP	Hazards		What	How	Frequen cy	Who	Action	Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1									
FPI 1999	Approve	:			Date				

Process	Critical						Record-		
Step and CCP	Hazards	Limits for each CCP	What	How	Frequen cy	Who	Action Corrective	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein								

Process		Critical		Monitorin	ıg				Record-
Step and CCP	Hazards	Limits for each CCP	What	How	Frequen cy	Who	Corrective Action	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein	Zero Toleranc e	Cleanout certificate certificate for carriers, Bill of Lading from supplier, Product labeling, Letter of Guarantee (LOG) from supplier, Presence of prohibited animal protein	Visual observation of documentation  Purchase only from approved supplier,  Use of Neogen test strips	Every load receive d into the facility	Receivin g employe e	Reject to adi in the shience of documentation, test failure, or non-approved supplier and documentation test failure, or non-approved supplier and documentation must be received at delivery must be received at delivery and proposed at delivery protection of supplier from Approved Supplier List Training of previousing personnel from non-approved supplier and appropriate disciplinary action of the proposed supplier and appropriate disciplinary action of the propriate disciplinary acti		

Process	Critical						Record-		
Step and CCP	Hazards	Limits for each CCP	What	How	Frequen cy	Who	Corrective Action	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein	Zero Toleranc e							

Product C	Category: Catt	Critical	neral medicated	f supplement Monitorin	18				Record-
Step and CCP	Hazards	Limits for each CCP	What	How	Frequen cy	Who	Corrective Action	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein	Zero Toleranc e	Cleanout certificate for carriers, Bill of Bill of Froduct labeling, Letter of Guarantee (LOG) from supplier. Product labeling, Letter of Guarantee (LOG) from supplier of the product labeling from the	Visual observation of documentation of documentation n  Purchase only from approved supplier,  Use of Neogen test strips	Every load receive d into the facility	Receivin g employe e	Reject to adi in the sheetine of documentation, test failure, or non-supported singletine of the supported singletine of supported singletine of supported singletine of supported singletine of supported singletine and appropriate disciplinary action	Daily review of receiving log and repartment by QA/QC department Operational audit performed by designated managemen t personnel to make sure Receiving Bulk Ingredients SOP is followed	
FPI 1999	Approve	d:			Date	к			

Process Step and CCP	Hazards	Critical Limits for each CCP	Monitoring						Record-
			What	How	Frequen cy	Who	Corrective Action	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein	Zero Toleranc e	Cleanout certificate certificate certificate Bill of supplier. Product labeling. Letter of Guarantee (LOG) from supplier, Presence of prohibited animal protein	Visual observation of documentation of documentation n  Purchase only from approved supplier,  Use of Neogen test strips	Every load receive d into the facility	Receivin g employe e			

Process Step and CCP	Hazards	Critical Limits for each CCP	Monitoring						Record-
			What	How	Frequen cy	Who	Corrective Action	Verification Activities	keeping procedure
Bulk Ing. Receiv- ing Pit, CCP#1	Prohibited animal protein	Zero Toleranc e	Cleanout certificate for carriers, Bill of consumplier, Product labeling, Letter of Guarantee (LOG) from supplier, Presence of prohibited animal protein	Visual observation of documentation  Purchase only from approved supplier,  Use of Neogen test strips	Every load receive d into the facility	Receivin g employe e	Reject toad in the sheece of documentation, test failure, or non-approved supplier to that documentation that documentation must be received at delivery and the supplier from Approved as University of the supplier from Approved Supplier List Training of pretraining presumed product pro	Daily review of receiving log and paperwork by QA/QC department Operational audit performed by designated management personnel to make sure Receiving Bulk lagredients SOP is followed	Receiving Bulk ingredients SOP. Cleasout certificate from carrier Bill of lading from supplier Froduct labeling Letter of Guarantee from Supplier Receiving log Receiving log Approved supplier list Recent of tenting (test salip). Training log (for purchasing personnel if product came

• The "HACCP Summary Form" must be completed and included as part of your HACCP Plan.