Dear Applicant,

A permit is required in New Mexico to operate a manufactured food facility. A manufactured food facility is a commercial operation that manufactures food products and provides those products to other business entities (wholesale sales). Manufactured Food facilities that operate in conjunction with another food facility are required to be permitted separately. For example, a person who operates a restaurant and also begins a manufacturing operation in the same facility would require a retail food permit and a manufactured food permit for the single facility.

Permit Applications must be completed as follows:

- 1. All sections of the application must be completed
- 2. All required attachments <u>must</u> be submitted as one "packet" with pages numbered and listed in the table of contents (it's preferred in a three-ring binder)
- 3. The packet **must** have a table of contents
- 4. Each section, corresponding to application section #, should be labeled and inserted in chronological order (i.e. Section 1, Section 2, Section 3)
 - a. Manufactured food Facility Application: Sections and subsections should be labeled to match the updated application. (i.e. Section 4 would be labeled as "Product Information" in the packet, Subsection 4.2.3 would be labeled "Manager and employee training" and the SOP or plan for manager and employee training would be labeled as 4.2.3.1 under that Subsection)
- 5. Clearly denote any portions of your application that you believe to be a trade secret. See "Trade Secrets" below for further details.
- 6. Submit a hard copy of the application is preferred. When ready to submit, email food.program@state.nm.us for submission instructions.

TRADE SECRETS

<u>PRIOR</u> to submitting the application, please <u>clearly denote</u> any portions of your application that you believe to be a trade secret under the meaning of Subsection 14-2-1(F) of the Inspection of Public Records Act.

"Trade Secret", as used in the Inspection of Public Records Act, means information, including a formula, pattern, compilation, program, device, method, technique or process, that:

- (1) derives independent economic value, actual or potential, from not being generally known to and not being readily ascertainable by proper means by other persons who can obtain economic value from its disclosure or use; and
 - (2) is the subject of efforts that are reasonable under the circumstances to maintain its secrecy.

If you have further questions, please email us at food.program@state.nm.us.



Application Date	•
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All fields must be completed. If a field does not apply, please indicate. Incomplete applications will not be processed.

General Information						
			Facility Information			
Name of Manu	factured Food Fac	ility:				
Street Address:	:			Phone:		
City:	Cou	nty:	Zip:	Fax:		
Mailing Addres	ss (if different than	n above):				
City:		State:	County:		Zip:	
		O	wnership Information	1		
Select on	e: Association	Corporation 🔲	Individual □ Partnership □	LLC Othe	er Legal Entity	
		d Reporting Sys	tem Identification Number (
	orporate Name:			Phone:		
Mailing Addres	S:	1		Fax:		
City:		State:		Zip:		
		Owne	ership Contact Informa	Ī	Same as "Ownership Information"	
Name and Title				Phone:		
Mailing Addres	S:			Cell:		
City:	1		T	Fax:		
State:	Zip:		Email:			
		Bill	ing Contact Informati	ı	Same as "Ownership Information"	
Name and Title	2:			Phone:		
Mailing Addres	S:			Cell:		
City:				Fax:		
State:	Zip:		Email:			
		Primary	Facility Contact Infor	mation	Same as "Ownership Information"	
Name and Title	2:			Phone:		
Mailing Addres	s:			Cell:		
City:				Fax:		
State:	Zip:		Email:			
	Addition	al Facility Co	ntact Information (attack	h additional she	eet if necessary) N/A	
Name and Title):			Phone:		
Mailing Addres	s:			Cell:		
City:				Fax:		
State:						
State.	Zip:		Email:			
State.	Zip:		Email: Regulations			



	Type of Construction (Check one)														
						Remo	del								
Facility Conversion to Manufactured Food Facility Op						penin	g or	Transfer of C	wnership of Ex	isting food	manufad	cturin	g Facility		
	Construction and Opening Details														
Plan	ned Co	nstruc	tion Start	Date:				Plan	nec	l Opening	Date:				
					Carron	E			Λ	a la catio	_				
			*If the fac	cility is in a						a Location which flo	n or each area i	s located.			
	Р	lease	indicate s	quare fo	otage in	eac	h area	`			Square Fo	otage		*Flo	or
											(ft. ²)			
			age of the												
			f the Man	•		<u> </u>	Area								
Squa	are Foc	tage o	f the Dry	Storage/v	varenous	e									
					•					peration					
		ı	f there is a							format:8ar second line	n to 8pm to insert add	itional hou	ırs.		
	Days	T	unday	Monda			day	Т		esday	Thursday	Frid		Sa	aturday
	ours		to	to	,	to)		to		to	to)	to	
Н	ours		to	to		to)	to			to to		to to		to
				Fors	seasona	l oı	perati	ons,	che	ck all th	at apply.				
Jan	□ Fe	eb 🗆	Mar 🗆	Apr □	May 🗆	Ţ	un 🗆	Jul		Aug □	Sept □	Oct 🗆	Nov	, _□	Dec □
				'							'				
Add	itional	inform	ation (if a	pplicable)	•										
			Тур	oe of Ma	nufactu	rec	d Foo	d Fac	ility	y (Check	all that ap	ply)			
	Acid I	ood							F	Refrigerat	ed Food				
			Acid Fo						_	Dry Mix F					
			w-Acid (ood					ams/Jelly	/				
			anned Fo	ood						erky					
	Seafo								١	Narehou	se				
	Shellf	ish							(Candy					
Bottled Water					F	ermente	d Food								
Bakery					1	Meat Pro	duct								
	Salsa								(Other:					
	Juice								(Other:					
	Raw F	ood							(Other:					
	Chile	Produ	ıct							Other:					
	Tortil	la								Other:					
	Froze	n Foo	d						(Other:					

		Below is a checklist of required inform	atio	n n	eeded to complete the plan review.				
	Please ensure all information is included.								
		Lack of complete information v	vill d	ela	y review and plan approval.				
checl	۲		check						
	1	1 Plans		3	Water & Sewage				
		 Floor, Mechanical, Electrical, and Site Plans 			Water supply and sewage disposal				
					Water test results				
	12	2 Equipment & Plumbing	4 Product Information						
		Floor Plan/Equipment Layout	Products manufactured						
		Equipment Specification Sheets		5	FDA and/or USDA Registration				
		Refrigeration		6	Additional Information & Other NMED Permits				
		 Handwashing sinks 			Signatures				
		 Warewashing 							
		 Plumbing Connections 			Applicant signatures				
		Water heaters & fixtures			NMED signatures				

Section 1 - Plans

1.1 FLOOR PLAN:

Submit floor plans drawn to scale that include the location and identification of all equipment including but not limited to, the items listed in Section 2.1 below. Number each item on the floor plan and provide a key identifying the equipment corresponding to each number on the floor plan.

1.2 MECHANICAL VENTILATION PLANS AND SCHEDULES (new construction, facility conversion, or ventilation change): Provide plans and schedules that indicate the location and specifications of ventilation hoods, fire suppression systems (as required by state or local fire authority), and restroom exhaust fans. Submit specification sheets for all ventilation hoods and fire suppression systems.

Provide make and model numbers and CFMs for each ventilation hood and exhaust fan in table below.

	Ventilation Information									
ID # on Plans or Location Make Model CFM										

1.3 ELECTRICAL PLANS AND SCHEDULES (new construction, facility conversion, or ventilation change):

Provide plans and schedules that indicate the locations and specifications of all lighting.

Note: All lights in processing areas, dry storage areas, dishwashing areas, inside equipment, and above areas where open products are held or displayed must be equipped with shatter proof bulbs or shields that will protect open product, utensils and equipment from broken glass if a bulb is broken.

1.4 SITE PLAN:

Provide a site plan which includes the following:

- 1) Dumpster enclosures and trash compactors
- 2) Outside walk-in coolers/freezers
- 3) Outside product storage areas
- 4) Location of well heads and well water supply lines servicing the building (if applicable).
- 5) Location of on-site liquid waste treatment systems and associated lines servicing the building (if applicable)
- 6) Grease interceptors/grease traps (if applicable)
- 7) Submit piping diagram of the disinfection system. Include size of holding tank(s), pressure tank(s), make and model number of treatment system, etc. (if applicable)



Section 2 - Equipment & Plumbing

2.1 Floor Plan/Equipment Layout

Check all that apply to your facilit	Check all that apply to your facility & add others not listed. When requested list ID #. If necessary, use another page.							
Hand Sink(s) (required in all processing area(s))	1 1 3 1							
Stoves	Dry Storage Areas	Other:						
Ovens	Chemical Dispensing Units	Other:						
Refrigerators	Laundry Facility Locations	Other:						
Freezers	Garbage/Recyclables Storage	Other:						
Ventilation Hoods	Toilet Facilities	Other:						
Utility Mop Sinks	Floor Sinks/Floor Drains	Other:						
Chemical Storage Areas	Hose Bibs/Hose Reels	Other:						
Personal Storage Areas	Grease Interceptor/Grease Trap	Other:						

2.2 EQUIPMENT SPECIFICATIONS:

Submit equipment specification sheets, including make and model numbers for all equipment listed in this section. If a specification sheet lists more than one piece of equipment, identify the specific equipment to be used. Number each specification sheet with the floor plan number assigned in 1.1 above.

2.3 REFRIGERATION:

	Refrigeration Capacities									
ID # on Plan										
or Location										
	Walk-in Cooler									
	Walk-in Freezer									
	Reach-in Cooler									
	Reach-in Freezer									
	Other:									

2.4 HANDWASHING SINKS:

Handwashing Sink (required in all processing areas)								
Do all sinks have a mixing valve or combination faucet with hot and cold running water under pressure?	YES□	NO*□						
Are enclosed paper towel dispensers and hand cleanser available at each sink?	YES□	NO*□						
*If the answer to either question above is "No", explain:								

2.5 WAREWASHING:

Manual Warewashing - Include the size of each compartment (*length x width x depth*) of the warewashing sinks, soiled and clean drain board lengths, and whether or not a pre-rinse spray hose will be installed for each warewashing area, including bars.

Note: Warewashing sinks must be large enough to accommodate the largest piece of equipment or utensils used.



	Manual Warewashing Information (required)							
ID # on Plans or Location	Length (inches) of Soiled Drain board (required)	Dimensions (inches) of Sink Compartments (L x W x D)	Length (inches) of Clean Drain board (required)	Spr	Rinse ayer s/No			
		х х		YES □	NO□			
		х х		YES □	NO□			

				х х				YES □	NO□		
				х х				YES □	NO□		
	Drain board Alternatives:										
If soiled and o	clean drainboa	ırds will not k	oe provided, ir	ndicate the method:	s that wi	ll be us	ed and provide	specificatio	n sheets :		
Mechanical warewashing				model numbers a pelow.	nd atta	ch spe	cification shee	ts for each			
			Mechanic	al Warewashing	Inform	ation			N/A□		
ID # on Plar	ns or Locatio	on									
Make	Model #	Sanitizin	g Method	Drain board Length (inches)	Pre-R	inse	Utensil Soak Sink Dim (inches)		ensions		
		Heat	Chemical		Yes	No	(L x	W x	D)		
							х	х			
							х	Х			
				Dirty Dishes							
	dirty dishes b										



Complete table below to indicate equipment that cannot be moved to be cleaned and sanitized and must be cleaned in place. If more room is necessary, attach an additional page.

	Clean-In-Place Equipment Installation List							Installation Method				
^	Clean-In-Place Equipment Installation List Note: Under "Installation Method", check all that apply. (attach additional sheet if necessary)						d		ounte Table lount	-		
ID # on Plan or Location	Equipment	Make/Model	New (N) / Used (U)	Plumbing Required Yes / No	Casters	Legs (at least 6 inches)	Sealed in Place	Portable	Legs (at least 4 inches)	Sealed in Place		

2.6 PLUMBING CONNECTIONS:

Complete table below for all related equipment and plumbing fixtures. Indicate if fixtures or equipment will be indirectly drained (e.g. floor sink or air gap), directly connected to the sewer (p-trap), and/or what method of backflow prevention will be used (if applicable).

ID # on Plan or Location	Fixture or Equipment		Indirect/Direct Drainage	Method of Backflow Prevention
	Warewashing Sinks	N/A□		
	Warewashing Machines	N/A□		
	Garbage Disposals	N/A□		
	Hand Sinks	N/A□		
	Chemical Dispensing Units	N/A□		
	Walk-in Refrigeration /	N/A□		
	Mop / Utility Sink	N/A□		
	Other:			
	Other:			

Note: Approved backflow protection is intended to protect the water supply. A vacuum breaker on water inlet lines for dishwashing machines, garbage disposals, or hose bibs is an example. Indirect drainage is intended to protect fixtures from sewage backup. An air gap at warewashing is an example.

2.7 WATER HEATER(S)

Provide type and capacity of all water heaters. **Provide specification sheet(s).**

Water Heater				
Туре	Capacity			
(Ex: Standard, Quick Recovery, Tankless)				



Booster Heater:		N/A□
Is a separate booster heater provided?	YES□	NO□

2.8 FIXURES REQUIRING HOT WATER

Provide the number of plumbing fixtures requiring hot water in the table below. This information will be used to determine the hot water demand for the facility and sizing criteria for the water heater.

Plumbing Fixtures Requiring Hot Water	# of Fixtures Throughout Facility	Plumbing Fixtures Requiring Hot Water	# of Fixtures Throughout Facility
3-compartment sinks		Garbage can washer	
Warewashing machines		Showers	
Pre-rinse sprayers		Hose bibs used for cleaning	
Utensil soak sinks		Other:	
Hand sinks include restrooms		Other:	
Mop sinks/Utility sinks		Other:	

Section 3 - Water & Sewage

3.1 WATER SOURCE, AVAILABILITY, & SAMPLING

3.1 WATER SOURCE, AVAILABILITY, & SAWF LING							
	Water Availability:						
	I acknowledge and understand that running water is required at <u>ALL</u> times and agree to discontinue all product-						
relate	ed activities, if wat	er is not av	ailable, un	til water s	service is restored,	, or an alternative plan is app	roved by NMED.
Signa	ature:						
Water Supply: (Select the type of water supply system that services the facility)							
□Pub	olic Water System - I	Name of mu	nicipality:				
☐ Priv	/ate (sampling req u	ired as outl	ined below	– if possib	ole, initial samples	should be submitted with appli	cation):
	Submit a copy of	the most re	cent water	sample te	st results that meet	t the drinking Water quality star	ıdards of a non-
	community wate	r system as	specified in	20.7.10 N	MAC.		
	Туре		Fre	quency		Limit	
	Total Coliform		Init	tial and M	onthly	Absent	
	Nitrate		Init	tial and Ar	nnual	10 ppm	
	Nitrite		Init	tial		<1.0 ppm	
	A list of certified labs can be located at: https://www.env.nm.gov/dwb/sampling/CertifiedLabs.htm						
Private Drinking Water Supply Information N/A□							
Well Depth (feet):			Setback to liquid	d waste drain field (feet):			
Disinfection YES NO		NO□	Туре:				

If yes, is a backflow device installed?

 $NO\square$

If a water treatment device is installed, how will the device be inspected and serviced? Attach separate page, if additional space is required.

YES□

NO

YES□

Is there a water treatment device?



3.2 SEWAGE DISPOSAL

Sewage Disposal:				
Select the type of sewage disposal system that services the facility				
□ Public - Name of municipality:				
☐ On-site liquid waste system – Permit number:				

Section 4 – Product Information

4.1 PRODUCT(S):

Provide a list of all products manufactured

Products Manufactured (list all products)			
Attach separate page, if additional space is required.			

4.2 OPERATIONAL PLAN(S):

Provide the	Provide the following information for all products manufactured.					
	Operational Plan Checklist					
Genera	General Information (one attachment needed for all products) REQUIRED					
4.2.1	Planned source of ingredients used in production (20.10.2.11.F)					
	4.2.1.1 Attach Standard Operating Procedure(s) OR describe in detail how you determine where ingredients will be purchased.					
	·					
4.2.2	Finished product testing 4.2.2.1 Attach Standard Operating Procedure(s) (SSOPs) OR testing plan for all products, including the product name, testing performed and frequency of testing. If product testing is not planned, list N/A.					
4.2.3	Manager and employee training					
	4.2.3.1 Attach Standard Operating Procedure(s) OR describe the manager and employee training plan and record					
	keeping system to track training.					
	4.2.3.2 Attach training log or record keeping system utilized to track training.					
4.2.4	Employee Health & Hygiene					
	4.2.4.1 Attach Standard Operating Procedure(s) OR Employee Illness Policy, describing:					
	 How permit holder will require employees report illness information to the person in charge. 					
	 How employees will report illnesses to the permit holder/person in charge. 					
	Specific illnesses and symptoms covered by the policy.					

- How to determine when employees will be excluded or restriction in work duties due to illness or infected cuts or lesions (See chapter 2, section 201 of the NMED Retail and Manufactured Food Field Guide for
- How to determine when employee exclusion or restriction will be removed.
- 4.2.4.2 Attach Standard Operating Procedure(s) OR describe how will bare hand contact with ready-to-eat products will be avoided during production and packing (i.e. gloves, utensils, dispensing equipment, others). **Helpful Resources**

The FDA Employee Health and Personal Hygiene Handbook is a great employee hygiene and illness resource to utilize.



4.2.5 | Standard Sanitation Operating Procedures (SSOPs)

4.2.5.1 Attach SSOPs that addresses sanitation conditions and practices before, during, and after processing. SSOPs to address, at a minimum, the following should be included:

- Practices
 - Safety of the water.
 - Monitoring backflow prevention devices.
 - Water sampling and limits (if private source).
 - Condition and cleanliness of product-contact surfaces, including equipment, work surfaces, utensils, gloves, and outer garments.
 - Prevention of cross contamination from insanitary objects, including chemicals and personal items, to product, product packaging material, and other product-contact surfaces, including equipment, work surfaces, utensils, gloves, and outer garments, and from raw product to processed product.
 - Prevention of allergen cross contact.
 - o Maintenance of hand washing, hand sanitizing, and toilet facilities.
 - Prevention of adulteration of product, product-packaging material, and product-contact surfaces from adulteration with lubricants, fuel, pesticides, cleaning compounds, sanitizing agents, condensate, and other chemical, physical, and biological contaminants.
 - o Proper labeling, storage, and use of toxic compounds. Include:
 - Type and concentration of sanitizer used for warewashing (i.e. chlorine, 100ppm).
 - Type and concentration of sanitizer used for food contact surfaces, such as tables/counters (i.e. chlorine, 100ppm).
 - Control of Employee health conditions that could result in the microbiological contamination of food products.
 - Exclusion of pests from the manufactured food facility.
- Monitoring Describe how the manufactured food facility will monitor the conditions and practices
 during manufacturing with sufficient frequency to ensure, at a minimum, conformance with those
 conditions and practices specified in the SSOPs are being met.
- Records Describe how the facility shall maintain SSOPs records that, at a minimum, document the monitoring and corrections of practices.

4.2.6 Pest Control Plan:

4.2.6.1 Attach proposed pest control plan.

4.2.7 Production Monitoring Equipment List

4.2.7.1 Attach a list and specification sheets for proposed equipment to measure and monitor product safety factors related to the production of food products. Examples include food safety thermometer, water activity meter, and pH meter.

4.2.8 Recall Plan

4.2.8.1 Attach a description of the firm's written product recall procedure, including:

- Plans for identifying products which may be adulterated or misbranded.
- Procedures for collecting, sampling, alerting consumers and businesses, warehousing, controlling, reworking, and/or disposal of recalled products.
- System for determining the effectiveness of recalls.
- Persons and governmental agencies to contact when implementing a recall, including the NMED.



Product Information (one attachment per product*) REQUIRED

*Product Information is required for each product that will be manufactured. Products or types of production methods may be grouped together, if the Product Hazard, Critical Control Points, Critical Limits, and procedures required to manufacture the products are essentially identical. The grouping of operational plans together must be approved by NMED.

• An example is beef jerky. If you produce multiple flavors of beef jerky using the same beef and production process, but only vary the dry flavorings added during the process (salt, chile, pepper), a single Operational Plan can be provided with all of the products listed (i.e. salt, red chile, lemon pepper) on the first page.

Prepare Product Information as a separate attachment and in the order outlined in the checklist below. This will make the review process more efficient.

4.2.9 <u>Product Information and Production</u>

- 4.2.9.1 Name of food product(s).
- 4.2.9.2 Names of the ingredient(s) listed in order by weight (largest quantity first).
- 4.2.9.3 Final product pH. (if applicable)
- 4.2.9.4 Final product water activity (a_w). (if applicable)
- 4.2.9.5 Names of any preservatives. (if none, write none)
- 4.2.9.6 Complete operational procedure for producing the product beginning with receiving incoming ingredients and continuing to final product distribution. <u>Include a flow chart.</u> <u>Identify critical control points on the operational procedure or flow chart.</u>
- 4.2.9.7 Type of packaging to be used and whether the packaging is integral to product.
- stability. Attach specification sheet for packaging.
- 4.2.9.8 Proposed product label(s) that comply with title 21, part 101 or title 9, 7.6.2.11.C NMAC and the New Mexico Food Act. Attached proposed label(s).

The FDA Food Labeling Guide is a great resource to assist with labeling requirements of 21 CFR 101.

- 4.2.9.9 Description of the batch / lot ID coding system, identifying the date and place of manufacture of each product and how/where it'll be placed on the package to be clearly visible on the product label or securely affixed to the body of the container.
- 4.2.9.10 Proposed shelf life. Provide supporting documentation to support proposal.
- 4.2.9.11 Product state during transportation (i.e. ambient temp., refrigerated, frozen).
- 4.2.9.12 Product care, including:
 - Condition of product (i.e. ready-to-eat, raw & must be cooked).
 - Product preparation steps required by the consumer.
 - Mishandling that may occur during storage, shipping, and in the hands of consumers.
 - Steps taken to address mishandling that may occur.
- 4.2.9.13 Intended distribution of product. List all that apply.

4.2.10 Proposed record keeping system to assure traceability of products from receiving to distribution

4.2.10.1 Attach Standard Operating Procedure(s) OR plan to describe what records will be kept, how they will be maintained, and how long they will be maintained.

4.2.10.2 Attach logs/records used to maintain traceability of all products.

4.2.10.3 Attach logs/records to monitor/document achievement of critical limits of critical control points. Examples of logs/records include, but are not limited to:

- Receiving: May contain the following information (note: terminology may vary): date received, product received, supplier, lot #, amount received, initial or signature of receiver.
- Storage: May include refrigeration temperature logs.
- Production: To monitor production requirements, including critical limits of critical control points.
- Analytical Lab Testing (if applicable): to verify compliance with testing SOPs or testing plan.
- Shipping: To maintain traceability in the event of a recall and to document critical limits of critical control points are met (if applicable during transportation).

4.2.11 | 4.2.11.1 HACCP Plan (if applicable)

- List all Food Hazards that are reasonably likely to occur and must be controlled for each product type
- List the Critical Control Points for each of the identified Food Hazards that is reasonably likely to occur, including as appropriate
- List the Critical Limits that shall be met at each of the Critical Control Points.
- List the procedures, and the frequency with which they are to be performed, that will be used to monitor each of the Critical Control Points to ensure compliance with the Critical Limits.
- Include any Corrective Action plans that have been developed and will be followed in response to deviations from critical limits at Critical Control Points.
- List the Validation and Verification procedures, and the frequency with which they are to be performed.
- Describe the recordkeeping system to document the monitoring of the Critical Control Points.
- Any additional scientific data or information supporting the determination that food safety is not compromised by the proposal.

Additional Requirements (if applicable)

4.2.12 4.2.11.1 Beef Jerky

 Documentation confirming a final water activity demonstrating that <u>each</u> final product is a non-TCS food in accordance with Table A or B under the definition of "Time/temperature control for safety food" in 7.6.2 NMAC.



Section 5 - FDA and/or USDA Registration

FDA or USDA Registration:				
Did you register with FDA or USDA?				
The FDA Food Safety Modernization Act (FSMA), enacted on January 4, 2011, amended section 415 of the Federal Food, Drug, and Cosmetic Act (FD&C Act), in relevant part, to require that facilities engaged in manufacturing, processing, packing, or holding food for consumption in the United States submit additional registration information to FDA, including an assurance that FDA will be permitted to inspect the facility at the times and in the manner permitted by the FD&C Act. Section 415 of the FD&C Act, as amended by FSMA, also requires food facilities required to register with FDA to renew such registrations every other year, and provides FDA with authority to suspend the registration of a food facility in certain circumstances.	YES	NO□		
Low-acid canned Foods and Acidified Foods Processors A commercial processor, when first engaging in the manufacture, processing, or packing of acidified foods (AF) or low-acid canned Foods (LACF) shall register and file with FDA. Registration and processing information forms are obtainable on request from: Food and Drug Administration, LACF Registration Coordinator (HFF-233), 200-C Street, SW, Washington, D.C. 20204	YES□	NO□		
Meat and Poultry Processors Meat, poultry products, or Siluriformes (i.e. catfish) inspected by USDA-FSIS or exempted.	YES□	NO□		
<u> Section 6 – Additional Information</u>				
Additional Information				
If you believe additional information would be helpful to clarify the process, please attach it in the application packet submitted. Please direct further questions to the NMED Food Program email				

Additional Information If you believe additional information would be helpful to clarify the process, please attach it in the application packet submitted. Please direct further questions to the NMED Food Program email food.program@state.nm.us. Other NMED Permits Held by Owner of this Facility Name of Facility Permit



<u>Section 7 – Signatures</u>

Applicant's Signature Page					
Comments:	s signature rage				
Comments.					
STATEMENT: I hereby certify that the above information is correct, and I fully understand that any deviation from the above without prior permission from the State of New Mexico Environment Department may nullify final approval. I agree to comply with 7.6.2 NMAC – Food Service and Food Processing Regulations and allow the regulatory authority access to the facility and records. I also certify that I have clearly denoted any portions of the application that I deem to be trade secret under the meaning of Subsection 14-2-1(F) of the Inspection of Public Records Act.					
Applicant or responsible representative(s) Signature / Title	Date				
	_				
Applicant or responsible representative(s) Signature / Title Approval of these plans and specifications by the State of New Mexico	Date	a not indicate compliance with any other			
code, law or regulation that may be requiredfederal, state, or local. It is facility (structure or equipment). A pre-opening inspection of the facility if it complies with 7.6.2 NMAC – Food Service and Food Processin approval to open is given, the fee remittance will be required and acce	y with equipment in place & o g Regulations . After the pre-	perational will be necessary to determine			
NME	O Use Only				
Signature:	Date:				
Approved □	Denied □				
Final reviewer's comments:					
ignature/Title: Date:					
Approved □	Denied □				
Office		Facility			
District:	Owner #:				
Field Office:	Permit #:				
Assigned Inspector:	Туре:				
Review Date:	Date Opened:	Date Closed:			