DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration Center for Veterinary Medicine

Expiration Date: November 30, 2024

See PRA Statement on page 3.

Form Approved: OMB No. 0910-0751

Qualified Facility Attestation for Animal Food Facility

If entering by hand, use blue or black ink only.

Section 1 – FACILITY INFORMATION					
Facility Registration Number					
Facilità Mana					
Facility Name					
Facility Address					
Address 1 (Street address, P.O. box, etc.)					
Address 2 (If applicable; apartment, suite, unit, building, floor, etc	:)				
rtaarese z (ir appricasie, apartirein, eane, arm, sanarig, neel, etc	•,				
City	State/Province	ate/Province/Territory			
		7ID as Darkel Oada			
Country		ZIP or Postal Code			
Telephone Number (Include area code)	FAX Nun	nber (Include area code)			
E-mail Address					
Section 2 – TYPE	OF NOTIFICA	ATION			
 a. Initial Submission (21 CFR 507.7(c)(2)(i)) – Complete Sec b. Biennial (Renewal) Submission (21 CFR 507.7(c)(2)(ii)) – 		·			
c. Status Change (21 CFR 507.7(c)(3)) – Complete Section	-				
Section 3 – QUALIFICATION FOR MODIFIED REQ	UIREMENTS	(Fill out only if Section 6 does not apply.)			
Animal food facilities may be exempt from the preventive controls regulations in 21 CFR part 507, primarily in subparts C and E, with associated requirements in subparts A, D, and F, under 21 CFR 507.5(d). Check the appropriate box to indicate the reason why your facility is a qualified facility.					
When including the sales by any subsidiary; affiliate; or subsidiar subsidiary or affiliate:	ries or affiliate	s, collectively, of any entity of which the facility is a			
The above-named facility qualifies for the exemption a during the preceding three calendar years, the busines \$2,500,000, adjusted for inflation, per year, in sales of manufactured, processed, packed, or held without sale	ss (including a animal food p	ny subsidiaries and affiliates) averaged less than lus the market value of animal food			
☐ The above-named facility qualifies for the exemption a	☐ The above-named facility qualifies for the exemption as a "qualified facility" as defined in 21 CFR 507.3 because;				
(1) during the preceding three calendar years, the average annual monetary value of the food manufactured, processed, packed, or held at the facility that was sold directly to qualified end-users (as defined in 21 CFR 507.3) exceeded the average annual monetary value of the food sold by the facility to all other purchasers; and					
(2) the average annual monetary value of all food sold during the preceding three calendar years was less than \$500,000, adjusted for inflation.					

Section 4 – COMPLIANCE WITH 21 CFR 507.7 (Fill out only if Section 6 does not apply.)					
Check the box to indicate how your facility is in compliance with 21 CFR 507.7(a)(2).					
I, as the owner, operator, or agent in charge of the above-named facility, (1) have identified the potential hazards associated with the animal food being produced, (2) am implementing preventive controls to address the hazards, and (3) am monitoring the performance of the preventive controls to ensure that such controls are effective. (21 CFR 507.7(a)(2)(i).) I understand that I am required to maintain records to support this attestation, but I am not required to submit those records with this attestation. (21 CFR 507.7(f).)					
The above-named facility is in compliance with State, local, county, tribal, or other applicable non-Federal food safety law including relevant laws and regulations of foreign countries. This is based on my knowledge, as the owner, operator, or agent in charge of the above-named facility, of the facility's licenses, inspection reports, certificates, permits, credentials, certification by an appropriate agency (such as a State department of agriculture), or other evidence of oversight. (21 CFR 507.7(a)(2)(ii).) I understand that I am required to maintain records to support this attestation, but I am not required to submit those records with this attestation. (21 CFR 507.7(f).)					
Section 5 – ATTESTATION STATEMENT (Fill out only if Section 6 does not apply.)					
I attest that, to the best of my knowledge and belief, the inform accurate and complete and that the above-named facility qualicowner, operator, or agent in charge of the above-named facility these attestations (21 CFR 507.7(f)) and make those records proceed by Secretary of Health and Human Services for official review and I also understand that under 18 U.S.C. 1001, anyone who know fraudulent statement to the U.S. Government is subject to crim	lifies for the exemption requested. I understand that, as the ty, I must maintain those records relied upon to support promptly available to a duly authorized representative of the d copying upon oral or written request (21 CFR 507.200(c)). Dewingly and willfully makes a materially false, fictitious, or				
Signature	Date				
Printed Name and Title:					
Please check one option below that best describes your relationship	to the facility.				
Owner Operator Agent in 0	Charge				
Please provide your contact information below if it differs from the fac	cility information provided in Section 1.				
Contact Address					
Address 1 (Street address, P.O. box, etc.)					
Address 2 (Apartment, suite, unit, building, floor, etc.)					
City	State/Province/Territory				
Country	ZIP or Postal Code				
Telephone Number (Include area code)	FAX Number (Include area code)				
E-mail Address					

Section 6 – STATUS CHANGE (If applicable)					
Animal food facilities that have changed status from a "qualified facility" to "not a qualified facility" must notify FDA of that change in status by July 31 of the applicable calendar year. Check the box below to indicate a status change.					
The above-named facility is no longer a "qualified facility" as defined in 21 CFR 507.3 based on the annual determination.					
Signature			Date		
Printed Name and Title:					
Please check one option below that best describes your relationship to the facility.					
Owner Operator Agent in Charge					
Please provide your contact information below if it differs from the facility information provided in Section 1.					
Contact Address					
Address 1 (Street address, P.O. box, etc.)					
Address 2 (Apartment, suite, unit, building, floor, etc.)					
City	State/Province/Territory				
Country		ZIP or Postal Code			
Country		ZIP OI POSIAI COGE	;		
Telephone Number (Include area code) FAX		AX Number <i>(Include area code)</i>			
E-mail Address					
If section 6 applied to you, refer to the FDA return address noted beneath Section 6.					
Return your completed Form FDA 3942b to the following FDA address:					
U.S. Food and Drug Administration					
(HFS-681)					
5001 Campus Drive					
College Park, MD 20740					

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

The burden time for this collection of information is estimated to average 30 minutes per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services Food and Drug Administration Office of Operations Paperwork Reduction Act (PRA) Staff PRAStaff@fda.hhs.gov

"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."