U.S. Food and Drug Administration Export Listing Module (ELM) Screenshots for OMB Approval https://www.access.fda.gov/

Table of Contents

Figure 1: Identification of Food Facility	
Figure 1: Identification of Food Facility	4
Figure 3: FEI selection	5
Figure 4: DUNS selection	
Figure 5: Select Product Type and Country/Region	
Figure 6: Business Information	8
Figure 7: Business Information (Facility Information and Inspection Details)	9
Figure 8: Additional Documents (Optional)	
Figure 9: Contact Information	11
Figure 10: Product Information	12
Figure 11: Product List	
Figure 12: Review screen (1 of 3)	14
Figure 13: Review screen (2 of 3)	
Figure 14: Review screen (3 of 3)	16
Figure 15: Signature Page	17

Figure 1: Identification of Food Facility

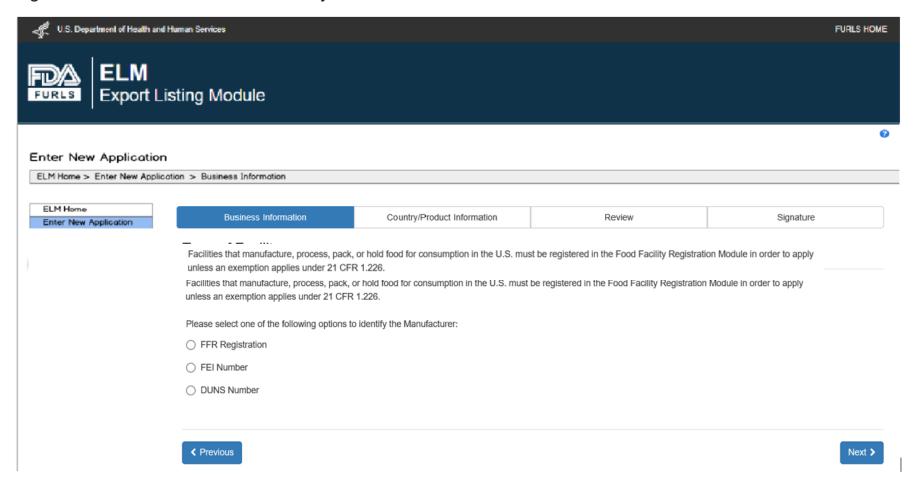


Figure 1: FFR selection

Signature **Business Information** Country/Product Information Review Type of Facility Facilities that manufacture, process, pack, or hold food for consumption in the U.S. must be registered in the Food Facility Registration Module in order to apply unless an exemption applies under 21 CFR 1.226. Please select one of the following options to identify the Manufacturer: FFR Registration FEI Number DUNS Number Food Facility Registration Information Please enter the Food Facility Registration Number and PIN for the manufacturing facility. If you represent the manufacturing facility, you may access information for your registered food facilities by logging into the Food Facility Registration Module from the FURLS home page. If you do not represent the manufacturing facility, you may wish to contact the facility to request this information. If you have any questions, you may contact the FDA Industry Systems Help Desk. FFR PIN Number FFR Registration Number

≺ Previous

Next >

Figure 2: FEI selection

Business Information	Country/Product Information	Review	Signature
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Type of Facility

O DUNS Number

Facilities that manufacture, process, pack, or hold food for consumption in the U.S. must be registered in the Food Facility Registration Module in order to apply unless an exemption applies under 21 CFR 1.226.

Please select one of the following options to identify the Manufacturer:
○ FFR Registration
FEI Number

FEI Number		

FEI Number	
FEI Number	



Next >

Figure 3: DUNS selection

Business Information	Country/Product Information	Review	Signature
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Type of Facility

Facilities that manufacture, process, pack, or hold food for consumption in the U.S. must be registered in the Food Facility Registration Module in order to apply unless an exemption applies under 21 CFR 1.226.

unless an exemption applies under 21 CFR 1.226.
Please select one of the following options to identify the Manufacturer:
○ FFR Registration
○ FEI Number
OUNS Number

DUNS Number	
DUNS Number	



Next :

Figure 4: Select Product Type and Country/Region

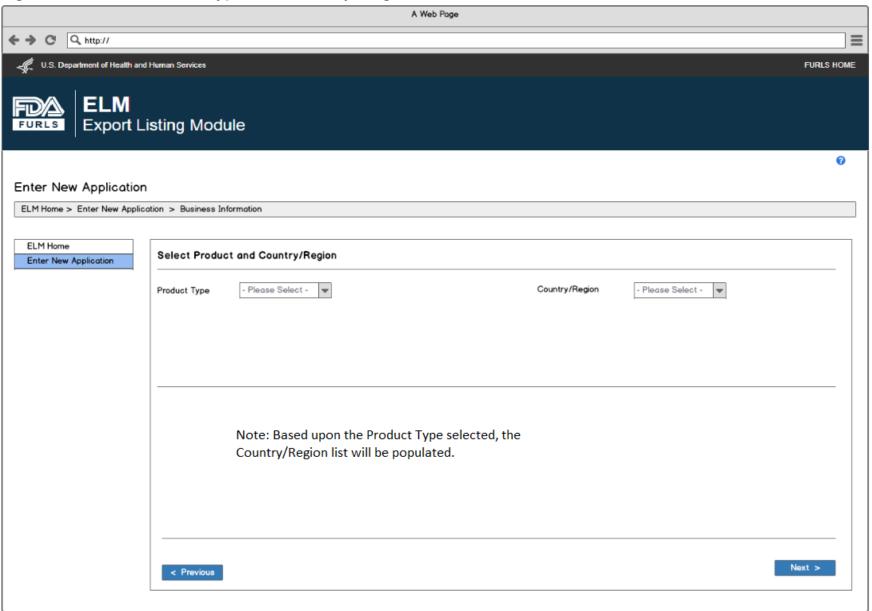


Figure 5: Business Information

Business Information	Country/Product Information	Review	Signature
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Business Information

Parent Company Information and Manufacturer Address are prepopulated from the Food Facility Registration Module. If you wish to update this information, you may log in to the Food Facility Registration Module from the FURLS home page.

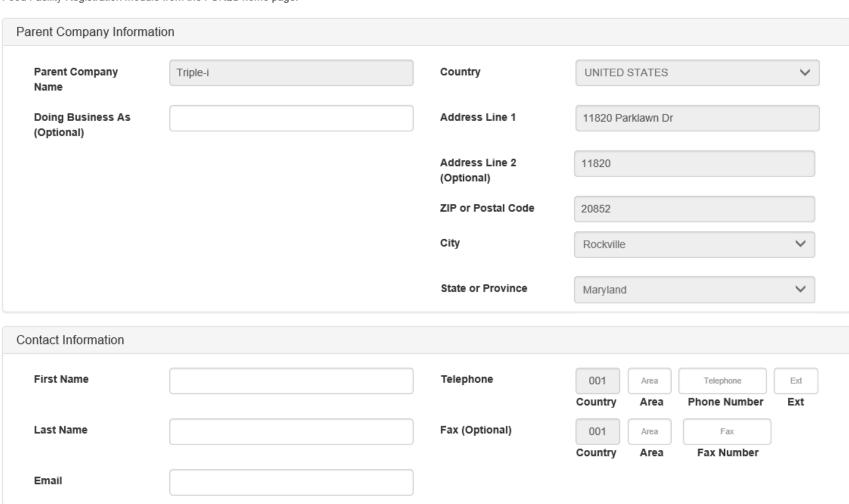


Figure 6: Business Information (Facility Information and Inspection Details)

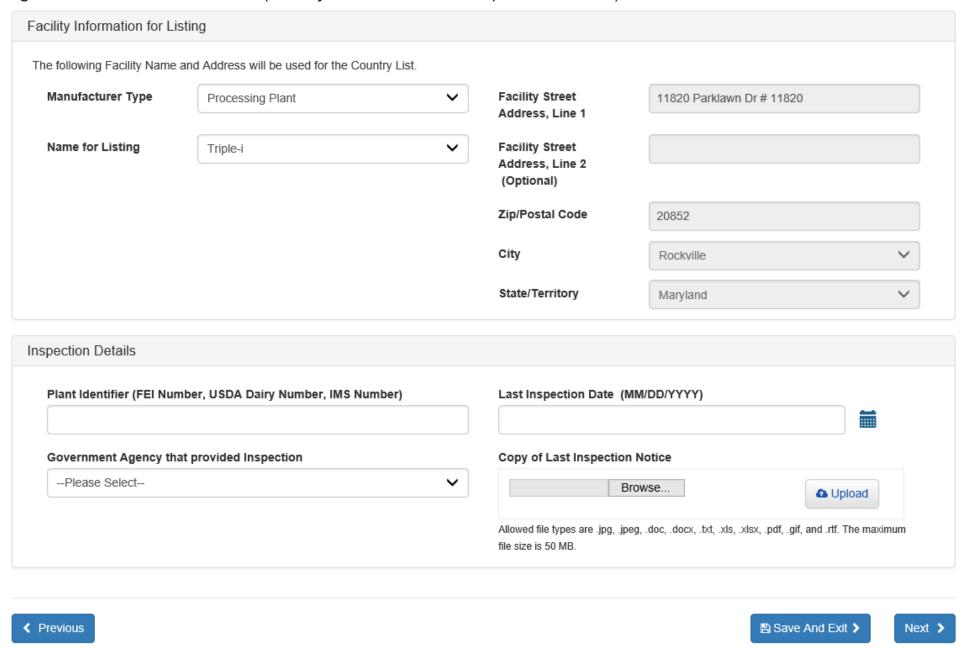


Figure 7: Additional Documents (Optional)

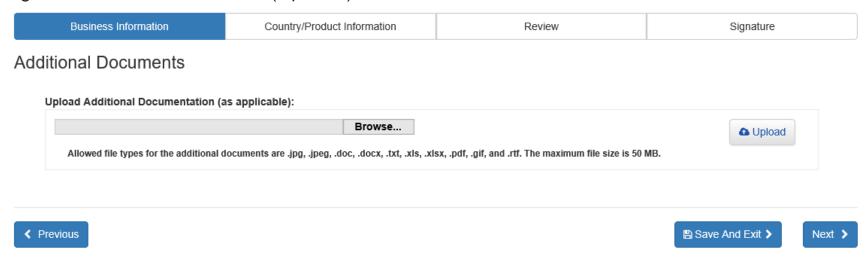


Figure 8: Contact Information

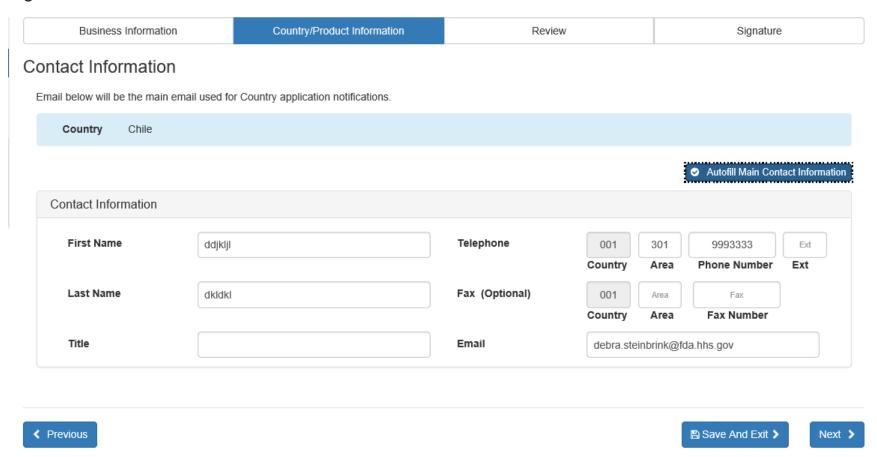
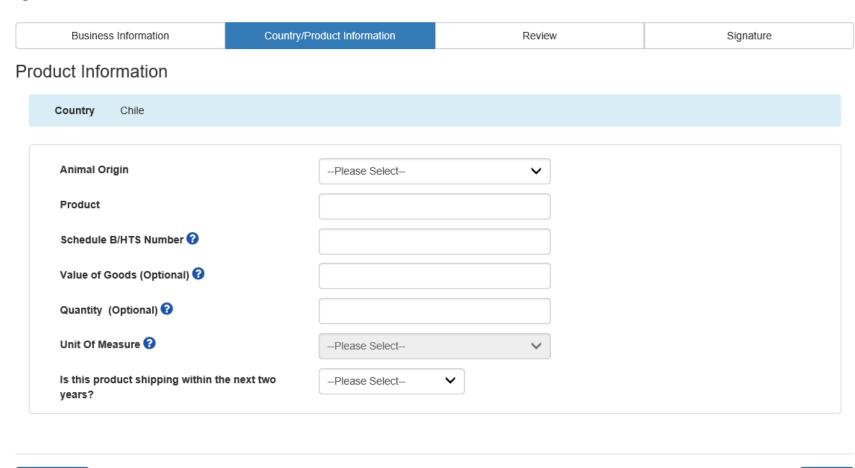


Figure 9: Product Information

Previous



> Next

Figure 10: Product List

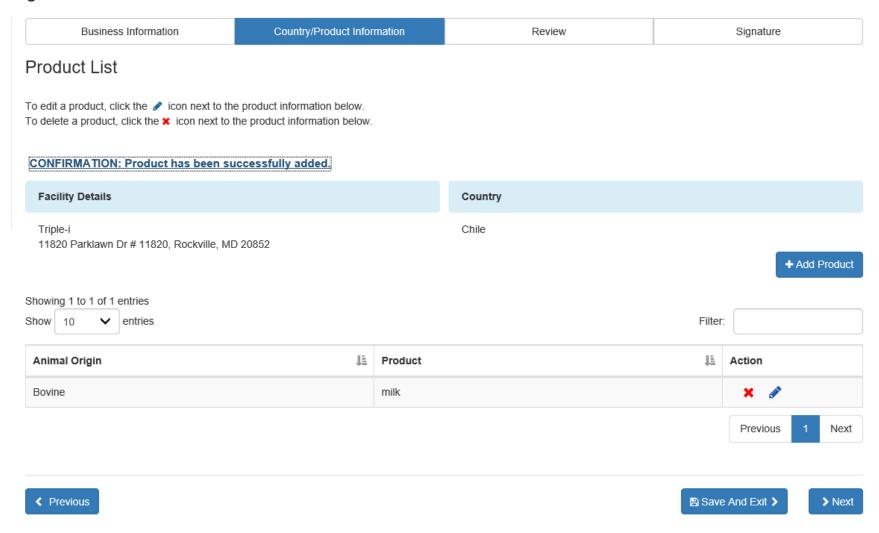


Figure 11: Review screen (1 of 3)

Business Information	Country/Product Information	Review	Signature	

Please review the information entered for the Dairy Facility. Please select the "Edit" buttons next to each section to update. Click "Next" to submit the application.



Figure 12: Review screen (2 of 3)

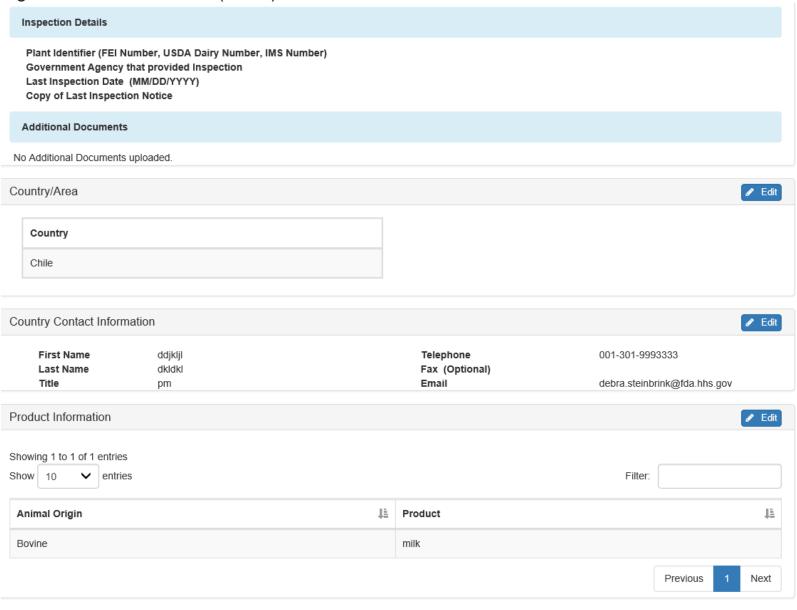


Figure 13: Review screen (3 of 3)

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Next >

Figure 14: Signature Page

Signature Page

As the responsible official or designee of the establishment listed on this application, I hereby certify to the United States Food and Drug Administration that the establishment and the products listed on this application are, to the best of my knowledge, in compliance with all applicable requirements of the Federal Food, Drug, and Cosmetic Act (the FD&C Act), and all applicable regulations and standards, including the following:

- 1. The establishment listed on this application is currently registered with the FDA, if required by the FD&C Act.
- 2. The products listed on this application for export are legally marketed within the United States or meet the requirements of Section 801(e)(1) of the FD&C Act.
- 3. The products listed on this application are not the subject of a pending judicial enforcement action (e.g., an injunction or seizure) or a pending administrative action (e.g., warning letter).
- 4. The products listed on this application are being exported from the United States.

The undersigned certifies that the information in this submission is complete and accurate. The undersigned understands that the information submitted is intended to assist FDA in establishing and maintaining a list of exporters. FDA considers the information on this list, which is provided voluntarily with the understanding that it will be communicated to the competent authority and posted on the Internet, to be information that is not protected from disclosure under 5 U.S.C § 552(b)(4).

I agree							
On Behalf Of (Optional)					Address, Line 1		
First Name					Address, Line 2 (Optional)		
Last Name					Zip/Postal Code		
Title					City	Please Select	~
Telephone	001 Country	Area Area	Telephone Phone Number	Ext Ext	State/Territory	Please Select	~
Email					Country/Area	United States	
Making or submitting false star penalties including up to \$250		-		-	itute violations of the United State	es Code Title 18, Chapter 47, Section 1001 with	
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