

Table 1A.--Prior Notice Information Required by Category

Information	Transshipment, Storage and Export, Manipulation and Export	Carried by or Accompanies an Individual	Food Not in Natural State	Food in Natural State	Mail	After Section 801(m) Refusal
§1.281 paragraph(s)	(a) and (c)	(a) and (c)	(a), (b), and (c)	(a), (b), and (c)	(b)	(c)
Submitter	Y	Y	Y	Y	Y	Y
Transmitter	Y	Y	Y	Y	Y	Y
Entry Type	Y	Y	Y	Y	Y	Y
Entry Identifier	Y	Y	Y	Y	Y	Y
FDA Product Code	Y	Y	Y	Y	Y	Y
Common, usual, or market name	Y	Y	Y	Y	Y	Y
Estimated Quantity	Y	Y	Y	Y	Y	Y ²
Lot/Code #	Y	Y	Y	N	Y	Y
Manufacturer	Y	Y	Y	N	Y/N	Y
Manufacturer Registration # ¹	N	Y	Y/N	N	Y/N	Y
Grower, if known	Y	Y	N	Y	Y	Y
Cty of Production	Y	Y	Y	Y	Y	Y
Shipper	Y	Y	Y	Y	Y	Y
Shipper Registration # ¹	N	Y	Y	Y	Y	Y
Country Shipped	Y	Y	Y	Y	Y	Y
Port of Arrival	Y	Y	Y	Y	N	Y ²
Date of Arrival	Y	Y	Y	Y	N	N
Time of Arrival	Y	Y	Y	Y	N	N
Date of Shipment	N	N	N	N	Y	N
Importer	N	Y	Y	Y	N	Y
Owner	N	Y	Y	Y	N	Y
Ultimate Consignee	N	Y	Y	Y	N	Y
U.S. Recipient	N	N	N	N	Y	Y
Mode of Transport	Y	N	Y	Y	N	Y
Carrier	Y	Y	Y	Y	N	Y
Airbill or Bill(s) of Lading	Y	N	Y	Y	N	Y ²
Vessel/Voyage	Y	Y	Y	Y	N	Y ²
Flight #	Y	Y	Y	Y	N	Y ²
Trip #	Y	Y	Y	Y	N	Y ²
Container #	Y	N	Y	Y	N	Y ²
Car #	Y	N	Y	Y	N	Y ²
License Plate #	Y	Y	Y	Y	N	Y ²
HTS code	Y	Y	Y	Y	N	Y
Hold Location	N	N	N	N	N	Y

¹ Registration numbers are required only if the firm is required to register for a facility associated with the article of food under section 415 of the FD&C Act, 21 U.S.C. 350d and 21 CFR part 1, subpart H; if registration number is provided, city and country can be provided instead of the full address.

² After arrival, therefore, no longer anticipated or planned.

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III. Comments on the Proposed Rule

FDA received approximately 470 timely responses containing one or more comments in response to the proposed rule. To make it easier to identify comments and responses to the comments, the word "Comments" will appear before the description of the comment, and the word "Response"

will appear before our response. A summary follows which includes a description of the appropriate section in the interim final rule.

A. General Comments and Outreach

(Comments) Some comments suggest revision of section 307 of the Bioterrorism Act. Other comments

recommend that FDA repropose the rule or not implement the rule.

(Response) Changes to the statute are beyond the scope of this rulemaking. Postponing implementation of or not implementing the rule is not viable under section 307(c) of the Bioterrorism Act, which not only directs the FDA to "promulgate proposed and final regulations for the requirement of