§ 1.1

REQUIREMENTS FOR NONTRANSPORTERS TO ESTABLISH AND MAINTAIN RECORDS TO IDENTIFY THE NONTRANSPORTER AND TRANSPORTER IMMEDIATE PREVIOUS SOURCES OF FOOD

1.337 What information must nontransporters establish and maintain to identify the nontransporter and transporter immediate previous sources of food?

REQUIREMENTS FOR NONTRANSPORTERS TO ESTABLISH AND MAINTAIN RECORDS TO IDENTIFY THE NONTRANSPORTER AND TRANSPORTER IMMEDIATE SUBSEQUENT RECEIVERS OF FOOD

1.345 What information must nontransporters establish and maintain to identify the nontransporter and transporter immediate subsequent recipients of food?

REQUIREMENTS FOR TRANSPORTERS TO ESTABLISH AND MAINTAIN RECORDS

1.352 What information must transporters establish and maintain?

GENERAL REQUIREMENTS

1.360 What are the record retention requirements?

1.361 What are the record availability requirements?

1.362 What records are excluded from this subpart?

1.363 What are the consequences of failing to establish or maintain records or make them available to FDA as required by this subpart?

COMPLIANCE DATES

1.368 What are the compliance dates for this subpart?

Subpart K—Administrative Detention of Food for Human or Animal Consumption

GENERAL PROVISIONS

1.377 What definitions apply to this subpart?

1.378 What criteria does FDA use to order a detention?

1.379 How long may FDA detain an article of food?

1.380 Where and under what conditions must the detained article of food be held?

1.381 May a detained article of food be delivered to another entity or transferred to another location?

1.382 What labeling or marking requirements apply to a detained article of food?

1.383 What expedited procedures apply when FDA initiates a seizure action against a detained perishable food?

1.384 When does a detention order terminate?

HOW DOES FDA ORDER A DETENTION?

1.391 Who approves a detention order?

1.392 Who receives a copy of the detention order?

1.393 What information must FDA include in the detention order?

WHAT IS THE APPEAL PROCESS FOR A DETENTION ORDER?

1.401 Who is entitled to appeal?

1.402 What are the requirements for submitting an appeal?

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1.404 Who serves as the presiding officer for an appeal, and for an informal hearing?

1.405 When does FDA have to issue a decision on an appeal?

1.406 How will FDA handle classified information in an informal hearing?


SOURCE: 42 FR 15553, Mar. 22, 1977, unless otherwise noted.
§ 1.20 Presence of mandatory label information.

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