**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2012–N–0001]

**Requirements for Importing Food and Drug Administration Regulated Products Into the United States**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of meeting.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the following meeting: “Requirements for Importing Food and Drug Administration Regulated Products Into the United States.” The topics to be discussed are FDA regulations with respect to importing pharmaceutical products, medical devices, food products, as well as technology which applies to brokers and forwarders.

**Date and Time:** The meeting will be held on July 24, 2012, from 8:30 a.m. to 5 p.m.

**Location:** The public workshop will be held at the Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Ave., Bethesda, MD 20814, 301–657–1234.

**Contact Person:** Mark Walderhaug, Center for Biologics Evaluation and Research (HFM–210), Food and Drug Administration, 1401 Rockville Pike, Suite 200N, Rockville, MD 20852–1448, 301–827–6028, email: Mark.Walderhaug@fda.hhs.gov.

**Registration:** Mail or email your registration information (including name, title, firm name, address, telephone, and fax numbers) to Mark Walderhaug (see Contact Person) by July 15, 2012. There is no registration fee for the public workshop. Early registration is recommended because seating is limited. Registration on the day of the public workshop will be provided on a space available basis beginning at 8 a.m.

If you need special accommodations due to a disability, please contact Mark Walderhaug (see Contact Person) at least 7 days in advance.

**SUPPLEMENTARY INFORMATION:** The public workshop presentations and panel discussions will:

(1) Discuss...