	Substances	CAS Nos.
217	1,2–DICHLOROETHYLENE	000540–59–0
218	1,2–DICHLOROETHENE, CIS-	000156–59–2
219	CARBON DISULFIDE	000075–15–0
220	PALLADIUM	007440–05–3
221	CHLOROETHANE	000075–00–3
222	ACETONE	000067–64–1
223	DIBENZOFURAN	000132–64–9
224	2,4–DIMETHYLPHENOL	000105–67–9
225	CHLOROMETHANE	000074–87–3
226	BIS(2–METHOXYETHYL) PHTHALATE	034006–76–3
227	BUTYL BENZYL PHTHALATE	000085–68–7
228	N–NITROSODIMETHYLAMINE	000062–75–9
229	1,2,4–TRICHLOROBENZENE	000120–82–1
230	N–NITROSODIPHENYLAMINE	000086–30–6
231	2–BUTANONE	000078–93–3
232	FLUORINE	007782–41–4
233	HYDROGEN FLUORIDE	007664–39–3
234	FLUORIDE ION	016984–48–8
235	AMMONIA	007664–41–7
236	COPPER	065357–62–2
237	CHLORINE DIOXIDE	010049–04–4
238	PBBs	059536–65–1
239	PBDEs	001163–19–5
240	SYNTHETIC VITREOUS FIBERS	NONE

Submission of Nominations for the Evaluation Set 24 Proposed Substances: Today's notice also invites voluntary public nominations for substances not listed in this notice. Nominations are most useful if they include the nominator, including full name, title, affiliation, e-mail address, and telephone number.

ATSDR will evaluate all data and information associated with nominated substances and will determine the final list of substances to be chosen for toxicological profile development. Substances will be chosen according to ATSDR's specific guidelines for selection, found in the *Selection Criteria* announced in the **Federal Register** on May 7, 1993 (87 FR 27288).

Submission of Comments: Submit comments via e-mail at: *tpcandidatecomments@cdc.gov*. Please include "Set 24" in the subject line of the e-mail. Or, mail to: Commander Jessilynn B. Taylor, Division of Toxicology and Environmental Medicine, 1600 Clifton Road, MS F–62, Atlanta, GA 30333; e-mail: *jbtaylor@cdc.gov*.

Please ensure that your comments are submitted within the specified nomination period. Nominations received after the closing date will be marked as late and may be considered only if time and resources permit.

Dated: March 26, 2010.

Ken Rose,

Associate Director, Office of Policy, Planning and Evaluation, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

[FR Doc. 2010–7169 Filed 3–30–10; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Pharmaceutical Supply Chain; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public workshop entitled "2010 PDA/FDA Pharmaceutical Supply Chain Workshop—Enough Talk: Let's Find and Implement Solutions." The workshop, cosponsored with the Parenteral Drug Association (PDA), will focus on solutions to reduce the risk to product quality in the pharmaceutical supply chain.

Date and Time: The conference will be held on Monday, April 26, 2010, from 8 a.m. to 6 p.m.; Tuesday, April 27, 2010, from 7:15 a.m. to 5:45 p.m.; and Wednesday, April 28, 2010, from 7:15 a.m. to 1:15 p.m.

Location: The public workshop will be held at the Hyatt Regency Bethesda, 7400 Wisconsin Ave., 1 Bethesda Metro Center, Bethesda, MD 20814; Phone: 301–657–1234; FAX: 301–657–6453.

Contact: Wanda Neal, Parenteral Drug Association, PDA Global Headquarters, Bethesda Towers, 4350 East-West Hwy., Suite 200, Bethesda, MD 20814; Phone: 301–656–5900, ext. 149.

Accommodations: Attendees are responsible for their own accommodations. To make reservations at the Hyatt Regency Bethesda, at the reduced conference rate, contact the Hyatt Regency Bethesda (see *Location*), citing meeting code "PDA." Room Rates are: Single: $209, plus 13% state and local taxes and Double: $234, plus 13% state and local taxes. Reservations can be made on a space and rate availability basis.

Registration: You are encouraged to register at your earliest convenience. The PDA registration fees cover the cost of facilities, materials, and breaks. Seats are limited; please submit your registration as soon as possible. Conference space will be filled in order of receipt of registration. Those accepted in to the conference will receive confirmation. Registration will close after applicable conference is filled. Onsite registration will be available on a space-available basis on the day of the public conference beginning at 7 a.m. on Monday, April 26, 2010.

The cost of registration is as follows:

PDA Members	$1850
PDA Nonmembers	$2099
PDA Member Government	$530
PDA Nonmember Government	$530
PDA Member Health Authority	$700
PDA Nonmember Health Authority	$800
PDA Member Academic	$700
PDA Nonmember Academic	$780

PDA Member Students	$280
PDA Nonmember Students	$310

If you need special accommodations due to a disability, please contact Wanda Neal, PDA (see *Contact*) at least 7 days in advance of the workshop.

Registration instructions: To register, please submit your name, affiliation, mailing address, phone, FAX number, and e-mail address, along with a check or money order payable to "PDA." Mail to: PDA, Global Headquarters, Bethesda Towers, 4350 East-West Hwy., Suite 200, Bethesda, MD 20814. To register via the Internet, go to See PDA Web site, *www.pda.org/supplychain2010* (FDA has verified the Web site address, but is not responsible for subsequent changes to the Web site after this document publishes in the **Federal Register**). The registrar will also accept payment by major credit cards (VISA/MasterCard only). For more information on the meeting, or for questions on registration, contact the PDA: Phone: 301–656–5900, FAX: 301–986–1093, or e-mail: *info@pda.org*.

SUPPLEMENTARY INFORMATION: A reliable supply of high quality, safe, and effective drug products and drug ingredients depends upon a series of controls across the entire supply chain from sourcing of incoming starting materials to distribution controls to marketing. Recent experiences in the market have highlighted the need for effective controls across the supply chain. There is a surge in global cooperation and efforts toward harmonization of good manufacturing practices (GMPs) and good distribution practices (GDPs) and controls pertaining to the supply chain among members of industry and regulatory agencies. Understanding and securing the entire ingredient manufacturing and distribution chain helps to ensure the quality and safety of medicines for patients.

Through a series of plenary sessions and working group breakout sessions, the workshop will provide participants the opportunity to:

- Hear from senior FDA personnel on the current regulatory environment.
- Share improvements in programs and technology.
- Identify any barriers to securing the entire ingredient manufacturing and distribution chain and associated actions to implement effective solutions.

Personnel with experience related to supply chain issues, including quality and technical functions, will find this level of information exchange with members of industry and regulatory agencies useful to their specific areas.

Dated: March 25, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–7151 Filed 3–30–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

United States Immigration and Customs Enforcement

Agency Information Collection Activities: Extension of an Existing Information Collection; Comment Request

ACTION: 30-Day notice of information collection for review; Form G–79A, Information Relating to Beneficiary of Private Bill; OMB Control No. 1653–0026.

The Department of Homeland Security, U.S. Immigration and Customs Enforcement (USICE), has submitted the following information collection request for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection was previously published in the **Federal Register** on January 19, 2010 Vol. 75 No. 11 2881, allowing for a 60 day public comment period. No comments were received on this information collection.

The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted for thirty days April 30, 2010. Written comments and suggestions from the public and affected agencies regarding items contained in this notice and especially with regard to the estimated public burden and associated response time should be directed to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to OMB Desk Officer, for U. S. Immigration and Customs Enforcement, Department of Homeland Security, and sent via electronic mail to *oira_submission@omb.eop.gov* or faxed to (202) 395–5806.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection:

(1) *Type of Information Collection:* Extension of a currently approved information collection.

(2) *Title of the Form/Collection:* Information Relating to Beneficiary of Private Bill.

(3) *Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* Form G–79A. U.S. Immigration and Customs Enforcement.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract: Primary:* Individuals or Households. The information collected on the Form G–79A is necessary for U.S. Immigration and Customs Enforcement to provide reports to Congress on Private Bills when requested.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 100 responses at 60 minutes (1 hour) per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 100 annual burden hours.

Requests for a copy of the proposed information collection instrument, with instructions; or inquiries for additional information should be requested via e-mail to: *forms.ice@dhs.gov* with "Form G–79A" in the subject line.

Dated: March 26, 2010.

Joseph M. Gerhart,

Chief, Records Management Branch, U.S. Immigration and Customs Enforcement, Department of Homeland Security.

[FR Doc. 2010–7186 Filed 3–30–10; 8:45 am]

BILLING CODE 9111–28–P