requirements of the applicable statutes and regulations.

III. Paperwork Reduction Act of 1995

Under the Paperwork Reduction Act of 1995 (the PRA) (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3 and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506 (c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information before submitting the collection to OMB for approval. To comply with this requirement, we are publishing a notice of the proposed collection of information set forth below.

With respect to the following collection of information, we invite comments on: (1) Whether the proposed collection of information is necessary for the proper performance of our functions, including whether the information will have practical utility; (2) the accuracy of our estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Title: Draft Compliance Policy Guide on Labeling and Marketing of Nutritional Products Intended for Use to Diagnose, Cure, Mitigate, Treat, or Prevent Disease in Dogs and Cats.

Description: The purpose of this CPG is to communicate FDA's strategy with respect to dog and cat food products that are labeled and/or marketed as intending to diagnose, cure, mitigate, treat, or prevent diseases and to provide nutrients in support of meeting the animal's total daily nutrient requirements.

Description of Respondents: Manufacturers of dog and cat foods that are labeled and/or marketed as intending to diagnose, cure, mitigate, treat, or prevent diseases and to provide nutrients in support of meeting the animal's total daily nutrient requirements.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Sections 402 and 403 of the FD&C Act	5	75	375	.25	94

[1] There are no operating costs or maintenance costs associated with this collection of information.

CVM estimates from its experience that approximately 5 manufacturers will be affected by the draft CPG, times 75 products produced annually equals 375 total annual responses. The hours per response are based on approximately .25 hour per response for respondents to look up the ingredient names in the AFFCO *Official Publication*.

This draft CPG also refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR part 571 (Food Additive Petitions and FAP Labeling) have been approved under OMB control number 0910–0546. The collection of information in 21 CFR 570.35 (GRAS) has been approved under OMB control number 0910–0342. The requirement for food facility registration has been approved under OMB control number 0910–0502.

IV. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) electronic or written comments regarding this document. It is only necessary to send one set of comments. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

V. Electronic Access

Copies of the CPG may be downloaded to a personal computer with access to the Internet. The Office of Regulatory Affairs home pages include this draft CPG and may be accessed at *http://www.fda.gov/ICECI/ComplianceManuals/* under ''Compliance Policy Guides.''

Dated: August 24, 2012.

Dara A. Corrigan,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 2012–22231 Filed 9–7–12; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0902]

Withdrawal of Approval of New Animal Drug Applications; Chorionic Gonadotropin; Naloxone; Oxymorphone; Oxytocin

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of four new animal drug applications (NADAs) at the sponsor's request because the products are no longer manufactured or marketed.

DATES: Withdrawal of approval is effective September 20, 2012.

FOR FURTHER INFORMATION CONTACT: David Alterman, Center for Veterinary Medicine (HFV–212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6843, email: *david.alterman@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The sponsors in table 1 of this document have requested that FDA withdraw approval of the four NADAs listed

because the products are no longer manufactured or marketed.

TABLE 1—WITHDRAWAL OF APPROVAL REQUESTS

NADA No.	Trade name (drug)	Applicant
030–525	NUMORPHAN (oxymorphone hydrochloride) Injection	Endo Pharmaceuticals Inc., 100 Painters Dr., Chadds Ford, PA 19317.
035–825	NARCAN (naloxone hydrochloride) Injection	Endo Pharmaceuticals Inc., 100 Painters Dr., Chadds Ford, PA 19317.
046–822	VETOCIN (oxytocin) Injection	United Vaccines, A Harlan Sprague Dawley, Inc., Co., P.O. Box 4220, Madison, WI 53711.
103–090	CHORTROPIN (chorionic gonadotropin) Injection	United Vaccines, A Harlan Sprague Dawley, Inc., Co., P.O. Box 4220, Madison, WI 53711.

Therefore, under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, and in accordance with § 514.116 *Notice of withdrawal of approval of application* (21 CFR 514.116), notice is given that approval of NADAs 030–525, 035–825, 046–822, and 103–090, and all supplements and amendments thereto, is hereby withdrawn, effective September 20, 2012.

Elsewhere in this issue of the **Federal Register**, FDA is amending the animal drug regulations to reflect the voluntary withdrawal of approval of these applications.

Dated: September 5, 2012.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. 2012–22195 Filed 9–7–12; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Prevention; Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given that the Substance Abuse and Mental Health Services Administration's (SAMHSA) Center for Substance Abuse Prevention (CSAP) Drug Testing Advisory Board (DTAB) will meet on September 24, 2012 from 9 a.m. to 5 p.m. and September 25, 2012 from 9 a.m. to 2 p.m. E.D.T.

The Board will discuss proposed revisions to the Mandatory Guidelines for Federal Workplace Drug Testing Programs. Therefore, this meeting is closed to the public as determined by the Administrator, SAMHSA, in accordance with 5 U.S.C. 552b(c)(9)(B) and 5 U.S.C. App. 2, Section 10(d).

Meeting information and a roster of DTAB members may be obtained by accessing the SAMHSA Advisory Committees' Web site, *http://www.nac.samhsa.gov/DTAB/meetings.aspx,* or by contacting Dr. Cook.

Committee Name: Substance Abuse and Mental Health Services Administration's Center for Substance Abuse Prevention, Drug Testing Advisory Board.

Dates/Time/Type: September 24, 2012 from 9 a.m. to 5 p.m. E.D.T.: CLOSED, September 25, 2012 from 9 a.m. to 2 p.m. E.D.T.: CLOSED.

Place: Sugarloaf Conference Room, SAMHSA Office Building, 1 Choke Cherry Road, Rockville, Maryland 20857.

Contact: Janine Denis Cook, Ph.D., Designated Federal Official, CSAP Drug Testing Advisory Board, 1 Choke Cherry Road, Room 7–1043, Rockville, Maryland 20857, Telephone: 240–276–2600, Fax: 240–276–2610, Email: *janine.cook@samhsa.hhs.gov*.

Janine Denis Cook,

Designated Federal Official, DTAB, Division of Workplace Programs, Center for Substance Abuse Prevention, Substance Abuse and Mental Health Services Administration.

[FR Doc. 2012–22167 Filed 9–7–12; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[Docket No. USCG–2012–0782]

Public Workshop on Marine Technology and Standards

AGENCY: Coast Guard, DHS.

ACTION: Notice.

SUMMARY: The American Society of Mechanical Engineers (ASME), in coordination with the United States Coast Guard (USCG), is sponsoring a two-day public workshop on marine technology and standards in Arlington, VA. This public workshop will provide a unique opportunity for classification societies, industry groups, standards development organizations, government organizations, and other interested members of the public to come together for a professional exchange of information on topics ranging from technological impacts on the marine industry, corresponding coverage in related codes and standards, and government regulations.

DATES: The two-day workshop will be held on Wednesday, July 24, 2013, and Thursday, July 25, 2013. The deadline for advance registration is Monday, July, 1, 2013. If you are interested in presenting a paper at the workshop, you must submit a 100 word abstract by email to *workshop@uscg.mil.* Abstracts are due on or before November 2, 2012.

See **SUPPLEMENTARY INFORMATION** below for other dates related to submission of abstracts, draft papers, and presentations, as well as more information on how to register for the workshop.

ADDRESSES: The workshop will be held at The Double Tree by Hilton Hotel, in the Crystal City neighborhood of Arlington VA. The hotel is located at 300 Army Navy Drive, Arlington, VA; the hotel phone number is (703) 416–4100. The hotel is located approximately three miles from Ronald Reagan Washington National Airport (DCA) and approximately four blocks from the Pentagon City Metro station. For registration information or to obtain further information about this workshop, visit the USCG Web site at *http://www.uscg.mil/marine_event.* The docket for this notice is available for inspection or copying at the Docket Management Facility (M–30), U.S. Department of Transportation, West Building Ground Floor, Room W12–140, 1200 New Jersey Avenue SE, Washington, DC 20590, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays. You may also find this docket on the Internet by going to *http://www.regulations.gov,* inserting USCG–2012–0782 in the ''Search'' box, and then clicking ''Search.''