MZ4339390–02X, MZ4306000–02X, MZ4339390–10X, or MZ4306000–10X as ''March 5, 2010,'' the calendar compliance time is April 14, 2011 (18 months after October 14, 2009 (the effective date of AD 2009–18–20, Amendment 39–16017 (74 FR 46313, September 9, 2009))).

(6) Where Note (6) of ''ATA 27–64–00 Flight Control—Spoiler Hydraulic Actuation,'' of Sub-part 4–2–1, ''Life Limits,'' of Sub-part 4–2, ''Systems Life Limited Components,'' of Airbus A330 ALS Part 4—Aging Systems Maintenance, Revision 04, dated August 27, 2013, defines a calendar date of ''September 5, 2008,'' as a date for the determination of accumulated flight cycles since the aircraft initial entry into service, the date is October 14, 2009 (the effective date of AD 2009–18–20, Amendment 39–16017 (74 FR 46313, September 9, 2009)).

(7) Where Note (6) of ''ATA 27–64–00 Flight Control—Spoiler Hydraulic Actuation,'' of Sub-part 4–2–1, ''Life Limits,'' of Sub-part 4–2, ''Systems Life Limited Components,'' of Airbus A330 ALS Part 4—Aging Systems Maintenance, Revision 04, dated August 27, 2013, defines a calendar compliance time as ''March 5, 2010,'' for the modification of affected servo controls, the calendar compliance time is April 14, 2011 (18 months after October 14, 2009 (the effective date of AD 2009–18–20, Amendment 39–16017 (74 FR 46313, September 9, 2009))).

(i) No Alternative Actions or Intervals

After accomplishing the revision required by paragraph (g) of this AD, no alternative actions (*e.g.,* inspections) or intervals may be used unless the actions or intervals are approved as an alternative method of compliance (AMOC) in accordance with the procedures specified in paragraph (j)(1) of this AD.

(j) Other FAA AD Provisions

The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs):* The Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA, has the authority to approve AMOCs for this AD, if requested using the procedures found in 14 CFR 39.19. In accordance with 14 CFR 39.19, send your request to your principal inspector or local Flight Standards District Office, as appropriate. If sending information directly to the International Branch, send it to ATTN: Vladimir Ulyanov, Aerospace Engineer, International Branch, ANM–116, Transport Airplane Directorate, FAA, 1601 Lind Avenue SW., Renton, WA 98057–3356; telephone 425–227–1138; fax 425–227–1149. Information may be emailed to: *9-ANM-116-ACO-AMOC-REQUESTS@faa.gov.* Before using any approved AMOC, notify your appropriate principal inspector, or lacking a principal inspector, the manager of the local flight standards district office/certificate holding district office. The AMOC approval letter must specifically reference this AD.

(2) *Contacting the Manufacturer:* For any requirement in this AD to obtain corrective actions from a manufacturer, the action must be accomplished using a method approved by the Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA; or the European Aviation Safety Agency (EASA); or Airbus's EASA Design Organization Approval (DOA). If approved by the DOA, the approval must include the DOA-authorized signature.

(k) Related Information

(1) Refer to Mandatory Continuing Airworthiness Information (MCAI) European Aviation Safety Agency (EASA) AD 2013–0268, dated November 7, 2013, for related information. You may examine the MCAI in the AD docket on the Internet at *http://www.regulations.gov/#!documentDetail;D=FAA-2013-0834-0003.*

(2) For service information identified in this AD, contact Airbus SAS, Airworthiness Office—EAL, 1 Rond Point Maurice Bellonte, 31707 Blagnac Cedex, France; telephone +33 5 61 93 36 96; fax +33 5 61 93 45 80; email *airworthiness.A330–A340@airbus.com;* Internet *http://www.airbus.com.* You may view this service information at the FAA, Transport Airplane Directorate, 1601 Lind Avenue SW., Renton, WA. For information on the availability of this material at the FAA, call 425–227–1221.

Issued in Renton, Washington, on February 20, 2015.

Victor Wicklund,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 2015–05031 Filed 3–6–15; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Part 702

[RIN 0694–AG17]

U.S. Industrial Base Surveys Pursuant to the Defense Production Act of 1950; Correction

AGENCY: Bureau of Industry and Security, Commerce.

ACTION: Proposed rule; correction.

SUMMARY: This rule corrects the preamble to a proposed rule published in the **Federal Register** of March 3, 2015, regarding U.S. Industrial Base Surveys by adding the inadvertently omitted **ADDRESSES** Caption.

DATES: March 9, 2015.

FOR FURTHER INFORMATION CONTACT: William Arvin, Bureau of Industry and Security Regulatory Policy Division, 202–482–2440 or *william.arvin@bis.doc.gov.*

Correction

In proposed rule FR Doc. 2015–04299, on page 11350 in the issue of March 3, 2015, in the first column, immediately following the **DATES** section, add the following:

ADDRESSES: Comments may be submitted:

- Via the Federal eRulemaking Portal: *http://www.regulations.gov.* Search for this rule using its regulations.gov docket number: BIS–2015–0010.
- By email directly to *publiccomments@bis.doc.gov.* Include ''RIN 0694–AG17'' in the subject line.
- By mail or delivery to Regulatory Policy Division, Bureau of Industry and Security, U.S. Department of Commerce, Room 2099B, 14th Street and Pennsylvania Avenue NW., Washington, DC 20230. Refer to ''RIN 0694–AG17.''

Dated: March 3, 2015.

Matthew S. Borman,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 2015–05324 Filed 3–6–15; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 201, 606, and 610

[Docket No. FDA–2007–N–0363]

RIN 0910–AG18

Electronic Distribution of Prescribing Information for Human Prescription Drugs, Including Biological Products; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed rule; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending the comment period for the proposed rule that appeared in the **Federal Register** of December 18, 2014. In the proposed rule, FDA requested comments on its proposal to amend its labeling regulations for human prescription drugs and biological products to require that the prescribing information intended for health care professionals that is on or within the package from which the product is dispensed be distributed electronically and not in paper form, except as provided by the proposed rule. The Agency is taking this action in response to a request for an extension to allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period on the proposed rule published on December 18, 2014 (79 FR 75506). Submit either electronic or written comments by May 18, 2015.

ADDRESSES: You may submit comments to the proposed rule by any of the following methods:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

Written Submissions

Submit written submissions in the following ways:

- *Mail/Hand delivery/Courier (for paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Instructions: All submissions received must include the Docket No. FDA–2007–N–0363 for this rulemaking. All comments received may be posted without change to *http://www.regulations.gov,* including any personal information provided. For additional information on submitting comments, see the ''Comments'' heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or comments received, go to *http://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Emily Gebbia, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6226, Silver Spring, MD 20993, 240–402–0980.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of December 18, 2014, FDA published a proposed rule to amend its labeling regulations for human prescription drugs and biological products to require that the prescribing information intended for health care professionals that is on or within the package from which the product is dispensed be distributed electronically and not in paper form, except as provided by the proposed rule. The proposed rule is intended to facilitate the distribution of updated prescribing information as new information becomes available and as changes in prescribing information are made. FDA is proposing the change to help ensure that the most current prescribing information will be available and readily accessible to health care professionals at the time of clinical decision making and dispensing. FDA provided a 90-day comment period (*i.e.,* until March 18, 2015) for the proposed rule.

The Agency has received a request for a 60-day extension of the comment period for the proposed rule. The request conveyed concern that the current 90-day comment period does not allow sufficient time for entities and individuals who will be most affected by a final rule to examine and to comment upon the proposed rule. The request suggested that FDA would benefit by granting stakeholders sufficient time to develop their comments and to address as many relevant issues as possible.

FDA has considered the request and is extending the comment period for the proposed rule for 60 days, until May 18, 2015. The Agency believes that a 60-day extension allows adequate time for interested persons to submit comments without significantly delaying rulemaking on this important issue.

II. Request for Comments

Interested persons may submit either electronic comments regarding the proposed rule to *http://www.regulations.gov* or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

Dated: March 3, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–05336 Filed 3–6–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 165

[Docket Number USCG–2015–0071]

RIN 1625–AA08

Safety Zone; 24 Mile Tampa Bay Marathon Swim, Tampa Bay, Tampa, FL

AGENCY: Coast Guard, DHS.

ACTION: Notice of proposed rulemaking.

SUMMARY: The Coast Guard is proposing to establish a temporary moving safety zone on the waters of the Tampa Bay in Tampa, Florida during the 24 Mile Tampa Bay Marathon Swim. The swim is scheduled to take place from 4 a.m. to 9 p.m. on April 25, 2015. Approximately 30 swimmers are anticipated to participate in the marathon swim event. No spectators are expected to be present during the event. The safety zone is necessary to provide for the safety of the participants, participant vessels, and the general public on the navigable waters of the United States during the event. The safety zone will establish a moving protective area around all swimmers involved in the race. Persons and vessels, except those participating in the event, will be prohibited from entering, transiting through, anchoring in, or remaining within the regulated area unless authorized by the Captain of the Port St. Petersburg or a designated representative.

DATES: Requests for public meetings, comments or related material must be received by the Coast Guard on or before March 16, 2015.

ADDRESSES: You may submit comments identified by docket number using any one of the following methods:

(1) *Federal eRulemaking Portal: http://www.regulations.gov*.

(2) *Fax:* (202) 493–2251.

(3) *Mail or Delivery:* Docket Management Facility (M–30), U.S. Department of Transportation, West Building Ground Floor, Room W12–140, 1200 New Jersey Avenue SE., Washington, DC 20590–0001. Deliveries accepted between 9 a.m. and 5 p.m., Monday through Friday, except federal holidays. The telephone number is (202) 366–9329.

See the ''Public Participation and Request for Comments'' portion of the **SUPPLEMENTARY INFORMATION** section below for further instructions on submitting comments. To avoid duplication, please use only one of these three methods.

FOR FURTHER INFORMATION CONTACT: If you have questions on this rule, call or email Tyrone J. Stafford, Sector St. Petersburg Prevention Department, Coast Guard; telephone (813) 228–2191, email *D07-SMB-Tampa-WWM@uscg.mil*. If you have questions on viewing or submitting material to the docket, call Cheryl Collins, Program Manager, Docket Operations, telephone (202) 366–9826.

SUPPLEMENTARY INFORMATION: