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.