

The FAA's authority to issue rules regarding aviation safety is found in title 49 of the U.S. Code. Subtitle 1, section 106, describes the authority of the FAA Administrator. Subtitle VII, Aviation Programs, describes in more detail the scope of the agency's authority. This rulemaking is promulgated under the authority described in subtitle VII, part A, subpart I, section 40103. Under that section, the FAA is charged with prescribing regulations to assign the use of airspace necessary to ensure the safety of aircraft and the efficient use of airspace. This regulation is within the scope of that authority as it creates controlled airspace at Hardin County Airport, Kenton, OH.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (Air).

Adoption of the Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—DESIGNATION OF CLASS A, B, C, D, AND E AIRSPACE AREAS; AIR TRAFFIC SERVICE ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40103, 40113, 40120; E. O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

■ 2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9U, Airspace Designations and Reporting Points, dated August 18, 2010, and effective September 15, 2010, is amended as follows:

Paragraph 6005 Class E airspace areas extending upward from 700 feet or more above the surface.

* * * * *

AGL OH E5 Kenton, OH [Amended]

Kenton, Hardin County Airport, OH
(Lat. 40°36'36" N., long. 83°38'39" W.)

That airspace extending upward from 700 feet above the surface bounded by a line beginning at lat. 40°43'34" N., long. 83°33'51" W., to lat. 40°38'16" N., long. 83°28'39" W., to lat. 40°30'37" N., long. 83°30'57" W., to lat. 40°24'00" N., long. 83°33'37" W., to lat. 40°13'31" N., long. 83°40'22" W., to lat. 40°11'47" N., long. 83°52'11" W., to lat. 40°16'44" N., long. 84°01'10" W., to lat. 40°24'31" N., long. 84°02'39" W., to lat. 40°31'30" N., long. 83°56'56" W., to lat. 40°32'13" N., long. 83°50'20" W., to lat. 40°34'45" N., long. 83°47'33" W., to lat. 40°38'56" N., long. 83°48'49" W., to lat. 40°43'49" N., long. 83°42'14" W., to the point of beginning.

Issued in Fort Worth, Texas, on March 15, 2011.

Richard J. Kervin, Jr.,

*Acting Manager Operations Support Group,
ATO Central Service Center.*

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DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Part 774

[Docket No. 110106012–1013–01]

RIN 0694–AF04

Implementation of the Understandings Reached at the 2010 Australia Group (AG) Plenary Meeting and Other AG-Related Clarifications and Corrections to the EAR

AGENCY: Bureau of Industry and Security, Commerce.

ACTION: Final rule.

SUMMARY: The Bureau of Industry and Security (BIS) publishes this final rule to amend the Export Administration Regulations (EAR) to implement the understandings reached at the June 2010 plenary meeting of the Australia Group (AG) and to make certain AG-related editorial clarifications and corrections to the EAR. Consistent with the June 2010 AG understandings, this rule amends the chemical manufacturing equipment entry on the Commerce Control List (CCL) of the EAR to reflect the addition of two parenthetical phrases that clarify the description of certain “materials” contained in items on the AG “Control List of Dual-Use Chemical Manufacturing Facilities and Equipment and Related Technology and Software.” In addition, this rule makes AG-related clarifications and corrections to the EAR. Specifically, this rule amends the human and zoonotic pathogens and toxins entry and the animal pathogens entry on the CCL by making an update and a clarification that are consistent with the description of items on the AG “List of Biological Agents for Export Control” and the AG “List of Animal Pathogens for Export Control,” respectively. Finally, this rule amends the listing for “valves” in the chemical manufacturing equipment entry on the CCL to clarify that it controls “valves” for the “production” of chemicals, as well as “valves” for the “processing” or “containment” of chemicals. The purpose of this rule is to ensure that the AG-related entries on the CCL conform with the wording in the AG Control Lists (as updated by the

understandings reached at the 2010 AG Plenary) and to clarify the meaning of terms used in these entries.

DATES: This rule is effective April 20, 2011.

ADDRESSES: Send comments regarding this collection of information, including suggestions for reducing the burden, to Jasmeet Sehra, Office of Management and Budget (OMB), by e-mail to Jasmeet.K.Sehra@omb.eop.gov, or by fax to (202) 395–7285; and to the Regulatory Policy Division, Bureau of Industry and Security, Department of Commerce, 14th Street & Pennsylvania Avenue, NW., Room 2705, Washington, DC 20230. Comments on this collection of information should be submitted separately from comments on the final rule (*i.e.*, RIN 0694–AF04)—all comments on the latter should be submitted by one of the three methods outlined above.

FOR FURTHER INFORMATION CONTACT: Elizabeth Sangine, Director, Chemical and Biological Controls Division, Office of Nonproliferation and Treaty Compliance, Bureau of Industry and Security, Telephone: (202) 482–3343.

SUPPLEMENTARY INFORMATION:

Background

The Bureau of Industry and Security (BIS) is amending the Export Administration Regulations (EAR) to implement the understandings reached at the annual plenary meeting of the Australia Group (AG) that was held in Paris, France, from May 31 through June 4, 2010. The AG is a multilateral forum consisting of 40 participating countries that maintain export controls on a list of chemicals, biological agents, and related equipment and technology that could be used in a chemical or biological weapons program. The AG periodically reviews items on its control list to enhance the effectiveness of participating governments' national controls and to achieve greater harmonization among these controls.

Consistent with the understandings reached at the 2010 AG Plenary, this final rule amends Export Control Classification Number (ECCN) 2B350 (Chemical manufacturing facilities and equipment) on the Commerce Control List (CCL) (Supplement No. 1 to Part 774 of the EAR) to reflect the addition of two parenthetical phrases that describe types of “materials” contained in items on the AG “Control List of Dual-Use Chemical Manufacturing Facilities and Equipment and Related Technology and Software.” Specifically, this rule adds a parenthetical phrase to ECCN 2B350.a.3, .b.3, .c.3, .d.3, .e.3, .g.3, .h.3, and .i.3 to clarify the meaning

of the term “fluoropolymers” in connection with the types of “materials” from which certain chemical manufacturing equipment is made. This parenthetical phrase describes “fluoropolymers” as “polymeric or elastomeric materials with more than 35% fluorine by weight.” This rule also adds a parenthetical phrase in ECCN 2B350.i.11 (under the listing for “pumps”) to clarify that the “material” “ferrosilicon” refers to “high silicon iron alloys.”

In addition to the AG Plenary changes described above, this rule amends ECCN 1C351 (human and zoonotic pathogens and “toxins”) and ECCN 1C352 (animal pathogens) on the CCL by updating ECCN 1C351 and clarifying ECCN 1C352 consistent with the controls described in the AG “List of Biological Agents for Export Control” and the AG “List of Animal Pathogens for Export Control,” respectively. Specifically, this rule revises the listing for “Chlamydia psittaci” in ECCN 1C351.c.7 by updating the name of the bacterium to read “Chlamydophilapsittaci (formerly known as Chlamydia psittaci).” This rule also revises the listing for the “Lyssa virus” in ECCN 1C352.a.8 by adding a parenthetical phrase to indicate that the virus is also known as “Rabies.”

This rule also makes two clarifications to ECCN 2B350, consistent with the controls described in the AG “Control List of Dual-Use Chemical Manufacturing Facilities and Equipment and Related Technology and Software.” First, this rule amends the introductory text of ECCN 2B350.g (“valves”) to clarify that the ECCN controls valves specified therein that are used in the “production” of chemicals, as well as valves that are used in the “processing” or “containment” of chemicals. Second, this rule revises the description of the material “Glass or glasslined (including vitrified or enameled coatings)” in ECCN 2B350.g.4 to read “Glass (including vitrified or enameled coating or glass lining)” to clarify the extent to which valves containing this type of material are controlled under this ECCN.

None of the changes made by this rule alters the scope of the controls in ECCNs 1C351, 1C352, or 2B350.

Although the Export Administration Act expired on August 20, 2001, the President, through Executive Order 13222 of August 17, 2001, 3 CFR, 2001 Comp., p. 783 (2002), as extended by the Notice of August 12, 2010, 75 FR 50681 (August 16, 2010), has continued the Export Administration Regulations in effect under the International Emergency Economic Powers Act.

Rulemaking Requirements

1. Executive Orders 13563 and 12866 direct agencies to assess all costs and benefits of available regulatory alternatives and, if regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety effects, distributive impacts, and equity). Executive Order 13563 emphasizes the importance of quantifying both costs and benefits, of reducing costs, of harmonizing rules, and of promoting flexibility. This rule has been designated a “significant regulatory action” although not economically significant, under section 3(f) of Executive Order 12866. Accordingly, the rule has been reviewed by the Office of Management and Budget.

2. Notwithstanding any other provision of law, no person is required to respond to, nor shall any person be subject to a penalty for failure to comply with, a collection of information subject to the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*) (PRA), unless that collection of information displays a currently valid Office of Management and Budget (OMB) Control Number. This rule contains a collection of information subject to the requirements of the PRA. This collection has been approved by OMB under Control Number 0694–0088 (Multi-Purpose Application), which carries a burden hour estimate of 58 minutes to prepare and submit form BIS–748. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to Jasmeet Sehra, Office of Management and Budget (OMB), and to the Regulatory Policy Division, Bureau of Industry and Security, Department of Commerce, as indicated in the **ADDRESSES** section of this rule.

3. This rule does not contain policies with Federalism implications as that term is defined in Executive Order 13132.

4. The provisions of the Administrative Procedure Act (5 U.S.C. 553) requiring notice of proposed rulemaking, the opportunity for public participation, and a delay in effective date, are inapplicable because this regulation involves a military and foreign affairs function of the United States (See 5 U.S.C. 553(a)(1)). Immediate implementation of these amendments is non-discretionary and fulfills the United States’ international obligation to the Australia Group (AG). The AG contributes to international

security and regional stability through the harmonization of export controls and seeks to ensure that exports do not contribute to the development of chemical and biological weapons. The AG consists of 40 member countries that act on a consensus basis and the amendments set forth in this rule implement agreements reached at the June 2010 plenary session of the AG and other changes that are necessary to ensure consistency with the controls maintained by the AG. Since the United States is a significant exporter of the items in this rule, implementation of this provision is necessary for the AG to achieve its purpose. Any delay in implementation will create a disruption in the movement of affected items globally because of disharmony between export control measures implemented by AG members, resulting in tension between member countries. Export controls work best when all countries implement the same export controls in a timely and coordinated manner.

Further, no other law requires that a notice of proposed rulemaking and an opportunity for public comment be given for this final rule. Because a notice of proposed rulemaking and an opportunity for public comment are not required to be given for this rule under the Administrative Procedure Act or by any other law, the analytical requirements of the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*) are not applicable. Therefore, this regulation is issued in final form.

List of Subjects in 15 CFR Part 774

Exports, Foreign trade, Reporting and recordkeeping requirements.

Accordingly, part 774 of the Export Administration Regulations (15 CFR parts 730–774) is amended as follows:

PART 774—[AMENDED]

■ 1. The authority citation for 15 CFR part 774 continues to read as follows:

Authority: 50 U.S.C. app. 2401 *et seq.*; 50 U.S.C. 1701 *et seq.*; 10 U.S.C. 7420; 10 U.S.C. 7430(e); 22 U.S.C. 287c, 22 U.S.C. 3201 *et seq.*; 22 U.S.C. 6004; 30 U.S.C. 185(s), 185(u); 42 U.S.C. 2139a; 42 U.S.C. 6212; 43 U.S.C. 1354; 15 U.S.C. 1824a; 50 U.S.C. app. 5; 22 U.S.C. 7201 *et seq.*; 22 U.S.C. 7210; E.O. 13026, 61 FR 58767, 3 CFR, 1996 Comp., p. 228; E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783; Notice of August 12, 2010, 75 FR 50681 (August 16, 2010).

Supplement No. 1 to Part 774—[Amended]

2. In Supplement No. 1 to Part 774 (the Commerce Control List), Category 1—Special Materials and Related Equipment, Chemicals,

“Microorganisms” & “Toxins,” ECCN 1C351 is amended by removing the name “Chlamydia psittaci”, where it appears in paragraph c.7 of the “Items” paragraph in the List of Items Controlled section, and adding in its place the name “Chlamyphilapsittaci (formerly known as Chlamydia psittaci)”.

3. In Supplement No. 1 to Part 774 (the Commerce Control List), Category 1—Special Materials and Related Equipment, Chemicals,

“Microorganisms” & “Toxins,” ECCN 1C352 is amended by removing the name “Lyssa virus”, where it appears in paragraph a.8 of the “Items” paragraph in the List of Items Controlled section, and adding in its place the name “Lyssa virus (a.k.a. Rabies)”.

4. In Supplement No. 1 to Part 774 (the Commerce Control List), Category 2—Materials Processing, ECCN 2B350 is amended under the “Items” paragraph in the List of Items Controlled section:

a. By adding the parenthetical phrase “(polymeric or elastomeric materials with more than 35% fluorine by weight)” immediately following the word “Fluoropolymers”, where it appears in paragraphs a.3, b.3, c.3, d.3, e.3, g.3, h.3, and i.3;

b. By removing the phrase “chemical(s) being processed or contained”, where it appears in the introductory text to paragraph g, and adding in its place the phrase “chemical(s) being produced, processed, or contained”;

c. By removing the phrase “Glass or glasslined (including vitrified or enameled coatings);”, where it appears in paragraph g.4, and adding in its place the phrase “Glass (including vitrified or enameled coating or glass lining);”;

d. By adding the parenthetical phrase “(high silicon iron alloys)” immediately following the word “Ferrosilicon”, where it appears in paragraph i.11.

Dated: April 12, 2011.

Kevin J. Wolf,

Assistant Secretary for Export Administration.

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CONSUMER PRODUCT SAFETY COMMISSION

16 CFR Part 1217

RIN 3041–AC79

Safety Standard for Toddler Beds

AGENCY: Consumer Product Safety Commission.

ACTION: Final rule.

SUMMARY: The Consumer Product Safety Improvement Act of 2008 (“CPSIA”) requires the United States Consumer Product Safety Commission (“Commission,” “CPSC”) to promulgate consumer product safety standards for durable infant or toddler products. These standards are to be “substantially the same as” applicable voluntary standards or more stringent than the voluntary standard if the Commission concludes that more stringent requirements would further reduce the risk of injury associated with the product. The Commission is issuing a safety standard for toddler beds in response to the CPSIA. The safety standard addresses entrapment in bed end structures, entrapment between the guardrail and side rail, entrapment in the mattress support system, and component failures of the bed support system and guardrails. The standard also addresses corner post extensions that can catch items worn by a child.

DATES: The rule will become effective on October 20, 2011, and apply to products manufactured or imported on or after that date. The incorporation by reference of the publications listed in this rule are approved by the Director of the Federal Register as of October 20, 2011.

FOR FURTHER INFORMATION CONTACT: Troy Whitfield, Office of Compliance and Field Operations, Consumer Product Safety Commission, Bethesda, MD 20814–4408; telephone (301) 504–7548; twhitfield@cpsc.gov.

SUPPLEMENTARY INFORMATION:

A. Background: Section 104(b) of the Consumer Product Safety Improvement Act

The Consumer Product Safety Improvement Act of 2008 (“CPSIA”, Pub. L. 110–314) was enacted on August 14, 2008. Section 104(b) of the CPSIA requires the Commission to promulgate consumer product safety standards for durable infant or toddler products. The law requires that these standards are to be “substantially the same as” applicable voluntary standards or more stringent than the voluntary standards if the Commission concludes that more stringent requirements would further reduce the risk of injury associated with the product. The term “durable infant or toddler product” is defined in section 104(f) of the CPSIA as a durable product intended for use, or that may be reasonably expected to be used, by children under the age of 5 years. Toddler beds are one of the products specifically identified in section 104(f)(2) of the CPSIA as a durable infant or toddler product.

In this document, the Commission is issuing a safety standard for toddler beds. The standard is largely the same as a voluntary standard developed by ASTM International (formerly the American Society for Testing and Materials), ASTM F 1821–09, *Standard Consumer Safety Specification for Toddler Beds*, but with several modifications that strengthen the standard.

In the **Federal Register** of April 28, 2010, the Commission published a notice of proposed rulemaking that proposed to incorporate by reference ASTM F 1821–09, *Standard Consumer Safety Specification for Toddler Beds*, with several modifications. 75 FR 22291. The final rule is very similar to the proposed rule. We summarize the proposed rule in section F of this preamble and discuss the final rule (including differences between the proposal and the final rule) in section G of this preamble. The information discussed in this preamble comes from CPSC staff’s briefing package for the toddler bed final rule, which is available on the CPSC’s Web site at <http://www.cpsc.gov/library/foia/foia11/brief/toddlerfinal.pdf>.

B. The Product

The ASTM voluntary standard defines a toddler bed as any bed sized to accommodate a full-size crib mattress having minimum dimensions of 51⁵/₈ inches in length and 27¹/₄ inches in width and that is intended to provide free access and egress to a child not less than 15 months of age and weighing no more than 50 pounds. The standard includes cribs that can be converted into a toddler bed using a full-size crib mattress.

CPSC staff estimates that there are currently at least 73 known manufacturers or importers supplying toddler beds and/or convertible cribs to the U.S. market. Approximately 48 suppliers are domestic manufacturers (66 percent); 13 are domestic importers (18 percent); 11 are foreign manufacturers (15 percent); and the remaining firm is a foreign supplier that imports from other countries and exports to the United States.

Based on information from a 2005 survey conducted by the American Baby Group, CPSC staff estimates annual convertible crib sales to number about 776,000 and annual sales of toddler beds to total about 819,000. Thus, a total of approximately 1.6 million units (convertible cribs and toddler beds) sold per year might be affected by the toddler bed standard.