after adjustment for inflation is $122 million, using the most current (2005) Implicit Price Deflator for the Gross Domestic Product. FDA does not expect this proposed rule to result in any 1-year expenditure that would meet or exceed this amount.

XII. Federalism

FDA has analyzed this proposed rule in accordance with the principles set forth in Executive Order 13132. FDA has determined that the proposed rule, if finalized, would not contain policies that would have substantial direct effects on the States, on the relationship between the National Government and the States, or on the distribution of power and responsibilities among the various levels of government. Accordingly, the agency tentatively concludes that the proposed rule does not contain policies that have federalism implications as defined in the Executive order and, consequently, a federalism summary impact statement has not been prepared.

XIII. Paperwork Reduction Act of 1995

FDA tentatively concludes that this proposed rule contains no new collections of information. Therefore, clearance by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA) (44 U.S.C. 3501–3520) is not required.

This proposed rule designates a guidance document as a special control. FDA also tentatively concludes that the draft special control guidance document does not contain new information collection provisions that are subject to review and clearance by OMB under the PRA. Elsewhere in this issue of the **Federal Register**, FDA is publishing a notice announcing the availability of that draft guidance document entitled ''Class II Special Controls Guidance Document: Tissue Adhesive for the Topical Approximation of Skin,'' which contains an analysis of the paperwork burden for the draft guidance.

XIV. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments regarding this document. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified 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.

XV. References

The following references have been placed on display in the Division of Dockets Management (see **ADDRESSES**) and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday.

1. Regulatory & Clinical Research Institute, Inc. (RCRI), reclassification petition, Docket No. 2006P–0071, Minneapolis MN, February 9, 2006.

2. Regulatory & Clinical Research Institute, Inc., reclassification petition, Docket No. 2006P–0071, Minneapolis MN, May 15, 2006.

3. Regulatory & Clinical Research Institute, Inc., reclassification petition, Docket No. 2006P–0071, Minneapolis MN, July 18, 2006.

4. General and Plastic Surgery Devices Panel, Transcript, pp. 199 to 207, August 25, 2006.

List of Subjects in 21 CFR Part 878

Medical devices.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, it is proposed that 21 CFR part 878 be amended as follows:

PART 878—GENERAL AND PLASTIC SURGERY DEVICES

1. The authority citation for 21 CFR part 878 continues to read as follows:

Authority: 21 U.S.C. 351, 360, 360c, 360e, 360j, 371.

2. Section 878.4010 is added to subpart E to read as follows:

§ 878.4010 Tissue adhesive.

(a) *Tissue adhesives for the topical approximation of skin*—(1) *Identification.* Tissue adhesives for the topical approximation of skin are intended for topical closure of surgical incisions, including laparoscopic incisions, and simple traumatic lacerations that have easily approximated skin edges. Tissue adhesives for the topical approximation of skin may be used in conjunction with, but not in place of, deep dermal stitches.

(2) *Classification.* Class II (special controls). The special control for this device isEFDA's ''Class II Special Controls Guidance Document: ''Tissue Adhesive for the Topical Approximation of Skin.'' See § 878.1(e) for the availability of this guidance document.

(b) *Tissue adhesives for non-topical use*—(1) *Identification.* A tissue adhesive for non-topical use, including adhesives intended for use in the embolization of brain arteriovenous malformation or for use in ophthalmic surgery, is a device used for adhesion of internal tissues and vessels.

(2) *Classification.* Class III (premarket approval). As of May 28, 1976, an approval under section 515 of the act is required before this device may be commercially distributed. See § 878.3.

Dated: June 22, 2007.

Linda S. Kahan,

Deputy Director, Center for Devices and Radiological Health.

[FR Doc. E7–12797 Filed 7–2–07; 8:45 am]

BILLING CODE 4160–01–S

ARCHITECTURAL AND TRANSPORTATION BARRIERS COMPLIANCE BOARD

36 CFR Parts 1193 and 1194

Telecommunications Act Accessibility Guidelines; Electronic and Information Technology Accessibility Standards

AGENCY: Architectural and Transportation Barriers Compliance Board.

ACTION: Notice of meeting.

SUMMARY: The Architectural and Transportation Barriers Compliance Board (Access Board) has established a Telecommunications and Electronic and Information Technology Advisory Committee (Committee) to assist it in revising and updating accessibility guidelines for telecommunications products and accessibility standards for electronic and information technology. This notice announces the dates, time, and location of the next committee meeting.

DATES: The meeting is scheduled for July 16–18, 2007 (beginning at 9 a.m. and ending at 5 p.m. each day).

ADDRESSES: The meeting will be held at the National Science Foundation. Report to the National Science Foundation, 4201 Wilson Boulevard, Arlington, VA 22230, to pick up security passes and then report to 4121 Wilson Boulevard, Stafford Place II, Room 555, Arlington, VA 22230 for the meeting.

FOR FURTHER INFORMATION CONTACT: Timothy Creagan, Office of Technical and Information Services, Architectural and Transportation Barriers Compliance Board, 1331 F Street, NW., suite 1000, Washington, DC 20004–1111. Telephone number: 202–272–0016 (Voice); 202–272–0082 (TTY). Electronic mail address: *creagan@access-board.gov.*

SUPPLEMENTARY INFORMATION: The Architectural and Transportation Barriers Compliance Board (Access Board) established the Telecommunications and Electronic and Information Technology Advisory Committee (Committee) to assist it in

revising and updating accessibility guidelines for telecommunications products and accessibility standards for electronic and information technology. The next committee meeting will take place on July 16–18, 2007. The meeting will focus on reports and discussion of recommendations from the following subcommittees:

- Software, Web, and Content
- General Interface Requirements and Functional Performance Criteria
- Computer Hardware
- Subpart A
- Documentation and Technical Support
- Telecommunications
- Audio/Visual
- Self Contained, Closed Products
- Editorial Working Group

The meeting will also discuss the status of the committee's work to date and when a final report may be ready for presentation to the Access Board. The full agenda for the July 16–18, 2007 meeting is available at *http://www.access-board.gov/sec508/refresh/agenda.htm.* Notices of future meetings will be published in the **Federal Register**.

Information about the committee, including future meeting dates is available at *http://www.access-board.gov/sec508/update-index.htm* or at a special Web site created for the committee's work (*http://teitac.org*). The site includes a calendar for subcommittee meetings, e-mail distribution lists, and a ''Wiki'' (*http://teitac.org/wiki/TEITAC_Wiki*) which provides interactive online work space.

Committee meetings are open to the public and interested persons can attend the meetings and communicate their views. Members of the public will have opportunities to address the committee on issues of interest to them and the committee during public comment periods scheduled on each day of the meeting. Members of groups or individuals who are not members of the committee are invited to participate on subcommittees; participation of this kind is very valuable to the advisory committee process.

The meeting site is accessible to individuals with disabilities. Sign language interpreters, an assistive listening system, and real-time captioning will be provided. For the comfort of other participants, persons attending committee meetings are requested to refrain from using perfume, cologne, and other fragrances. Due to security measures at the National Science Foundation, all attendees must notify the Access Board's receptionist at 202–272–0007 or *receptionist@access-board.gov* by July 11, 2007 of their intent to attend the meeting. This notification is required for expeditious entry into the facility and will enable the Access Board to provide additional information as needed.

Lawrence W. Roffee,

Executive Director.

[FR Doc. E7–12811 Filed 7–2–07; 8:45 am]

BILLING CODE 8150–01–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R03–OAR–2007–0451; FRL–8333–5]

Approval and Promulgation of Air Quality Implementation Plans; Delaware; Control of VOC Emissions from Crude Oil Lightering Operations

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: EPA is proposing to approve a State Implementation Plan (SIP) revision submitted by the State of Delaware. This SIP revision pertains to the control of volatile organic compound (VOC) emissions from crude oil lightering operations. This action is being taken under the Clean Air Act (CAA).

DATES: Written comments must be received on or before August 2, 2007.

ADDRESSES: Submit your comments, identified by Docket ID Number EPA–R03–OAR–2007–0451 by one of the following methods:

A. *http://www.regulations.gov.* Follow the on-line instructions for submitting comments.

B. *E-mail: cripps.christopher@epa.gov.*

C. *Mail*: EPA–R03–OAR–2007–0451, Christopher Cripps, Acting Chief, Air Quality Planning Branch, Mailcode 3AP21, U.S. Environmental Protection Agency, Region III, 1650 Arch Street, Philadelphia, Pennsylvania 19103.

D. *Hand Delivery*: At the previously-listed EPA Region III address. Such deliveries are only accepted during the Docket's normal hours of operation, and special arrangements should be made for deliveries of boxed information.

Instructions: Direct your comments to Docket ID No. EPA–R03–OAR–2007–0451. EPA's policy is that all comments received will be included in the public docket without change, and may be made available online at *www.regulations.gov*, including any personal information provided, unless the comment includes information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you consider to be CBI or otherwise protected through *www.regulations.gov* or e-mail. The *www.regulations.gov* Web site is an ''anonymous access'' system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an e-mail comment directly to EPA without going through *www.regulations.gov*, your e-mail address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the Internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD–ROM you submit. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses.

Docket: All documents in the electronic docket are listed in the *www.regulations.gov* index. Although listed in the index, some information is not publicly available *i.e.*, CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either electronically in *www.regulations.gov* or in hard copy during normal business hours at the Air Protection Division, U.S. Environmental Protection Agency, Region III, 1650 Arch Street, Philadelphia, Pennsylvania 19103. Copies of the State submittal are available at the Delaware Department of Natural Resources & Environmental Control, 89 Kings Highway, P.O. Box 1401, Dover, Delaware 19901.

FOR FURTHER INFORMATION CONTACT: Rose Quinto, (215) 814–2182, or by e-mail at *quinto.rose@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

On May 2, 2007, the Delaware Department of Natural Resources and Environmental Control (DNREC) submitted a revision to its SIP for Regulation No. 1124, Section 46—Control of VOC Emissions from Crude Oil Lightering Operations. Lightering is the transfer at anchorage for some of the