U.S.C. 3507 (d)), The Department of Health and Human Services is submitting a copy of the revised State plan preprint page to the Office of Management and Budget (OMB) for its review.

The State plan preprint and amendments serve as a contract with OCSE in outlining the activities the States will perform as required by law in order for States to receive federal funds to meet the costs of these activities. We are asking for approval of the revised State plan preprint page for Periodic Reporting to Consumer Reporting Agencies to reflect new Federal requirements. Procedures to Improve Program Effectiveness, is amended by adding a new section 7, Periodic Reporting to Consumer Reporting Agencies, which requires the State to have procedures, (1) to periodically report information regarding the amount of overdue support owed by an absent parent to consumer reporting agencies when such amount exceeds $1,000 and is at least two months in arrears in accordance with section 666(a)(7) of the Act; and 92) for making absent parent information available to Consumer Reporting Agencies in accordance with Sec. 302.70(d). The information collected on the State plan pages is necessary to enable OCSE to monitor compliance with the requirements in Title IV–D of the Social Security Act and implementing regulations.

Respondents: State governments.

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Plan ..... Estimated Total Annual Burden Hours: 1,316.	54	1836	.717	1,316

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer, Larry Guerrero.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street NW., Washington, DC 20503, Attn: Ms. Wendy Taylor.

Dated: April 25, 1997.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 97–11234 Filed 4–30–97; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97C–0171]

Closure Medical Corp.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Closure Medical Corp. filed a petition proposing that the color additive regulations be amended to provide for the safe use of D&C Violet No. 2 to color 2-octyl cyanoacrylate topical tissue adhesives.

DATES: Written comments on the petitioner's environmental assessment by June 2, 1997.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Ellen M. Waldron, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3089.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 721(d)(1) (21 U.S.C. 379e(d)(1)), notice is given that a color additive petition (CAP 7C0250) has been filed by Closure Medical Corp., 5265 Capital Blvd., Raleigh, NC 27616. The petition proposes to amend the color additive regulations in § 74.3602 *D&C Violet No. 2* (21 CFR 74.3602) to provide for the safe use of D&C Violet No. 2 to color 2-octyl cyanoacrylate topical tissue adhesives.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before June 2, 1997, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one 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 office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: April 16, 1997.

Alan M. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 97–11238 Filed 4–30–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA–R–0097]

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration