

II. Significance of the Guidance

This draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the agency's current thinking on oxygen pressure regulators and oxygen conserving devices with a built-in oxygen pressure regulator. It does not create or confer any rights for or on any person and would not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute and regulations.

III. Electronic Access

Persons interested in obtaining a copy of the draft guidance may do so by using the Internet. To receive "Class II Special Controls Guidance Document: Pressure Regulators For Use With Medical Oxygen and Oxygen Conserving Devices," you may either send an e-mail request to dsmica@fda.hhs.gov to receive an electronic copy of the document, or send a fax request to 240-276-3151 to receive a hard copy. Please use the document number 1227 to identify the guidance you are requesting.

CDRH maintains an entry on the Internet for easy access to information including text, graphics, and files that may be downloaded to a personal computer with Internet access. Updated on a regular basis, the CDRH home page includes device safety alerts, **Federal Register** reprints, information on premarket submissions (including lists of approved applications and manufacturers' addresses), small manufacturer's assistance, information on video conferencing and electronic submissions, Mammography Matters, and other device-oriented information. The CDRH Web site may be accessed at <http://www.fda.gov/cdrh>. A search capability for all CDRH guidance documents is available at <http://www.fda.gov/cdrh/guidance.html>. Guidance documents are also available on the Division of Dockets Management Internet site at <http://www.fda.gov/ohrms/dockets>.

IV. Paperwork Reduction Act of 1995

This draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA) (44 U.S.C. 3501-3520). The collections of information in 21 CFR part 807, subpart E (premarket notification procedures) have been

approved under OMB Control number 0910-0120. The labeling statements that would be required by this regulation are "public disclosure[s] of information originally supplied by the Federal government to the recipient for the purpose of disclosure to the public * * *" (5 CFR 1320.3(c)(2)).

Accordingly, FDA concludes that the labeling requirements in this proposed rule are not subject to review by OMB under the PRA.

V. 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. The draft guidance and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: February 8, 2007.

Linda S. Kahan,

Deputy Director, Center for Devices and Radiological Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104-13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to Office of Management and Budget under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443-1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including

whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: The Nurse Faculty Loan Program (NFLP): Annual Operating Report (AOR) Form—NEW

Under Title VIII of the Public Health Service Act, as amended by Public Law 107-205, Section 846A, the Secretary of Health and Human Services (HHS) enters into an agreement with a school of nursing to establish and operate the NFLP fund. HHS makes an award to the school in the form of a Federal Capital Contribution (FCC). The award is used to establish a distinct account for the NFLP loan fund at the school or is deposited into an existing NFLP fund. The school of nursing makes loans from the NFLP fund to eligible students enrolled full-time in a master's or doctoral nursing education program that will prepare them to become qualified nursing faculty. Following graduation from the NFLP lending school, loan recipients may receive up to 85 percent NFLP loan cancellation over a consecutive four-year period in exchange for service as full-time faculty at a school of nursing. The NFLP lending school collects any portion of the loan that is not cancelled and any loan in repayment and deposits these monies into the NFLP loan fund to make additional NFLP loans. The school of nursing must keep records of all NFLP loan fund transactions.

The NFLP Annual Operating Report will be used to collect information relating to the NFLP loan fund operations and financial activities for a specified reporting period (July 1 through June 30 of the academic year). Participating schools will complete and submit the AOR annually to provide the Federal Government with current and cumulative information on: (1) The number and amount of loans made, (2) the number of NFLP recipients and graduates, (3) the number and amount of loans collected, (4) the number and amount of loans in repayment, (5) the number of NFLP graduates employed as nurse faculty, (6) NFLP loan fund receipts, disbursements and other related cost. The NFLP loan fund balance is used to determine future awards to the school.

The estimate of burden for this form is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per responses	Total burden hours
Nurse Faculty Loan Program Annual Operating Report (AOR)	150	1	150	8	1200
Total Burden	150	1	150	8	1200

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 10-33, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: February 21, 2007.

Alexandra Huttinger,
Acting Director, Division of Policy Review and Coordination.

[FR Doc. E7-3406 Filed 2-26-07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995 Pub. L. 104-13), the Health Resources and Services Administration (HRSA) will

publish periodic summaries of proposed projects being developed for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain the data collection plans, call HRSA Reports Clearance Office on (301) 443-1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Scholarships for Disadvantaged Students Program—(OMB No. 0915-0149) Application—Reinstatement

The Scholarships for Disadvantaged Students (SDS) Program has as its

purpose the provision of funds to eligible schools to provide scholarships to full-time, financially needy students from disadvantaged backgrounds enrolled in health professions and nursing programs.

To qualify for participation in the SDS program, a school must be carrying out a program for recruiting and retaining students from disadvantaged backgrounds, including students who are members of racial and ethnic minority groups (section 737(d)(1)(B) of the PHS Act). A school must meet the eligibility criteria to demonstrate that the program has achieved success based on the number and/or percentage of disadvantaged students who graduate from the school. In awarding SDS funds to eligible schools, funding priorities must be given to schools based on the proportion of graduating students going into primary care, the proportion of underrepresented minority students, and the proportion of graduates working in medically underserved communities (section 737(c) of the PHS Act).

The estimated response burden is as follows:

Form	Number of respondents	Responses per respondent	Hours per response	Total hour burden
Application	500	1	20	10,000
Report	500	1	1	1,500
Total	500	1	21	11,500

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 10-33, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Written comments should be received within 60 days of this notice.

Dated: February 21, 2007.

Alexandra Huttinger,
Acting Director, Division of Policy Review and Coordination.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Notice of Availability of Draft Policy Document for Comment

AGENCY: Health Resources and Services Administration (HRSA), HHS.

ACTION: Notice of availability and solicitation of comments.

SUMMARY: This is a Notice of Availability and request for comments on the draft policy document, "Emergency Management Program Expectations," prepared by HRSA. This document is currently posted on the Internet at <http://bphc.hrsa.gov>. Comments will be reviewed and analyzed, and a summary and general response will be published as soon as possible after the deadline for receipt of comments.