recommendations and vote on a premarket approval application, sponsored by SyntheMed, Inc., for the REPEL-CV, which is a surgical adjuvant indicated for reducing the incidence, severity and extent of post-operative adhesion formation in patients undergoing cardiac surgery.

On September 20, 2007, the committee will discuss and make recommendations regarding clinical trial designs for cardiac ablation devices designed to treat patients with medically refractory atrial fibrillation.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/ohrms/dockets/ac/acmenu.htm*, click on the year 2007 and scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before September 5, 2007. Oral presentations from the public will be scheduled for approximately 30 minutes at the beginning of committee deliberations on each day and for approximately 30 minutes near the end of the deliberations on each day. Those desiring to make formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before August 28, 2007. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by August 29, 2007.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, at 240–276–8932, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: July 23, 2007.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E7–14600 Filed 7–26–07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: The National Health Service Corps Loan Repayment Program (OMB No. 0915–0127)—Extension

The National Health Service Corps (NHSC) Loan Repayment Program (LRP) was established to assure an adequate supply of trained primary care health care professionals to provide services in the neediest Health Professional Shortage Areas (HPSAs) of the United States. Under this program, the Department of Health and Human Services agrees to repay the educational loans of the primary care health professionals. In return, the health professionals agree to serve for a specified period of time in a federally-designated HPSA approved by the Secretary for LRP participants.

The NHSC LRP forms provide information that is needed for selecting participants and making determinations regarding repayment of qualifying loans for education. The LRP forms include the following: The NHSC LRP Application; the Loan Information and Verification form; the Community Site Information form; the Applicant Checklist; the Payment Information form; and the Authorization to Release Information form.

The estimated annual burden is as follows:

Type of form	Number of respondents	Responses per respondent	Total number of responses	Hours per response	Total burden hours
NHSC LRP Application	1920	1	1920	.5	960
Community Site Information form	1920	1	1920	.25	480
Loan Information and Verification form	1920	3	5760	.25	1440
Authorization to Release Information	1920	1	1920	.1	192
Applicant Checklist	1920	1	1920	.2	384
Lenders	80	1	80	.25	20
Total	2000	.....	9680	1.55	3476

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the ''attention of the desk officer for HRSA.''

Dated: July 5, 2007.

Alexandra Huttinger,

Acting Director, Division of Policy Review and Coordination.

[FR Doc. E7–14525 Filed 7–26–07; 8:45 am]

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