Accordingly, pursuant to section 564(g)(2) of the Act, FDA revokes the EUA issued on April 28, 2020.

Notice of this revocation will be published in the Federal Register, pursuant to section 564(h)(1) of the Act.

Sincerely,

/s/

RADM Denise M. Hinton
Chief Scientist
Food and Drug Administration


Lauren K. Roth,
Associate Commissioner for Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting; Postponement

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of meeting: postponement.

SUMMARY: The meeting of the Circulatory System Devices Panel of the Medical Devices Advisory Committee was originally announced in the Federal Register of August 20, 2020 (85 FR 51453), and was initially scheduled for October 7, 2020. FDA has decided to postpone this public meeting until further notice.


Lauren K. Roth,
Associate Commissioner for Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval: Public Comment Request; Information Collection Request Title: Application for Deemed Health Center Program Award Recipients To Sponsor Volunteer Health Professionals for Deemed PHS Employment

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with of the Paperwork Reduction Act of 1995, HRSA has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period. OMB may act on HRSA’s ICR only after the 30 day comment period for this notice has closed.

DATES: Comments on this ICR must be received no later than November 4, 2020.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443–1984.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the information request collection title for reference.

Information Collection Request Title: Application for Deemed Health Center Program Award Recipients To Sponsor Volunteer Health Professionals for Deemed PHS Employment

Agency: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

SUMMARY: The Food and Drug Administration (FDA) is postponing the meeting of the Circulatory System Devices Panel of the Medical Devices Advisory Committee scheduled from October 7, 2020, to a later date to be determined. The meeting was announced in the Federal Register of August 20, 2020. A future meeting date will be announced in the Federal Register.

FOR FURTHER INFORMATION CONTACT: Aden Asefa, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5214, Silver Spring, MD 20993–0002, Aden.Asefa@fda.hhs.gov, 301–796–0400, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area) and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting.