in 5 U.S.C. App. 2, section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). Grant applications for the ''AHRQ RFAHS20–003 Novel, High-Impact Studies Evaluating Health System and Healthcare Professional Responsiveness to COVID–19 (R01).'' are to be reviewed and discussed at this meeting. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: July 17, 2020.

Virginia L. Mackay-Smith,

Associate Director.

[FR Doc. 2020–15936 Filed 7–22–20; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) announces a Special Emphasis Panel (SEP) meeting on ''HCRT SEP 2020/10 ZHS1 HSR A (01).'' This SEP meeting will be closed to the public.

DATES: August 14, 2020.

ADDRESSES: Agency for Healthcare Research and Quality, (Video Assisted Review), 5600 Fishers Lane, Rockville, Maryland 20850.

FOR FURTHER INFORMATION CONTACT: Jenny Griffith, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, Agency for Healthcare Research and Quality, (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20850, Telephone: (301) 427–1557.

SUPPLEMENTARY INFORMATION: A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by AHRQ, and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

The SEP meeting referenced above will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2, section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). Grant applications for the ''HCRT SEP 2020/10 ZHS1 HSR A (01).'' are to be reviewed and discussed at this meeting. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: July 17, 2020.

Virginia L. Mackay-Smith,

Associate Director.

[FR Doc. 2020–15935 Filed 7–22–20; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

AGENCY: Agency for Healthcare Research and Quality, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) announces a Special Emphasis Panel (SEP) meeting on ''HSQR ZHS1 HSR X-(01).'' This SEP meeting will be closed to the public.

DATES: August 19, 2020.

ADDRESSES: Agency for Healthcare Research and Quality, (Video Assisted Review), 5600 Fishers Lane, Rockville, Maryland 20850.

FOR FURTHER INFORMATION CONTACT: Jenny Griffith, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, Agency for Healthcare Research and Quality, (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20850, Telephone: (301) 427–1557.

SUPPLEMENTARY INFORMATION: A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the AHRQ, and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

The SEP meeting referenced above will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2, section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). Grant applications for the ''AHRQ-HSQR ZHS1 HSR X-(01).'' are to be reviewed and discussed at this meeting. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy. Agenda items for this meeting are subject to change as priorities dictate.

Dated: July 17, 2020.

Virginia L. Mackay-Smith,

Associate Director.

[FR Doc. 2020–15934 Filed 7–22–20; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Health Statistics (NCHS), ICD–10 Coordination and Maintenance (C&M) Committee Meeting

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: The CDC, National Center for Health Statistics (NCHS), Classifications and Public Health Data Standards Staff, announces the following meeting of the ICD–10 Coordination and Maintenance (C&M) Committee meeting. This meeting is open to the public, limited only by audio and web conferences lines available. Online Registration is not required.

DATES: The meeting will be held on September 8, 2020, from 9:00 a.m. to 5:00 p.m., EDT, and September 9, 2020, from 9:00 a.m. to 5:00 p.m., EDT.

ADDRESSES: This is a virtual meeting. Information will be provided on each of our respective web pages when it becomes available. For CDC/NCHS *https://www.cdc.gov/nchs/icd/icd10cm_maintenance.htm.*

For CMS *https://www.cms.gov/Medicare/Coding/ICD9ProviderDiagnosticCodes/meetings.*

FOR FURTHER INFORMATION CONTACT: Traci Ramirez, Program Specialist, CDC, 3311 Toledo Road, Hyattsville, Maryland 20782, Telephone (301) 458–4454; *TRamirez@cdc.gov.*

SUPPLEMENTARY INFORMATION: *Purpose:* The ICD–10 Coordination and Maintenance (C&M) Committee is a public forum for the presentation of proposed modifications to the International Classification of Diseases, Tenth Revision, Clinical Modification and ICD–10 Procedure Coding System.

Matters To Be Considered: The tentative agenda will include discussions on ICD–10–CM and ICD–10–PCS topics listed below. Agenda items are subject to change as priorities dictate.

Please refer to the posted agenda for updates one month prior to the meeting.

ICD–10–PCS Topics

Vertebral Body Tethering
Removal of a Transplanted/Rejected Kidney
Isotope Administration
Administration of Lifileucel
Administration of Narsoplimab
Insertion of Implantable Bone Void Filler
Single-Use Duodenoscope
Administration of Immune Effector Cell Therapy
Spinal Stabilization
Administration of Idecabtagene Vicleucel (ide-cel)
Restriction of Coronary Sinus Embolic Protection

ICD–10–CM Topics:

Complications of immune effector cellular (IEC) therapy
Endometriosis
Immune Effector Cell Associated Neurotoxicity Syndrome (ICANS)
Addenda

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2020–16000 Filed 7–22–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0490]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Cosmetic Labeling Regulations and Voluntary Cosmetic Registration Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by August 24, 2020.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://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. The OMB control number for this information collection is 0910–0599. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Cosmetic Labeling Regulations—21 CFR part 701 and Voluntary Cosmetic Registration Program—21 CFR parts 710 and 720

OMB Control Number 0910–0599—Revision

The Federal Food, Drug, and Cosmetic Act (FD&C Act) and the Fair Packaging and Labeling Act (FPLA) require that cosmetic manufacturers, packers, and distributors disclose information about themselves or their products on the labels or labeling of their products. Sections 201, 301, 502, 601, 602, 603, 701, and 704 of the FD&C Act (21 U.S.C. 321, 331, 352, 361, 362, 363, 371, and 374) and sections 4 and 5 of the FPLA (15 U.S.C. 1453 and 1454) provide authority to FDA to regulate the labeling of cosmetic products. Failure to comply with the requirements for cosmetic labeling may render a cosmetic adulterated under section 601 of the FD&C Act or misbranded under section 602 of the FD&C Act.

I. Cosmetic Labeling Regulations

FDA's cosmetic labeling regulations are codified in part 701 (21 CFR part 701). Section 701.3 requires the label of a cosmetic product to bear a declaration of the ingredients in descending order of predominance. Section 701.11 requires the principal display panel of a cosmetic product to bear a statement of the identity of the product. Section 701.12 requires the label of a cosmetic product to specify the name and place of business of the manufacturer, packer, or distributor. Section 701.13 requires the label of a cosmetic product to declare the net quantity of contents of the product. The information collection provisions found in part 701 are currently approved under OMB control number 0910–0027. To improve the efficiency of Agency operations, we are consolidating these information collection elements into OMB control number 0910–0599.

II. Voluntary Cosmetic Registration Program

Information collection associated with our Voluntary Cosmetic Registration Program (VCRP) are found in parts 710 and 720 (21 CFR parts 710 and 720). Participants have the option of submitting information via paper forms or via an online interface. The use of the term ''form'' refers to both the paper form and the online system.

Pursuant to part 710, we request that establishments that manufacture or package cosmetic products voluntarily register with us using Form FDA 2511 entitled ''Registration of Cosmetic Product Establishment.'' The online version of Form FDA 2511 is available on our VCRP website at *https://www.fda.gov/cosmetics/voluntary-cosmetic-registration-program/online-registration-voluntary-cosmetic-registration-program-vcrp.* We encourage online registration of Form FDA 2511 because it is faster and more efficient for the filer and the Agency. A registering facility will receive confirmation of online registration, including a registration number by