notice that the oncology drugs at issue were unapproved and that the medical practice discontinued ordering those drugs when he learned of that regulatory status, as discussed above, it is undisputed that the offense to which he pled guilty led to his administering foreign, unapproved drug products to his patients. Even assuming Dr. Tahsildar's representations with respect to his reduced role as a manager in the practice to be true, the Chief Scientist also cannot conclude that his managerial role is a favorable consideration, given his status as a partner and a physician in that practice. Balancing the applicable considerations—including his voluntary steps in mitigation under section 306(c)(3)(C) of the FD&C Act and the absence of previous criminal convictions related to matters within the jurisdiction of FDA under section 306(c)(3)(F)—the Chief Scientist has determined that a 2-year debarment period is appropriate. Inasmuch as there are no material factual disputes for resolution at a hearing, the Chief Scientist is also denying Dr. Tahsildar's hearing request.

Separately, Dr. Tahsildar requests that, in lieu of debarment by FDA, he enter into a settlement agreement with FDA whereby he would voluntarily agree to the terms of the proposed debarment for the proposed period of debarment and to not provide services in any capacity to a person that has an approved or pending drug product application. Dr. Tahsildar appears to be proposing an informal resolution of this debarment matter. However, his request is now moot given that the foregoing findings support debarment for a 2-year period.

III. Findings and Order

Therefore, the Chief Scientist, under section 306(b)(2)(B)(i)(I) of the FD&C Act and authority delegated to her by the Commissioner of Food and Drugs, finds that Dr. Tahsildar has been convicted of a misdemeanor under Federal law for conduct related to the regulation of drugs under the FD&C Act and that the type of conduct underlying the conviction undermines the regulation of drugs. FDA has considered the relevant factors listed in section 306(c)(3) of the FD&C Act and determined that a 2-year debarment is appropriate.

As a result of the foregoing findings, Dr. Tahsildar is debarred for 2 years from providing services in any capacity to a person with an approved or pending drug product application under sections 505, 512, or 802 of the FD&C Act (21 U.S.C. 355, 360b, or 382), or under section 351 of the Public Health Service Act (42 U.S.C. 262), effective February 8, 2023, (see 21 U.S.C. 335a(c)(1)(B) and (c)(2)(A)(iii) and 21 U.S.C. 321(dd))). Any person with an approved or pending drug application who knowingly uses the services of Dr. Tahsildar, in any capacity during his debarment, will be subject to civil money penalties (section 307(a)(6) of the FD&C Act (21 U.S.C. 335b(a)(6))). If Dr. Tahsildar, during his period of debarment, provides services in any capacity to a person with an approved or pending drug product application, he will be subject to civil money penalties (section 307(a)(7) of the FD&C Act). In addition, FDA will not accept or review any abbreviated new drug applications submitted by or with the assistance of Dr. Tahsildar during his period of debarment (section 306(c)(1)(B) of the FD&C Act).

Dated: February 2, 2023.

Namandjé N. Bumpus,

Chief Scientist.

[FR Doc. 2023–02634 Filed 2–7–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

[OMB No. 0915–0318—Revision]

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request; Ryan White HIV/AIDS Program: Allocations Forms

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 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 should be received no later than March 10, 2023.

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 Samantha Miller, the HRSA Information Collection Clearance Officer, at *paperwork@hrsa.gov* or call 301–594–4394.

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

Information Collection Request Title: Ryan White HIV/AIDS Program: Allocations Forms, OMB No. 0915–0318—Revision.

Abstract: HRSA administers the Ryan White HIV/AIDS Program (RWHAP) authorized under Title XXVI of the Public Health Service Act. The RWHAP Allocations and Expenditures Reports (A&E Reports) allow HRSA to monitor and track the use of grant funds for compliance with program and grants policies, and requirements as outlined in the legislation. To avoid duplication and reduce recipient reporting burden, HRSA created an electronic grantee contract management system (GCMS) that includes data required for various reports, including the Allocations Reports and other HRSA data reports, such as the RWHAP Services Report. Recipients can access GCMS year-round to upload or manually enter data on their service provider contractors or subrecipients, the RWHAP core medical and support services provided, and their funding amounts. Data required for Allocations Reports and other reports are automatically prepopulated from GCMS. Expenditures Report data are not auto-populated in the GCMS, and are still manually entered into the data reporting system.

Allocations and Expenditures (A&E) Reports

Recipients funded under RWHAP Parts A, B, C, and D are required to report financial data to HRSA at the beginning (Allocations Report) and at the end (Expenditures Report) of their grant budget period. The A&E Reports request information recipients already collect, including the use of RWHAP grant funds for core medical and support services; and on various program components, such as administration, planning and evaluation, and clinical quality management (CQM). RWHAP Parts A and B recipients funded under the

Ending the HIV Epidemic in the U.S. (EHE) initiative are also required to report allocations and expenditures of the grant budget period in the EHE A&E Reports. This allows HRSA to track and report progress toward meeting the EHE goals.

The reports are similar in content; however, in the first report, recipients document the allocation of their RWHAP or EHE grant award at the beginning of their grant budget period. In the second report, recipients document actual expenditures of their RWHAP or EHE grant award (including any carryover dollars) at the end of their grant budget period.

HRSA proposes the following updates to the RWHAP Allocations Reports.

RWHAP Part A Allocations Report

- Revising row and column headers and other language for clarity and alignment with RWHAP requirements;
- Combining the columns for RWHAP Part A Formula and Supplemental Allocation amounts and updating the title;
- Moving the RWHAP Part A Minority AIDS Initiative (MAI) Award Amount row after the RWHAP Part A Supplemental Award Amount row;
- Changing the calculation for Service Allocation Subtotal percent in the Total RWHAP Part A Allocation Amounts column;
- Blacking out the percent columns for the RWHAP Part A Formula and Supplemental Allocation Amounts, RWHAP Part A MAI Allocation Amounts, and selected cells in the Total RWHAP Part A Allocation Amounts column; and
- Adding the Legislative Requirements Checklist.

RWHAP Part B Allocations Report

- Revising row and column headers and other language for clarity and alignment with RWHAP requirements;
- Adding the following rows to Table 1: 4c. Part B HIV Care Consortia Planning & Evaluation/Emerging Communities (EC) HIV Care Consortia Planning & Evaluation and 4d. Part B HIV Care Consortia CQM/EC HIV Care Consortia CQM except for the AIDS Drug Assistance Program (ADAP) Earmark + ADAP Supplemental Award cells;
- Removing row 11. Total Part B X07 Allocations;
- Allowing users to enter data in Table 2 for 1d. Health Insurance Premium & Cost Sharing and 1e. Home and Community-based Health Services;
- Blacking out selected cells in the following rows, columns, or tables:
  - 2. Part B Health Insurance Premium & Cost Sharing Assistance for Low-Income Individuals (Table 1) as this information is also reported in Table 2
  - 3. Part B Home and Community-based Health Services (Table 1) as this information is also reported in Table 2
  - 4. Total Column (Table 1)
  - 1a. ADAP Treatments (Table 2) as this information is also reported in Table 1
  - MAI Award (Table 3)
- Updating calculations and language in the Legislative Requirements Checklist; and
- Removing the following services under the Legislative Requirements Checklist's Core Medical Services:
  - Health Insurance Premium & Cost Sharing Assistance, and
  - Home and Community-based Health Services.

RWHAP Part C Allocations Report

- There are no proposed changes to the RWHAP Part C Allocations Report.

RWHAP Part D Allocations Report

- There are no proposed changes to the RWHAP Part D Allocations Report.

HRSA EHE Initiative A&E Reports

- There are no proposed changes to the HRSA EHE Allocations Reports. A 60-day notice published in the **Federal Register**, 87 FR pp. 71339–40 (November 22, 2022). There were no public comments.

Need and Proposed Use of the Information: Accurate allocation, expenditure, and service contract records of the recipients receiving RWHAP and EHE funding are critical to the implementation of the RWHAP legislation and EHE initiative appropriation language and thus are necessary for HRSA to fulfill its monitoring and oversight responsibilities.

Likely Respondents: RWHAP Part A, Part B, Part C, and Part D recipients.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Part A Allocations Report	52	1	52	4	208
Part B Allocations Report	54	1	54	6	324
Part C Allocations Report	346	1	346	4	1,384
Part D Allocations Report	116	1	116	4	464
EHE Allocations Reports	47	1	47	4	188
Total	615				2,568

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information

technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2023–02686 Filed 2–7–23; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Addiction Risks and Mechanisms Study Section, February 23, 2023, 9:00 a.m. to February 24, 2023, 7:30 p.m., Darcy Hotel, 1515 Rhode Island Avenue, Washington, DC 20005 which was published in the **Federal Register** on January 26, 2023, 88 FR 5010.

This meeting is being amended to change the name of the hotel from Darcy Hotel, 1515 Rhode Island Avenue, Washington, DC 20005 to the Hotel George, 15 E Street NW, Washington, DC 20001. The meeting is closed to the public.

Dated: February 3, 2023.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–02675 Filed 2–7–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. 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.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Cell Biology, Developmental Biology, and Bioengineering.

Date: March 7–8, 2023.

Time: 8:30 a.m. to 7:30 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, 7400 Wisconsin Ave., Bethesda, MD 20814.

Contact Person: Alexander Gubin, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4196, MSC 7812, Bethesda, MD 20892, 301–435–2902, *gubina@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Topics in Health Services Research.

Date: March 7–8, 2023.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Mary Kate Baker, DRPH, Scientific Review Officer, The Center for Scientific Review, The National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, 301–594–5117, *katie.baker2@nih.gov.*

Name of Committee: Center for Scientific Review, Special Emphasis Panel; Special: Modern Equipment for Shared-use Biomedical Research Facilities.

Date: March 9–10, 2023.

Time: 9:00 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Frederique Yiannikouris, Ph.D., Scientific Review Officer, The Center for Scientific Review, The National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, 301–594–3313, *frederique.yiannikouris@nih.gov.*

Name of Committee: Center for Scientific Review, Special Emphasis Panel; Therapeutic Immune Regulation.

Date: March 9–10, 2023.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Yue Wu, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 803C, Bethesda, MD 20892, (301) 867–5309, *wuy25@csr.nih.gov.*

Name of Committee: Center for Scientific Review, Special Emphasis Panel; Fellowships: Epidemiology and Population Sciences.

Date: March 9–10, 2023.

Time: 9:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Canopy by Hilton, 940 Rose Avenue, North Bethesda, MD 20852.

Contact Person: Erin Harrell, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 1000G, Bethesda, MD 20892, (301) 594–4935, *harreller@csr.nih.gov.*

Name of Committee: Center for Scientific Review, Special Emphasis Panel; Small Business: The Cardiovascular and Hematological Sciences.

Date: March 9–10, 2023.

Time: 9:00 a.m. to 9:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Dmitri V Gnatenko, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 867–5309, *gnatenkod2@nih.gov.*

Name of Committee: Center for Scientific Review, Special Emphasis Panel; Fellowships: Brain Disorders and Related Neurosciences.

Date: March 9–10, 2023.

Time: 9:30 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Vilen A Movsesyan, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4040M, MSC 7806, Bethesda, MD 20892, 301–402–7278, *movsesyanv@csr.nih.gov.*

Name of Committee: Molecular, Cellular and Developmental Neuroscience Integrated Review Group; Cellular and Molecular Biology of Neurodegeneration Study Section.

Date: March 9–10, 2023.

Time: 9:30 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Laurent Taupenot, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4188, MSC 7850, Bethesda, MD 20892, 301–435–1203, *laurent.taupenot@nih.gov.*

Name of Committee: Biological Chemistry and Macromolecular Biophysics Integrated Review Group; Maximizing Investigators' Research Award B Study Section.

Date: March 9–10, 2023.

Time: 9:30 a.m. to 8:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Sudha Veeraraghavan, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4166, MSC 7846, Bethesda, MD 20892, (301) 827–5263, *sudha.veeraraghavan@nih.gov.*

Name of Committee: Risk, Prevention and Health Behavior Integrated Review Group; HIV/AIDS Intra- and Inter-personal Determinants and Behavioral Interventions Study Section.

Date: March 9–10, 2023.

Time: 10:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.