information in 21 CFR 310.305(c) and 21 CFR 314.80(c)(2)(iii) and (e) for submitting postmarketing safety reports have been approved under OMB control number 0910–0230. The collections of information for submitting postmarketing safety reports under MedWatch have been approved under OMB control number 0910–0291. The collections of information in 21 CFR 201.56 and 201.57, including 21 CFR 201.57(c)(9)(i)(A) for preparing human prescription drug labeling to include pregnancy registries and relevant contact information under the subheading Pregnancy Exposure Registry have been approved under OMB control number 0910–0572. The collections of information contained in the guidance for clinical trial sponsors entitled ''Establishment and Operation of Clinical Trial Data Monitoring Committees'' (available at *https://www.fda.gov/downloads/RegulatoryInformation/Guidances/ucm127073.pdf*) have been approved under OMB control number 0910–0581. The collections of information in the ''Content and Format of Labeling for Human Prescription Drug and Biological Products; Requirements for Pregnancy and Lactation Labeling'' final rule have been approved under OMB control number 0910–0624. The collections of information in 21 CFR 50.25 for the elements of informed consent have been approved under OMB control number 0910–0755.

III. Electronic Access

Persons with access to the internet may obtain the document at *https://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm, https://www.fda.gov/BiologicsBloodVaccines/GuidanceComplianceRegulatoryInformation/default.htm,* or *https://www.regulations.gov.*

Dated: May 3, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Enrollment and Re-Certification of Entities in the 340B Drug Pricing Program, OMB Number 0915–0327—Revision

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

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Before submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than July 8, 2019.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 14N136B, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at (301) 443–1984.

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

Information Collection Request Title: Enrollment and Re-Certification of Entities in the 340B Drug Pricing Program, OMB No. 0915–0327—Revision.

Abstract: Section 602 of Public Law 102–585, the Veterans Health Care Act of 1992, enacted section 340B of the Public Health Service (PHS) Act, which instructs HHS to enter into a Pharmaceutical Pricing Agreement (PPA) with manufacturers of covered outpatient drugs. Manufacturers are required by section 1927(a)(5)(A) of the Social Security Act to enter into agreements with the Secretary of HHS (the Secretary) that comply with section 340B of the PHS Act if they participate in the Medicaid Drug Rebate Program. When a drug manufacturer signs a PPA, it is opting into the 340B Drug Pricing Program (340B Program), and it agrees to the statutory requirement that prices charged for covered outpatient drugs to covered entities will not exceed statutorily defined 340B ceiling prices. When an eligible covered entity voluntarily decides to enroll and participate in the 340B Program, it accepts responsibility for ensuring compliance with all provisions of the 340B Program, including all associated costs. Covered entities that choose to participate in the 340B Program must comply with the requirements of section 340B(a)(5) of the PHS Act. Section 340B(a)(5)(A) of the PHS Act prohibits a covered entity from accepting a discount for a drug that would also generate a Medicaid rebate. Further, section 340B(a)(5)(B) of the PHS Act prohibits a covered entity from reselling or otherwise transferring a discounted drug to a person who is not a patient of the covered entity.

Need and Proposed Use of the Information: To ensure its ongoing responsibility to administer the 340B Program while maintaining efficiency, transparency, and integrity, HRSA developed a process of registration for covered entities to address specific statutory mandates. Specifically, section 340B(a)(9) of the PHS Act requires HRSA to notify manufacturers of the identities of covered entities and of their status pertaining to certification and annual recertification in the 340B Program pursuant to section 340B(a)(7) and the establishment of a mechanism to prevent duplicate discounts as outlined at section 340B(a)(5)(A)(ii) of the PHS Act.

Also, section 340B(a)(1) of the PHS Act requires each participating manufacturer to enter into an agreement with the Secretary to offer covered outpatient drugs to 340B covered entities.

Finally, section 340B(d)(1)(B)(i) of the PHS Act requires the development of a system to enable the Secretary to verify the accuracy of ceiling prices calculated by manufacturers under subsection (a)(1) and charged to covered entities.

HRSA is requesting approval for existing information collections. HRSA notes that the previously approved collections are mostly unchanged, except that HRSA has transitioned completely to online versus hardcopy forms. In doing so, some of the forms have been revised to increase program efficiency and integrity. Below are descriptions of each of the forms and any resulting revisions in both the registration and pricing component of

the 340B Office of Pharmacy Affairs Information System (OPAIS).

Enrollment/Registration

To enroll and certify the eligible federally funded grantees and other safety net health care providers, HRSA requires entities to submit administrative information (*e.g.,* shipping and billing arrangements, Medicaid participation), certifying information (*e.g.,* Medicare Cost Report information, documentation supporting the hospital's selected classification), and attestation from appropriate grantee level or entity level authorizing officials and primary contacts. The purpose of this registration information is to determine eligibility for the 340B Program. To maintain accurate records, HRSA requests entities to submit modifications to any administrative information they submitted when initially enrolling in the Program. 340B covered entities have an ongoing responsibility to immediately notify HRSA of any change in eligibility for the 340B Program. No less than on an annual basis, entities must certify the accuracy of the information provided and continued maintenance of their eligibility and comply with statutory mandates of the Program.

Registration and annual recertification information are entered into the 340B OPAIS by entities and verified by HRSA staff according to 340B Program requirements. The following forms are being revised:

1. 340B Program Registrations & Certifications for Hospitals (applies to all hospital types): With the launch of 340B OPAIS in September 2017, HRSA removed the requirement for a Government Official to attest to the hospital classification of a parent hospital. HRSA would like to require parent hospitals to attach documents supporting the hospital classification that they select during registration. This is a more accurate and efficient way to determine the eligibility of parent hospital registrations, without increasing the burden, since the Government Official attestation has been removed.

2. 340B Program Registrations for STD/TB Clinics: HRSA is requesting that any STD entity provide its Notice of Funding Opportunity (NOFO) number at the time of registration. HRSA is also requesting that an entity describe the type of in-kind funding it receives, as well as the time period of the funding. This will assist HRSA in accurately determining the eligibility of the covered entity registration. This requirement would impose minimal burden on the public, as the NOFO number correlates to the Federal Grant Number, which is already required during registration.

3. 340B Registrations for Ryan White Entities: HRSA is requesting that any Ryan White entity provide its NOFO number at the time of registration. HRSA is also requesting that an entity provide the time period of the assistance. This will assist HRSA to accurately determine the eligibility of the registration. This requirement would impose minimal burden on the public, as the NOFO number correlates to the Federal Grant Number, which is already required during registration.

4. Medicaid Billing: HRSA is making a minor change to clarify the question about Medicaid billing.

Accurate records are critical to the prevention of drug diversion to non-eligible individuals as well as duplicate discounts in the 340B Program. To maintain accurate records, HRSA also requires that covered entities recertify eligibility annually, and that they notify the program of updates to any administrative information that they submitted when initially enrolling in the program. HRSA expects that the burden imposed by these processes is low for recertification and minimal for submitting change requests.

Contract Pharmacy Self-Certification

To ensure that drug manufacturers and drug wholesalers recognize contract pharmacy arrangements, covered entities that elect to use one or more contract pharmacies are required to submit general information about the arrangements and certify that signed agreements are in place with those contract pharmacies.

Pharmaceutical Pricing Agreement and Addendum

In accordance with the 340B Program guidance issued in the May 7, 1993, **Federal Register**, section 340B(a)(1) of the PHS Act provides that a manufacturer who sells covered outpatient drugs to eligible entities must sign a Pharmaceutical Pricing Agreement (the ''Agreement'') with the Secretary of HHS (the ''Secretary'') in which the manufacturer agrees to charge a price for covered outpatient drugs that will not exceed the average manufacturer price decreased by a rebate percentage. Also, section 340B(a)(1) of the PHS Act includes specific required components of the PPA with manufacturers of covered outpatient drugs. In particular, section 340B(a)(1) includes the following requirements:

I. ''Each such agreement shall require that the manufacturer furnish the Secretary with reports, on a quarterly basis, of the price for each covered outpatient drug subject to the agreement that, according to the manufacturer, represents the maximum price that covered entities may permissibly be required to pay for the drug (referred to in this section as the ''ceiling price'') and

II. ''. . . shall require that the manufacturer offer each covered entity covered outpatient drugs for purchase at or below the applicable ceiling price if such drug is made available to any other purchaser at any price.''

The burden imposed on manufacturers by submission of the PPA and PPA Addendum is low as the information is readily available.

Pricing Data Submission, Validation and Dissemination

To implement section 340B(d)(1)(B)(i)(II) of the PHS Act, HRSA developed a system to calculate 340B ceiling prices prospectively from data obtained from the Centers for Medicare & Medicaid Services as well as a third party commercial database. However, to conduct the comparison required under the statute, manufacturers must submit the quarterly pricing data as required by section 340B(d)(1)(B)(i)(II). The 340B OPAIS securely collects the following data from manufacturers on a quarterly basis: Average manufacturer price, unit rebate amount, package size, case pack size, unit type, national drug code, labeler code, product code, period of sale (year and quarter), FDA product name, labeler name, wholesale acquisition cost, and the manufacturer determined ceiling price for each covered outpatient drug produced by a manufacturer subject to a PPA. The burden imposed on manufacturers is low because the information requested is readily available and used by manufacturers in other areas.

Likely Respondents: Drug manufacturers and covered entities.

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	Hours per respondent	Total burden hours
Hospital Enrollment, Additions & Recertifications					
340B Program Registrations & Certifications for Hospitals *	248	1	248	2.00	496
Certifications to Enroll Hospital Outpatient Facilities	665	8	5,320	0.50	2,660
Hospital Annual Recertifications	2,481	10	24,810	0.25	6,202
Registrations and Recertifications for Entities Other Than Hospitals					
340B Registrations for Community Health Centers *	360	3	1,080	1.00	1,080
340B Registrations for STD/TB Clinics *	535	1	535	1.00	535
340B Registrations for Various Other Eligible Entity Types *	392	1	392	1.00	392
Community Health Center Annual Recertifications	1,277	7	8,939	0.25	2,235
STD & TB Annual Recertifications	4,033	1	4,033	0.25	1,008
Annual Recertification for entities other than Hospitals, Community Health Centers, and STD/TB Clinics	4,472	1	4,472	0.25	1,118
Contracted Pharmacy Services Registration & Recertifications					
Contracted Pharmacy Services Registration	2,048	11	22,528	1.00	22,528
Other Information Collections					
Submission of Administrative Changes for any Covered Entity	19,322	1	19,322	** 0.25	4,831
Submission of Administrative Changes for any Manufacturer	350	1	350	0.50	175
Pharmaceutical Pricing Agreement and PPA Addendum	200	1	200	1.00	200
Total	36,383		92,229		43,460

* Revised since last OMB submission, but burden was not affected.
** Burden changed from .5 to .25 due to the 340B OPAIS improvement.

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.

Amy McNulty,

Acting Director, Division of the Executive Secretariat.

[FR Doc. 2019–09601 Filed 5–8–19; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Tick-Borne Disease Working Group

AGENCY: Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) is hereby giving notice that the Tick-Borne Disease Working Group (Working Group) will hold a meeting. The meeting will be open to the public. For this meeting, Working Group members will focus on plans to develop the next report to the HHS Secretary and Congress on federal tick-borne activities and research, taking into consideration the 2018 report. The 2020 report will also address a wide range of federal activities and research related to tick-borne diseases, such as, surveillance, prevention, diagnosis, diagnostics, and treatment; identify gaps in tick-borne disease research; and provide recommendations to the HHS Secretary regarding changes or improvements to such activities and research. In developing the report, the Working Group will solicit stakeholder input.

DATES: The meeting will be held on June 4, 2019, from 8:30 a.m. to 5:00 p.m. ET (times are tentative and subject to change). The confirmed times and agenda items for the meeting will be posted on the website for the Working Group at *https://www.hhs.gov/ash/advisory-committees/tickbornedisease/meetings/2019-6-4/index.html* when this information becomes available.

ADDRESSES: The meeting will be held at the Holiday Inn Washington-Capitol, 550 C Street SW, Washington, DC 20024. Members of the public may also attend the meeting via webcast. Instructions for attending via webcast will be posted one week prior to the meeting at *https://www.hhs.gov/ash/advisory-committees/tickbornedisease/meetings/2019-6-4/index.html.*

FOR FURTHER INFORMATION CONTACT: James Berger, Designated Federal Officer for the Working Group; Office of HIV/AIDS and Infectious Disease Policy, Office of the Assistant Secretary for Health, Department of Health and Human Services, Mary E Switzer Building, 330 C Street SW, Suite L100, Washington, DC 20024. Email: *tickborne disease@hhs.gov;* Phone: 202–795–7697.

SUPPLEMENTARY INFORMATION: In-person attendance at the meeting is limited to space available; therefore, preregistration for public members is advisable and can be accomplished by