

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN<sup>1</sup>—Continued

21 CFR section; activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Use of veterinary master files during all phases of product development (including product development that precedes the establishment of an INAD file or the submission of a NADA) .....	15	1	15	20	300
Total .....			1,022		22,083

<sup>1</sup> There are no capital costs or operating and maintenance costs associated with this collection of information.

<sup>2</sup> NADAs and supplements regarding antimicrobial animal drugs that use a recommended approach to assessing antimicrobial concerns as part of the overall preapproval safety evaluation.

Based on a review of the information collection since our last request for OMB approval, we have made no adjustments to our previous estimates. However, as discussed, we have separately estimated the burden of the “Use of veterinary master files during all phases of product development (including product development that precedes the establishment of an INAD file or the submission of a NADA)” in table 1, row 10. We base our estimate of the total annual responses for the use of veterinary master files on such uses initiated during calendar year 2018. We base our estimate of the hours per response upon our experience with the respondents’ use of veterinary master files. We estimate that the time it takes to compile information and submit it to a veterinary master file will vary from 1 to 50 hours, depending on the complexity of the information; therefore, we are estimating on average the burden per response to be 20 hours. Accordingly, we report an additional 300 burden hours and 15 total annual responses in row 10. We are also correcting several rounding errors that were made in our last request for OMB approval. Correcting these rounding errors reduces our previously reported total burden hours and total responses. Thus, our estimated burden for the information collection reflects a net overall increase of 124 hours and a corresponding increase of 14 responses.

Dated: February 11, 2019.

**Lowell J. Schiller,**

*Acting Associate Commissioner for Policy.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

#### Advisory Council on Blood Stem Cell Transplantation

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

**ACTION:** Notice of charter renewal.

**SUMMARY:** HHS is hereby giving notice that the Advisory Council on Blood Stem Cell Transplantation (ACBSCT) has been renewed. The effective date of the renewed charter is February 19, 2019.

#### FOR FURTHER INFORMATION CONTACT:

Robert Walsh, Executive Secretary, ACBSCT, HRSA, 5600 Fishers Lane, Rockville, Maryland 20857. Phone: 301–443–6839; email: [rwalsh@hrsa.gov](mailto:rwalsh@hrsa.gov).

**SUPPLEMENTARY INFORMATION:** Relevant statutes are Public Law 109–129 as amended by Public Law 111–264; 42 U.S.C. 274k; and Section 379 of the Public Health Service Act. The Council is governed by the provisions of Public Law 92–463, as amended (5 U.S.C. appendix 2), which sets forth standards for the formation and use of advisory committees.

ACBSCT advises and makes recommendations to the Secretary of Health and Human Services (Secretary) on matters related to the activities of the C.W. Bill Young Cell Transplantation Program and the National Cord Blood Inventory Program. One of its principal functions shall be to provide consolidated, comprehensive sources of expert, unbiased analysis and recommendations to the Secretary on the latest advances in the science of blood stem cell transplantation.

ACBSCT may meet up to three times during the fiscal year. The charter renewal for ACBSCT was approved on February 7, 2019. The filing date is February 19, 2019. Renewal of the

ACBSCT charter authorizes the Council to operate until February 19, 2021.

A copy of the ACBSCT charter is available on the ACBSCT website at: [https://bloodcell.transplant.hrsa.gov/about/advisory\\_council/index.html](https://bloodcell.transplant.hrsa.gov/about/advisory_council/index.html). A copy of the charter can also be obtained by accessing the FACA database that is maintained by the Committee Management Secretariat under the General Services Administration. The website address for the FACA database is <http://www.facadatabase.gov/>.

**Amy P. McNulty,**

*Acting Director, Division of the Executive Secretariat.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Retail Pharmacy Interest in Utilization of Innovative Educational Technology To Increase Human Papillomavirus (HPV) Vaccination Rates in Rural Areas

**AGENCY:** National Vaccine Program Office, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** This request for information (RFI) is issued for informational and planning purposes only. This RFI is not a solicitation; nor does it commit the Department of Health and Human Services (HHS) to issue a solicitation, make any award, or pay any costs associated with responding to this announcement.

The RFI is being issued by the National Vaccine Program Office (NVPO) of the U.S. Department of Health and Human Services. The NVPO is located in the Office of the Assistant Secretary for Health (ASH), Office of the Secretary (OS), U.S. Department of Health and Human Services (HHS). The NVPO provides strategic leadership and