including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *regulations.gov,* approximately two-to-three business days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Mr. Curtis E. Glover, Sr., Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA 202–501–1448 or via email at *curtis.glover@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

FAR Part 47 contains policies and procedures for applying transportation and traffic management considerations in the acquisition of supplies. The FAR part also contains policies and procedures when acquiring transportation or transportation-related services. Generally, contracts involving transportation require information regarding the nature of the supplies, method of shipment, place and time of shipment, applicable charges, marking of shipments, shipping documents and other related items. Contractors are required to provide the information in accordance with the following FAR Part 47 clauses: 52.247–29 through 52.247–44, 52.247–48, 52.247–52, and 52.247–64. The information is used to ensure that: (1) Acquisitions are made on the basis most advantageous to the Government and; (2) supplies arrive in good order and condition, and on time at the required place.

B. Annual Reporting Burden

Respondents: 65,000.
Responses per Respondent: 22.
Annual Responses: 1,430,000.
Hours per Response: .05.
Total Burden Hours: 71,500.

C. Public Comments

A 60-day notice was published in the **Federal Register** at 83 FR 15571 on April 11, 2018. No comments were received. Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, telephone 202–501–4755.

Please cite OMB Control No. 9000–0061, Transportation Requirements, in all correspondence.

William Clark,

Director, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2018–17929 Filed 8–17–18; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Office on Trafficking in Persons; Notice of Meeting

AGENCY: Administration for Children and Families (ACF), Department of Health and Human Services.

ACTION: Announcement of meeting.

SUMMARY: Notice is hereby given, pursuant to the provisions of the Federal Advisory Committee Act (FACA) and the Preventing Sex Trafficking and Strengthening Families Act, that a meeting of the National Advisory Committee (NAC) on the Sex Trafficking of Children and Youth in the United States (Committee) will be held on September 13–14, 2018. The purpose of the meeting is for the Committee to discuss its duties and information for a draft report on recommended best practices for states to follow in combating the sex trafficking of children and youth based on multidisciplinary research and promising, evidence-based models and programs.

DATES: The meeting will be held on Thursday, September 13, 2018, from 9:00 a.m. to 5:00 p.m. ET and on Friday, September 14, 2018, from 9:00 a.m. to 1:00 p.m. ET.

ADDRESSES: The meeting will be held at 200 Independence Ave. SW, Washington, DC 20201. Space is limited. Identification will be required at the entrance of the facility (*e.g.,* passport, state ID, or federal ID).

To attend the meeting virtually, please register for this event online: *https://www.acf.hhs.gov/otip/resource/nacagenda0918.*

FOR FURTHER INFORMATION CONTACT: Katherine Chon, Director, Office on Trafficking in Persons, Designated Federal Officer (DFO) at *EndTrafficking@acf.hhs.gov* or (202) 205–4554 or 330 C Street SW, Washington, DC 20201. Additional information is available at *https://www.acf.hhs.gov/otip/partnerships/the-national-advisory-committee.*

SUPPLEMENTARY INFORMATION: The formation and operation of the NAC are governed by the provisions of Public Law 92–463, as amended (5 U.S.C. App. 2), which sets forth standards for the formation and use of federal advisory committees.

Purpose of the NAC: The purpose of the NAC is to advise the Secretary and the Attorney General on practical and general policies concerning improvements to the nation's response to the sex trafficking of children and youth in the United States. The NAC is established pursuant to Section 121 of the Preventing Sex Trafficking and Strengthening Families Act of 2014 (P.L. 113–183).

Tentative Agenda: The agenda can be found at *https://www.acf.hhs.gov/otip/resource/nacagenda0918.*

Public Accessibility to the Meeting: Pursuant to 5 U.S.C. 552b and 41 CFR 102–3.140 through 102–3.165, and subject to the availability of space, this meeting is open to the public. Seating is on a first to arrive basis. Because the meeting of the NAC will be held at 200 Independence Ave. SW, Washington, DC 20201, security screening is required. Attendees are requested to register by submitting their name, affiliation, email address, and daytime phone number 10 business days prior to the meeting by email to: *adonald@nhttac.org.* A photo ID is required to enter the premises. Please note that space and parking is limited. The building is fully accessible to individuals with disabilities.

Written Comments or Statements: Pursuant to 41 CFR 102–3.105(j) and 102–3.140 and section 10(a)(3) of the FACA, the public or interested organizations may submit written comments or statements to the NAC in response to the stated agenda of the meeting or in regard to the committee's mission in general. Written comments or statements should be addressed to Katherine Chon, the NAC DFO, via email, the preferred mode of submission, at *adonald@nhttac.org.* The DFO will review all submitted written comments or statements and provide them to members of the NAC for consideration in advance of the meeting. Written comments or statements submitted in response to the agenda set forth in this notice must be received by the DFO at least 15 business days prior

to the meeting to be considered by the NAC. Written comments or statements received after this date may not be provided to the NAC until its next meeting.

Verbal Comments or Statements: Pursuant to 41 CFR 102–3.140d, the NAC is not obligated to allow a member of the public to speak or otherwise address the NAC during the meeting. Members of the public are invited to provide verbal comments or statements during the NAC meeting only at the time and manner described below. All requests to speak or otherwise address the NAC during the meeting must be submitted to the NAC's DFO at least 15 days prior to the meeting, via email, the preferred mode of submission, at *adonald@nhttac.org.* The request should include a brief statement of the subject matter to be addressed by the comment and should be relevant to the stated agenda of the meeting or in regard to the NAC's mission in general. The DFO will log each request in the order received. A period near the end of the meeting will be available for verbal public comments. The time allotted will depend on the number of public comments or statements received and the NAC's agenda items. To provide time for as many people to speak as possible, speaking time for each individual will be limited to 3 minutes.

Minutes: The minutes of this meeting will be available for public review and copying within 90 days at: *https://www.acf.hhs.gov/otip/partnerships/the-national-advisory-committee.*

Dated: August 14, 2018.

Steven Wagner,

Acting Assistant Secretary for Children and Families.

[FR Doc. 2018–17891 Filed 8–17–18; 8:45 am]

BILLING CODE 4184–40–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–1558]

Food and Drug Administration's Evaluation of Approaches To Demonstrate Effectiveness of Heartworm Preventatives for Dogs; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is extending the comment period for the request for comments that appeared in the **Federal Register** of May 24, 2018. In the request for comments, FDA requested comments on the design of studies intended to generate data to support substantial evidence of effectiveness for investigational new animal drugs intended for the prevention of heartworm disease in dogs. The Agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period on the request for comments published May 24, 2018 (83 FR 24122). Submit either electronic or written comments by November 20, 2018.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before November 20, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of November 20, 2018. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov*.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2018–N–1558 for ''FDA's Evaluation of Approaches to Demonstrate Effectiveness of Heartworm Preventatives for Dogs.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the