Secretary, 600 Pennsylvania Avenue NW., Suite CC–5610 (Annex D), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610 (Annex D), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at *http://www.ftc.gov* to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before April 27, 2015. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission (''FTC'') has accepted, subject to final approval, an agreement containing a consent order from TT of Longwood, Inc., also doing business as Cory Fairbanks Mazda. The proposed consent order has been placed on the public record for thirty (30) days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After thirty (30) days, the FTC will again review the agreement and the comments received, and will decide whether it should withdraw from the agreement and take appropriate action or make final the agreement's proposed order.

The respondent is a motor vehicle dealer. According to the FTC's complaint, the respondent has misrepresented: (1) Vehicle purchase prices; (2) that advertised prices, discounts, rebates, bonuses, and incentives are available to all consumers; (3) the prices for added features such as spoilers and sunroofs; (4) that vehicles are available for sale or lease for zero down, zero payments, or zero interest; (5) that vehicles are available for $99; and (6) that consumers can pay $0 at the inception of a lease to lease the advertised vehicle for the advertised monthly payment amount. The complaint alleges therefore that the representations are false and misleading in violation of Section 5 of the FTC Act.

In addition, the complaint alleges the respondent violated the Consumer Leasing Act (''CLA'') and Regulation M for failing to disclose or to disclose clearly and conspicuously certain costs and terms when advertising vehicles for lease.

The proposed order is designed to prevent the respondent from engaging in similar deceptive practices in the future. Part I.A of the proposed order prohibits the respondent from misrepresenting the cost of: (1) Purchasing a vehicle with financing, including but not necessarily limited to the amount or percentage of the down payment, the number of payments or period of repayment, the amount of any payment, and the repayment obligation over the full term of the loan, including any balloon payment; or (2) leasing a vehicle, including but not limited to the total amount due at lease inception, the down payment, amount down, acquisition fee, capitalized cost reduction, any other amount required to be paid at lease inception, and the amounts of all monthly or other periodic payments. Part I.B prohibits the respondent from misrepresenting any other material fact about the price, sale, financing, or leasing of any vehicle.

Part II.A of the proposed order prohibits respondent from representing that a discount, rebate, bonus, incentive or price is available unless: (1) It is available to all consumers, and for all vehicles advertised; or (2) the representation clearly and conspicuously discloses all qualifications or restrictions on: (a) A consumer's ability to obtain the discount, rebate, bonus, incentive, or price and (b) the vehicles available at the discount, rebate, bonus incentive, or price. Part II.B prohibits respondent from misrepresenting any of the following: (1) The existence or amount of any discount, rebate, bonus, incentive, or price; (2) the existence, price, value, coverage, or features of any product or service associated with the motor vehicle purchase; (3) the number of vehicles available at particular prices; or (4) any other material fact about the price, sale, financing, or leasing of motor vehicles.

Part III of the proposed order addresses the CLA allegations. Part III.A prohibits the respondent from stating the amount of any payment or that any or no initial payment is required at lease inception without disclosing clearly and conspicuously: (1) That the transaction advertised is a lease; (2) the total amount due at lease signing or delivery; (3) whether or not a security deposit is required; (4) the number, amounts, and timing of scheduled payments; and (5) that an extra charge may be imposed at the end of the lease term. Part III.B prohibits the respondent from violating any provision of the CLA or Regulation M.

Part IV of the proposed order requires the respondent to keep copies of relevant advertisements and materials substantiating claims made in the advertisements. Part V requires the respondent to provide copies of the order to certain of its personnel. Part VI requires notification to the Commission regarding changes in corporate structure that might affect compliance obligations under the order. Part VII requires the respondent to file compliance reports with the Commission. Finally, Part VIII is a provision ''sunsetting'' the order after twenty (20) years, with certain exceptions.

The purpose of this analysis is to aid public comment on the proposed order. It is not intended to constitute an official interpretation of the complaint or proposed order, or to modify in any way the proposed order's terms.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2015–07407 Filed 3–31–15; 8:45 am]

BILLING CODE 6750–01–P

GENERAL SERVICES ADMINISTRATION

[Notice–MVC–2015–01, Docket No. 2015–0054, Sequence 1]

National Dialogue and Pilot To Reduce Reporting Compliance Costs for Federal Contractors and Grantees

AGENCY: General Services Administration (GSA) and Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Chief Acquisition Officers Council, Department of Health and Human Services, and the General Services Administration (GSA) are conducting a national dialogue to discuss ideas on how to reduce the costs (compliance and other) associated with reporting compliance under Federal awards (contracts, subcontracts, grants, subgrants, and cooperative agreements). This dialogue is part of an effort to improve the economy and efficiency of the federal award system by identifying impactful steps that can be taken to streamline, reporting, reduce burden, and reduce costs.

DATES: Interested parties may participate in the national dialogue through an online platform by reviewing the information and participation dates posted at *www.cao.gov.* The dialogue will open on May 30, 2015 and close on May 30, 2017.

ADDRESSES: Interested parties may participate in the dialogue through the online platform by reviewing the information and participation dates posted at *www.cao.gov.*

FOR FURTHER INFORMATION CONTACT: Christopher Zeleznik at *dataactpmo@hhs.gov* or 202–205–3514 or Emily Gartland at *IAEOutreach@gsa.gov* or 703–605–2532.

SUPPLEMENTARY INFORMATION:

This notice announces a dialogue to explore opportunities to streamline processes and reduce or eliminate burden in federal procurement and grants processes. This dialogue furthers the goals of the President's Management Agenda, which lays the foundation for creating a 21st century government that delivers better results to the American people, and addresses requirements in the Digital Accountability and Transparency Act of 2014 (Public Law 113–101) to gain a better understanding of the costs of compliance with Federal contracting and grants awards as well as recommendations to standardize data, eliminate unnecessary duplication, and reduce compliance costs.

During last year's Open Dialogue on Federal Procurement, published in the **Federal Register** at 79 FR 22682, on April 23, 2014, many commenters pointed to the potential reduction of redundant reporting and related processes as one way to improve the efficiency and effectiveness of the government's acquisition practices. This feedback is helping to support ongoing efforts to modernize the IT infrastructure supporting Federal procurement data collection and display, which will include development of a single Web site for Federal contractors to use for federal contract reporting requirements.

Management of federal contract and grant business arrangements requires multiple layers of reporting across multiple agencies. In some cases, lack of standardization results in slight (or significant) variations in reports that create additional administrative and burdensome requirements for awardees that could be readily rectified. This dialogue is intended to continue the conversation begun last year in the context of federal procurement and expand it to cover federal grants by identifying opportunities for reducing burden, discussing ideas for standardizing processes and forms, and identifying recommended actions to reduce costs and eliminate duplication for awardees. The open dialogue focuses on three topics (campaigns). Each campaign focuses on a unique aspect of the Federal contracting and grants process for which we welcome your insight, ideas, and feedback.

- Campaign 1—Reporting compliance requirements shared by prime and sub-awardees of Federal procurements and grants.
- Campaign 2—Procurement practices, processes, and reporting.
- Campaign 3—Grants practices and processes.

Note—We are looking for ideas to reduce your burden through data standards and changes to reporting procedures. We are interested in hearing about proposed changes that can be accomplished through executive (regulatory, administrative, or management) action, as well as potential legislative proposals where requirements are based in statute.

To facilitate a national dialogue, an online platform will be launched in May 2015 so that interested parties may submit ideas, comment on others, respond to questions posed by moderators, and vote to indicate which ideas they think are most promising and impactful. Information on the platform, and the dates for participating in the dialogue, will be posted at *www.cao.gov.* A separate notice will beposted to address additional dialogue topics on federal procurement for conversation later in the spring and summer.

Dated: March 27, 2015.

William Clark,

Director, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2015–07441 Filed 3–31–15; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–D–0838]

Procedures for Meetings of the Medical Devices Advisory Committee; Draft Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the draft guidance entitled ''Procedures for Meetings of the Medical Devices Advisory Committee.'' The Center for Devices and Radiological Health (CDRH) is issuing this guidance to provide additional information regarding the processes for meetings of the Medical Devices Advisory Committee panels other than the Dispute Resolution Panel (DRP). This guidance describes the general circumstances in which CDRH consults with a panel, the process for exchange of information between CDRH, the members of the panel, industry, and the public, and the conduct of panel meetings. This guidance supplements existing FDA Agency-wide guidance on the conduct of Advisory Committee meetings. This draft guidance is not final nor is it in effect at this time.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment of this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by June 1, 2015.

ADDRESSES: An electronic copy of the guidance document is available for download from the Internet. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance. Submit written requests for a single hard copy of the draft guidance document entitled ''Procedures for Meetings of the Medical Devices Advisory Committee'' to the Office of the Center Director, Guidance and Policy Development, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5431, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request.

Submit electronic comments on the draft guidance to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. Identify comments with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: James Swink, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1609, Silver Spring, MD 20993–0002, 301–796–6313.

SUPPLEMENTARY INFORMATION:

I. Background

CDRH is issuing this draft guidance to provide additional information regarding the processes for meetings of the Medical Devices Advisory Committee panels other than the DRP. The term ''panel,'' as used in this guidance, refers to the panels established under the Medical Devices Advisory Committee charter excluding the DRP. This guidance describes the