app stores. Game developers relied on Tapjoy to generate revenue for themselves and offer gamers a way to earn currency to enhance their play. However, Tapjoy’s failure to screen fraudulent offers left both gamers and developers holding the bag. The settlement proposed today should help reverse the lax policing practices that led hundreds of thousands of gamers to file complaints. But when it comes to addressing the deeper structural problems in this marketplace that threaten both gamers and developers, the Commission will need to use all of its tools—competition, consumer protection, and data protection—to combat middlemen mischief, including by the largest gaming gatekeepers.

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SUPPLEMENTARY INFORMATION:
Procedures for Attendance and Public Comment
Contact Dr. Ken Sandler at ken.sandler@gsa.gov or 202–219–1121 to register to attend the Committee meeting. To attend, submit your full name, organization, email address, and phone number. Requests to attend the meeting must be received by 5:00 p.m. ET, on Monday, January 25, 2021. (GSA will be unable to provide technical assistance to any listener experiencing technical difficulties. Testing access to the Web meeting site before the calls is recommended.)
Contact Dr. Sandler to register to comment during the meeting public comment period. Registered speakers/organizations will be allowed a maximum of five minutes each and will need to provide written copies of their presentations. Requests to comment at the meeting must be received by 5:00 p.m., ET, on Monday, January 25, 2021.

Background
The Administrator of GSA established the Committee on June 20, 2011 (Federal Register / Vol. 76, No. 118) pursuant to Section 494 of the Energy Independence and Security Act of 2007 (EISA, 42 U.S.C. 17123). Under this authority, the Committee provides independent policy advice and recommendations to GSA to advance federal building innovations in planning, design, and operations to reduce costs, enable agency missions, enhance human health and performance, and minimize environmental impacts.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration
[Docket No. FDA–2008–N–0312]
Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Extralabel Drug Use in Animals
AGENCY: Food and Drug Administration, HHS.
ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by February 12, 2021.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to https://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. The OMB control number for this information collection is 0910–0325. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, PRASStaff@fda.hhs.gov.