for small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid OMB control number.

DATES: Written Paperwork Reduction Act (PRA) comments should be submitted on or before March 15, 2013. If you anticipate that you will be submitting PRA comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the FCC contact listed below as soon as possible.

ADDRESSES: Submit your PRA comments to Nicholas A. Fraser, Office of Management and Budget, via fax at 202–395–5167 or via Internet at Nicholas_A._Fraser@omb.eop.gov and to Judith B. Herman, Federal Communications Commission, via the Internet at Judith.herman@fcc.gov. To submit your PRA comments by email send them to: PRA@fcc.gov.

FOR FURTHER INFORMATION CONTACT: Judith B. Herman, Office of Managing Director, (202) 418–0214.

SUPPLEMENTARY INFORMATION:
OMB Control Number: 3060–0952.
Title: Proposed Demographic Information and Notifications, Second Further Notice of Proposed Rulemaking (FNPRM), CC Docket No. 98–147 and Fifth NPRM (NPRM), CC Docket No. 96–98.
Form Number: N/A.
Type of Review: Extension of a currently approved collection.
Respondents: Business or other for-profit entities.
Number of Respondents: 1,200 respondents; 1,200 responses.
Estimated Time per Response: 2 hours.
Frequency of Response: On occasion reporting requirements and third party disclosure requirement.
Total Annual Burden: 4,800 hours.
Total Annual Cost: N/A.
Privacy Impact Assessment: N/A.
Nature and Extent of Confidentiality: The Commission is not requesting respondents to submit confidential information to the FCC. If the applicants wish to submit information which they believe is confidential, they may request confidential treatment of such information under 47 CFR 0.459 of the Commission’s rules.
Needs and Uses: The Commission is seeking an extension of this information collection in order to obtain the full three year approval from OMB. There is no change to the reporting and third party disclosure requirements.
The Commission asked whether physical collocation in remote terminals presents technical or security concerns, and if so, whether these concerns warrant modification of its collocation rules. The Commission asked whether incumbent LECs should be required to provide requesting carriers with demographic and other information regarding particular remote terminals similar to the information available regarding incumbent LEC central offices. Requesting carriers use demographic and other information obtained from incumbent LECs to determine whether they wish to collocate at particular remote terminals.
This proposed information collection in the Second Further Notice of Proposed Rulemaking, FCC 98–147, will be used by the Commission, state commissions, and competitive carriers to facilitate the deployment of advanced services and other telecommunications services in implementation of section 251(c)(6) of the Communications Act of 1934, as amended.
Federal Communications Commission.
Marlene H. Dorch, Secretary.
[FR Doc. 2013–00554 Filed 1–11–13; 8:45 am] BILLING CODE 6712–01–P

FEDERAL TRADE COMMISSION
Revised Jurisdictional Thresholds of the Clayton Act

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: The Federal Trade Commission announces the revised thresholds for interlocking directorates required by the 1990 amendment of Section 8 of the Clayton Act. Section 8 prohibits, with certain exceptions, one person from serving as a director or officer of two competing corporations if two thresholds are met. Competitor corporations are covered by Section 8 if each one has capital, surplus, and undivided profits aggregating more than $10,000,000, with the exception that no corporation is covered if the competitive sales of either corporation are less than $1,000,000. Section 8(a)(5) requires the Federal Trade Commission to revise those thresholds annually, based on the change in gross national product. The new thresholds, which take effect immediately, are $28,883,000 for Section 8(a)(1), and $2,888,300 for Section 8(a)(2)(A).

DATES: Effective Date: January 14, 2013.

FOR FURTHER INFORMATION CONTACT: James F. Mongoven, Bureau of Competition, Office of Policy and Coordination, (202) 326–2879.


By direction of the Commission.

Richard C. Donohue,
Acting Secretary.

[FR Doc. 2013–00482 Filed 1–11–13; 8:45 am] BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Mandatory Guidelines for Federal Workplace Drug Testing Programs

AGENCY: Substance Abuse and Mental Health Services Administration (SAMHSA), Department of Health and Human Services.

ACTION: HHS Approval of Entities that Certify Medical Review Officers (MRO).

SUMMARY: The current version of the Department of Health and Human Services (HHS) Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines), effective on October 1, 2010, addresses the role and qualifications of Medical Review Officers (MROs) and HHS approval of entities that certify MROs.

Subpart M-Medical Review Officer (MRO), Section 13.1(b) states as follows: “Nationally recognized entities that certify MROs or subspecialty boards for physicians performing a review of Federal employee drug testing results that seek approval by the Secretary must submit their qualifications and a sample examination. Based on an annual objective review of the qualifications and content of the examination, the Secretary shall publish a list in the Federal Register of those entities and boards that have been approved.”

HHS has completed its review of entities that train and certify MROs, in accordance with requests submitted by such entities to HHS.

(1) The HHS Secretary approves the following MRO certifying entities that offer both MRO training and certification through examination: American Association of Medical Review Officers (AAMRO), P.O. Box 12873, Research Triangle Park, NC 27709, Phone: (800) 489–1839, Fax:...
Deterrent Opioids—Evaluation and Labeling.

The draft guidance describes how abuse-deterrent properties of opioid analgesic products should be studied and evaluated, and what claims regarding such properties may be suitable for inclusion in labeling. In addition to general input on this draft guidance, FDA is seeking input on the research topics outlined in the final section of the draft guidance. FDA also intends to hold a public meeting to solicit additional input from affected stakeholders on the draft guidance.

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 on 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 March 15, 2013.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 2201, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. See the SUPPLEMENTARY INFORMATION section for electronic access to the draft guidance document.

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, 10903 New Hampshire Ave., Bldg. 38, Room 1–25, Silver Spring, MD 20993–0082.

FOR FURTHER INFORMATION CONTACT: Jennifer Fan, Pharm.D., J.D., Division of Substance Abuse Prevention (CSAP), Substance Abuse and Mental Health Services Administration (SAMHSA), 1 Choke Cherry Road, Room 7–1038, Rockville, MD 20857; Telephone: (240) 276–1759; Email: jennifer.fan@samhsa.hhs.gov.


Kathleen Sebelius,
Secretary.

[FR Doc. 2013–00476 Filed 1–11–13; 8:45 am]

BILLING CODE 4160–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
[Docket No. FDA–2013–D–0045]

Draft Guidance for Industry on Abuse-Deterrent Opioids—Evaluation and Labeling; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled “Abuse-Deterrent Opioids—Evaluation and Labeling.” This draft guidance is intended to provide industry with a framework for evaluating and labeling abuse-deterrent opioid products. The draft guidance discusses how the potentially abuse-deterrent properties of opioid analgesics intended to deter abuse should be evaluated and what claims regarding such properties may be suitable for inclusion in labeling. In addition to general input on this draft guidance, FDA is seeking input on the research topics outlined in the final section of the draft guidance. FDA also intends to hold a public meeting to solicit additional input from affected stakeholders on the draft guidance.

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 on 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 March 15, 2013.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 2201, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. See the SUPPLEMENTARY INFORMATION section for electronic access to the draft guidance document.

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, 10903 New Hampshire Ave., Bldg. 38, Room 1–25, Silver Spring, MD 20993–0082.

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Kathleen Sebelius,
Secretary.

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BILLING CODE 4160–20–P