

information. Please allow at least five days' advance notice; last-minute requests will be accepted, but may be impossible to fill.

Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. 2014-04459 Filed 2-27-14; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL ELECTION COMMISSION

Sunshine Act Meetings

AGENCY: Federal Election Commission.

DATE AND TIME: Tuesday March 4, 2014 at 10 a.m.

PLACE: 999 E Street NW., Washington, DC.

STATUS: This meeting will be closed to the public.

ITEMS TO BE DISCUSSED: Compliance matters pursuant to 2 U.S.C. 437g.

Information the premature disclosure of which would be likely to have a considerable adverse effect on the implementation of a proposed Commission action.

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PERSON TO CONTACT FOR INFORMATION:
Judith Ingram, Press Officer, Telephone:

copy can be obtained from the FTC Public Reference Room, Room 130–H, 600 Pennsylvania Avenue NW., Washington, DC 20580, either in person or by calling (202) 326–2222.

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before March 24, 2014. Write “N.E.W. Plastics Corp.,—Consent Agreement; File No. 132 3126” on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission Web site, at <http://www.ftc.gov>.

As a matter of discretion, the Commission tries to remove individuals’ home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, like anyone’s Social Security number, date of birth, driver’s license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any “[t]rade secret or any commercial or financial information which . . . is privileged or confidential,” as discussed in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you have to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c).¹ Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to

heightened security screening. As a result, we encourage you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at <http://www.ftc.gov>.

by following the instructions on the web-based form. If this Notice appears at <http://www.ftc.gov>, you also may file a comment through that Web site.

If you file your comment on paper, write “N.E.W. Plastics Corp.,—Consent Agreement; File No. 132 3126” on your comment and on the envelope, and mail or deComas disc2.itQhGeneral nonsel,Tny coders your , OffTj /15 U.S. number, dagraary, Rore H T*6113 (Annex D Tw 600s C <http://www.ftc.gov> “N.ov/os/

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¹ In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. FTC Rule 4.9(c), 16 CFR 4.9(c).

instrumentalities to make any false, unsubstantiated, or otherwise misleading representation of material fact regarding any product or package.

Part IV requires N.E.W. to deliver a letter to its distributors and retailers that instructs them to stop using Evolve and Trimax plastic lumber advertising and marketing materials provided by N.E.W. prior to December 2013. This requirement seeks to ensure that deceptive claims will be entirely removed from the market.

Parts V through IX are reporting and compliance provisions. Part V requires Respondent to keep (and make available to the Commission on request): Copies of advertisements and promotional materials containing the representations covered by the order; materials relied upon in disseminating those representations; evidence that contradicts, qualifies, or calls into question the representations, or the basis relied upon for the representations. Part VI requires dissemination of the order now and in the future to principals, officers, directors, and managers, and to all current and future employees, agents, and representatives having responsibilities relating to the subject matter of the order. It also requires Respondent to maintain and make available to the FTC all acknowledgments of receipt of the order. Part VII requires notification to the FTC of changes in corporate status. Part VIII mandates that Respondent submit an initial compliance report to the FTC and make available to the FTC subsequent reports. Part IX is a provision terminating the order after twenty (20) years, with certain exceptions.

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

By direction of the Commission.

Donald S. Clark,

[FR Doc. 2014-04380 Filed 2-27-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10203]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by 29, 2014.

ADDRESSES: When commenting, please reference the document identifier or OMB control number (OCN). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. You may send your comments electronically to www.reg.gov. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.
2. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at www.cms.gov / 1995.
2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to cms.comments@hhs.gov.
3. Call the Reports Clearance Office at (410) 786-1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786-1326

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS-10203 Medicare Health Outcomes Survey

Under the PRA (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. **Revision of a currently approved collection:** **CMS-10203 Medicare Health Outcomes Survey (HOS);** The collection of Medicare Health Outcomes Survey (HOS) is necessary to hold Medicare managed care contracts accountable for the quality of care they deliver to beneficiaries. This reporting requirement allows us to obtain the information necessary for proper oversight of the Medicare Advantage program. It is critical to our mission that