O (August 27, 2013 Notice [1]). Due to the federal government shutdown (October 1, 2013–October 16, 2013), the FTC extended the comment period to compensate for that interval and ensure that interested persons had a full opportunity to file comments.[2] No comments were received in either instance. Pursuant to the OMB regulations, 5 CFR Part 1320, that implement the PRA, 44 U.S.C. 3501 et seq., the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for the Rule. All comments should be filed as prescribed herein, and must be received on or before January 10, 2014.

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5167.

Burden Statement

The FTC is seeking clearance for its assumed share of the estimated PRA burden regarding the disclosure requirements under the FTC and CFPB Rules. The FTC's assumed share of estimated PRA burden, explained in the August 27, 2013 Notice, is 32,500 hours, $1,866,975 for labor costs, and $250,000 for non-labor costs.[3]

Request for Comment

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before January 10, 2014. Write ''Regulation O PRA Comment, FTC File No. P134812'' 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 :// . . / / . 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 obtained from any person and which is privileged or confidential,'' as provided 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).[4] Your comment will be kept confidential only if the FTC General Counsel 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 :// . / / 2, by following the instructions on the web-based form. If this Notice appears at :// . /#! , you also may file a comment through that Web site.

If you file your comment on paper, write ''Regulation O PRA Comment, FTC File No. P134812'' on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex J), 600 Pennsylvania Avenue NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

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 January 10, 2014. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at :// . . / / .

David C. Shonka,

P .

[FR Doc. 2013–29589 Filed 12–10–13; 8:45 am]

BILLING CODE 6750–01–P

[1] 78 FR 52915.

[2] 78 FR 75649, 75650 (November 1, 2013) (comment period for Regulation O extended to November 13, 2013).

[3] 78 FR at 52917.

[4] 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).

FEDERAL TRADE COMMISSION

[File No. 132 3087]

Goldenshores Technologies, LLC and Erik M. Geidl; Analysis of Proposed Consent Order To Aid Public Comment

AGENCY: Federal Trade Commission.

ACTION: Proposed consent agreement.

SUMMARY: The consent agreement in this matter settles alleged violations of federal law prohibiting unfair or deceptive acts or practices or unfair methods of competition. The attached Analysis to Aid Public Comment describes both the allegations in the draft complaint and the terms of the consent order—embodied in the consent agreement—that would settle these allegations.

DATES: Comments must be received on or before January 6, 2014.

ADDRESSES: Interested parties may file a comment at :// . / / online or on paper, by following the instructions in the Request for Comment part of the **SUPPLEMENTARY INFORMATION** section below. Write ''Goldenshores, File No. 132 3087'' on your comment and file your comment online at :// . / / by following the instructions on the web-based form. If you prefer to file your comment on paper, mail or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex D), 600 Pennsylvania Avenue NW., Washington, DC 20580.

FOR FURTHER INFORMATION CONTACT: Kerry O'Brien (415–848–5189), FTC, Western Region, San Francisco, 600

or refuse the terms of the Brightest Flashlight EULA.

The proposed consent order contains provisions designed to prevent respondents from engaging in similar acts or practices in the future. Specifically, Part I prohibits respondent from misrepresenting (1) the extent to which ''covered information'' is collected, used, disclosed, or shared and (2) the extent to which users may exercise control over the collection, use, disclosure, or sharing of ''covered information'' collected from or about them, their computers or devices, or their online activities. ''Covered information'' is defined as ''(a) a first and last name; (b) a home or other physical address, including street name and name of city or town; (c) an email address or other online contact information, such as an instant messaging user identifier or a screen name; (d) a telephone number; (e) a Social Security number; (f) a driver's license or other state-issued identification number; (g) a financial institution account number; (h) credit or debit card information; (i) a persistent identifier, such as a customer number held in a ''cookie,'' a static Internet Protocol (''IP'') address, a mobile device ID, or processor serial number; (j) precise geolocation data of an individual or mobile device, including but not limited to GPS-based, WiFi-based, or cell-based location information (''geolocation information''); (k) an authentication credential, such as a username and password; or (l) any other communications or content stored on a consumer's mobile device.''

Part II requires respondents to give users of their mobile applications a clear and prominent notice and to obtain express affirmative consent prior to collecting their geolocation information. Part III requires respondents to delete any ''covered information'' in their possession, custody, or control that they collected from users of the Brightest Flashlight App prior to the entry of the order.

Parts IV, V, VI, VII, and VIII of the proposed order require respondent to keep copies of relevant advertisements and materials substantiating claims made in the advertisements; to provide copies of the order to its personnel; to notify the Commission of changes in corporate structure that might affect compliance obligations under the order; and to file compliance reports with the Commission. Part IX provides that the order will terminate after twenty (20) years, with certain exceptions.

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

By direction of the Commission.

Donald S. Clark,

[FR Doc. 2013–29531 Filed 12–10–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–14–0739]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–7570 or send comments to Kim Lane, 1600 Clifton Road, MS D–74, Atlanta, GA 30333 or send an email to .

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

CDC Oral Health Management Information System (OMB No. 0920–0739, exp. 4/30/2014)—Revision—National Center for Chronic Disease Prevention and Public Health Promotion (NCDDPHP), Centers for Disease Control and Prevention (CDC).

The CDC works with state health departments to improve the oral health of the nation. Targeted efforts include building and/or maintaining effective public health capacity for the implementation, evaluation, and dissemination of best practices in oral disease prevention and advancement of oral health. Through a cooperative agreement program (Program Announcement DP13–1307), CDC will provide funding to 21 states over a five-year period. New cooperative agreements went into effect in September 2013 and build on previous funded collaborations involving CDC and state programs. Of the 21 awardees, 3 are funded at the Basic level (Component 1, infrastructure) and 18 are funded at the Enhanced level (Component 2) which includes additional activities. The cooperative agreement funding will be used to strengthen state-based oral health infrastructure and capacity, implement and expand evidence-based interventions that increase community-clinical linkages, such as school-based dental sealant programs; increase and maintain environmental systems level changes that support healthy behaviors, such as community water fluoridation; implement strategies that improve the delivery of targeted clinical preventive services; and promote beneficial health systems changes. CDC funding will also help states reduce health disparities among high-risk populations including, but not limited to, those of lower socio-economic status, rural populations, Hispanic, African American and other ethnic groups.

CDC is currently approved to collect annual progress and activity reports from state-based oral health programs. An electronic reporting system has been in place since 2007 and was enhanced in 2008 to capture information about grantees' success stories and environmental scanning activities. The information collected in the management information system (MIS) improved CDC's ability to disseminate information about successful public health approaches that can be replicated or adapted for use in other states.

CDC plans to implement changes to the existing information collection. Through a Revision request, CDC will increase the number of awardees from 20 to 21; describe changes in the MIS platform and data elements that will align the monitoring and evaluation framework for oral health awardees with the framework used for a number of other programs in the National Center for Chronic Disease Prevention and