

every year in order to retain their status as “current” members of the Safe Harbor Framework.

California Skate-Line sells skating-related lessons and clothing, hosts events, and sponsors live performances. According to the Commission’s complaint, since at least January 2015, California Skate-Line set forth on its Web site, <http://caliskateline.com/index.php?col=3&page=privacy>, privacy policies and statements about its practices, including statements related to its participation in the U.S.-EU Safe Harbor Framework.

The Commission’s complaint alleges that California Skate-Line falsely represented that it was a participant in the U.S.-EU Safe Harbor Framework when, in fact, California Skate-Line was never a participant in the U.S.-EU Safe Harbor Framework. Commerce has never included the company on its public Web site.

Part I of the proposed order prohibits California Skate-Line from making misrepresentations about its membership in any privacy or security program sponsored by the government or any other self-regulatory or standard-setting organization, including, but not limited to, the U.S.-EU Safe Harbor Framework and U.S.-Swiss Safe Harbor Framework.

Parts II through VI of the proposed order are reporting and compliance provisions. Part II requires California Skate-Line to retain documents relating to its compliance with the Order for a five-year period. Part III requires dissemination of the order now and in the future to persons with responsibilities relating to the subject matter of the order. Part IV ensures the notification to the FTC of changes in corporate status. Part V mandates that California Skate-Line submit an initial compliance report to the FTC, and make available to the FTC subsequent reports. Part VI is a provision “sunsetting” the order 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 proposed complaint or order or to modify the order’s terms in any way.

By direction of the Commission.

**April J. Tabor,**

*Acting Secretary.*

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## GENERAL SERVICES ADMINISTRATION

[Notice–2015–PM–03; Docket No. 2015–0002; Sequence No. 18]

### Notice of Public Meeting for the Supplemental Draft Environmental Impact Statement for the Federal Bureau of Investigation Central Records Complex in Winchester County, Virginia

**AGENCY:** General Services  
Administration (GSA).

**ACTION:** Meeting notice.

**SUMMARY:** Pursuant to the National Environmental Policy Act (NEPA) of 1969, as implemented by the Council on Environmental Quality regulations, the GSA has prepared and filed with the Environmental Protection Agency (EPA), a Supplement to the Final Environmental Impact Statement (EIS), from May 2007, analyzing the environmental impacts of site acquisition and development of the Federal Bureau of Investigation (FBI), Central Records Complex (CRC), in Winchester County, Virginia.

**DATES:** *Effective Date:* August 21, 2015. The public may submit comments on the Supplemental Draft EIS during a 45-day public review and comment period beginning Friday, August 21, 2015, and ending on Monday, October 5, 2015. Instructions for submitting comments may be found under the heading **SUPPLEMENTAL INFORMATION** in this notice.

**Public Meeting:** A public information meeting is scheduled for Thursday, September 10, 2015 between 6:00 p.m. and 8:00 p.m., Eastern Standard Time (EST), at the War Memorial Building Social Hall at Jim Barnett Park, located at 1001 East Cork Street, Winchester, VA 22601.

**FOR FURTHER INFORMATION CONTACT:** Ms. Courtenay Hoernemann, Project Environmental Planner, 20 N 8th Street, Philadelphia PA 19107 at 215–446–4710.

**ADDRESSES:** Send written comments by email to [frederick.va.siteacquisition@gsa.gov](mailto:frederick.va.siteacquisition@gsa.gov), or U.S. Postal Service to Courtenay Hoernemann, Project Environmental Planner, 20 N 8th Street, Philadelphia, PA 19107.

#### **SUPPLEMENTARY INFORMATION:**

**Background:** The proposed FBI facility would consolidate the FBI’s records currently housed within the Washington DC area, in addition to field offices and information technology centers nationwide. The project requirements are for an overall square footage of 256,425 gross square feet, and will

include the records storage building, support area, visitor’s screening facility, service center, and guard booth. Parking is proposed at 427 spaces. A Notice of Intent to prepare a Supplemental Draft EIS was published in the **Federal Register** at 80 FR 8311 on February 17, 2015. A public scoping comment period was held for 30 days following publication of the Notice of Intent.

The alternatives fully evaluated in the Supplemental Draft EIS include the No Action Alternative, the Arcadia Route 50 property, and Whitehall Commerce Center.

The Supplemental Draft EIS incorporates by reference and builds upon the analyses presented in the 2007 Final EIS, and documents the Section 106 process under the National Historic Preservation Act of 1966, as amended (36 CFR part 800). The Supplemental Draft EIS addresses changes to the proposed action relevant to environmental concerns and assesses any new circumstances or information relevant to potential environmental impacts.

The Supplemental Draft EIS has been distributed to various federal, state, and local agencies. The Supplemental Draft EIS is available for review on the project Web site <http://www.fbicrc-seis.com>. A printed copy of the Supplemental Draft EIS is available for viewing at the following libraries:

- Handley Library, 100 West Piccadilly Street, P.O. Box 58, Winchester, VA 22604
- Bowman Library, 871 Tasker Road, P.O. Box 1300, Stephens City, VA 22655
- Smith Library, Shenandoah University, 718 Wade Miller Drive, Winchester, VA 22601

Federal, state, and local agencies, and other interested parties, are invited and encouraged to be present or represented at the public meeting on Thursday, September 10, 2015. All formal comments will become part of the public record and substantive comments will be responded to in the Final Supplemental EIS.

**Public Comments:** Comments on the Supplemental Draft EIS can be submitted three ways: (1) Submit comments via the project email address: [frederick.va.siteacquisition@gsa.gov](mailto:frederick.va.siteacquisition@gsa.gov), (2) provide written comments during the public meeting, or (3) mail a comment form or letter to: Ms. Courtenay Hoernemann, Project Environmental Planner, 20 N. 8th Street, Philadelphia, PA 19107. Written comments postmarked by October 5, 2015 will become part of the official public record.

**Public Meeting:** The format will be open house with informational posters

on display, and representatives from GSA and FBI will be available to explain the proposed project, answer questions, and receive comments from the public. Comment forms will be available for the public to provide formal written comments.

Dated: August 14, 2015.

**John Hofmann,**

*Division Director, Facilities Management & Services Programs Division, General Services Administration, Mid-Atlantic Region.*

[FR Doc. 2015-20532 Filed 8-19-15; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Agency for Healthcare Research and Quality, HHS.

**ACTION:** Notice.

**SUMMARY:** This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: “*Online Application Order Form for Products from the Healthcare Cost and Utilization Project (HCUP)*.”

In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection.

**DATES:** Comments on this notice must be received by October 19, 2015.

**ADDRESSES:** Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at [doris.lefkowitz@AHRQ.hhs.gov](mailto:doris.lefkowitz@AHRQ.hhs.gov).

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

**FOR FURTHER INFORMATION CONTACT:** Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427-1477, or by email at [doris.lefkowitz@AHRQ.hhs.gov](mailto:doris.lefkowitz@AHRQ.hhs.gov).

#### SUPPLEMENTARY INFORMATION:

##### Proposed Project

*Online Application Order Form for Products From the Healthcare Cost and Utilization Project (HCUP)*

The Healthcare Cost and Utilization Project (HCUP) is a vital resource helping the Agency achieve its mission to produce evidence to make health care

safer, higher quality, more accessible, equitable, and affordable. HCUP is a family of health care databases and related software tools and products developed through a Federal-State-Industry partnership and sponsored by AHRQ. HCUP includes the largest collection of longitudinal hospital care data in the United States, with all-payer, encounter-level information beginning in 1988. The HCUP databases are annual files that contain anonymous information from hospital discharge records for inpatient care and certain components of outpatient care, such as emergency care and ambulatory surgeries. The project currently releases seven types of databases created for research use on a broad range of health issues, including cost and quality of health services, medical practice patterns, access to health care programs, and outcomes of treatments at the national, State, and local market levels. HCUP also produces a large number of software tools to enhance the use of administrative health care data for research and public health use. Software tools use information available from a variety of sources to create new data elements, often through sophisticated algorithms, for use with the HCUP databases.

HCUP's objectives are to:

- Create and enhance a powerful source of national, state, and all-payer health care data.
- Produce a broad set of software tools and products to facilitate the use of HCUP and other administrative data.
- Enrich a collaborative partnership with statewide data organizations (that voluntarily participate in the project) aimed at increasing the quality and use of health care data.
- Conduct and translate research to inform decision making and improve health care delivery.

This project is being conducted by AHRQ through its primary contractor and subcontractor, Truven Health Analytics and Social & Scientific Systems, Inc., pursuant to AHRQ's statutory authority to conduct and support research on health care and on systems for the delivery of such care, including activities with respect to the outcomes, cost, cost-effectiveness, and use of health care services and access to such services. 42 U.S.C. 299a(a)(3).

##### Method of Collection

The HCUP releases seven types of databases for public research use:

(1) The National Inpatient Sample (NIS) is the largest all-payer inpatient care database in the United States, yielding national estimates of hospital inpatient stays. The NIS approximates

20 percent of the discharges from all U.S. community hospitals and contains data from approximately 8 million hospital stays each year. NIS data releases are available for purchase from the HCUP Central Distributor for data years beginning in 1988.

(2) The Kids' Inpatient Database (KID) is the only all-payer inpatient care database for children in the United States. The KID was specifically designed to permit researchers to study a broad range of conditions and procedures related to child health issues. The KID contains a sample of 2 to 3 million discharges for children age 20 and younger from more than 3,500 U.S. community hospitals. KID data releases are available every third year starting in 1997.

(3) The Nationwide Emergency Department Sample (NEDS) is the largest all-payer Emergency Department (ED) database in the United States. It is constructed to capture information both on ED visits that do not result in an admission and on ED visits that result in an admission to the same hospital. The NEDS contains more than 25 million unweighted records for ED visits at about 1,000 U.S. community hospitals and approximates a 20-percent stratified sample of U.S. hospital-based EDs. NEDS data releases are available beginning with data year 2006.

(4) The State Inpatient Databases (SID) contain the universe of inpatient discharge abstracts from data organizations in 46 States and the District of Columbia that currently participate in the SID. Together, the SID encompass approximately 96 percent of all U.S. community hospital discharges. Most States that participate in the SID make their data available for purchase through the HCUP Central Distributor. Files are available beginning with data year 1990.

(5) The State Ambulatory Surgery and Services Databases (SASD) contain encounter-level data from ambulatory surgery and other outpatient services from hospital-owned facilities. In addition, some States provide data for ambulatory surgery and outpatient services from nonhospital-owned facilities. Currently, 34 States participate in the SASD. Files are available beginning with data year 1997.

(6) The State Emergency Department Databases (SEDD) contain data from hospital-owned EDs for visits that do not result in a hospitalization. Currently, 32 States participate in the SEDD. Files are available beginning with data year 1999.

(7) A new database called the Nationwide Readmissions Database (NRD) is planned for release in late