ACTION: Notice of a revision of a currently approved information collection (1024–0224).

SUMMARY: We (National Park Service) will ask the Office of Management and Budget (OMB) to approve the information collection request (ICR) described below. As required by the Paperwork Reduction Act of 1995 and as part of our continuing efforts to reduce paperwork and respondent burden, we invite the general public and other Federal agencies to take this opportunity to comment on this ICR which is an extension of a currently approved collection of information (OMB #1024–0224). We may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

DATES: Public comments will be accepted on or before July 29, 2011.

ADDRESSES: Please submit written comments on this ICR to the OMB Office of Information and Regulatory Affairs, Attention: Desk Officer for the Department of the Interior via e-mail to oira.docket@omb.eop.gov or fax at 202–395–5806; and reference Information Collection 1024–0224 in the subject line. Please also submit a copy of your comments to Phadrea Ponds, Information Collection Coordinator, National Park Service, 1201 Oakridge Drive, Fort Collins, CO 80525 (mail); or phadrea_ponds@nps.gov (e-mail); and reference Information Collection 1024–0224 in the subject line.

FOR FURTHER INFORMATION CONTACT: Dr. Bruce Peacock, Chief, NPS Social Science Division, 1201 Oakridge Drive, Fort Collins, CO 80525 (phone); 970–225–3597 (Fax); or Bruce_Peacock@nps.gov (e-mail). To see a copy of the entire ICR submitted to OMB, go to http://www.reginfo.gov (Information Collection Review; Currently under Review).

SUPPLEMENTARY INFORMATION:

I. Abstract

The NPS needs information concerning park visitors and visitor services, potential park visitors, and residents of communities near parks to provide National Park Service (NPS) managers with usable knowledge for improving the quality and utility of agency programs, services, and planning efforts. Since many of the NPS surveys are similar in terms of the populations being surveyed, the types of questions being asked, and research methodologies, the NPS proposes to renew its clearance from OMB for a generic Information Collection (1024–0224) of NPS-sponsored surveys. Since 1999, the benefits of this generic approval program have been significant to the NPS, Department of the Interior, OMB, NPS cooperators, and the public. Significant time and cost savings have been incurred and 514 surveys have been conducted in units throughout the National Park System. Approval was typically granted in 60 days or less from the date the Principal Investigator (PI) first submitted the survey package for review. This is a significant reduction over the approximately 6–8 months involved in the regular OMB review process. From FY 1999 through FY 2010, the generic ICR process has produced an estimated cost savings to the Federal government and PI’s of $1,017,495.

II. Data


III. Request for Comments

Comments are invited on: (1) The practical utility of the information being gathered; (2) the accuracy of the burden hour estimate; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden to respondents, including use of automated information techniques or other forms of information technology. Please note that the comments submitted in response to this notice are a matter of public record. Before including your address, phone number, e-mail address, or other personal identifying information in your comment, you should be aware that your entire comment, including your personal identifying information—may be made publicly available at any time. While you can ask OMB in your comment to withhold your personal identifying information from public review, we cannot guarantee that it will be done.

Dated: June 24, 2011.

Robert M. Gordon, Information Collection Clearance Officer, National Park Service.

[FR Doc. 2011–16321 Filed 6–28–11; 8:45 am]

BILLING CODE 4312–52–P

DEPARTMENT OF THE INTERIOR

National Park Service


National Register of Historic Places; Notification of Pending Nominations and Related Actions Nominations for the following properties being considered for listing or related actions in the National Register were received by the National Park Service before June 11, 2011. Pursuant to section 60.13 of 36 CFR part 60, written comments are being accepted concerning the significance of the nominated properties under the National Register criteria for evaluation. Comments may be forwarded by United States Postal Service, to the National Register of Historic Places, National Park Service, 1849 C St., NW., MS 2280, Washington, DC 20240; by all other carriers, National Register of Historic Places, National Park Service, 1201 Eye St., NW., 8th floor, Washington, DC 20005; or by fax, 202–371–6447. Written or faxed comments should be submitted by July 14, 2011. Before including your address, phone number, e-mail address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

James Gabbert, Acting Chief, National Register of Historic Places, National Historic Landmarks Program.

ARIZONA

Maricopa County

Koonz, Kinter K., (North Central Phoenix Farmhouses and Rural Estate Homes, 1895–1959) 7620 N. 7th St., Phoenix, 11000463

ARKANSAS

Clark County

Arkadelphia Commercial Historic District, Roughly Main St. between 5th & 7th Sts.,
DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to §1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 14, 2011, Pharmagra Labs, Inc., 158 McLean Road, Brevard, North Carolina 28712, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed substances for analytical research and clinical trials. Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a). Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 29, 2011.

Dated: June 22, 2011.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 11, 2011, and published in the Federal Register on April 19, 2011, 76 FR 21915, Meda Pharmaceuticals, Inc., 705 Eldorado Street, Decatur, Illinois 62523, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance as a finished drug product in dosage form only for distribution to its customers. The company does not import the listed controlled substance in bulk active pharmaceutical ingredient (API) form.

There are no domestic sources of Nabilone in finished drug product form available in the United States. The U.S. Food and Drug Administration has approved this product for medical use in the United States.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and §952(a), and determined that the registration of Meda Pharmaceuticals Inc. to import the basic class of controlled substance is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Meda Pharmaceuticals Inc. to ensure that the company’s registration is consistent with the public interest. The investigation has included inspection and testing of the company’s physical security systems, verification of the company’s compliance with state and local laws, and a review of the company’s background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: June 22, 2011.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Office of Justice Programs

Meeting of the Department of Justice’s (DOJ’s) National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee

AGENCY: Bureau of Justice Assistance, Justice.

ACTION: Notice of meeting.

SUMMARY: This is an announcement of a meeting of DOJ’s National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee to discuss the role of the NMVTIS Federal Advisory Committee Members and various issues relating to the operation and implementation of NMVTIS.