

Information Collection

1. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Healthy Indiana Program (HIP) 2.0 Beneficiaries Survey; *Use:* Approval for testing and developing the survey is vital to adequately inform CMS decision making regarding Section 1115 Waivers, in particular the State's upcoming NEMT waiver due for renewal by December 1, 2016. The NEMT benefit provides transportation for Medicaid beneficiaries who otherwise have no means of transportation to get to and from medical services. The Healthy Indiana Program (HIP) 2.0 demonstration provides authority for the State to not offer NEMT for the new adult group during the first year of the demonstration (except for pregnant women and individuals determined to be medically frail). CMS may extend the State's authority, subject to evaluation of the impact of this policy on access to care. *Form Number:* CMS-10615 (OMB control number: 0938-1300); *Frequency:* Once; *Affected Public:* Individuals and households; *Number of Respondents:* 36; *Total Annual Responses:* 36; *Total Annual Hours:* 36. (For policy questions regarding this collection contact Teresa DeCaro at 202-384-6309).

Written comments and recommendations will be considered from the public if received by the date and address noted above.

Dated: March 22, 2016.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Community Living

Proposed Information Collection Activity; Comment Request; State Developmental Disabilities Council 5-Year State Plan

AGENCY: Administration on Intellectual and Developmental Disabilities, Administration on Community Living, HHS.

ACTION: Notice.

SUMMARY: A plan developed by the State Council on Developmental Disabilities is required by federal statute. Each State Council on Developmental Disabilities must develop the plan, provide for public comments in the State, provide for approval by the State's Governor, and finally submit the plan on a five-year basis. On an annual basis, the Council must review the plan and make any amendments. The State Plan will be used (1) by any amendments. The State Plan will be used (2) by the Council as a planning document; (3) by the citizenry of the State as a mechanism for commenting on the plans of the Council; (4) by the Department as a stewardship tool, for ensuring compliance with the Developmental Disabilities Assistance and Bill of Rights Act, as one basis for providing technical assistance (e.g., during site visits), and as a support for management decision making.

DATES: Submit written comments on the collection of information by May 31, 2016.

ADDRESSES: Submit written comments on the collection of information by email to: *Valerie.Bond@acl.hhs.gov*.

FOR FURTHER INFORMATION CONTACT:

Valerie Bond, Administration on Community Living, Administration on Intellectual and Developmental Disabilities, Office of Program Support, 330 C Street SW., Room 1139-C, Washington, DC 20201, (202) 795-7311.

SUPPLEMENTARY INFORMATION: In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration on Community Living is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to: Valerie Bond, Administration on Community Living, Administration on Intellectual and Developmental Disabilities, Office of Program Support, 330 C Street NW., Room 1139-C, Washington, DC 20201.

The Department specifically requests comments on: (a) Whether the proposed Collection of information is necessary for the proper performance of the function 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) the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden information to be collected; and (e) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection technique comments and or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Respondents: 56 State Developmental Disabilities Councils.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Developmental Disabilities Council 5-Year State Plan	56	1	367	20,552

Estimated Total Annual Burden Hours: 20,552.

Dated: March 22, 2016.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-N-3037]

Pediatric Studies of Lorazepam; Establishment of Public Docket

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of docket.

SUMMARY: The Food and Drug Administration (FDA) is establishing a public docket to make available to the public a report of the pediatric studies of Lorazepam that were conducted in accordance with the Public Health Service Act (PHS Act) and submitted to the Director of the National Institutes of Health (NIH) and the Commissioner of Food and Drugs.

DATES: Submit either electronic or written comments by April 28, 2016.