up survey will be low-income parents and their children from the Rhode Island site currently participating in the HtE Project. As described in the prior OMB submission, these parents are Medicaid recipients between the ages of 18 and 45 receiving Medicaid through the managed care provider United Behavioral Health (UBH) in Rhode Island who meet study criteria with regard to their risk for depression. Children are the biological, adopted, and stepchildren of these parents, between 1 and 18 years of age.

The annual burden estimates are detailed below, and the substantive content of each component will be detailed in the supporting statement attached to the forthcoming 30-day notice.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
RI 15-month, parent physiological component	400	8	5 minutes or .08 hrs	266.66
RI 15-month, young child physiological component	160	8	5 minutes or .08 hrs	106.66
RI 15-month, youth physiological component	242	8	5 minutes or .08 hrs	161.33

Estimated Total Annual Burden Hours: 534.65.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families 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 the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: September 13, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–7763 Filed 9–19–06; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Office of Family Assistance; Single-Source Program Expansion Supplement

AGENCY: Office of Family Assistance, Administration for Children and Families, HHS.

CFDA#: 93.575.

Legislative Authority: Child Care and Development Block Grant Act of 1990, as amended.

Amount of Award: $101,774.00 for one year.

Project Period: 09/30/2006–09/29/2007.

Justification for the Exception to Competition: Oregon State University (the grantee) is currently conducting data analyses with funding from a research grant awarded in FY 2004 to validate methodologies used to conduct State market rate surveys on the price for child care and early education programs at the State and local levels. The supplemental funds will allow the grantee to include additional datasets in the ongoing analyses representing sampling methodologies that include a more diverse care provider sample, a broader geographical coverage, and several additional data collection methods, and will in turn make the findings from the project more generalizable to States, Tribes and Territories implementing the Child Care and Development Fund program.

CONTACT FOR FURTHER INFORMATION: Ivelisse Martinez-Beck, Research Coordinator, Child Care Bureau, Portals Building, Suite 800, 1250 Maryland Avenue, SW., Washington, DC 20024.

Telephone: 202–690–7885.

Dated: September 1, 2006.

Sidonie Squier,

Director, Office of Family Assistance.

[FR Doc. E6–15559 Filed 9–19–06; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004E–0040]

Determination of Regulatory Review Period for Purposes of Patent Extension; CYDECTIN

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for CYDECTIN and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of Patents and Trademarks, Department of Commerce, for the extension of a patent which claims that animal drug product.

ADDRESSES: Submit written comments and petitions to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments.*

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy (HFD–7), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane,Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: The Drug Price Competition and Patent Term Restoration Act of 1984 (Public Law 98–417) and the Generic Animal Drug and