

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 14-45, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Written comments should be received within 60 days of this notice.

Dated: January 23, 2003.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 03-2215 Filed 1-30-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104-13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To

request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443-1129.

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.

Proposed Project: Uniform Data System (OMB No. 0915-0193)—Revision

This is a request for a revision of approval of the Uniform Data System (UDS), which contains the annual reporting requirements for the cluster of primary care grantees funded by the Bureau of Primary Health Care (BPHC), Health Resources and Services Administration (HRSA). Authorizing Legislation is Section 330 of the Public Health Service Act. The UDS includes reporting requirements for grantees of the following primary care programs:

Community Health Centers, Migrant Health Centers, Health Care for the Homeless, Outreach and Primary Health Services for Homeless Children and Public Housing Primary Care, and Healthy Schools Healthy Communities. BPHC collects data on its programs to ensure compliance with legislative mandates and to report to Congress and policy makers on program accomplishments. To meet these objectives, BPHC requires a core set of information collected annually that is appropriate for monitoring and evaluating performance and reporting on annual trends. The UDS includes two components: the Universal Report, completed by all grantees, provides data on services, staffing, and financing; and the Grant Report, completed by grantees funded under the Homeless, Public Housing Program or Healthy Schools Healthy Communities as well as one of the other programs, provides data on characteristics of users whose services fall within the scope of the Homeless, Public Housing Program, Healthy Schools Healthy Communities grant. Grantees are also asked to provide information on the charges, collections, bad debt write off and contractual disallowances by payor sources (Medicaid, Medicare, self pay and private insurance).

Estimated annualized reporting burden are as follows:

| Type of report | Number of respondents | Hours per response | Total burden hours |
|------------------------|-----------------------|--------------------|--------------------|
| Universal Report | 982 | 27 | 26,514 |
| Grant Report | 184 | 18 | 3,312 |
| Total | 982 | | 29,826 |

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 14-45, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: January 27, 2003.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 03-2349 Filed 1-30-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443-1129.

The following request has been submitted to the Office of Management

and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: Voluntary Partner Surveys To Implement Executive Order 12862 in the Health Resources and Services Administration—(OMB 0915-0212)—Extension

In response to Executive Order 12862, the Health Resources and Services Administration (HRSA) is proposing to conduct voluntary customer surveys of its "partners" to assess strengths and weaknesses in program services. A generic approval is being requested from OMB to conduct the partner surveys. HRSA partners are typically State or local governments, health care facilities, health care consortia, health care providers, and researchers.

Partner surveys to be conducted by HRSA might include, for example, mail or telephone surveys of grantees to

determine satisfaction with a technical assistance contractor, or in-class evaluation forms completed by providers who receive training from HRSA grantees, to measure satisfaction with the training experience. Results of these surveys will be used to plan and redirect resources and efforts as needed

to improve service. Focus groups may also be used to gain partner input into the design of mail and telephone surveys. Focus groups, in-class evaluation forms, mail surveys, and telephone surveys are expected to be the preferred methodologies.

A generic approval will permit HRSA to conduct a limited number of partner

surveys without a full-scale OMB review of each survey. If generic approval is granted, information on each individual partner survey will not be published in the **Federal Register**.

The estimated response burden is as follows:

| Type of survey | Number of respondents | Responses per respondent | Hours per response | Total hour burden |
|------------------------------|-----------------------|--------------------------|--------------------|-------------------|
| In-class evaluations | 40,000 | 1 | .05 | 2,000 |
| Mail/Telephone surveys | 2,000 | 1 | .25 | 3,000 |
| Focus groups | 50 | 1 | 1.5 | 75 |
| Total | 52,050 | | | 5,075 |

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Morrall, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: January 27, 2003.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 03-2348 Filed 1-30-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Environmental Factors in the Development of Polycystic Ovary Syndrome

Summary: Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Environmental Health Sciences (NIEHS), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on October 25, 2002, pages 56690-56691 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or

after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: Environmental Factors in the Development of Polycystic Ovary Syndrome. **Type of Information Collection Request:** Revision of OMB No. 0925-0483 and expiration date 2/28/2003. **Need and Use of Information Collection:** The purpose of this study is to identify a cohort of living female twin pairs in which at least one member is likely to have Polycystic Ovary Syndrome (PCOS) for future study. Potential participants (~3,700) will come from the Mid-Atlantic Twin Registry (MATR) and were chosen based on their answers to several questions (in a preliminary MATR survey) concerning irregular periods and a history of polycystic cystic ovaries. The instrument to be used here will be administered by telephone by professional interviewers at the MATR. It contains 15 simple and direct questions and will take about 10 minutes to complete. Its contents deal with the frequency of menstrual periods, a history of polycystic ovaries, obesity, excess facial hair and other evidence of hyperandrogenism. Since this is such a short telephone survey, participants will receive no prior notification. Informed consent will be asked for verbally over the phone at the time of the interview. All participants will be asked about their willingness to participate in future studies if their answers meet certain criteria. The major objectives of future studies using this cohort are to determine more reliable concordance rates for PCOS in monozygotic and dizygotic twins, establish baseline heritability estimates, and develop hypotheses concerning possible pathogenetic and/or environmental factors. The findings from this study will aid in developing:

(1) Genetic tests to identify high risk women; (2) preventative strategies; and (3) more effective therapies for PCOS and related syndromes such as type 2 diabetes, obesity, idiopathic, hyperandrogenism, and male pattern baldness. **Frequency of Response:** One time. **Affected Public:** Individuals or households. **Type of Respondents:** Adult women. The annual reporting burden is as follows: **Estimated Number of Respondents:** 3,700; **Estimated Number of Responses per Respondent:** 1; **Average Burden Hours Per Response:** 0.167; and **Estimated Total Annual Burden Hours Requested:** 205.9 hours. The annualized cost to respondents is estimated at \$3,449.94. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated