Methodist Hospital of Indiana, Inc., Department of Pathology and Laboratory Medicine, 1701 N. Senate Blvd., Indianapolis, IN 46202, 317–929–3587
Methodist Medical Center Toxicology Laboratory, 221 N.E. Glen Oak Ave., Peoria, IL 61636, 800–752–1835/309–671–5199
MetroLab-Legacy Laboratory Services, 235 N. Graham St., Portland, OR 97227, 503–413–4512 800–237–7808 (x4512)
National Toxicology Laboratories, Inc., 1100 California Ave., Bakersfield, CA 93304, 805–322–4250
Northwest Toxicology, Inc., 1141 E. 3900 South, Salt Lake City, UT 84124, 800–322–3361
Oregon Medical Laboratories, P.O. Box 972, 722 East 11th Ave., Eugene, OR 97440–0972, 503–687–2134
Pathology Associates Medical Laboratories, East 11604 Indiana, Spokane, WA 99206, 509–926–2400
PDLA, Inc. (Princeton), 100 Corporate Court, So. Plainfield, NJ 07080, 908–769–8500/800–237–7352
PharmChem Laboratories, Inc., 1505–A O'Brien Dr., Menlo Park, CA 94025, 415–328–6200/800–446–5177
PharmChem Laboratories, Inc., Texas Division, 7606 Pebble Dr., Fort Worth, TX 76118, 817–595–0294 (formerly: Harris Medical Laboratory)
Physicians Reference Laboratory, 7800 West 110th St., Overland Park, KS 66210, 913–338–4070/800–821–3627
Poisonlab, Inc., 7272 Clairemont Mesa Rd., San Diego, CA 92111, 619–279–2600/800–882–7272
Premier Analytical Laboratories, 15201 I–10 East, Suite 125, Channelview, TX 77530, 713–457–3784 (formerly: Drug Labs of Texas)
Presbyterian Laboratory Services, 1851 East Third Street, Charlotte, NC 28204, 800–473–6640
Puckett Laboratory, 4200 Mamie St., Hattiesburgh, MS 39402, 601–264–3856/800–844–8378
Scientific Testing Laboratories, Inc., 463 Southlake Blvd., Richmond, VA 23236, 804–378–9130
Scott & White Drug Testing Laboratory, 600 S. 25th St., Temple, TX 76504, 800–749–3788
S.E.D. Medical Laboratories, 500 Walter NE, Suite 500, Albuquerque, NM 87102, 505–244–8800, 800–999–LABS
Sierra Nevada Laboratories, Inc., 888 Willow St., Reno, NV 89502, 800–648–5472
SmithKline Beecham Clinical Laboratories, 7600 Tyrone Ave., Van Nuys, CA 91045, 818–989–2520
SmithKline Beecham Clinical Laboratories, 801 East Dixie Ave., Leesburg, FL 34748, 904–787–9006 (formerly: Doctors & Physicians Laboratory)
SmithKline Beecham Clinical Laboratories, 3175 Presidential Dr., Atlanta, GA 30340, 770–452–1590 (formerly: SmithKline Bio-Science Laboratories)
SmithKline Beecham Clinical Laboratories, 506 E. State Pkwy., Schaumburg, IL 60173, 708–885–2010 (formerly: International Toxicology Laboratories)
SmithKline Beecham Clinical Laboratories, 400 Egypt Rd., Norristown, PA 19403, 800–523–5447 (formerly: SmithKline Bio-Science Laboratories)
SmithKline Beecham Clinical Laboratories, 8000 Sovereign Row, Dallas, TX 75247, 214–638–1301 (formerly: SmithKline Bio-Science Laboratories)
SmithKline Beecham Clinical Laboratories, 1737 Airport Way South, Suite 200, Seattle, WA 98134, 206–623–8100
South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 219–234–4176
Southwest Laboratories, 2727 W. Baseline Rd., Suite 6, Tempe, AZ 85283, 602–438–8507
St. Anthony Hospital (Toxicology Laboratory), P.O. Box 205, 1000 N. Lee St., Oklahoma City, OK 73102, 405–272–7052
Toxicology & Drug Monitoring Laboratory, University of Missouri Hospital & Clinics, 2703 Clark Lane, Suite B, Lower Level, Columbia, MO 65202, 314–882–1273
Toxicology Testing Service, Inc., 5426 N.W. 79th Ave., Miami, FL 33166, 305–593–2260
TOXWORX Laboratories, Inc., 6160 Variel Ave., Woodland Hills, CA 91367, 818–226–4373, (formerly: Laboratory Specialists, Inc.; Abused Drug Laboratories; MedTox Bio-Analytical, a Division of MedTox Laboratories, Inc.)
UNILAB, 18408 Oxnard St., Tarzana, CA 91356, 800–492–0800/818–343–8191 (formerly: MetWest-BPL Toxicology Laboratory)

The following laboratory withdrew from the National Laboratory Certification Program on March 29, 1996:

National Psychopharmacology Laboratory, Inc., 9320 Park W. Blvd., Knoxville, TN 37923, 800–251–9492

Richard Kopanda,

Acting Executive Officer Substance Abuse and Mental Health Services. Administration.

[FR Doc. 96–8382 Filed 4–3–96; 8:45 am]

BILLING CODE 4160–20–U

Dietary Supplement Labels Commission; Meeting; Correction

AGENCY: Office of Disease Prevention and Health Promotion, HHS.

ACTION: Correction.

SUMMARY: In notice document 96–7639 beginning on page 14102 in the issue of Friday, March 29, 1996, make the following corrections:

On page 14102 in the third column line 3 in **SUMMARY**, the notice refers to the second meeting. This should be changed to read third meeting.

On page 14102 in the third column in **FOR FURTHER INFORMATION CONTACT** change the telephone number to read (202) 690–7102.

On page 14102 in the third column in Announcement of Meeting, the meeting date, time, and location are listed incorrectly. The sentences should read the Commission's third meeting will be April 26, 1996, 8:30 a.m. to 4:30 p.m. Pacific Standard Time. The meeting will be held in the Sausalito Room, at the Holiday Inn Fisherman's Wharf, 1300 Columbus Avenue, San Francisco, California 94133.

On page 14103 in the first column in Public Participation at Meeting, the last two sentences should read please request the opportunity to present oral comments in writing and provide nine (9) copies of the written comments from which the oral presentation is abstracted to the address above by April 19, 1996. If you will require a sign language interpreter, please call Sandra Saunders (202) 690–7102 by 4:30 p.m. E.S.T. on April 19, 1996.

Dated: April 1, 1996.

Claude Earl Fox,

Deputy Assistant Secretary for Health (Disease Prevention and Health Promotion), U.S. Department of Health and Human Services.

[FR Doc. 96–8270 Filed 4–3–96; 8:45 am]

BILLING CODE 4160–17–M

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Early Head Start Evaluation.

OMB No: New Request.

Description: The Head Start Reauthorization Act of 1994 established a special initiative creating funding for services for families with infants and toddlers. In response the Administration on Children, Youth and Families (ACYF) designed the Early Head Start (EHS) program. In September, 1995, ACYF awarded grants to 68 local programs to serve families with infants and toddlers.

EHS programs are designed to produce outcomes in four domains: (1) child development, (2) family development, (3) staff development and (4) community development. The Reauthorization required that this new initiative be evaluated. To study the effect of the initiative ACYF awarded a contract through a competitive procurement to Mathematica Policy Research, Inc. (MPR) with a subcontract to Columbia University Center for Young Children and Families. Evaluation will be carried out from October 1, 1995, through September 30, 2000. Data collection activities that are the subject of this Federal Register notice are intended for the first phase of the EHS evaluation.

The sample for the child and family assessments will be approximately

3,400 families who include a pregnant woman or a child under 12 months of age, in 17 EHS study sites. Each family will be randomly assigned to a treatment group or a control group. The sample for the child care assessments will include the primary child care provider for the focal child in each of the 3,400 study sample families. The sample for the staff assessments will be all EHS staff who have contact with the study children and families. The surveys and assessments will be conducted through computer assisted telephone interviewing, pencil and paper self-administered questionnaires, structured observations and videotaping. All data collection instruments have been designed to minimize the burden on respondents by minimizing interviewing and assessment time. Participation in the study is voluntary and confidential.

The information will be used by government managers, Congress and others to identify the features and evaluate the effectiveness of the EHS program.

Respondents: Applicants to the Early Head Start program, child care providers for Early Head Start families and Early Head Start staff.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
14-Month Parent Interview, Child Assessment Videotaping Protocol	3,230	1	2.5	8,075
6-Month Parent Services Follow-Up Interview	3,298	1	.75	2,474
Child Care Provider Interview	1,259	1	.50	630
Child Care Provider Observation Protocol	1,259	1	2	2,518
Staff Questionnaire	170	1	.5	85
Estimated Total Annual Burden Hours: 13,782.				

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 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 Information Services, Division of Information Resource Management, 370 L'Enfant Promenade, S.W., Washington, D.C. 20047, Attn.: ACF Reports Clearance Officer. All requests should be identified by title.

In addition, requests for copies may be made and comments forwarded to the Reports Clearance Officer over the Internet by sending a message to rkatson@acf.dhhs.gov. Internet messages must be submitted as an ASCII file without special characters or encryption.

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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: March 28, 1996.

Roberta Katson,

Director, Division of Information Resource Management Services.

[FR Doc. 96–8203 Filed 4–3–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0089]

Establishment of Lists of U.S. Firms/Processors Exporting Shell Eggs, Dairy Products, Game Meat and Game Meat Products to the European Community; Request for Information From Such Firms

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that it intends to establish lists of U.S. firms/processors exporting shell eggs, dairy products, game meat and game meat products to the European Community (EC) that manufacture products in compliance with U.S. food laws and regulations. FDA is taking this action in response to current changes in the EC legislation that will require countries trading with any of the EC member countries to provide lists of firms exporting certain animal derived commodities to the EC. FDA is requesting that U.S. firms presently exporting, or who anticipate exporting these commodities to the EC, provide the agency with information for inclusion on the appropriate list. This list will be updated on a quarterly basis and will be submitted to the EC. This notice is intended to alert all U.S. exporters to the EC requirement for lists of companies processing animal derived commodities that are exported to the EC member states. The agency is also requesting comments on the best mechanisms to be used in obtaining future changes and additions to the EC lists for these commodities.

DATES: Written comments and information for inclusion on the EC list by April 30, 1996. Written comments on the information collection requirements by May 6, 1996.

ADDRESSES: Submit written information for inclusion on the EC list to Marilyn F. Balmer, Center for Food Safety and Applied Nutrition (HFS–306), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, FAX 202–205–4422 or E-mail MFB@FDACF.SSW.DHHS.GOV.

Submit written comments on the best mechanisms to be used in obtaining future changes and additions to the EC lists for dairy products, shell eggs, game meat and game meat products to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the Docket number found in brackets in the heading of this document.

Submit written comments on the information collection requirements to DHHS Reports Clearance Officer, Paperwork Reduction Project (0910–0320), Hubert Humphrey Bldg., 200 Independence Ave. SW., rm. 531–H, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Marilyn F. Balmer, Center for Food