

that sets out the areas of noncompliance and the steps that must be taken to correct the noncompliance. By regulation, OCR must attempt to secure voluntary compliance through informal means. In practice, OCR has been quite successful in securing voluntary compliance and will continue these efforts. If the matter cannot be resolved informally, OCR must secure compliance through (a) the termination of Federal assistance after the recipient/covered entity has been given an opportunity for an administrative hearing, (b) referral to DOJ for injunctive relief or other enforcement proceedings, or (c) any other means authorized by law.

13. Q. Does issuing this guidance mean that OCR will be changing how it enforces compliance with Title VI?

A. No. How OCR enforces Title VI is governed by the Title VI implementing regulations. The methods and procedures used to investigate and resolve complaints, and conduct compliance reviews, have not changed.

14. Q. What is HHS doing to ensure it is following the guidance it is giving to States and others?

A. Although legally, federally conducted programs and activities are not subject to Title VI, HHS recognizes the importance of ensuring that its programs and services are accessible to LEP persons. To this end, HHS has established a working group to assess how HHS itself is providing language access. Currently, agencies across HHS have taken a number of important steps to ensure that their programs and services are accessible to LEP persons. For example, a number of agencies have translated important consumer materials into languages other than English. Also, several agencies have launched Spanish language web sites. In order to ensure that all HHS federally conducted programs and activities are accessible to LEP persons, the Secretary has directed the working group to develop and implement a Department-wide plan for ensuring LEP persons meaningful access to HHS programs. This internal HHS initiative was begun prior to the President's August 11, 2000, Executive Order 13166, "Improving Access to Services for Persons with Limited English Proficiency". The Executive Order requires Federal Agencies to develop and implement a system for ensuring LEP persons meaningful access to their federally-conducted programs. It also requires agencies to issue guidance to their recipients on the recipients' obligations to provide LEP persons meaningful access to their federally-assisted programs. HHS is a step ahead on each of the obligations outlined in the Executive Order.

Appendix B—Selected Federal and State Laws and Regulations Requiring Language Assistance

Federal Laws and Regulations

Federal laws that recognize the need for language assistance include:

1. The Voting Rights Act, which bans English-only elections and prescribes other remedial devices to ensure

nondiscrimination against language minorities;¹

2. The Food Stamp Act of 1977, which requires states to provide written and oral language assistance to LEP persons under certain circumstances;²

3. Judicial procedure laws that require the use of certified or otherwise qualified interpreters for LEP parties and witnesses, at the government's expense, in certain proceedings;³

4. The Older Americans Act, which requires state planning agencies to use outreach workers who are fluent in the languages of older LEP persons, where there is a substantial number of such persons in a planning area;⁴

5. The Substance Abuse and Mental Health Administration Reorganization Act, which requires services provided with funds under the statute to be bilingual if appropriate;⁵

6. The Disadvantaged Minority Health Improvement Act, which requires the Office of Minority Health (OMH) to enter into contracts to increase the access of LEP persons to health care by developing programs to provide bilingual or interpreter services;⁶

7. The Equal Educational Opportunities Act of 1974, which requires educational agencies to take appropriate action to accommodate the language differences that impede equal participation by students in instructional programs;⁷ and

8. Regulations issued by the Health Care Financing Administration (HCFA) which require that evaluations for the mentally ill and mentally retarded be adapted to the cultural background, language, ethnic origin and means of communication of the person being evaluated.⁸

State Laws and Regulations

Many states have recognized the seriousness of the language access challenge and have enacted laws that require providers to offer language assistance to LEP persons in many service settings.⁹ States that require language assistance include:

1. California, which provides that intermediate care facilities must use interpreters and other methods to ensure adequate communication between staff and patients;¹⁰

¹ 42 U.S.C. Section 1973b(f)(1).

² 7 U.S.C. Section 2020(e)(1) and (2)(A).

³ 28 U.S.C. Section 1827(d)(1)(A).

⁴ 42 U.S.C. Section 3027(a)(20)(A).

⁵ 42 U.S.C. Section 290aa(d)(14).

⁶ 42 U.S.C. Section 300u-6(b)(7).

⁷ 20 U.S.C. Section 1703(f).

⁸ 42 CFR section 483.128(b).

⁹ At least twenty six (26) states and the District of Columbia have enacted legislation requiring language assistance, such as interpreters and/or translated forms and other written materials, for LEP persons.

¹⁰ 22 California Code of Regulations, Section 73501. California has a wide array of other laws and regulations that require language assistance, including those that require: (a) intermediate nursing facilities to use interpreters and other methods to ensure adequate communication with patients, (b) adult day care centers to employ ethnic and linguistic staff as indicated by participant characteristics, (c) certified interpreters for non-English speaking persons at administrative

2. New Jersey, which provides that drug and alcohol treatment facilities must provide interpreter services if their patient population in non-English speaking;¹¹

3. Pennsylvania, which provides that a patient who does not speak English should have access, where possible, to an interpreter;¹² and

4. Massachusetts, which in April 2000, enacted legislation that requires every acute care hospital to provide competent interpreter services to LEP patients in connection with all emergency room services.¹³

Medical Accreditation Organizations

1. The Joint Committee on Accreditation of Healthcare Organizations (JCAHO), which accredits hospitals and other health care institutions, requires language assistance in a number of situations. For example, its accreditation manual for hospitals provides that written notice of patients' rights must be appropriate to the patient's age, understanding and language.¹⁴

2. The National Committee for Quality Assurance (NCQA), which provides accreditation for managed care organizations, also requires language assistance in a variety of settings. As part of its evaluation process, the NCQA assesses managed care member materials to determine whether they are available in languages, other than English, spoken by major population groups.¹⁵

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Procedures for Requests to use Child Care and Development Funds for Construction or Major Renovation of Child Care Facilities.

OMB No.: 0970-0160.

Description: The Child Care and Development Block Grant Act, as amended, allows Indian Tribes to use Child Care and Development Fund (CCDF) grant awards for construction and renovation of child care facilities. A tribal grantee must first request and receive approval from the Administration for Children and

hearings, and (d) health licensing agencies to translate patients rights information into every language spoken by 1% or more of the nursing home population.

¹¹ New Jersey Administrative Code Section 42A-6.7.

¹² 28 Pennsylvania Administrative Code Section 103.22(b)(14).

¹³ M.G.L.A. 111, Section 25J

¹⁴ JCAHO, 1997 Accreditation Manual for Hospitals, Section R1.1.4.

¹⁵ NCQA, 1997 Accreditation Standards, RR 6.2.

Families (ACF) before using CCDF funds for construction or major renovation. This information collection contains the statutorily-mandated uniform procedures for the solicitation and consideration of requests, including

instructions for preparation of environmental assessments in conjunction with the National Environmental Policy Act. The proposed draft procedures update and clarify the original procedures that were

issued in August 1997. Respondents will be CCDF tribal grantees requesting to use CCDF funds for construction or major renovation.

Respondents: Tribal Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number or responses per respondent	Average burden hours per response	Total burden hours
Construction and Renovation	25	1	20	500
Estimated Total Annual Burden Hours				500

Additional Information: 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 may be obtained by writing to The Administration for Children and Families, Office of Information Services, 370 L'Infant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Desk Officer for ACF.

Dated: August 25, 2000.

Bob Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Nonprescription Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee

of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Nonprescription Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on October 19, 2000, 8 a.m. to 5 p.m.

Location: Holiday Inn, The Ballroom, Two Montgomery Ave., Gaithersburg, MD.

Contact Person: Sandra L. Titus, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093) Rockville, MD 20857, 301-827-7001, or e-mail Tituss@cder.fda.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12541. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will consider safety issues regarding the use of Phenylpropanolamine (PPA) in over-the-counter (OTC) drug products. The discussion will focus on the reported results of an epidemiological study designed to assess the risk of hemorrhagic stroke associated with the use of PPA. The Consumer Health Products Association (CHPA) commissioned the study which was conducted by Yale University. The material which the committee will review will be available at least 1 business day before the meeting at: <http://www.fda.gov/ohrms/dockets/ac/acmenu.htm>. Click on the year 2000 and then locate the Nonprescription Drugs Advisory Committee meeting for October 20, 2000.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending

before the committee. Written submissions may be made to the contact person by October 9, 2000. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before October 9, 2000, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: August 21, 2000.

Linda A. Suydam,

Senior Associate Commissioner.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Joint Meeting of the Nonprescription Drugs Advisory Committee and the Gastrointestinal Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committees: Nonprescription Drugs Advisory Committee and the Gastrointestinal Drugs Advisory Committee.

General Function of the Committees: To provide advice and