to the Web sites after this document publishes in the **Federal Register**.)

1. Malam, S., S. Clegg, S. Kirwan, and S. McGinigal, ''Comprehension and Use of UK Nutrition Signpost Labelling Schemes,'' report prepared for Food Standards Agency, May 2009.

2. Borgmeier, I, and J. Westenhoefer, ''Impact of Different Food Label Formats on Healthiness Evaluation and Food Choice of Consumers: a Randomized-Controlled Study,'' *BMC Public Health*, 9: 184, 2009, accessed online at *http://www.biomedcentral.com/content/pdf/1471-2458-9-184.pdf*.

3. Kelly, B, C. Hughes, K. Chapman, J.C.-Y. Louie, H. Dixon, J. Crawford, L. King, M. Daube, T. Slevin, ''Consumer Testing of the Acceptability and Effectiveness of Front-of-Pack Food Labelling Systems for the Australian Grocery Market,'' Health Promotion International 24(2):120–9, 2009.

4. Feunekes, G.I.J., I.A. Gortemaker, A.A. Willems, R. Lion, and M. van den Kommer, ''Front-of-pack Nutrition Labelling: Testing Effectiveness of Different Nutrition Labelling Formats Front-of-pack in Four European Countries,'' *Appetite*, 50:57–70, 2008.

Dated: November 24, 2009.

David Horowitz,

Assistant Commissioner for Policy.

[FR Doc. E9–28699 Filed 11–30–09; 8:45 am]

BILLING CODE 4160–01–S

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, e-mail *paperwork@hrsa.gov* or 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: National Health Service Corps Travel Request Worksheet (OMB No. 0915–0278)—Extension

Clinicians participating in the HRSA National Health Service Corps (NHSC) Scholarship Program use the online Travel Request Worksheet to receive travel funds from the Federal Government to perform pre-employment interviews at sites on the NHSC's Opportunities List.

The travel approval process is initiated when a scholar notifies the NHSC of an impending interview at one or more NHSC approved practice sites. The Travel Request Worksheet is also used to initiate the relocation process after a NHSC scholar has successfully been matched to an approved practice site. Upon receipt of the Travel Request Worksheet, the NHSC will review and approve or disapprove the request and promptly notify the scholar and the NHSC logistics contractor regarding travel arrangements and authorization of the funding for the site visit or relocation.

The estimated annual burden is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Travel Request Worksheet	140	2	280	.06	16.8

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the ''attention of the desk officer for HRSA.''

Dated: November 24, 2009.

Alexandra Huttinger,

Director, Division of Policy Review and Coordination.

[FR Doc. E9–28698 Filed 11–30–09; 8:45 am]

BILLING CODE 4165–15–P

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, e-mail *paperwork@hrsa.gov* or 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: The Health Education Assistance Loan (HEAL) Program: Forms (OMB No. 0915–0043 Extension)

The Health Education Assistance Loan (HEAL) program continues to administer and monitor outstanding loans which were provided to eligible students to pay for educational costs in a number of health professions. HEAL forms collect information that is required for responsible program management. The HEAL Repayment Schedule, Fixed and Variable, provides the borrower with the cost of a HEAL loan, the number and amount of payments, and the Truth-in-Lending disclosures. The Lender's Report on HEAL Student Loans Outstanding (Call Report), provides information on the status of loans outstanding by the number of borrowers and total number of loans whose loan payments are in various stages of the loan cycle, such as student education and repayment, and the corresponding dollar amounts. These forms are needed to provide borrowers with information on the cost of their loan(s) and to determine which

lenders may have excessive delinquencies and defaulted loans.

The annual estimate of burden is as follows:

Instrument	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Disclosure: Repayment Schedule HRSA 502–1,2	8	396	3,168	0.50	1,584
Reporting: Call Report HRSA 512	13	4	52	0.75	39
Total Reporting and Disclosure	21		3,220		1,623

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the ''attention of the desk officer for HRSA.''

Dated: November 24, 2009.

Alexandra Huttinger,

Director, Division of Policy Review and Coordination.

[FR Doc. E9–28696 Filed 11–30–09; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2007–D–0369] (formerly Docket No. 2007D–0168)

Draft and Revised Draft Guidances for Industry Describing Product-Specific Bioequivalence Recommendations; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of additional draft and revised draft product-specific bioequivalence (BE) recommendations. The recommendations provide product-specific guidance on the design of BE studies to support abbreviated new drug applications (ANDAs). In the **Federal Register** of May 31, 2007, FDA announced the availability of a draft guidance for industry entitled ''Bioequivalence Recommendations for Specific Products,'' explaining the process that would be used to make product-specific BE recommendations available to the public on FDA's Web site. The BE recommendations identified in this notice were developed using the process described in that guidance.

DATES: Submit written or electronic comments on the draft and revised draft product-specific BE recommendations listed in this notice by February 1, 2010.

ADDRESSES: Submit written requests for single copies of the individual BE guidances to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 2201, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. Submit written comments on the draft product-specific BE recommendations 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.regulations.gov*. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance recommendations.

FOR FURTHER INFORMATION CONTACT: Doan T. Nguyen, Center for Drug Evaluation and Research (HFD–600), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9314.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of May 31, 2007 (72 FR 30388), FDA announced the availability of a draft guidance for industry entitled ''Bioequivalence Recommendations for Specific Products,'' that explained the process that would be used to make product-specific BE recommendations available to the public on FDA's Web site at *http://www.fda.gov/CDER/GUIDANCE/bioequivalence/default.htm*. As described in that draft guidance, FDA adopted this process as a means to develop and disseminate product-specific BE recommendations and provide a meaningful opportunity for the public to consider and comment on those recommendations. Under that process, draft recommendations are posted on FDA's Web site and announced periodically in the **Federal Register**. The public is encouraged to submit comments on those recommendations within 60 days of their announcement in the **Federal Register**. FDA considers any comments received and either publishes final recommendations, or publishes revised draft recommendations for comment. Recommendations were last announced in the **Federal Register** of June 8, 2009 (74 FR 27146). This notice announces draft product-specific recommendations, either new or revised, that have been posted on FDA's Web site in the period from November 1, 2008, through December 1, 2009.

II. Drug Products for Which New Draft Product-Specific BE Recommendations Are Available

FDA is announcing draft BE product-specific recommendations for drug products containing the following active ingredients:

A

Adapalene (multiple reference listed drugs (RLDs))
Adapalene; Benzoyl Peroxide
Alendronate Sodium; Cholecalciferol
Aliskiren Hemifumarate
Aliskiren Hemifumarate; Hydrochlorothiazide
Allopurinol
Ambrisentan
Amlodipine Besylate; Atorvastatin Calcium
Atenolol

B

Bromfenac Sodium
Bromocriptine
Budesonide

C

Calcium Acetate
Cephalexin
Chlorpheniramine Polistirex; Hydrocodone Polistirex
Ciprofloxacin
Clonidine
Clotrimazole (multiple RLDs)

D

Desmopressin Acetate
Desogestrel; Ethinyl Estradiol (multiple RLDs)
Desvenlafaxine Succinate
Dextroamphetamine Sulfate
Dextromethorphan Hydrobromide; Guaifenesin
Diclofenac Sodium (multiple RLDs)
Doxycycline Hyclate
Drospirenone; Ethinyl Estradiol

E

Eletriptan Hydrobromide