forward a dose reconstruction report to DOL and to the claimant, and closes the record on data used for the dose reconstruction. The dose reconstruction results will be supplied to the claimant and to the DOL which will factor them into its determination whether the claimant is eligible for compensation under the Act.

On October 31, 2001, the Office of Management and Budget approved DHHS' request for emergency Paperwork Reduction Act clearance, so that NIOSH could begin its dose reconstruction duties under the Act. That emergency clearance expires on April 30, 2002. This notice pertains to DHHS request for normal Paperwork Reduction Act clearance to permit NIOSH to continue conducting dose reconstruction activities after April 30, 2002. The total annual burden for this data collection is 16,250 hours.

Respondents	Number of respondents	Number of responses	Average burden per response (in hrs)
Initial interview	15,000	1	60/60
Conclusion form	15,000	1	5/60

Dated: February 1, 2002.

Julie Fishman,

Acting Deputy Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 02–3151 Filed 2–8–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services

[Document Identifier: CMS–10051]

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

AGENCY: Centers for Medicare and Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: New Collection; *Title of Information Collection:* Evaluation of the MassHealth Insurance Partnership; *Form No.:* CMS–10051 (OMB# 0938–NEW); *Use:* This collection will be used to evaluate the Massachusetts' 1115 Waiver Demonstration, including Insurance Partnership program, offering subsidies to small employers to encourage them to offer health insurance coverage to employees. The purpose of the survey is to determine the factors influencing an employer's decision to participate or not, in the IP program and their respective characteristics.; *Frequency:* Other: One-time; *Affected Public:* Business or other for-profit, Not-for-profit institutions, and Farms; *Number of Respondents:* 2,016; *Total Annual Responses:* 2,016; *Total Annual Hours:* 336.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS' Web Site address at *http://www.hcfa.gov/regs/prdact95.htm*, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov*, or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, D.C. 20503.

Dated: January 8, 2002.

Dawn M. Willinghan,

Acting, CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–3252 Filed 2–8–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0036]

Aventis Pharmaceuticals et al.; Withdrawal of Approval of 12 New Drug Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of 12 new drug applications (NDAs). The holders of the applications notified the agency in writing that the drug products were no longer marketed and requested that the approval of the applications be withdrawn.

DATES: Effective March 13, 2002.

FOR FURTHER INFORMATION CONTACT: Florine P. Purdie, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: The holders of the applications listed in the table in this document have informed FDA that these drug products are no longer marketed and have requested that FDA withdraw approval of the applications. The applicants have also, by their request, waived their opportunity for a hearing.

NDA No.	Drug	Applicant
8–102	Tace (chlorotrianisene).	Aventis Pharmaceuticals, 399 Interpace Pkwy., P.O. Box 663, Parsippany, NJ 07054.
9–925	Dyclone (dyclonine hydrochloride (HCl)) Topical Solution, 0.5% and 1%.	AstraZeneca LP, 1800 Concord Pike, P.O. Box 8355, Wilmington, DE 19803–8355.
11–444	Tace (chlorotrianisene) Capsules, 25 milligrams (mg).	Aventis Pharmaceuticals
14–322	Meprobamate Tablets, 200 mg and 400 mg.	IMPAX Laboratories, Inc., 30831 Huntwood Ave., Hayward, CA 94544.
16–235	Tace (chlorotrianisene) Capsules, 72 mg.	Aventis Pharmaceuticals
17–829	Diprosone (betamethasone dipropionate) Aerosol.	Schering Corp., 2000 Galloping Hill Rd., Kenilworth, NJ 07033.
19–188	Gastrocrom (cromolyn sodium) Capsules.	Celltech Pharmaceuticals, Inc., 755 Jefferson Rd., P.O. Box 31710, Rochester, NY 14603–1710.
19–399	Total Parenteral Nutrition Electrolytes.	Abbott Laboratories, D–389 Bldg. AP30, 200 Abbott Park Rd., Abbott Park, IL 60064–3537.
20–227	Normiflo (ardeparin sodium) Injection.	Pharmacia & Upjohn Co., 7000 Portage Rd., Kalamazoo, MI 49001–0199.
50–370	Ilotycin Gluceptate (erythromycin gluceptate).	Eli Lilly and Co., Lilly Corp. Center, Indianapolis, IN 46285.
50–579	Monocid (cefonicid sodium) Injection.	SmithKline Beecham Pharmaceuticals, One Franklin Plaza, P.O. Box 7929, Philadelphia, PA 19101–7929.
50–581	Mefoxin (cefoxitin sodium) Premixed IV Solution.	Merck & Co., Inc., P.O. Box 4, BLA-20, West Point, PA 19486.

Therefore, under section 505(e) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(e)) and under authority delegated to the Director, Center for Drug Evaluation and Research (21 CFR 5.82), approval of the applications listed in the table in this document, and all amendments and supplements thereto, is hereby withdrawn, effective March 13, 2002.

Dated: January 18, 2002.

Steven K. Galson,

Deputy Director, Center for Drug Evaluation and Research.

[FR Doc. 02–3199 Filed 2–8–02; 8:45 am]

BILLING CODE 4160–02–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02F–0042]

Ecolab, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Ecolab, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of a mixture of peroxyacetic acid, octanoic acid, acetic acid, hydrogen peroxide, peroxyoctanoic acid, and 1-hydroxyethylidene-1,1-diphosphonic acid as an antimicrobial agent on meat parts, trim, and organs.

DATES: Submit written comments on the petitioner's environmental assessment by March 13, 2002.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Andrew D. Laumbach, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 202–418–3071.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 2A4731) has been filed by Ecolab, Inc., Ecolab Center, 370 Wabasha St., St. Paul, MN 55102. The petition proposes to amend the food additive regulations in Part 173 *Secondary Direct Food Additives Permitted in Food for Human Consumption* (21 CFR part 173) to provide for the safe use of a mixture of peroxyacetic acid, octanoic acid, acetic acid, hydrogen peroxide, peroxyoctanoic acid, and 1-hydroxyethylidene-1,1-diphosphonic acid as an antimicrobial agent on meat parts, trim, and organs.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may submit to the Dockets Management Branch written comments by March 13, 2002. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.51(b).

Dated: January 22, 2002.

L. Robert Lake,

Director of Regulations and Policy, Center for Food Safety and Applied Nutrition.

[FR Doc. 02–3139 Filed 2–8–02; 8:45 am]

BILLING CODE 4160–01–S