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.