

disease in a defined geographic area, the contribution of chronic viral hepatitis to the burden of disease, and the influence of etiologic agent(s) and other factors on mortality, and to monitor the incidence of and mortality from chronic liver disease over time. The information gathered will be analyzed in conjunction with data collected from

other sources to address these questions. The results of the project will assist the Hepatitis Branch, Division of Viral and Rickettsial Diseases, National Center for Infectious Diseases in accomplishing the part of its mission related to preparing recommendations for the prevention and control of all types of viral hepatitis and their sequella. In order to focus on

prevention efforts and resource allocation, a representative view of the overall burden of chronic liver disease, its natural history, and the relative contribution of viral hepatitis is needed. The total annualized estimated cost to respondents is \$5,000.

Respondents	No. of respondents	No. of responses per respondent	Average burden per response in hours	Total burden in hours
All Consenting adults with physician-diagnosed chronic liver disease living in catchment areas	500	1	1	500
Total				500

Dated: February 22, 2001.

Nancy Cheal,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1501]

Agency Information Collection Activities; Announcement of OMB Approval; Threshold of Regulation for Substances Used in Food-Contact Articles

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Threshold of Regulation for Substances Used in Food-Contact Articles" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 26, 2000 (65 FR 81530), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control

number. OMB has now approved the information collection and has assigned OMB control number 0910-0298. The approval expires on February 29, 2004. A copy of the supporting statement for this information collection is available on the Internet at <http://www.fda.gov/ohrms/dockets>.

Dated: February 23, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01-5017 Filed 3-1-01; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1379]

Agency Information Collection Activities; Announcement of OMB Approval; Procedures for the Safe Processing and Importing of Fish and Fishery Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Procedures for the Safe Processing and Importing of Fish and Fishery Products" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 21, 2000 (65 FR

45382), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0354. The approval expires on February 29, 2004. A copy of the supporting statement for this information collection is available on the Internet at <http://www.fda.gov/ohrms/dockets>.

Dated: February 23, 2001.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 01-5018 Filed 3-1-01; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1521]

Agency Information Collection Activities; Announcement of OMB Approval; Food Labeling Regulations (21 CFR Parts 101, 102, 104, and 105)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Food Labeling Regulations (21 CFR Parts 101, 102, 104, and 105)" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.