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Certifier	J. Woodward

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. 98N-1110]

**Agency Information Collection Activities; Announcement of OMB Approval; Current Good Manufacturing Practice Regulations for Finished Pharmaceuticals**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "CGMP Regulations for Finished Pharmaceuticals" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

**FOR FURTHER INFORMATION CONTACT:** Karen L. Nelson, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1482.

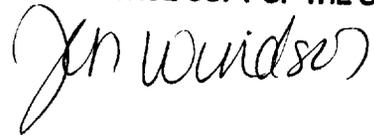
**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of April 19, 1999 (64 FR 19180), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond 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 0910-0139. The approval expires on June 30, 2002. A copy of the supporting statement for this information collection is available on the Internet at "<http://www.fda.gov/ohrms/dockets>".

Dated: 6/30/99  
June 30, 1999

  
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Margaret M. Dotzel  
Acting Associate Commissioner for  
Policy Coordination

**CERTIFIED TO BE A TRUE COPY OF THE ORIGINAL**



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