TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section/Form No.	No. of Respondents	Number of Responses per Respondent	Total Annual Responses	Hours per Response	Total Hours
FDA–2656 (Registration of Drug Establishment) 207.21 207.22 207.25 207.26 207.40	18,430	.36	6,700	2.50	16,750
FDA–2656 (Annual Update of Drug Establishment) 207.21 207.22 207.25 207.26 207.40	8,382	.82	6,859	2.50	17,147.50
FDA–2657 (Drug Product Listing) 207.21 207.22 207.25 207.30 207.31 207.40	15,530	3	46,713	2.50	116,782.50
FDA–2658 (Registered Establishments' Report of Private Label Distributors) 207.21 207.22 207.25 207.30 207.31	7,216	2.14	15,415	2.50	38,537.50
Total Reporting Burden					189,217.50

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: March 29, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–7907 Filed 4–7–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0463]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Infant Formula Requirements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Infant Formula Requirements'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

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

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 13, 2004 (69 FR 1985), 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 0910–0256. The approval expires on March 31, 2007.

A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: April 2, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–8024 Filed 4–7–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0507]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Study of Trans Fat Claims on Food

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Experimental Study of Trans Fat Claims on Food'' has been approved by the Office of Management and Budget