shell eggs, game meat and game meat products, gelatin, and collagen.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Products	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Shell Eggs	10	1	10	0.25 (15 minutes)	3
Game Meat and Game Meat Products	5	1	5	0.25 (15 minutes)	1
Gelatin	7	1	7	0.25 (15 minutes)	2
Collagen	18	1	18	0.25 (15 minutes)	5
Total					11

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

We base our estimates of the number of respondents and total annual responses on the submissions that we have received in the past 3 years for each product type. To calculate the estimate for the hours per response values, we assumed that the information requested is readily available to the submitter. We expect that the submitter will need to gather information from appropriate persons in the submitter's company and to prepare this information for submission. We believe that this effort should take no longer than 15 minutes (0.25 hour) per response. We estimate that we will receive 1 submission from 10 shell egg producers annually, for a total of 10 annual responses. Each submission is estimated to take 0.25 hour per response for a total of 2.5 hours, rounded to 3 hours. This collection has previously covered information collected to maintain lists of eligible exporters of dairy products; dairy products will be covered under OMB control number 0910–0509, so the estimated burden has been removed from this collection. We estimate that we will receive one submission from five game meat and game meat product producers annually, for a total of five annual responses. Each submission is estimated to take 0.25 hour per response for a total of 1.25 hours, rounded to 1 hour. We estimate that we will receive one submission from seven gelatin producers annually, for a total of seven annual responses. Each submission is estimated to take 0.25 hour per response for a total of 1.75 hours, rounded to 2 hours. We estimate that we will receive one submission from 18 collagen producers annually, for a total of 18 annual responses. Each submission is estimated to take 0.25 hour per response for a total of 4.5 hours, rounded to 5 hours. The estimated burden for collagen producers includes animal casings, which have been listed separately in previous notices. Therefore, the proposed annual burden for this information collection is 11 hours.

Dated: September 28, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–23930 Filed 10–3–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2013–N–0115; FDA–2013–N–0717]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, Three White Flint North, 11601 Landsdown St., North Bethesda, MD 20852, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507). The OMB control number and expiration date of OMB approval for each information collection are shown in table 1. Copies of the supporting statements for the information collections are available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.* 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.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control No.	Date approval expires
Manufactured Food Regulatory Program Standards	0910–0601	9/30/2019
Evaluation of the Food and Drug Administration's General Market Youth Tobacco Prevention Campaign	0910–0753	9/30/2019

Dated: September 27, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–23898 Filed 10–3–16; 8:45 am]

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