reading the news article, participants will complete a questionnaire assessing their emotional response, appraisals, attribution of responsibility, perceptions about the safety of the affected produce, intentions to grow, sell, or buy the affected produce, perceived probability of a repeat event, and a measure of their innate ability to effectively respond to the information in the article.

To help design and refine the questionnaire, we will recruit 25 participants in order to conduct 10 cognitive interviews. We estimate cognitive interview recruitment will take 5 minutes (0.083 hours), for a total of 2 hours. The cognitive interviews are estimated at 1 hour per response for a total of 10 hours for the cognitive interview activities. We expect to send screeners to 800 members of a consumer panel, each taking 2 minutes (0.03 hours) to complete, for a total of 24 hours for the consumer panel screener activity. We also expect to administer 360 screeners to growers and retailers, each taking 2 minutes (0.033 hours) to complete, for a total of 24 hours (11 + 11 = 22). Twenty-four participants (20 consumers, 2 growers, 2 retailers) will complete the pre-test. Each pre-test will take 10 minutes (0.17 hours) for a total of 5 hours for the pre-test activity. We estimate that 900 individuals (540 consumers, 180 growers, and 180 retailers) will complete the questionnaire for the experiment, each taking 10 minutes (0.17 hours) for a total of 153 hours for the experimental study activities. The estimated total hour burden of the collection of information is 215 hours.

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

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Portion of study	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response (in hours) [2]	Total hours
Cognitive Interview Recruitment	25	1	25	5/60	2
Cognitive Interviews	10	1	10	1	10
Consumer Panel Screener	800	1	800	2/60	24
Grower Screener	360	1	360	2/60	11
Retailer Screener	360	1	360	2/60	11
Pre-tests	24	1	24	10/60	5
Experiment	900	1	900	10/60	153
Total					216

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] Burden estimates of less than 1 hour are expressed as a fraction of an hour in the format "[number of minutes per response]/60".

II. References

11. Olsen, S., L. MacKinon, *et al.,* "Surveillance for Foodborne Disease Outbreaks—United States, 1993 to1997," *Morbidity and Mortality Weekly Report* 49(SS01), pp. 1 through 51, 2000.
2. FDA 101: Product Recalls—From First Alert to Effectiveness Checks, Available at *http://www.fda.gov/ForConsumers/ConsumerUpdates/ucm049070.htm.*
3. Calvin, L., "Outbreak Linked to Spinach Forces Reassessment of Food Safety Practices," *Amber Waves* 5(3), pp. 24 through 31, 2007.
4. Lucier, G. and R. Dettmann, "Vegetables and Melons Outlook," A Report From the United States Department of Agriculture, Economic Research Service, VGS–327, June 26, 2008.

Dated: April 11, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–9155 Filed 4–14–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–E–0241]

Determination of Regulatory Review Period for Purposes of Patent Extension; ATRYN; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of March 21, 2011 (76 FR 15323). The document announced the determination of the regulatory review period for ATRYN. The document was published with an incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Planning and Budget, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3208, Silver Spring, MD 20993–0002, 301–796–9138.

SUPPLEMENTARY INFORMATION: In FR Doc. 2011–6509, appearing on page 15323, in the **Federal Register** of Monday, March 21, 2011, the following correction is made:

1. On page 15323, in the first column, in the Docket No. heading, "[Docket No. FDA–2010–E–0241]" is corrected to read "[Docket No. FDA–2009–E–0241]".

Dated: April 8, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–9153 Filed 4–14–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0002]

Pediatric Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Pediatric Advisory Committee.