TABLE 2—ESTIMATED ANNUAL REPORTING BURDEN [1]—Continued

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Pretest	640	1	640	0.75 (45 minutes)	480
Main Study Screener	2,112	1	2,112	0.08 (5 minutes)	169
Main Study	1,056	1	1,056	0.75 (45 minutes)	792
Total	5,193	1	5,193		1,560

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

V. References

The following references are on display in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, and are available for viewing by interested persons between 9 a.m. and 4 p.m., Monday through Friday; they are also available electronically at *http://www.regulations.gov.* FDA has verified the Web site addresses, as of the date this document publishes in the **Federal Register**, but Web sites are subject to change over time.

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Dated: December 15, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–32251 Filed 12–22–15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0873]

Agency Information Collection Activities: Proposed Collection; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice entitled ''Agency Information Collection Activities: Proposed Collection; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products'' that appeared in the **Federal Register** of December 15, 2015 (80 FR 77637). The document solicited comments on the bar code label requirements for human drug and biological products. The document was published with an incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Lisa Granger, Office of Policy and Planning, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3330, Silver Spring, MD 20993–0002, 301–796–9115.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of Tuesday, December 15, 2015, in FR Doc. 2015–31402, the following correction is made:

1. On page 77637, in the second column, the docket number is corrected to read FDA–2012–N–0873.

Dated: December 15, 2015.

Leslie Kux,

Associate Commissioner for Policy.

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