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.


Leslie Kux.
Associate Commissioner for Policy.

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.


Leslie Kux.
Associate Commissioner for Policy.

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