4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 310, 314, 329, and 600[Docket No. FDA-2008-N-0334]

RIN 0910-AF96

Postmarketing Safety Reports for Human Drug and Biological Products; Electronic Submission Requirements; Corrections

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; corrections.

SUMMARY: The Food and Drug Administration (FDA) is correcting a document entitled "Postmarketing Safety Reports for Human Drug and Biological Products; Electronic Submission Requirements; Correction" that appeared in the Federal Register of August 14, 2014 (79 FR 47655). The document published without the required RIN number and in the Notice category. This document corrects those errors.

DATES: Effective [INSERT DATE OF PUBLICATION IN THE FEDERAL REGISTER].

FOR FURTHER INFORMATION CONTACT: Jean Chung, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, rm. 4466, Silver Spring, MD 20993-0002, 301-796-1874; or Stephen Ripley, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, rm. 7268, Silver Spring, MD 20993-0002, 240-402-7911.

SUPPLEMENTARY INFORMATION:

In the Federal Register of August 14, 2014, in FR Doc. 2014-19255, the following correction is made:

1. On page 47655, in the first column, add the heading "RIN 0910-AF96" between the Docket No. and the title of the document.

Dated: September 2, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

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