4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 864

[Docket No. FDA-2018-N-0399]

Medical Devices; Hematology and Pathology Devices; Classification of Lynch Syndrome Test Systems; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final order; correction.

SUMMARY: The Food and Drug Administration is correcting a final order entitled "Medical Devices; Hematology and Pathology Devices; Classification of Lynch Syndrome Test Systems" that appeared in the *Federal Register* of February 27, 2018. The document was published with the incorrect docket number. This document corrects that error.

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

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 February 27, 2018 (83 FR 8355), in FR Doc. 2018-03924, on page 8355, the following correction is made:

1. On page 8355, in the third column, in the header of the document, the docket number is corrected to read "FDA-2018-N-0399".

Dated: March 8, 2018.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2018-05115 Filed: 3/13/2018 8:45 am; Publication Date: 3/14/2018]