

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 807

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Certifier J. Hawkins

**Medical Device Reports; Reports of Corrections and Removals;
Establishment Registration and Device Listing: Premarket Approval
Supplements; Quality System Regulation; Importation of Electronic
Products; Technical Amendment; Correction**

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of April 8, 2004 (69 FR 18472). That document corrected a final rule that appeared in the **Federal Register** of March 10, 2004 (69 FR 11310). The April 8, 2004, document published with inadvertent errors. This document corrects those errors.

DATES: This rule is effective [*insert date of publication in the Federal Register*].

FOR FURTHER INFORMATION CONTACT: Joyce A. Strong, Office of Policy and Planning (HF-27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 04-8022, appearing on page 18472 in the **Federal Register** of Thursday, April 8, 2004, the following corrections are made:

1. On page 18472, in the third column, under the **FOR FURTHER INFORMATION CONTACT** heading, the address is corrected to read "Joyce A. Strong, Office of

Policy and Planning (HF-27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-7010.”

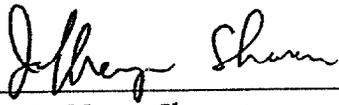
§ 807.22 [Corrected]

2. On page 18473, in the first column, in § 807.22, in paragraph (a), the

first sentence is corrected to read "The first registration of a device establishment shall be on Form FDA-2891 (Initial Registration of Device Establishment)."

Dated: 4-29-04

April 29, 2004.



Jeffrey Shuren,
Assistant Commissioner for Policy.

[FR Doc. 04-????? Filed ??-??-04; 8:45 am]

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Dawn P. Hawkins