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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 807

[Docket No. 98N-0520]

Medical Devices; Establishment Registration and Device Listing for Manufacturers and Distributors of Devices; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a proposed rule that appeared in the **Federal Register** of September 29, 1998 (63 FR 51874). The document proposed to amend certain regulations governing establishment registration and device listing by domestic distributors. The document was published with an error. This document corrects that error.

EFFECTIVE DATE: February 11, 1999.

FOR FURTHER INFORMATION CONTACT: Walter W. Morgenstern, Center for Devices and Radiological Health (HFZ-305), Food and Drug Administration, 2094 Gaither Rd., Rockville, MD 20857, 301-594-4699.

SUPPLEMENTARY INFORMATION: In FR Doc. 98-25797 appearing on page 51874 in the **Federal Register** of September 29, 1998, the following correction is made:

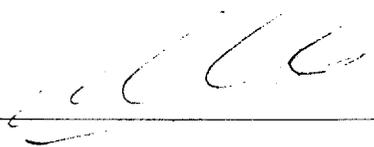
On page 51875, in the third column, amendatory paragraph four is corrected to read:

4. Section 807.20 is amended by revising paragraph (a)(4), by removing paragraph (c), by redesignating paragraph (d) as paragraph (c), and by adding paragraph (c)(3) to read as follows:

* * * * *

Dated: November 19, 1998

November 19, 1998



William K. Hubbard
Associate Commissioner for Policy Coordination

[FR Doc. 98-???? Filed ??-??-98; 8:45 am]

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