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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

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Certifier D. Hawkins

[Docket No. FDA-2007-P-0300] (formerly 2007P-0326)

**Determination That SANOREX (Mazindol) Tablets 1 and 2 Milligrams Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness; Correction**

**AGENCY:** Food and Drug Administration. HHS.

**ACTION:** Notice; correction.

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**SUMMARY:** The Food and Drug Administration is correcting a notice that appeared in the **Federal Register** of July 15, 2008 (73 FR 40582). The document announced the determination that SANOREX (mazindol) Tablets, 1 and 2 milligrams (mg), were not withdrawn from sale for reasons of safety or effectiveness. The document was published with an incorrect docket number. This document corrects that error.

**FOR FURTHER INFORMATION CONTACT:** Joyce 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. E8-15998, appearing on page 40582 in the **Federal Register** of Tuesday, July 15, 2008, the following correction is made:

1. On page 40582, in the third column, in the headings section of the document, “[Docket No. FDA-2007-P-0326]” is corrected to read “[Docket No. FDA-2007-P-0300] (formerly 2007P-0326)”.

Dated: JUL 22 2008

July 22, 2008.

Jeffrey Shuren

Jeffrey Shuren,  
Associate Commissioner for Policy and Planning.

[FR Doc. 08-????? Filed ??-??-08; 8:45 am]

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