

K800364 ULTRASONIC DIAGNOSTIC SYSTEM, MOD. 202Mar 12, 1980
21 days to decisionK800364 · Product code: **DKX** · Toxicology
Source: <https://www.510kdatabase.net/k800364/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thin Layer Chromatography, Barbiturate (DKX)
Date received	Feb 20, 1980
Decision date	Mar 12, 1980
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Hoffrel Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1978-1987

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Device record: <https://www.510kdatabase.net/k800364/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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