

**K925434 SECTOR & TRANSVAGINAL MODEL #'S
SW4/5B/A,EW5/7K/A**Sep 30, 1993
337 days to decisionK925434 · Product code: ITX · Radiology
Source: <https://www.510kdatabase.net/k925434/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Ultrasonic, Diagnostic (ITX)
Date received	Oct 28, 1992
Decision date	Sep 30, 1993
Days to decision	337 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kretz Technik
Location	Washington, DC, US
Contact	PATRICIA B SHRADER
510(k) history	1 submissions · 1 cleared · 1993-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925434/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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