FDA 510(k) Premarket Notification

K925434 SECTOR & TRANSVAGINAL MODEL #'S SW4/5B/A,EW5/7K/A

CLEARED

Sep 30, 1993
337 days to decision

K925434 · 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