

K962142 SONOLINE VERSA 555 DIAGNOSTIC ULTRASOUND SYSTEM

May 29, 1997
360 days to decision

K962142 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k962142/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jun 3, 1996
Decision date	May 29, 1997
Days to decision	360 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	STEVE HESLER
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k962142/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026