FDA 510(k) Premarket Notification

K951695 OPERAND* ULTRASOUND TRANSMISSION GEL

CLEARED

May 12, 1995
30 days to decision

K951695 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k951695/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Apr 12, 1995
Decision date	May 12, 1995
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Redi Products, Inc.**
Location	Prichard, WV, US
Contact	JOSEPH LANGDON
510(k) history	1 submissions · 1 cleared · 1995-1995

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k951695/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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