FDA 510(k) Premarket Notification

K810133 ULTRAPROBE

CLEARED

Jun 9, 1981
141 days to decision

K810133 · Product code: **IYN** · Radiology
Source: https://www.510kdatabase.net/k810133/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Doppler, Ultrasonic (IYN)
Date received	Jan 19, 1981
Decision date	Jun 9, 1981
Days to decision	141 days
Third-party review	No

APPLICANT

Company	**Ars Magna**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810133/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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