FDA 510(k) Premarket Notification

K863387 MULTIPLE

CLEARED

Sep 4, 1987
366 days to decision

K863387 · Product code: **DXK** · Cardiovascular
Source: https://www.510kdatabase.net/k863387/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Sep 3, 1986
Decision date	Sep 4, 1987
Days to decision	366 days
Third-party review	No

APPLICANT

Company	**Del Mar Avionics**
Location	Mchenry, IL, US
Contact	DAVID R JOHNSON
510(k) history	54 submissions · 54 cleared · 1977-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k863387/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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