FDA 510(k) Premarket Notification

K811638 DATASCOPE INTERNAL DEFIBRILLATOR PADDLES

CLEARED

Jul 10, 1981
30 days to decision

K811638 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k811638/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jun 10, 1981
Decision date	Jul 10, 1981
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019