FDA 510(k) Premarket Notification

K844103 CARDIOPAC DEFIBRILLATOR 444

CLEARED

Jan 18, 1985
87 days to decision

K844103 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k844103/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 23, 1984
Decision date	Jan 18, 1985
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Birtcher Corp.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1988