FDA 510(k) Premarket Notification

K880575 CARDIOLIFE TEC-7300 W/OPTIONAL ACCESSORIES

CLEARED

Aug 2, 1988
174 days to decision

K880575 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k880575/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 10, 1988
Decision date	Aug 2, 1988
Days to decision	174 days
Third-party review	No

APPLICANT

Company	**Nihon Kohden America, Inc.**
Location	Foothill Ranch, CA, US
Contact	MIKE DASHEFSKY
510(k) history	166 submissions · 163 cleared · 1979-2012