FDA 510(k) Premarket Notification

K863405 CARDIOLIFE TEC 7200

CLEARED

Oct 31, 1986
58 days to decision

K863405 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k863405/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 3, 1986
Decision date	Oct 31, 1986
Days to decision	58 days
Third-party review	No

APPLICANT

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