FDA 510(k) Premarket Notification

CLEARED

K981737 K-DEFIB/PACE ADULT ELECTRODE MODEL NUMBER KDP-60

Dec 9, 1998
205 days to decision

K981737 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k981737/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	May 18, 1998
Decision date	Dec 9, 1998
Days to decision	205 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Katecho, Inc.**
Location	Des Moines, IA, US
Contact	LORNE S CHARNBERG
510(k) history	26 submissions · 25 cleared · 1984-2001