

**K981737 K-DEFIB/PACE ADULT ELECTRODE MODEL NUMBER
KDP-60**Dec 9, 1998
205 days to decisionK981737 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k981737/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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