

K863899 K-STIK DEFIBRILLATION PADSNov 12, 1986
36 days to decisionK863899 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k863899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 7, 1986
Decision date	Nov 12, 1986
Days to decision	36 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k863899/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026