

K103159 CARDIO MED MODEL A102, H104, O101, P103, AND Z100Jan 11, 2011
77 days to decisionK103159 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k103159/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 26, 2010
Decision date	Jan 11, 2011
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardio Medical Products, Inc.
Location	Rockaway, NJ, US
Contact	NICK MENDISE
510(k) history	3 submissions · 3 cleared · 2011-2011

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