

**K914971 TEC-8251A DEFIBRILLATOR MONITOR,
MODIFICATION**Dec 4, 1991
47 days to decisionK914971 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k914971/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 18, 1991
Decision date	Dec 4, 1991
Days to decision	47 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914971/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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