

K890988 MODEL 3110 HEART WIRE & 3110P PEDIATRIC HEART WIREAug 25, 1989
179 days to decisionK890988 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k890988/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Feb 27, 1989
Decision date	Aug 25, 1989
Days to decision	179 days
Third-party review	No

APPLICANT

Company	Pace Medical
Location	Waltham, MA, US
Contact	RALPH E HANSON
510(k) history	19 submissions · 19 cleared · 1987-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890988/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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