

**K883930 SEAMED MODEL 3300 DUAL CHAMBER PACING
SYSTEM ANALY**Nov 9, 1988
55 days to decisionK883930 · Product code: **DTE** · Cardiovascular
Source: <https://www.510kdatabase.net/k883930/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Sep 15, 1988
Decision date	Nov 9, 1988
Days to decision	55 days
Third-party review	No

APPLICANT

Company	Seamed Corp.
Location	Walker, MI, US
Contact	MARCIA A W.-PAGE
510(k) history	18 submissions · 18 cleared · 1982-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883930/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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