

K823048 SEA MED MODEL 880Dec 15, 1982
61 days to decisionK823048 · Product code: **DTE** · CardiovascularSource: <https://www.510kdatabase.net/k823048/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Oct 15, 1982
Decision date	Dec 15, 1982
Days to decision	61 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k823048/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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