

K831381 RAPID ATRIAL PACERAug 31, 1983
125 days to decisionK831381 · Product code: **DTE** · CardiovascularSource: <https://www.510kdatabase.net/k831381/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Apr 28, 1983
Decision date	Aug 31, 1983
Days to decision	125 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/k831381/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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