FDA 510(k) Premarket Notification

K831381 RAPID ATRIAL PACER

CLEARED

Aug 31, 1983
125 days to decision

K831381 · Product code: **DTE** · Cardiovascular
Source: 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