FDA 510(k) Premarket Notification

K823048 SEA MED MODEL 880

CLEARED

Dec 15, 1982
61 days to decision

K823048 · Product code: **DTE** · Cardiovascular
Source: 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