FDA 510(k) Premarket Notification

K813174 BRADYCARDIA ALARM

CLEARED

Jan 22, 1982
77 days to decision

K813174 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k813174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 6, 1981
Decision date	Jan 22, 1982
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Douglas Scientific Products**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1982