FDA 510(k) Premarket Notification

K790644 SUBMAX

CLEARED

Apr 26, 1979
27 days to decision

K790644 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k790644/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 30, 1979
Decision date	Apr 26, 1979
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Chase Mfg. Co.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1977-1979