FDA 510(k) Premarket Notification

K820745 DIASCOPE 2

CLEARED

Apr 1, 1982
13 days to decision

K820745 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k820745/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Mar 19, 1982
Decision date	Apr 1, 1982
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Puritan Bennett Corp.**
Location	Mchenry, IL, US
510(k) history	110 submissions · 101 cleared · 1976-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820745/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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