FDA 510(k) Premarket Notification

K872443 COMPANION 50 REGULATOR/FLOWMETER

CLEARED

Jul 21, 1987
28 days to decision

K872443 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k872443/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 23, 1987
Decision date	Jul 21, 1987
Days to decision	28 days
Third-party review	No

APPLICANT

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