FDA 510(k) Premarket Notification

K900682 PHANTOM RADIOTRANSLUCENT ECG MONITORING ELECTRODE

CLEARED

Jul 31, 1990
168 days to decision

K900682 · Product code: **DRY** · Cardiovascular
Source: https://www.510kdatabase.net/k900682/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Feb 13, 1990
Decision date	Jul 31, 1990
Days to decision	168 days
Third-party review	No

APPLICANT

Company	**Contour Electrodes, Inc.**
Location	Herrin, IL, US
Contact	PARIS WALKER
510(k) history	4 submissions · 4 cleared · 1987-1990