FDA 510(k) Premarket Notification

K810154 LIFEPAK 7 ACUTE CARDIAC CARE SYSTEM

CLEARED

Mar 26, 1981
64 days to decision

K810154 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k810154/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jan 21, 1981
Decision date	Mar 26, 1981
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Physio-Control Corp.**
Location	Mchenry, IL, US
510(k) history	80 submissions · 78 cleared · 1976-1999