FDA 510(k) Premarket Notification

K760676 LIFEPAK 5 CARDIOSCOPE-RECORDER MODULE

CLEARED

Oct 13, 1976
23 days to decision

K760676 · Product code: **DSF** · Cardiovascular
Source: https://www.510kdatabase.net/k760676/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Sep 20, 1976
Decision date	Oct 13, 1976
Days to decision	23 days
Third-party review	No

APPLICANT

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