FDA 510(k) Premarket Notification

K760563 ECG MODULATOR

CLEARED

Sep 10, 1976
8 days to decision

K760563 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k760563/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Sep 2, 1976
Decision date	Sep 10, 1976
Days to decision	8 days
Third-party review	No

APPLICANT

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