FDA 510(k) Premarket Notification

K771447 PREGELLED DISP. EKG MONITORING ELECT-

CLEARED

Aug 4, 1977
2 days to decision

K771447 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k771447/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 2, 1977
Decision date	Aug 4, 1977
Days to decision	2 days
Third-party review	No

APPLICANT

Company	**Hayes Products, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1979