FDA 510(k) Premarket Notification

K791140 PREGELLED DISPOSABLE EELCTRO. SERIES 600

CLEARED

Jul 10, 1979
22 days to decision

K791140 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k791140/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 18, 1979
Decision date	Jul 10, 1979
Days to decision	22 days
Third-party review	No

APPLICANT

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