

K771447 PREGELLED DISP. EKG MONITORING ELECT-Aug 4, 1977
2 days to decisionK771447 · Product code: **DRX** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k771447/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026