FDA 510(k) Premarket Notification

K832429 CARDIO TRACE ELECTRODE

CLEARED

May 9, 1984
292 days to decision

K832429 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k832429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 22, 1983
Decision date	May 9, 1984
Days to decision	292 days
Third-party review	No

APPLICANT

Company	**Poly-Med Splint Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k832429/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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