

K813176 GEL ELECTRODEDec 29, 1981
47 days to decisionK813176 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k813176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Nov 12, 1981
Decision date	Dec 29, 1981
Days to decision	47 days
Third-party review	No

APPLICANT

Company	Dyn-A-Med Products
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1984

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Device record: <https://www.510kdatabase.net/k813176/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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