

K781518 PREGELLED DISPOSABLE ELECTRODEOct 6, 1978
36 days to decisionK781518 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k781518/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 31, 1978
Decision date	Oct 6, 1978
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Vente Medical Products, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k781518/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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