

K801846 ELECTROCARDIOGRAPH ELECTRODESep 9, 1980
36 days to decisionK801846 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k801846/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 4, 1980
Decision date	Sep 9, 1980
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Phoenix Glove Co.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1978-1983

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Device record: <https://www.510kdatabase.net/k801846/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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