

K821846 LDS MONITORING ELECTRODEJul 30, 1982
38 days to decisionK821846 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k821846/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 22, 1982
Decision date	Jul 30, 1982
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Life Design Systems, Inc.
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1982-1992

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Device record: <https://www.510kdatabase.net/k821846/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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