

K820122 PIONEER II BACK PLATE ELECTRODEMar 1, 1982
42 days to decisionK820122 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k820122/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 18, 1982
Decision date	Mar 1, 1982
Days to decision	42 days
Third-party review	No

APPLICANT

Company	Life Science Instrumentation, Inc.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

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Device record: <https://www.510kdatabase.net/k820122/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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