FDA 510(k) Premarket Notification

K820122 PIONEER II BACK PLATE ELECTRODE

CLEARED

Mar 1, 1982
42 days to decision

K820122 · Product code: **DRX** · Cardiovascular
Source: 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