FDA 510(k) Premarket Notification

K821846 LDS MONITORING ELECTRODE

CLEARED

Jul 30, 1982
38 days to decision

K821846 · Product code: **DRX** · Cardiovascular
Source: 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