FDA 510(k) Premarket Notification

K792305 A SLEEP ASSESSMENT DEVICE

CLEARED

Dec 20, 1979
34 days to decision

K792305 · Product code: **LEL** · Neurology
Source: https://www.510kdatabase.net/k792305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Sleep Assessment (LEL)
Date received	Nov 16, 1979
Decision date	Dec 20, 1979
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Farrall Instruments, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1976-1995