

**K792305 A SLEEP ASSESSMENT DEVICE**Dec 20, 1979  
34 days to decisionK792305 · Product code: **LEL** · Neurology  
Source: <https://www.510kdatabase.net/k792305/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k792305/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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