

K832999 RESPIRATION MONITORFeb 13, 1984
179 days to decisionK832999 · Product code: **FLS** · AnesthesiologySource: <https://www.510kdatabase.net/k832999/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Apnea, Facility Use (FLS)
Date received	Aug 18, 1983
Decision date	Feb 13, 1984
Days to decision	179 days
Third-party review	No

APPLICANT

Company	Litton Medical Electronics
Location	Walker, MI, US
510(k) history	38 submissions · 38 cleared · 1982-1985

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Device record: <https://www.510kdatabase.net/k832999/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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