

K832017 LME BLOOD PRESSURE MONITOROct 31, 1983
130 days to decisionK832017 · Product code: **DSK** · CardiovascularSource: <https://www.510kdatabase.net/k832017/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Blood-pressure (DSK)
Date received	Jun 23, 1983
Decision date	Oct 31, 1983
Days to decision	130 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/k832017/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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