FDA 510(k) Premarket Notification

# K832017 LME BLOOD PRESSURE MONITOR

**CLEARED**

Oct 31, 1983
130 days to decision

K832017 · Product code: **DSK** · Cardiovascular
Source: https://www.510kdatabase.net/k832017/

## SUBMISSION DETAILS

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|---|---|
| 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

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