FDA 510(k) Premarket Notification

# K810153 VENTILATORY EFFORT MONITOR

**CLEARED**

Feb 9, 1981
19 days to decision

K810153 · Product code: **FLS** · Anesthesiology
Source: https://www.510kdatabase.net/k810153/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Monitor, Apnea, Facility Use (FLS) |
| Date received | Jan 21, 1981 |
| Decision date | Feb 9, 1981 |
| Days to decision | 19 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Medicon, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 14 submissions · 14 cleared · 1981-1997 |