

K810153 VENTILATORY EFFORT MONITORFeb 9, 1981
19 days to decisionK810153 · Product code: **FLS** · AnesthesiologySource: <https://www.510kdatabase.net/k810153/>**SUBMISSION DETAILS**

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

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Device record: <https://www.510kdatabase.net/k810153/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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