

K831877 ARF 850 ENov 3, 1983
143 days to decisionK831877 · Product code: **CBK** · Anesthesiology
Source: <https://www.510kdatabase.net/k831877/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jun 13, 1983
Decision date	Nov 3, 1983
Days to decision	143 days
Third-party review	No

APPLICANT

Company	W.E. Leder Enterprises, Inc.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k831877/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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