FDA 510(k) Premarket Notification

K832508 VOLUME VENTILATOR

CLEARED

Dec 27, 1983
153 days to decision

K832508 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k832508/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jul 27, 1983
Decision date	Dec 27, 1983
Days to decision	153 days
Third-party review	No

APPLICANT

Company	**Sechrist Industries, Inc.**
Location	Mchenry, IL, US
510(k) history	23 submissions · 22 cleared · 1978-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k832508/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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