FDA 510(k) Premarket Notification

K833982 MODEL IV 100B-DA INFANT VENTILATOR

CLEARED

Mar 5, 1984
109 days to decision

K833982 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k833982/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Nov 17, 1983
Decision date	Mar 5, 1984
Days to decision	109 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/k833982/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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