FDA 510(k) Premarket Notification

K850563 SECHRIST VENTILATOR/NAUTILUS CHAMBER COMBINATION

CLEARED

Mar 12, 1985
28 days to decision

K850563 · Product code: **CBF** · Anesthesiology
Source: https://www.510kdatabase.net/k850563/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Hyperbaric (CBF)
Date received	Feb 12, 1985
Decision date	Mar 12, 1985
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Nautilus Environmedical Systems, Inc.**
Location	Walker, MI, US
Contact	& EGBE
510(k) history	2 submissions · 2 cleared · 1983-1985

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k850563/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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