FDA 510(k) Premarket Notification

K770249 VENTILATOR SET - CODE 2C7109

CLEARED

Feb 14, 1977
6 days to decision

K770249 · Product code: **BTT** · Anesthesiology
Source: https://www.510kdatabase.net/k770249/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Respiratory Gas, (direct Patient Interface) (BTT)
Date received	Feb 8, 1977
Decision date	Feb 14, 1977
Days to decision	6 days
Third-party review	No

APPLICANT

Company	**Travenol Laboratories, S.A.**
Location	Mchenry, IL, US
Website	https://www.baxter.com
510(k) history	206 submissions · 206 cleared · 1976-1988

Travenol Laboratories, S.A. is a medical device manufacturer based in McHenry, US. The company specializes in infusion, dialysis, and hospital care devices. Travenol Laboratories received FDA 510(k) clearances from total submissions between 1976 and 1988. The company's cleared devices span general hospital and gastroenterology/urology categories, including infusion systems, dialysis equipment, and administration sets. This regulatory record reflects the company's historical focus on critical care and renal therapy technologies. The company is inactive and represents a his...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k770249/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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