FDA 510(k) Premarket Notification

CLEARED

K770571 TRAY, ANESTHESIA, EPIDURAL, CONTINUOUS

Mar 30, 1977
5 days to decision

K770571 · Product code: **CAZ** · Anesthesiology
Source: https://www.510kdatabase.net/k770571/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anesthesia Conduction Kit (CAZ)
Date received	Mar 25, 1977
Decision date	Mar 30, 1977
Days to decision	5 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/k770571/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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