FDA 510(k) Premarket Notification

K870940 SOLUTION ADMINI. SETS- ALTER. DRIP CHAMBER/CONNEC.

CLEARED

Mar 17, 1987
8 days to decision

K870940 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k870940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 9, 1987
Decision date	Mar 17, 1987
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Travenol Laboratories, S.A.**
Location	Mchenry, IL, US
Contact	JULIA A MEYER
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...