

K780223 PARENTERAL INFUSION CONTROLLERApr 10, 1978
60 days to decisionK780223 · Product code: **FLN** · General Hospital
Source: <https://www.510kdatabase.net/k780223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Electric For Gravity Flow Infusion Systems (FLN)
Date received	Feb 9, 1978
Decision date	Apr 10, 1978
Days to decision	60 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...
