

**K860353 SYRINGE INFUSION PUMP**Sep 5, 1986  
218 days to decisionK860353 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k860353/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jan 30, 1986
Decision date	Sep 5, 1986
Days to decision	218 days
Third-party review	No

**APPLICANT**

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Company	<b>Travenol Laboratories, S.A.</b>
Location	McHenry, IL, US
Contact	ROGER A LEROUX
Website	<a href="https://www.baxter.com">https://www.baxter.com</a>
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...

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