

**K853881 THE TRAVENOL 5 ML/HR INFUSOR**Dec 2, 1985  
74 days to decisionK853881 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k853881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Sep 19, 1985
Decision date	Dec 2, 1985
Days to decision	74 days
Third-party review	No

**APPLICANT**

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Company	<b>Travenol Laboratories, S.A.</b>
Location	McHenry, IL, US
Contact	PATRICIA S BARSANTI
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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Device record: <https://www.510kdatabase.net/k853881/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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