

**K854586 HOLLOW FIBER DIALYZER**Dec 13, 1985  
25 days to decisionK854586 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k854586/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Nov 18, 1985
Decision date	Dec 13, 1985
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Travenol Laboratories, S.A.</b>
Location	Mchenry, IL, US
Contact	JULIA A MEYER
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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