

K832663 GAMMA STERIL. CAPILLARY FLOW DIALY-G15Sep 12, 1983
35 days to decisionK832663 · Product code: **FKQ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k832663/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Central Multiple Patient (FKQ)
Date received	Aug 8, 1983
Decision date	Sep 12, 1983
Days to decision	35 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...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k832663/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026