

K832284 SOLUTION ADMIN. SET 2C9096 W/10 PVCDec 29, 1983
169 days to decisionK832284 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k832284/>**SUBMISSION DETAILS**

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| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received | Jul 13, 1983 |
| Decision date | Dec 29, 1983 |
| Days to decision | 169 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...
