

**K780668 SOLUTIONS FOR RESPIRATORY THERAPY**Jun 13, 1978  
53 days to decisionK780668 · Product code: **CAF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k780668/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Apr 21, 1978
Decision date	Jun 13, 1978
Days to decision	53 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k780668/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026