

K923265 FUNNEL ADAPTORMar 18, 1993
259 days to decisionK923265 · Product code: **CAD** · AnesthesiologySource: <https://www.510kdatabase.net/k923265/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vaporizer, Anesthesia, Non-heated (CAD)
Date received	Jul 2, 1992
Decision date	Mar 18, 1993
Days to decision	259 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Southmedic, Inc.
Location	Ancaster, Ontario, CA
Contact	PAUL E DRYDEN
510(k) history	10 submissions · 10 cleared · 1984-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923265/> 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 25, 2026