FDA 510(k) Premarket Notification

K923265 FUNNEL ADAPTOR

CLEARED

Mar 18, 1993
259 days to decision

K923265 · Product code: **CAD** · Anesthesiology
Source: 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