FDA 510(k) Premarket Notification

K973544 VACU*MED, GOLD EDITION CARBON DIOXIDE GAS ANALYZER

CLEARED

May 26, 1998
250 days to decision

K973544 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k973544/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Sep 18, 1997
Decision date	May 26, 1998
Days to decision	250 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vacumed, Div. of Vacumetrics, Inc.**
Location	Salt Lake City, UT, US
Contact	JACQUILINE E FRAILEY
510(k) history	2 submissions · 2 cleared · 1998-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k973544/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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