FDA 510(k) Premarket Notification

K980325 MICROSTREAM NASAL CANNULA FILTERLINE

CLEARED

Apr 16, 1998
78 days to decision

K980325 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k980325/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Jan 28, 1998
Decision date	Apr 16, 1998
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Oridion Medical , Ltd.**
Location	Jerusalem, IL
Contact	SANFORD BROWN
510(k) history	4 submissions · 4 cleared · 1998-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k980325/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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