

**K060065 CAPNOSTREAM20 TWO PARAMETER BEDSIDE
MONITOR & CAPNOSTREAM10 ONE PARAMETER BEDSIDE
MONITOR**

May 4, 2006
115 days to decision

K060065 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k060065/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 9, 2006
Decision date	May 4, 2006
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Oridion Capnography, Inc.
Location	Jerusalem, IL
Contact	SANFORD BROWN
510(k) history	5 submissions · 5 cleared · 2006-2011

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k060065/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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