

K072323 MASTERSCREEN & OXYCON CPXOct 10, 2007
51 days to decisionK072323 · Product code: **BZC** · Anesthesiology
Source: <https://www.510kdatabase.net/k072323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Calculator, Pulmonary Function Data (BZC)
Date received	Aug 20, 2007
Decision date	Oct 10, 2007
Days to decision	51 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Viasys Healthcare GmbH
Location	Yorba Linda, CA, US
Contact	Yvette Lloyd
510(k) history	14 submissions · 14 cleared · 2003-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072323/> 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 24, 2026