FDA 510(k) Premarket Notification

K072323 MASTERSCREEN & OXYCON CPX

CLEARED

Oct 10, 2007
51 days to decision

K072323 · 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