

K011518 VITALSAT SERIESNov 15, 2002
547 days to decisionK011518 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k011518/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	May 17, 2001
Decision date	Nov 15, 2002
Days to decision	547 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Beta Biomed Services, Inc.
Location	Crofton, MD, US
Contact	E J SMITH
510(k) history	2 submissions · 2 cleared · 2001-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k011518/> 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 25, 2026