FDA 510(k) Premarket Notification

K011518 VITALSAT SERIES

CLEARED

Nov 15, 2002
547 days to decision

K011518 · 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