FDA 510(k) Premarket Notification

K092549 OXIPROBE, MODELS BM-100, BM-200, BM-400, BM-300, BM-300S, BM-600P

CLEARED

Nov 16, 2009
89 days to decision

K092549 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k092549/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Aug 19, 2009
Decision date	Nov 16, 2009
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bio Medical Technologies Co., Ltd.**
Location	Dunwoody, GA, US
Contact	CATHRYN CAMBRIA
510(k) history	2 submissions · 2 cleared · 2009-2014