

**K092549 OXIPROBE, MODELS BM-100, BM-200, BM-400,
BM-300, BM-300S, BM-600P**Nov 16, 2009
89 days to decisionK092549 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092549/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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