

**K081712 FINGERTIP PULSE OXIMETER, MODEL M70, AND
HANDHELD PULSE OXIMETER, MODEL M700**Sep 12, 2008
87 days to decisionK081712 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k081712/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 17, 2008
Decision date	Sep 12, 2008
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Guangdong Biolight Meditech Co., Ltd.
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	21 submissions · 21 cleared · 2008-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081712/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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