FDA 510(k) Premarket Notification

K090671 PULSE OXIMETER, MODELS CMS50E, CMS50F, CMS60C, CMS60D

CLEARED

Jun 11, 2009
90 days to decision

K090671 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k090671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 13, 2009
Decision date	Jun 11, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Contec Medical System Co., Ltd.**
Location	Zhong Shan, Shanghai, CN
Contact	Diana Hong
510(k) history	12 submissions · 12 cleared · 2008-2019