FDA 510(k) Premarket Notification

K160268 Fingertip Pulse Oximeter MD300CG11/MD300CG51

CLEARED

Oct 14, 2016
255 days to decision

K160268 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k160268/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Feb 2, 2016
Decision date	Oct 14, 2016
Days to decision	255 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Beijing Choice Electronic Technology Co., Ltd.**
Location	Beijing, CN
Contact	LEI CHEN
510(k) history	14 submissions · 14 cleared · 2016-2024