FDA 510(k) Premarket Notification

K071642 MEDIAID M960 SERIES, VITAL SIGNS MONITOR AND MEDIAID M900 SERIES, PULSE OXIMETER

CLEARED

May 2, 2008
319 days to decision

K071642 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k071642/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 18, 2007
Decision date	May 2, 2008
Days to decision	319 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mediaid, Inc.**
Location	Plano, TX, US
Contact	JAYESH PATEL
510(k) history	4 submissions · 4 cleared · 2004-2008