FDA 510(k) Premarket Notification

CLEARED

K002939 MODIFICATION TO MASIMO SET RADICAL PULSE OXIMETER WITH SATSHARE AND LNOP SERIES OF SENORS OXIMETER

Oct 6, 2000
15 days to decision

K002939 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k002939/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Oximeter (DQA)
Date received	Sep 21, 2000
Decision date	Oct 6, 2000
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Masimo Corp.**
Location	Irvine, CA, US
Contact	JAMES J CRONIN
Website	http://www.masimo.com/
510(k) history	28 submissions · 28 cleared · 1997-2004

Masimo Corp. is an American health technology company headquartered in Irvine, California. The company develops patient monitoring devices, non-invasive sensors, and related software platforms for hospital and home settings. Masimo has received FDA 510(k) clearances from total submissions since its first clearance in 1997. The company's regulatory submissions have focused exclusively on Anesthesiology devices, including pulse oximetry sensors, adaptor cables, and monitoring modules. The latest clearance on record dates to 2004, reflecting the company's historical regulato...