

**K062605 PHILIPS SPO2 REUSABLE SENSOR, MODELS M1196A
AND M1196T**Nov 9, 2006
65 days to decisionK062605 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k062605/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Oximeter (DQA)
Date received	Sep 5, 2006
Decision date	Nov 9, 2006
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems
Location	Seattle, WA, US
Contact	RICH MCCLEARY
510(k) history	107 submissions · 105 cleared · 2002-2021

Philips Medical Systems is a Dutch multinational health technology company headquartered in Amsterdam with U.S. operations based in Seattle. The company evolved from a consumer electronics conglomerate founded in 1891 to a healthcare-focused organization. Philips Medical Systems has received FDA 510(k) clearances from total submissions between 2002 and 2021. The company's regulatory focus centered on Cardiovascular devices, which represented 79% of all submissions. This historical record reflects the company's significant presence in diagnostic ultrasound systems and pati...

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Device record: <https://www.510kdatabase.net/k062605/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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