

K811836 PO-200 BLOOD OXYGEN MONITORJul 27, 1981
27 days to decisionK811836 · Product code: **KLK** · AnesthesiologySource: <https://www.510kdatabase.net/k811836/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Jun 30, 1981
Decision date	Jul 27, 1981
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Sumitomo Electric Industries, Inc.
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1985

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Device record: <https://www.510kdatabase.net/k811836/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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