

**K870805 MODEL 840 TRANSCUTAN. OXYGEN/CARBON DIOX.
MONITOR**May 27, 1987
85 days to decisionK870805 · Product code: **KLK** · Anesthesiology
Source: <https://www.510kdatabase.net/k870805/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Mar 3, 1987
Decision date	May 27, 1987
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Novametrix Medical Systems, Inc.
Location	Mchenry, IL, US
Contact	ROBERT SCHIFFMAN
510(k) history	45 submissions · 45 cleared · 1978-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k870805/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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