FDA 510(k) Premarket Notification

K870805 MODEL 840 TRANSCUTAN. OXYGEN/CARBON DIOX. MONITOR

CLEARED

May 27, 1987
85 days to decision

K870805 · 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