FDA 510(k) Premarket Notification

K813350 NOVAMETRIX MODEL 807 TRANSCUT. MONITOR

CLEARED

Feb 12, 1982
77 days to decision

K813350 · Product code: **KLK** · Anesthesiology
Source: https://www.510kdatabase.net/k813350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Nov 27, 1981
Decision date	Feb 12, 1982
Days to decision	77 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k813350/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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