FDA 510(k) Premarket Notification

K792009 TC02 M OXYGEN ELECTRODE CONTACT

CLEARED

Oct 26, 1979
21 days to decision

K792009 · Product code: **KLK** · Anesthesiology
Source: https://www.510kdatabase.net/k792009/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Oxygen, Cutaneous, For Infant Not Under Gas Anesthesia (KLK)
Date received	Oct 5, 1979
Decision date	Oct 26, 1979
Days to decision	21 days
Third-party review	No

APPLICANT

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