

**K792009 TC02 M OXYGEN ELECTRODE CONTACT**Oct 26, 1979  
21 days to decisionK792009 · Product code: **KLK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k792009/>**SUBMISSION DETAILS**

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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**

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Company	<b>Novamatrix Medical Systems, Inc.</b>
Location	Mchenry, IL, US
510(k) history	45 submissions · 45 cleared · 1978-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792009/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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