

**K904039 REFLECTANCE SENSOR (RS-10)**Dec 27, 1990  
114 days to decisionK904039 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k904039/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Sep 4, 1990
Decision date	Dec 27, 1990
Days to decision	114 days
Third-party review	No

**APPLICANT**

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Company	<b>Nellcor, Inc.</b>
Location	Mchenry, IL, US
Contact	KENNETH MICHAEL
510(k) history	18 submissions · 18 cleared · 1982-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904039/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026