

**K970368 SC9000/SC9015 MEDICAL INFORMATION BUS (MIS)  
 PROTOCOL CONVERTER**

May 6, 1997  
 95 days to decision

K970368 · Product code: **DQA** · Anesthesiology  
 Source: <https://www.510kdatabase.net/k970368/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 31, 1997
Decision date	May 6, 1997
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	JAQUELINE EMERY
510(k) history	778 submissions · 778 cleared · 1980-2026

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

Device record: <https://www.510kdatabase.net/k970368/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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