

**K030640 SIEMENS MICRO2+ WITH VAI SOFTWARE**Jun 3, 2003  
95 days to decisionK030640 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k030640/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Oximeter (DQA)
Date received	Feb 28, 2003
Decision date	Jun 3, 2003
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030640/> 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 15, 2026