FDA 510(k) Premarket Notification

K030640 SIEMENS MICRO2+ WITH VAI SOFTWARE

CLEARED

Jun 3, 2003
95 days to decision

K030640 · Product code: **DQA** · Anesthesiology
Source: 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	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	PENELOPE H GRECO
510(k) history	778 submissions · 778 cleared · 1980-2026