FDA 510(k) Premarket Notification

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

**CLEARED**

May 6, 1997
95 days to decision

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

## SUBMISSION DETAILS

| | |
|---|---|
| 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

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