FDA 510(k) Premarket Notification

# K863042 SIRECUST 888/888R NON-INVASIVE BLOOD PRESS MONITOR

**CLEARED**

Jan 14, 1987
155 days to decision

K863042 · Product code: **DSK** · Cardiovascular
Source: https://www.510kdatabase.net/k863042/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Computer, Blood-pressure (DSK) |
| Date received | Aug 12, 1986 |
| Decision date | Jan 14, 1987 |
| Days to decision | 155 days |
| Third-party review | No |

## APPLICANT

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