FDA 510(k) Premarket Notification

# K972379 PRECISA N

**CLEARED**

Sep 24, 1997
90 days to decision

K972379 · Product code: **DXQ** · Cardiovascular
Source: https://www.510kdatabase.net/k972379/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Blood Pressure Cuff (DXQ) |
| Date received | Jun 26, 1997 |
| Decision date | Sep 24, 1997 |
| Days to decision | 90 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Rudolf Riester GmbH & Co. KG** |
| Location | D-72417 Jungingen, DE |
| Contact | PATRICIA RIESTER-FREUDENMANN |
| 510(k) history | 30 submissions · 26 cleared · 1993-2000 |