FDA 510(k) Premarket Notification

# K832171 SELF CHECK DIGITAL BLOOD PRESS. MONITOR

**CLEARED**

Oct 4, 1983
90 days to decision

K832171 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k832171/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Measurement, Blood-pressure, Non-invasive (DXN) |
| Date received | Jul 6, 1983 |
| Decision date | Oct 4, 1983 |
| Days to decision | 90 days |
| Third-party review | No |

## APPLICANT

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|---|---|
| Company | **Ritter Co.** |
| Location | Mchenry, IL, US |
| 510(k) history | 10 submissions · 10 cleared · 1976-1983 |