FDA 510(k) Premarket Notification

# K923344 ELECTRONIC DIGITAL BLOOD PRESSURE MONITOR

**CLEARED**

Oct 7, 1993
476 days to decision

K923344 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k923344/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Measurement, Blood-pressure, Non-invasive (DXN) |
| Date received | Jun 18, 1992 |
| Decision date | Oct 7, 1993 |
| Days to decision | 476 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Citizen Watch Co., Ltd.** |
| Location | Los Angeles, CA, US |
| Contact | WILLIAM ANDROLIA |
| Website | http://www.citizen.fi/ |
| 510(k) history | 8 submissions · 8 cleared · 1991-2006 |