

K923344 ELECTRONIC DIGITAL BLOOD PRESSURE MONITOROct 7, 1993
476 days to decisionK923344 · Product code: **DXN** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923344/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026