

K760930 BLOOD PRESSURE & PULSE CHECKERNov 9, 1976
11 days to decisionK760930 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k760930/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Oct 29, 1976
Decision date	Nov 9, 1976
Days to decision	11 days
Third-party review	No

APPLICANT

Company	Bio Mega Diagnostic, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1976-1987

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k760930/> 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 29, 2026