FDA 510(k) Premarket Notification

K760930 BLOOD PRESSURE & PULSE CHECKER

CLEARED

Nov 9, 1976
11 days to decision

K760930 · Product code: **DXN** · Cardiovascular
Source: 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