FDA 510(k) Premarket Notification

K170666 The Wrist-Type Fully Automatic Digital Blood Pressure Monitors

CLEARED

Nov 22, 2017
261 days to decision

K170666 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k170666/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Mar 6, 2017
Decision date	Nov 22, 2017
Days to decision	261 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Joytech Healthcare Co. , Ltd.**
Location	Hangzhou, CN
Contact	Ren Yunhua
Website	https://www.joytechhealthcare.com
510(k) history	22 submissions · 22 cleared · 2017-2026