FDA 510(k) Premarket Notification

CLEARED

K152770 ProBP 2400 Digital Blood Pressure Device

Oct 21, 2015
26 days to decision

K152770 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k152770/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Sep 25, 2015
Decision date	Oct 21, 2015
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microlife Intellectual Property GmbH**
Location	Great Neck, NY, US
Contact	GERHARD FRICK
510(k) history	54 submissions · 54 cleared · 2003-2023