FDA 510(k) Premarket Notification

K093205 PADCHECK

CLEARED

Apr 1, 2010
170 days to decision

K093205 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k093205/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Oct 13, 2009
Decision date	Apr 1, 2010
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Padtest, LLC**
Location	Los Angeles, CA, US
Contact	JAMES R GREENWOOD
510(k) history	1 submissions · 1 cleared · 2010-2010