FDA 510(k) Premarket Notification

K053453 HOMMED CENTRAL STATION, VERSION 3.5

CLEARED

May 5, 2006
144 days to decision

K053453 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k053453/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 12, 2005
Decision date	May 5, 2006
Days to decision	144 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Honeywell Hommed, LLC**
Location	Houston, TX, US
Contact	TOMMIE J MORGAN
510(k) history	7 submissions · 7 cleared · 2006-2016