

**K053453 HOMMED CENTRAL STATION, VERSION 3.5**May 5, 2006  
144 days to decisionK053453 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k053453/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k053453/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 3, 2026