

**K040032 CENTRAL STATION, VERSION 3.3H**Feb 27, 2004  
50 days to decisionK040032 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k040032/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                      |
| Submission type       | Traditional   |
| Device classification | System, Measurement, Blood-pressure, Non-invasive (DXN) |
| Date received         | Jan 8, 2004   |
| Decision date         | Feb 27, 2004  |
| Days to decision      | 50 days   |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Hommed, LLC</b>                    |
| Location       | Houston, TX, US                       |
| Contact        | TOMMIE J MORGAN                       |
| 510(k) history | 8 submissions · 8 cleared · 2000-2004 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040032/> 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 May 25, 2026