

K040799 HOMMED GENESIS PATIENT MONITOR SYSTEM WITH OPTIONSAug 18, 2004
142 days to decisionK040799 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k040799/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Mar 29, 2004
Decision date	Aug 18, 2004
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040799/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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