FDA 510(k) Premarket Notification

CLEARED

K040799 HOMMED GENESIS PATIENT MONITOR SYSTEM WITH OPTIONS

Aug 18, 2004
142 days to decision

K040799 · 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