

**K912138 HOMEVUE MONITORING SYS.MODEL TD 2000**Mar 19, 1992  
310 days to decisionK912138 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k912138/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	May 14, 1991
Decision date	Mar 19, 1992
Days to decision	310 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Homevue Health Systems, Inc.</b>
Location	Laguna Beach, CA, US
Contact	GARY R.MOUNTS
510(k) history	2 submissions · 2 cleared · 1992-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912138/> 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 20, 2026