

**K032252 MODEL 120 SERIES MATERNAL/FETAL MONITOR**Aug 21, 2003  
30 days to decisionK032252 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k032252/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Jul 22, 2003
Decision date	Aug 21, 2003
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Information Technologies</b>
Location	Mchenry, IL, US
Contact	MELISSA ROBINSON
510(k) history	136 submissions · 132 cleared · 1978-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032252/> 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 26, 2026