

**K021135 MODEL 2120IS SERIES MATERNAL/FETAL MONITOR**May 9, 2002  
30 days to decisionK021135 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k021135/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Apr 9, 2002
Decision date	May 9, 2002
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	JOELLE NEIDER
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/k021135/> 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