

**K220732 Mural Perinatal Surveillance**Jun 23, 2022  
101 days to decisionK220732 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k220732/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Mar 14, 2022
Decision date	Jun 23, 2022
Days to decision	101 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Information Technologies, Inc.</b>
Location	Milwaukee, WI, US
Contact	Brandon O' Shea
510(k) history	31 submissions · 31 cleared · 2010-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220732/> 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 14, 2026