

**K813458 FETASCAN 1000, 1000A OBSTETRICAL DATA**Jan 28, 1982  
49 days to decisionK813458 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k813458/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Dec 10, 1981
Decision date	Jan 28, 1982
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Cranbury Medical Labs, Inc.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1982-1982

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k813458/> 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 21, 2026