

**K823188 M2000 INTRAPARTUM FETAL MONITOR**Jan 26, 1983  
92 days to decisionK823188 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k823188/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Oct 26, 1982
Decision date	Jan 26, 1983
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Huntleigh Technology, Inc.</b>
Location	Walker, MI, US
510(k) history	23 submissions · 23 cleared · 1981-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k823188/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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