

**K902104 ORION WATCHGUARD**Oct 25, 1990  
170 days to decisionK902104 · Product code: **HFM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k902104/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Uterine Contraction, External (for Use In Clinic) (HFM)
Date received	May 8, 1990
Decision date	Oct 25, 1990
Days to decision	170 days
Third-party review	No

**APPLICANT**

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Company	<b>Orion Industries</b>
Location	Fort Myers, FL, US
Contact	BOB GRANTHAM
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902104/>; 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 June 30, 2026