

**K854267 HUNTLEIGH DOPCORD, D140**Jul 9, 1986  
259 days to decisionK854267 · Product code: **KNG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k854267/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Oct 23, 1985
Decision date	Jul 9, 1986
Days to decision	259 days
Third-party review	No

**APPLICANT**

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Company	<b>Huntleigh Technology, Inc.</b>
Location	Walker, MI, US
Contact	JAMES BRITTON
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/k854267/>; 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 20, 2026