

**K860233 5 MHZ INTRACAVITARY PROBE, MODEL H4222V**Jun 2, 1986  
131 days to decisionK860233 · Product code: **HEM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k860233/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Imager, Ultrasonic Obstetric-gynecologic (HEM)
Date received	Jan 22, 1986
Decision date	Jun 2, 1986
Days to decision	131 days
Third-party review	No

**APPLICANT**

---

Company	<b>General Electric Co.</b>
Location	Mchenry, IL, US
Contact	HOWARD II
510(k) history	254 submissions · 254 cleared · 1976-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860233/>; 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 17, 2026