

**K871383 FETALERT ADR FETAL MONITOR**Jun 1, 1987  
55 days to decisionK871383 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k871383/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Apr 7, 1987
Decision date	Jun 1, 1987
Days to decision	55 days
Third-party review	No

**APPLICANT**

---

Company	<b>Advanced Technology Laboratories, Inc.</b>
Location	Mchenry, IL, US
Contact	JOHN DELLINGER
510(k) history	61 submissions · 59 cleared · 1980-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871383/> 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 19, 2026