

**K902190 FETAL PULSE DETECTOR MODEL DOP-1**Oct 22, 1990  
159 days to decisionK902190 · Product code: **HEK** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k902190/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Heart Sound, Fetal, Ultrasonic (HEK)
Date received	May 16, 1990
Decision date	Oct 22, 1990
Days to decision	159 days
Third-party review	No

**APPLICANT**

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Company	<b>Advanced Medical Systems, Inc.</b>
Location	Hamden, CT, US
Contact	ANTHONY CALDERONI
510(k) history	10 submissions · 10 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902190/>; 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 21, 2026