

**K870692 PERINATAL COMPUTERS
2015/2025/2040/2045/2080/2085**May 4, 1987
74 days to decisionK870692 · Product code: **HGM** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k870692/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Feb 19, 1987
Decision date	May 4, 1987
Days to decision	74 days
Third-party review	No

APPLICANT

Company	Peritronics Research and Development, Inc.
Location	Arcadia, CA, US
Contact	KOH, MD
510(k) history	2 submissions · 2 cleared · 1986-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k870692/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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