FDA 510(k) Premarket Notification

# K870692 PERINATAL COMPUTERS 2015/2025/2040/2045/2080/2085

**CLEARED**

May 4, 1987
74 days to decision

K870692 · 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 |