

K831371 MODEL FM5Jun 30, 1983
64 days to decisionK831371 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k831371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Apr 27, 1983
Decision date	Jun 30, 1983
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Sonicaid, Inc.
Location	Mchenry, IL, US
510(k) history	18 submissions · 17 cleared · 1977-1988

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831371/> Data sourced from FDA 510(k) public records (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. - Generated May 21, 2026