

K780487 FETAL MONITORING SYSTEMMay 9, 1978
43 days to decisionK780487 · Product code: **HFO** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k780487/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Pressure, Intrauterine (HFO)
Date received	Mar 27, 1978
Decision date	May 9, 1978
Days to decision	43 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k780487/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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