

**K812256 MODEL AP 180 MINI-MONITOR**Oct 20, 1981  
69 days to decisionK812256 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k812256/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Aug 12, 1981
Decision date	Oct 20, 1981
Days to decision	69 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k812256/> 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