

**K833227 SONICAID MODEL D206E AIR EMBOLI-DETECT**Oct 27, 1983  
72 days to decisionK833227 · Product code: **LBA** · Toxicology  
Source: <https://www.510kdatabase.net/k833227/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Procainamide Control Materials (LBA)
Date received	Aug 16, 1983
Decision date	Oct 27, 1983
Days to decision	72 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833227/> 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