

**K840359 POST-AURICULAR HEARING AID**Feb 21, 1984  
26 days to decisionK840359 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k840359/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 26, 1984
Decision date	Feb 21, 1984
Days to decision	26 days
Third-party review	No

**APPLICANT**

---

Company	<b>Phonic Ear, Inc.</b>
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1982-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k840359/> 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 July 3, 2026