

**K093715 EVOK 900 SERIES HEARING AID/TINNITUS MASKER  
OPTION DEVICE**Dec 22, 2010  
386 days to decisionK093715 · Product code: **KLW** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k093715/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Masker, Tinnitus (KLW)
Date received	Dec 1, 2009
Decision date	Dec 22, 2010
Days to decision	386 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Magnatone Hearing Aid Corp. Db a Persona Medical</b>
Location	Casselberry, FL, US
Contact	DON CAMPBELL
510(k) history	1 submissions · 1 cleared · 2010-2010

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093715/> 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 June 20, 2026