

**K982432 PILLOW MASKER, C2007M, C2008M, CE2000,  
WONDER EAR, MINI WONDER EAR, PT-2SM, PT-3SM, PT-3LFM,  
PT-3HFM, PT3CM, PT5-SM, PT5-**

Jan 25, 1999  
196 days to decision

K982432 · Product code: **KLW** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k982432/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Masker, Tinnitus (KLW)
Date received	Jul 13, 1998
Decision date	Jan 25, 1999
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Tinnitus Treatment Centers, Inc.</b>
Location	Dallas, TX, US
Contact	DAVID W HOLMES
510(k) history	2 submissions · 2 cleared · 1999-1999

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

Device record: <https://www.510kdatabase.net/k982432/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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