

K982451 TTCGHI-T AND TTCTN3-T-TMay 7, 1999
296 days to decisionK982451 · Product code: **KLW** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k982451/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Masker, Tinnitus (KLW)
Date received	Jul 15, 1998
Decision date	May 7, 1999
Days to decision	296 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982451/> 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 July 5, 2026