FDA 510(k) Premarket Notification

K982451 TTCGHI-T AND TTCTN3-T-T

CLEARED

May 7, 1999
296 days to decision

K982451 · Product code: **KLW** · Ear, Nose, Throat
Source: 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