

K790064 TINNITUS DEVICESFeb 16, 1979
32 days to decisionK790064 · Product code: **ETS** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k790064/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Electronic Noise (for Audiometric Testing) (ETS)
Date received	Jan 15, 1979
Decision date	Feb 16, 1979
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Vicon Instrument Co.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k790064/>; 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 6, 2026