

K901609 ELECTROGLOTTOGRAPHJun 28, 1990
83 days to decisionK901609 · Product code: **KLX** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k901609/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electroglottograph (KLX)
Date received	Apr 6, 1990
Decision date	Jun 28, 1990
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Glottal Ent., Inc.
Location	Washington, DC, US
Contact	& JAWORSKI
510(k) history	1 submissions · 1 cleared · 1990-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k901609/>; 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