

K010981 GSA GENITO SENSORY ANALYZERSep 20, 2001
171 days to decisionK010981 · Product code: **LLN** · Neurology
Source: <https://www.510kdatabase.net/k010981/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vibration Threshold Measurement (LLN)
Date received	Apr 2, 2001
Decision date	Sep 20, 2001
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medoc Ltd. Advanced Medical Systems
Location	Minnetonka, MN, US
Contact	ANN QUINLAN-SMITH
510(k) history	4 submissions · 4 cleared · 2001-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010981/> 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 June 29, 2026