FDA 510(k) Premarket Notification

CLEARED

K010981 GSA GENITO SENSORY ANALYZER

Sep 20, 2001
171 days to decision

K010981 · 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