FDA 510(k) Premarket Notification

K131568 SUDO PATH

CLEARED

Jun 28, 2013
29 days to decision

K131568 · Product code: **GZO** · Neurology
Source: https://www.510kdatabase.net/k131568/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Galvanic Skin Response Measurement (GZO)
Date received	May 30, 2013
Decision date	Jun 28, 2013
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ld Technology, LLC**
Location	Miami, FL, US
Contact	ALBERT MAAREK
510(k) history	14 submissions · 14 cleared · 2009-2020