FDA 510(k) Premarket Notification

K103370 RT200

CLEARED

Apr 5, 2011
139 days to decision

K103370 · Product code: **GZI** · Neurology
Source: https://www.510kdatabase.net/k103370/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Nov 17, 2010
Decision date	Apr 5, 2011
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Restorative Therapies, Inc.**
Location	Baltimore, MD, US
Contact	ANDREW BARRISKILL
510(k) history	10 submissions · 10 cleared · 2005-2017