FDA 510(k) Premarket Notification

K153704 Strive

CLEARED

Jun 3, 2016
163 days to decision

K153704 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k153704/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 23, 2015
Decision date	Jun 3, 2016
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Djo, LLC**
Location	Vista, CA, US
Contact	Rand Daoud
510(k) history	12 submissions · 12 cleared · 2009-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k153704/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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