FDA 510(k) Premarket Notification

K120691 BRAIN PORT

CLEARED

Jun 5, 2012
90 days to decision

K120691 · Product code: **GZT** · Neurology
Source: https://www.510kdatabase.net/k120691/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor, Self-retaining, For Neurosurgery (GZT)
Date received	Mar 7, 2012
Decision date	Jun 5, 2012
Days to decision	90 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Nico Corporation**
Location	Indianapolis, IN, US
Contact	JAY DITTMAN
510(k) history	9 submissions · 9 cleared · 2012-2024