

K120691 BRAIN PORTJun 5, 2012
90 days to decisionK120691 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120691/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026