

K121264 TRITON ELECTRIC HIGH TORQUE HANDPIECE AND SOFTWARE MODULE

Aug 27, 2012
123 days to decision

K121264 · Product code: **HBE** · Neurology
Source: <https://www.510kdatabase.net/k121264/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drills, Burrs, Trephines & Accessories (simple, Powered) (HBE)
Date received	Apr 26, 2012
Decision date	Aug 27, 2012
Days to decision	123 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Surgical Technologies
Location	Fort Worth, TX, US
Contact	RISHI SINHA
510(k) history	1 submissions · 1 cleared · 2012-2012

510k Database - www.510kdatabase.net
 Device record: <https://www.510kdatabase.net/k121264/> 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 14, 2026