

K150371 Diros OWL Sterile Single Use Trident™ R.F. Insulated Cannulae, Models DTR and DTRHJul 30, 2015
167 days to decisionK150371 · Product code: **GXI** · Neurology
Source: <https://www.510kdatabase.net/k150371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Feb 13, 2015
Decision date	Jul 30, 2015
Days to decision	167 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Diros Technology, Inc.
Location	Charleston, SC, US
Contact	GEORGE DARMOS
510(k) history	8 submissions · 8 cleared · 2002-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150371/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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