

**K062946 TOP NEUROPOLE NEEDLES, MODELS ST, X, RC, XE AND TL**

Feb 15, 2007  
140 days to decision

K062946 · Product code: **GXI** · Neurology  
Source: <https://www.510kdatabase.net/k062946/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Sep 28, 2006
Decision date	Feb 15, 2007
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Top Corp.</b>
Location	Littleton, CO, US
Contact	CATHRYN N CAMBRIA
510(k) history	3 submissions · 3 cleared · 1986-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k062946/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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