

**K222281 Intracept Intraosseous Nerve Ablation System**Oct 26, 2022  
89 days to decisionK222281 · Product code: **GXI** · Neurology  
Source: <https://www.510kdatabase.net/k222281/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Jul 29, 2022
Decision date	Oct 26, 2022
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Relievant Medsystems, Inc.</b>
Location	Hayward, CA, US
Contact	Thomas A. Slater
510(k) history	4 submissions · 4 cleared · 2007-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222281/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026