

**K180369 Intracept Intraosseous Nerve Ablation System  
(component Intracept RF Probe)**Sep 14, 2018  
214 days to decisionK180369 · Product code: **GXI** · Neurology  
Source: <https://www.510kdatabase.net/k180369/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Feb 12, 2018
Decision date	Sep 14, 2018
Days to decision	214 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Relievant Medsystems</b>
Location	Redwood City, CA, US
Contact	Alex DiNello
510(k) history	6 submissions · 6 cleared · 2010-2019

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180369/> 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 June 21, 2026