

**K213836 Intracept Intraosseous Nerve Ablation System**Mar 11, 2022  
92 days to decisionK213836 · Product code: **GXI** · Neurology  
Source: <https://www.510kdatabase.net/k213836/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Dec 9, 2021
Decision date	Mar 11, 2022
Days to decision	92 days
Third-party review	No
Combination product	No
PCCP authorized	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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