FDA 510(k) Premarket Notification

K213836 Intracept Intraosseous Nerve Ablation System

CLEARED

Mar 11, 2022
92 days to decision

K213836 · Product code: **GXI** · Neurology
Source: https://www.510kdatabase.net/k213836/

SUBMISSION DETAILS

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

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