FDA 510(k) Premarket Notification

K222281 Intracept Intraosseous Nerve Ablation System

CLEARED

Oct 26, 2022
89 days to decision

K222281 · Product code: **GXI** · Neurology
Source: https://www.510kdatabase.net/k222281/

SUBMISSION DETAILS

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

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