FDA 510(k) Premarket Notification

K100353 SPINEWAND SURGICAL DEVICE

CLEARED

May 13, 2010
90 days to decision

K100353 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k100353/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 12, 2010
Decision date	May 13, 2010
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Arthrocare Corp.**
Location	Mountain View, CA, US
Contact	VALERIE DEFIESTA-NG
Website	http://www.arthrocare.com/
510(k) history	112 submissions · 112 cleared · 1995-2016