FDA 510(k) Premarket Notification

K091228 SPINEWAND SURGICAL DEVICE

CLEARED

Sep 28, 2009
154 days to decision

K091228 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k091228/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 27, 2009
Decision date	Sep 28, 2009
Days to decision	154 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