K132099 ARTHROCARE COBLATOR IQ DLR SPINEWAND, ARTHROCARE COBLATOR IQ DLG SPINEWAND

CLEARED

Aug 22, 2013
45 days to decision

K132099 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k132099/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 8, 2013
Decision date	Aug 22, 2013
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**ArthroCare Corporation**
Location	Irvine, CA, US
Contact	SHIRLEY HYINK
Website	http://www.arthrocare.com/
510(k) history	36 submissions · 36 cleared · 2011-2026