FDA 510(k) Premarket Notification

K123353 CIQ CONTROLLER, CIQ WANDS

CLEARED

Apr 5, 2013
156 days to decision

K123353 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k123353/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 31, 2012
Decision date	Apr 5, 2013
Days to decision	156 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k123353/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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