

K183346 FLOW 90 WandJan 24, 2019
52 days to decisionK183346 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k183346/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 3, 2018
Decision date	Jan 24, 2019
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k183346/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026