

K000778 ENTEC REFLEX WAND 55, MODEL E4055-01, E4045-01May 3, 2000
54 days to decisionK000778 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k000778/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 10, 2000
Decision date	May 3, 2000
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k000778/> 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