

**K122092 MEDRANGE ELECTROSURGICAL EXPANSION
SYSTEM**Oct 23, 2012
99 days to decisionK122092 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k122092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 16, 2012
Decision date	Oct 23, 2012
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medrange Corporation
Location	Los Angeles, CA, US
Contact	KUOFANG HUANG
510(k) history	2 submissions · 2 cleared · 2010-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122092/> 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 June 15, 2026