

K820902 MULTIPERFECTEMPApr 26, 1982
26 days to decisionK820902 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820902/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 31, 1982
Decision date	Apr 26, 1982
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Kay Medical, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k820902/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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