

K790293 CIMARRON 1000 OR 1000-2Apr 12, 1979
63 days to decisionK790293 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k790293/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 8, 1979
Decision date	Apr 12, 1979
Days to decision	63 days
Third-party review	No

APPLICANT

Company	Cimarron Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 3 cleared · 1979-1984

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Device record: <https://www.510kdatabase.net/k790293/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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