

K812760 HI-TEMP CARDIOVASCULAR CAUTERYJan 18, 1982
109 days to decisionK812760 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k812760/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 1, 1981
Decision date	Jan 18, 1982
Days to decision	109 days
Third-party review	No

APPLICANT

Company	Key Medical, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k812760/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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