

K821356 DESERET CAPACITY PLUS ELECTROSURG. PADJul 14, 1982
69 days to decisionK821356 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k821356/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 6, 1982
Decision date	Jul 14, 1982
Days to decision	69 days
Third-party review	No

APPLICANT

Company	Warner-Lambert Co.
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003

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Device record: <https://www.510kdatabase.net/k821356/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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