

**K792600 MODEL III HANDSWITCH**Jan 10, 1980  
23 days to decisionK792600 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k792600/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 18, 1979
Decision date	Jan 10, 1980
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Sara Dan, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1980

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792600/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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