

**K780978 SINGLE USE ACTIVE ELECTRODE HANDLE**Jun 22, 1978  
13 days to decisionK780978 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k780978/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 9, 1978
Decision date	Jun 22, 1978
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Aspen Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1976-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k780978/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 22, 2026