

**K780975 SINGLE USE NEEDLE ELECTRODE**Jun 22, 1978  
13 days to decisionK780975 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k780975/>**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/k780975/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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