

**K902035 FLEXIBLE EXTENDER**Jul 19, 1990  
77 days to decisionK902035 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k902035/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 3, 1990
Decision date	Jul 19, 1990
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Beacon Laboratories, Inc.</b>
Location	Westminster, CO, US
Contact	RICHARD P FLEENOR
510(k) history	18 submissions · 18 cleared · 1989-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902035/>; 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 June 28, 2026