

**K930243 FLEXIBLE EXTENDER**Mar 11, 1994  
416 days to decisionK930243 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k930243/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 19, 1993
Decision date	Mar 11, 1994
Days to decision	416 days
Third-party review	No
Summary / Statement	Summary

**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/k930243/> 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