

**K952470 ACTI-FLEX LOW, MEDIUM, RESCARE I, RESCARE II.**Nov 28, 1995  
186 days to decisionK952470 · Product code: **GCD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k952470/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Connector, Catheter (GCD)
Date received	May 26, 1995
Decision date	Nov 28, 1995
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>R-Group Intl.</b>
Location	Gainesville, FL, US
Contact	KERRY ANNE KAHN
510(k) history	7 submissions · 7 cleared · 1992-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952470/> 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 26, 2026