

**K844164 REACTIFS, IBF ULTROGELS ACA & ULTROGELS A**Dec 10, 1984  
46 days to decisionK844164 · Product code: **DII** · Chemistry  
Source: <https://www.510kdatabase.net/k844164/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Columns, Glc (DII)
Date received	Oct 25, 1984
Decision date	Dec 10, 1984
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Lkb Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	ED ZIOR
510(k) history	52 submissions · 52 cleared · 1976-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k844164/>; 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 14, 2026