

**K821834 MODEL 100 BICARBONATE CONDUCTIVITY**Jul 2, 1982  
10 days to decisionK821834 · Product code: **FKQ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k821834/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Central Multiple Patient (FKQ)
Date received	Jun 22, 1982
Decision date	Jul 2, 1982
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Western Laboratories Corp.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1978-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k821834/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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