

**K884179 TURBOX FIBRINOGEN**Dec 20, 1988  
81 days to decisionK884179 · Product code: **KQJ** · Hematology  
Source: <https://www.510kdatabase.net/k884179/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Fibrinogen Determination (KQJ)
Date received	Sep 30, 1988
Decision date	Dec 20, 1988
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Unipath , Ltd.</b>
Location	Bedford Mk41 Oqg, GB
Contact	MR. G ZAJICEK
510(k) history	73 submissions · 73 cleared · 1986-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884179/>, 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 29, 2026