

**K893803 FEVER LINE AND FEVER LINE PLUS**Aug 21, 1989  
91 days to decisionK893803 · Product code: **KPD** · General Hospital  
Source: <https://www.510kdatabase.net/k893803/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Strip, Temperature, Forehead, Liquid Crystal (KPD)
Date received	May 22, 1989
Decision date	Aug 21, 1989
Days to decision	91 days
Third-party review	No

**APPLICANT**

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Company	<b>Liquid Crystal Sciences, Inc.</b>
Location	Atlanta, GA, US
Contact	L BRETT
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893803/>; 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 July 8, 2026