

**K032017 LIQUICHEK D-DIMER CONTROL LEVEL 1, 2 AND 3**Aug 8, 2003  
39 days to decisionK032017 · Product code: **GGN** · Hematology  
Source: <https://www.510kdatabase.net/k032017/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasma, Coagulation Control (GGN)
Date received	Jun 30, 2003
Decision date	Aug 8, 2003
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bio-Rad</b>
Location	Mchenry, IL, US
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510(k) history	319 submissions · 319 cleared · 1976-2017

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032017/>; 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