

**K092170 STA LIATEST FREE PS**Dec 22, 2009  
154 days to decisionK092170 · Product code: **GGP** · Hematology  
Source: <https://www.510kdatabase.net/k092170/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Test, Qualitative And Quantitative Factor Deficiency (GGP)
Date received	Jul 21, 2009
Decision date	Dec 22, 2009
Days to decision	154 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Diagnostica Stago, Inc.</b>
Location	Parsippany, NJ, US
Contact	UMBERTO V PARROTTA
Website	<a href="https://www.stago-us.com">https://www.stago-us.com</a>
510(k) history	16 submissions · 16 cleared · 2001-2026

Diagnostica Stago, Inc. is an industry leader in hemostasis and thrombosis analysis. The company provides coagulation instruments, reagent kits, and related systems for clinical and research laboratories. With a manufacturing facility in Parsippany, US, the company has served the hemostasis laboratory community for over 25 years. Diagnostica Stago, Inc. has received FDA 510(k) clearances from total submissions since 2001. All submissions focus on Hematology devices. The company's latest clearance in 2026 demonstrates continued innovation and active regulatory engagement i...

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