

**K954437 THROMBELASTOGRAPH COAGULATION ANALYZER  
3000S**Jan 30, 1996  
130 days to decisionK954437 · Product code: **GKP** · Hematology  
Source: <https://www.510kdatabase.net/k954437/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Coagulation, Automated (GKP)
Date received	Sep 22, 1995
Decision date	Jan 30, 1996
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Haemoscope Corp.</b>
Location	Morton Grove, IL, US
Contact	CHARMAINE SUTTON
510(k) history	7 submissions · 7 cleared · 1989-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954437/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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