

**K895844 THROMBELASTOGRAPH(R)**Oct 24, 1989  
21 days to decisionK895844 · Product code: **GKP** · Hematology  
Source: <https://www.510kdatabase.net/k895844/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Coagulation, Automated (GKP)
Date received	Oct 3, 1989
Decision date	Oct 24, 1989
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>Haemoscope Corp.</b>
Location	Morton Grove, IL, US
Contact	ZUCKERMAN, PHD
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/k895844/>; 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 28, 2026