

**K032203 HEMOCUE HEMOGLOBIN 201 + SYSTEM ARTICLE  
121704**Aug 25, 2003  
38 days to decisionK032203 · Product code: **GKR** · Hematology  
Source: <https://www.510kdatabase.net/k032203/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Hemoglobin, Automated (GKR)
Date received	Jul 18, 2003
Decision date	Aug 25, 2003
Days to decision	38 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Hemocue, Inc.</b>
Location	Potomac, MD, US
Contact	R. J SLOMOFF
510(k) history	9 submissions · 9 cleared · 1991-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032203/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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