

**K802266 HGB3 (1:251)**Oct 23, 1980  
37 days to decisionK802266 · Product code: **GKR** · Hematology  
Source: <https://www.510kdatabase.net/k802266/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Hemoglobin, Automated (GKR)
Date received	Sep 16, 1980
Decision date	Oct 23, 1980
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>R&amp;D Systems, Inc.</b>
Location	Mchenry, IL, US
510(k) history	79 submissions · 79 cleared · 1978-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802266/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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