

**K842921 REES & ECKER DILUTION FLUID**Sep 7, 1984  
44 days to decisionK842921 · Product code: **GLG** · Hematology  
Source: <https://www.510kdatabase.net/k842921/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Platelet Counting, Manual (GLG)
Date received	Jul 25, 1984
Decision date	Sep 7, 1984
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>E K Ind., Inc.</b>
Location	Mchenry, IL, US
510(k) history	50 submissions · 49 cleared · 1984-1984

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

Device record: <https://www.510kdatabase.net/k842921/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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