

**K781969 BLOOD, HEMATOLOGY REFERENCE CONTROL**Jan 3, 1979  
40 days to decisionK781969 · Product code: **JPK** · Hematology  
Source: <https://www.510kdatabase.net/k781969/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Nov 24, 1978
Decision date	Jan 3, 1979
Days to decision	40 days
Third-party review	No

**APPLICANT**

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Company	<b>Coulter Electronics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	101 submissions · 101 cleared · 1976-1994

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Device record: <https://www.510kdatabase.net/k781969/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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