

**K053571 MICROGEL ALKALINE HEMOGLOBIN  
ELECTROPHORESIS TEST SYSTEM, MICROGEL ACID  
HEMOGLOBIN ELECTROPHORESIS TEST SYSTEM**Jun 30, 2006  
190 days to decisionK053571 · Product code: **JBD** · Hematology  
Source: <https://www.510kdatabase.net/k053571/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                 |
| Submission type       | Traditional  |
| Device classification | System, Analysis, Electrophoretic Hemoglobin (JBD) |
| Date received         | Dec 22, 2005                                       |
| Decision date         | Jun 30, 2006                                       |
| Days to decision      | 190 days   |
| Third-party review    | No   |
| Summary / Statement   | Statement  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Interlab S.R.L.</b>                |
| Location       | East Stroudsburg, PA, US              |
| Contact        | Gary Lehnus                           |
| 510(k) history | 4 submissions · 4 cleared · 2003-2006 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k053571/>; 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