

**K831026 INDIRECT FLUORESCENT ANTIBODY REAGENTS**Jun 22, 1983  
84 days to decisionK831026 · Product code: **LHL** · Microbiology  
Source: <https://www.510kdatabase.net/k831026/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                                  |
| Submission type       | Traditional   |
| Device classification | Reagents, Antibody, Legionella, Direct & Indirect Fluorescent (LHL) |
| Date received         | Mar 30, 1983  |
| Decision date         | Jun 22, 1983  |
| Days to decision      | 84 days   |
| Third-party review    | No  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Zeus Technologies</b>              |
| Location       | Mchenry, IL, US                       |
| 510(k) history | 4 submissions · 4 cleared · 1983-1984 |

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

Device record: <https://www.510kdatabase.net/k831026/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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