

**K040111 ZEUS SCIENTIFIC, INC., ATHENA MULTI-LYTE  
MPO/PR3 IGG TEST SYSTEM**Apr 5, 2004  
76 days to decisionK040111 · Product code: **MOB** · Immunology  
Source: <https://www.510kdatabase.net/k040111/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                              |
| Submission type       | Traditional   |
| Device classification | Test System, Antineutrophil Cytoplasmic Antibodies (anca) (MOB) |
| Date received         | Jan 20, 2004  |
| Decision date         | Apr 5, 2004   |
| Days to decision      | 76 days   |
| Third-party review    | No  |
| Summary / Statement   | Statement   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Zeus Scientific, Inc.</b>  |
| Location       | Mchenry, IL, US   |
| Contact        | MARK J KOPNITSKY  |
| Website        | <a href="https://www.zeusscientific.com">https://www.zeusscientific.com</a> |
| 510(k) history | 135 submissions · 135 cleared · 1976-2022                                   |

Zeus Scientific, Inc. is a chemistry and immunology device manufacturer based in McHenry, US. The company specializes in flexible autoimmune and infectious disease testing solutions. Zeus Scientific has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company's regulatory portfolio spans microbiology devices and immunology testing systems, including ELISA-based assays and immunofluorescence platforms. The latest clearance on record dates to 2022, reflecting the company's historical contribution to diagnostic device development. ...

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

Device record: <https://www.510kdatabase.net/k040111/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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