

**K891782 HSV-2 ELISA TEST SYSTEM**Jun 20, 1989  
88 days to decisionK891782 · Product code: **GQL** · Microbiology  
Source: <https://www.510kdatabase.net/k891782/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, Herpesvirus Hominis 1,2 (GQL)
Date received	Mar 24, 1989
Decision date	Jun 20, 1989
Days to decision	88 days
Third-party review	No

**APPLICANT**

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Company	<b>Zeus Scientific, Inc.</b>
Location	McHenry, IL, US
Contact	JERRY W PICKERING
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/k891782/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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