

**K914022 SERIOLOGICAL TEST KIT/IGM ANTIBODY VARI-ZOST VIRUS**Jan 27, 1992  
140 days to decisionK914022 · Product code: **LFY** · Microbiology  
Source: <https://www.510kdatabase.net/k914022/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Varicella-zoster (LFY)
Date received	Sep 9, 1991
Decision date	Jan 27, 1992
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Pharmacia Diagnostics, Inc.</b>
Location	Columbia, MD, US
Contact	JUDITH SMITH
510(k) history	7 submissions · 7 cleared · 1990-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914022/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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