

TEST CHANGE

Varicella-Zoster Virus DFA with Reflex to Varicella-Zoster Virus Culture 0060282, VZVFAC

0000202, 1211710		
Specimen Requirements:		
Patient Preparation:	Collect vesicle specimen during first three days of rash.	
Collect:	Vesicle fluid or swab, tissue, or skin scrapings. Swab should be rolled in base of fresh vesicle to obtain cells.	
Specimen Preparation:	Immediately transfer specimen to viral transport media (ARUP Supply #12884) and refrigerate (extremely temperature sensitive). Available online through eSupply using ARUP Connect(TM) or contact ARUP Client Services at (800) 522-2787.	
Transport Temperature:	Refrigerated.	
Unacceptable Conditions:	Bone marrow, CSF, or whole blood. Calcium alginate, eSwab, dry or wood swabs.	
Remarks:	Specimen source preferred.	
Stability:	Ambient: Unacceptable; Refrigerated: 48 hours; Frozen: Unacceptable	
Methodology:	Immunofluorescent Direct Fluorescent Antibody (DFA) Stain / Cell Culture	
Note:	If DFA is negative or inadequate, then a VZV culture will be added. Additional charges apply.	
	Sensitivity of DFA methodology is dependent upon adequacy of the specimen. If specimen contains fewer than 20 cells, then the DFA result will be reported as "inadequate."	
CPT Codes:	87290; if reflexed, add 87252; 87254	
New York DOH Approval Status:	This test is New York DOH approved.	
Interpretive Data:		
Approximately 20% of samples su	bmitted for V7V are positive for herpes simplex virus. Testing	

Effective Date: January 20, 2026

Approximately 20% of samples submitted for VZV are positive for herpes simplex virus. Testing for herpes simplex virus is NOT included in this test.

Reference Interval:

<u>Test</u> <u>Number</u>	<u>Components</u>	Reference Interval
	Varicella-Zoster Virus, DFA	<u>Negative</u>
Negative		

HOTLINE NOTE: There is a component change associated with this test. One or more components have been added or removed. Refer to the Hotline Test Mix for interface build information.



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