

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108 EXAMPLE, REPORT W WX0000003827 M 07/08/1978 45 Y

	Referral	Testing					
		cted: 09/25/2023	3 15:38	Received:	09/25/2023	15:38	
<u>Test Name</u>	<u>Result</u>	Flag	<u>Ref-Range</u>	<u>s l</u>	<u>Jnits</u>	<u>Site</u>	
Diphtheria/Tetanus ABS IgG Diphtheria Ab IgG	7.4			I	U/mL	ARRL	
INTERPRETIVE INFORMATION:	Diphtheria Ab, I	gG					
Antibody concentration of greater than 0.1 IU/mL is usually considered protective.							
Responder status is determ one month post-vaccination concentrations of Diphther	sample to pre-v	accination	of a				
	 If the one month post-vaccination concentration is less than 1.0 IU/mL, the patient is considered to be a non-responder. 						
 If the post-vaccination equal to 1.0 IU/mL, a p 1.5 is a non-responder, a weak responder, and a responder. 	atient with a ra a ratio of 1.5	tio of less to less than	than 3.0,				
 If the pre-vaccination 1.0 IU/mL, it may be di based on a ratio alone. above 2.5 IU/mL in this 	fficult to asses A post-vaccinat	s the respon ion concentr	se				
This test was developed an determined by ARUP Laborat approved by the US Food an was performed in a CLIA ce intended for clinical purp	ories. It has no d Drug Administr rtified laborato	t been clear ation. This	ed or				
Tetanus Ab IgG	12.6			I	U/mL	ARRL	
INTERPRETIVE INFORMATION:	Tetanus Ab, IgG						
Antibody concentration of considered protective.	greater than 0.1	IU/mL is us	ually				
Responder status is determ one-month post-vaccination concentration of Tetanus I	sample to pre-v	accination	of a				
1. If the one month post-v	accination conce	ntration is					

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED

Ordered By: KAJAL SITWALA, MD, PhD WX00000000002365



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Referral Testing								
		Collected: 09/25/2023 15:38 Received: 09/25/2023 15:38						
<u>Test Name</u>		ResultFlagRef-RangesUnitsSiteless than 1.0 IU/mL, the patient is considered a non-responder.						
	2.	If the post-vaccination concentration is greater than or equal to 1.0 IU/mL, a patient with a ratio of less than 1.5 is a non-responder, a ratio of 1.5 to less than 3.0, a weak responder, and a ratio of 3.0 or greater, a good responder.						
	3.	If the pre-vaccination concentration is greater than 1.0 IU/mL, it may be difficult to assess the response based on a ratio alone. A post-vaccination concentration above 2.5 IU/mL in this case is usually adequate.						
	de ap wa in Pe 50 Sa	is test was developed and its performance characteristics termined by ARUP Laboratories. It has not been cleared or proved by the US Food and Drug Administration. This test s performed in a CLIA certified laboratory and is tended for clinical purposes. rformed By: ARUP Laboratories 0 Chipeta Way lt Lake City, UT 84108 boratory Director: Jonathan R. Genzen, MD, PhD						

 Performing Site:

 ARRL: ARUP REFERENCE LAB 500 Chipeta Way Salt Lake City UT 841081221

 Reported Date:
 2023.09.26
 15:38
 DITET

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED