



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988 34 Y

Referral Testing

Collected: 09/28/2023 07:54

Received: 09/28/2023 07:54

Test Name	Result	Flag	Ref-Ranges	Units	Site
Haemophilus influenza B Ab IgG	1.6			ug/mL	ARRL

INTERPRETIVE INFORMATION: H. Influenzae b Ab, IgG

Less than 1.0 ug/mL Antibody concentration not protective.
 1.0 ug/mL or greater Antibodies to H. Influenzae b detected. Suggestive of protection.

Responder status is determined according to the ratio of post-vaccination concentration to pre-vaccination concentration of Haemophilus influenza b antibody, IgG as follows:

1. If the post-vaccination concentration is less than 3.0 ug/mL, the patient is considered to be a non-responder.
2. If the post-vaccination concentration is greater than or equal to 3.0 ug/mL, a patient with a ratio of greater than or equal to 4 is a good responder, a ratio of 2-4 is a weak responder, and a ratio of less than 2 is considered a non-responder.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Performed By: ARUP Laboratories
500 Chipeta Way
Salt Lake City, UT 84108
Laboratory Director: Jonathan R. Genzen, MD, PhD
CLIA Number: 46D0523979

Performing Site:

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

Reported Date: 2023.09.28 7:54

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

F328000003
WX0000003826
Printed D&T: 09/28/23 07:55

Ordered By: KAJAL SITWALA, MD, PhD
WX0000000002353

Kajal V. Sitwala, MD, PhD - Medical Director
Form: MM RL1
PAGE 1 OF 1