

Client: Example Client ABC123
123 Test Drive
Salt Lake City, UT 84108
UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB: 6/14/1948
Gender: Female
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
Collection Date: 00/00/0000 00:00

Bordetella pertussis Antibodies, IgA, IgG, and IgM by ELISA with Reflex to Immunoblot

ARUP test code 2001775

B. pertussis Ab, IgG by ELISA

0.94 IV (Ref Interval: <=1.04)

When Bordetella pertussis Antibodies by ELISA testing is negative, no further testing is performed.

INTERPRETIVE INFORMATION: B. pertussis Ab, IgG

- 0.94 IV or less: Negative - No significant level of detectable B. pertussis IgG antibody.
- 0.95-1.04 IV: Equivocal - Repeat testing in 10-14 days may be helpful.
- 1.05 IV or greater: Positive - IgG antibody to B. pertussis detected, which may indicate a current or recent exposure/immunization to B. pertussis

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.

B. pertussis Ab, IgM by ELISA

1.0 IV (Ref Interval: <=1.1)

When Bordetella pertussis Antibodies by ELISA testing is negative, no further testing is performed.

INTERPRETIVE INFORMATION: B. pertussis Ab, IgM w/Reflex

- 0.9 IV or less: Negative - No significant level of detectable B. pertussis IgM antibody.
- 1.0-1.1 IV: Equivocal - Repeat testing in 10-14 days may be helpful.
- 1.2 IV or greater: Positive - IgM antibody to B. pertussis detected, which may indicate a current or recent exposure/immunization to B. pertussis.

Recommend that treatment decisions be based on the result of the B. pertussis IgM immunoblot test instead of the ELISA test. B. pertussis IgM test by ELISA may produce false-positive results.

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.

H=High, L=Low, *=Abnormal, C=Critical

B. pertussis Ab, IgA by ELISA

0.4 IV

(Ref Interval: <=1.1)

When Bordetella pertussis Antibodies by ELISA testing is negative, no further testing is performed.

INTERPRETIVE INFORMATION: B. pertussis Ab, IgA w/Reflex

- 0.9 IV or less: Negative - No significant level of detectable B. pertussis IgA antibody.
- 1.0-1.1 IV: Equivocal - Repeat testing in 10-14 days may be helpful.
- 1.2 IV or greater: Positive - IgA antibody to B. pertussis detected, which may indicate a current or past exposure/immunization to B. pertussis.

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.

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
B. pertussis Ab, IgG by ELISA	23-018-133139	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
B. pertussis Ab, IgM by ELISA	23-018-133139	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
B. pertussis Ab, IgA by ELISA	23-018-133139	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at: