MyQuest™

BAGER,ARWA

DOB: 03/31/1981
Sex: F
Phone: (305) 788-5319
Patient ID: 69797

Age: 41
Fasting:

Specimen: MR797267K
Requisition: 1955200
Lab Reference ID: 575436
Report Status: FINAL / SEE REPORT

Collected: 07/06/2022 10:45
Received: 07/06/2022 23:42
Reported: 07/11/2022 22:45

Client #: 11046
EISERMANN,JUERGEN
S FL INST FOR REPROD MED
7300 SW 62ND PL FL 4
SOUTH MIAMI, FL 33143-4800
Phone: (305) 662-7901 ext. 7718
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HIV 1/2 ANTIGEN/ANTIBODY,FOURTH GENERATION W/RFL

HIV 1/2 ANTIGEN/ANTIBODY,FOURTH GENERATION W/RFL

Analyte	Value	
HIV AG/AB, 4TH GEN	**NON-REACTIVE**	Reference Range: NON-REACTIVE

HIV-1 antigen and HIV-1/HIV-2 antibodies were not detected. There is no laboratory evidence of HIV infection.

PLEASE NOTE: This information has been disclosed to you from records whose confidentiality may be protected by state law. If your state requires such protection, then the state law prohibits you from making any further disclosure of the information without the specific written consent of the person to whom it pertains, or as otherwise permitted by law. A general authorization for the release of medical or other information is NOT sufficient for this purpose.

⚠ TSH

Analyte	Value	
⚠ TSH	**5.05 H**	mIU/L

Reference Range

> or = 20 Years 0.40-4.50

Pregnancy Ranges
First trimester 0.26-2.66
Second trimester 0.55-2.73
Third trimester 0.43-2.91

VITAMIN D,25-OH,TOTAL,IA

Analyte	Value	
VITAMIN D,25-OH,TOTAL,IA	**64**	Reference Range: 30-100 ng/mL

Vitamin D Status 25-OH Vitamin D:

Deficiency: <20 ng/mL
Insufficiency: 20 - 29 ng/mL
Optimal: > or = 30 ng/mL

For 25-OH Vitamin D testing on patients on D2-supplementation and patients for whom quantitation of D2 and D3 fractions is required, the QuestAssureD(TM) 25-OH VIT D, (D2,D3), LC/MS/MS is recommended: order code 92888 (patients >2yrs).

COMMENT

See Note 1

Note 1

For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ199 (This link is being provided for informational/ educational purposes only.)

CHLAMYDIA TRACHOMATIS AB (IGG,IGA,IGM)

Analyte	Value

C. TRACHOMATIS AB (IGG)	**<1:64**	Reference Range: <1:64 titer
C. TRACHOMATIS AB (IGA)	**<1:16**	Reference Range: <1:16 titer
C. TRACHOMATIS AB (IGM)	**<1:10**	Reference Range: <1:10 titer

INTERPRETATION **see note**

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Antibody Not Detected

Reference Range:
          IgM  <1:10
          IgG  <1:64
          IgA  <1:16

The immunofluorescent detection of specific antibodies
to Chlamydia trachomatis may be complicated by
crossreactive antibodies, non-specific antibody
stimulation, or past exposure to similar organisms
such as Chlamydophila pneumoniae and C. psittaci. IgM
titers of 1:10 or greater usually indicate recent
infection and any IgG titer may indicate past exposure.
IgA is typically present at low titers during primary
infection, but may be elevated in recurrent exposures
or in chronic infection.


This test was developed and its analytical performance
characteristics have been determined by Quest
Diagnostics Nichols Institute, Chantilly, VA. It has
not been cleared or approved by the U.S. Food and Drug
Administration. This assay has been validated pursuant
to the CLIA regulations and is used for clinical
purposes.
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ABO GROUP AND RH TYPE

Analyte	Value
ABO GROUP	**A**
RH TYPE	**RH(D) POSITIVE**

COMMENT

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For additional information, please refer to
http://education.QuestDiagnostics.com/faq/FAQ111
(This link is being provided for informational/
educational purposes only.)
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HEMOGLOBIN A1c

Analyte	Value	
HEMOGLOBIN A1c	**5.4**	Reference Range: <5.7 % of total Hgb

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For the purpose of screening for the presence of
diabetes:

<5.7%       Consistent with the absence of diabetes
5.7-6.4%    Consistent with increased risk for diabetes
            (prediabetes)
> or =6.5%  Consistent with diabetes

This assay result is consistent with a decreased risk
of diabetes.

Currently, no consensus exists regarding use of
hemoglobin A1c for diagnosis of diabetes in children.

According to American Diabetes Association (ADA)
guidelines, hemoglobin A1c <7.0% represents optimal
control in non-pregnant diabetic patients. Different
metrics may apply to specific patient populations.
Standards of Medical Care in Diabetes(ADA).
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T4 (THYROXINE), TOTAL

Analyte	Value	
T4 (THYROXINE), TOTAL	**11.4**	Reference Range: 5.1-11.9 mcg/dL

T4, FREE

Analyte	Value	
T4, FREE	**1.5**	Reference Range: 0.8-1.8 ng/dL

THYROGLOBULIN ANTIBODIES

Analyte	Value	
THYROGLOBULIN ANTIBODIES	**<1**	Reference Range: < or = 1 IU/mL

THYROID PEROXIDASE ANTIBODIES

Analyte	Value	
THYROID PEROXIDASE ANTIBODIES	**<1**	Reference Range: <9 IU/mL

HEPATITIS B SURFACE ANTIGEN W/REFL CONFIRM

Analyte	Value	
HEPATITIS B SURFACE ANTIGEN	**NON-REACTIVE**	Reference Range: NON-REACTIVE

HEPATITIS C AB W/REFL TO HCV RNA, QN, PCR

Analyte	Value	
HEPATITIS C ANTIBODY	**NON-REACTIVE**	Reference Range: NON-REACTIVE
INDEX	**0.01**	Reference Range: <1.00

HCV antibody was non-reactive. There is no laboratory evidence of HCV infection.

In most cases, no further action is required. However, if recent HCV exposure is suspected, a test for HCV RNA (test code 35645) is suggested.

For additional information please refer to http://education.questdiagnostics.com/faq/FAQ22v1 (This link is being provided for informational/ educational purposes only.)

RUBELLA AB (IGG), IMMUNE STATUS

Analyte	Value	
RUBELLA AB (IGG), IMMUNE STATUS	**6.32**	Index

Index	Interpretation
<0.90	Not consistent with immunity
0.90-0.99	Equivocal
> or = 1.00	Consistent with immunity

The presence of rubella IgG antibody suggests immunization or past or current infection with rubella virus.

VARICELLA ZOSTER VIRUS ANTIBODY (IGG)

Analyte	Value

Analyte	Value	
VARICELLA ZOSTER VIRUS ANTIBODY (IGG)	**1557.00**	index

Index	Interpretation
<135.00	Negative - Antibody not detected
135.00 - 164.99	Equivocal
> or = 165.00	Positive - Antibody detected

A positive result indicates that the patient has antibody to VZV but does not differentiate between an active or past infection. The clinical diagnosis must be interpreted in conjunction with the clinical signs and symptoms of the patient. This assay reliably measures immunity due to previous infection but may not be sensitive enough to detect antibodies induced by vaccination. Thus, a negative result in a vaccinated individual does not necessarily indicate susceptibility to VZV infection. A more sensitive test for vaccination-induced immunity is Varicella Zoster Virus Antibody Immunity Screen, ACIF.

PROLACTIN

Analyte	Value	
PROLACTIN	**5.2**	ng/mL

	Reference Range
Females	
Non-pregnant	3.0-30.0
Pregnant	10.0-209.0
Postmenopausal	2.0-20.0

RPR (DX) W/REFL TITER AND CONFIRMATORY TESTING

Analyte	Value	
RPR (DX) W/REFL TITER AND CONFIRMATORY TESTING	**NON-REACTIVE**	Reference Range: NON-REACTIVE

Performing Sites

AMD Quest Diagnostics/Nichols Chantilly-Chantilly VA, 14225 Newbrook Dr, Chantilly, VA 20151-2228 Laboratory Director: Patrick W Mason M.D.,PhD

MI Quest Diagnostics-Miami, 10200 Commerce Pkwy, Miramar, FL 33025-3938 Laboratory Director: Julie L Friedman, MD

Key

Priority Out of Range Out of Range