Laboratory Report - Harvard University Health Services

GREEN, TESSA D MCGINTY, SHANNON - 1710271523 Name: Ordered by: 70834480 L1649055-06 (WC558463Y) ld: Order #: 30 yrs at result time **Collected:** 4/14/2023 1:03 PM Age: Received: 4/14/2023 1:06 PM DOB: 3/26/1993 **EST** Timezone: Sex: Spec. Type:

36336 - CELIAC DISEASE COMP PANEL W/GLIADIN AB (IGG)

Final Status:

Reported:

QUEST DIAGNOSTICS Lab Name: MASSACHUSETTS LLC

Test Name Result

4/27/2023 9:00 PM

INTERPRETATION SEE BELOW No serological evidence for celiac disease is present.

Reference Range

Reference Range

<10.4 umol/L

Flags

U/mL

Flags

tTg may normalize in individuals with celiac disease who maintain a gluten free diet. If high suspicion of celiac disease, consider HLA DQ2 and DQ8 testing to rule out celiac disease.

Antibody not detected

Antibody detected

TISSUE TRANSGLUTAMINASE

AB, IGA Value Interpretation

<15.0

IMMUNOGLOBULIN A 109 47-310 mg/dL

30517 - METHYLMALONIC ACID AND HOMOCYSTEINE

TEST PERFORMED AT: QUEST Lab Name:

Status: Final

HOMOCYSTEINE

Reported:

> or = 15.0

DIAGNOSTICS NICHOLS INSTITUTE -CHANTILLY

4/27/2023 9:00 PM

Test Name

METHYLMALONIC ACID **72*** 87-318 nmol/L

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.

6.0

Result

Homocysteine is increased by functional deficiency of folate or vitamin B12. Testing for methylmalonic acid differentiates between these deficiencies. Other causes of increased homocysteine include renal failure, folate antagonists such as methotrexate and phenytoin, and exposure to nitrous oxide. Selhub J, et al. Ann Intern Med. 1999;131(5):331-9.

568 - INTRINSIC FACTOR BLOCKING ANTIBODY

4/27/2023 9:00 PM Reported: Final Status: Lab Name: TEST PERFORMED AT: QUEST DIAGNOSTICS NICHOLS INSTITUTE -

CHANTILLY

Test Name Reference Range Result **Flags** INTRINSIC FACTOR BLOCKING Negative Negative **ANTIBODY**

For additional information, please refer to

http://education.questdiagnostics.com/faq/IFAB (This link is being provided for informational/ educational purposes only.)

15114 - PARIETAL CELL AB, ELISA

Reported: 4/27/2023 9:00 PM Status: Final

TEST PERFORMED AT: QUEST Lab Name: DIAGNOSTICS NICHOLS INSTITUTE -

Positive

CHANTILLY

Test Name Reference Range Result PARIETAL CELL AB, ELISA 31.5* <=20.0 Unit Н

Reference Range:

>= 25.0

<= 20.0 Negative 20.1 - 24.9Equivocal

Anti-gastric parietal cell antibodies (Anti-GPA) were previously tested for by indirect immunofluorescence (IF) using mouse stomach as a substrate.

Identification of the specific antibody target as H+/K+ ATPase protein (a gastric proton pump) has led to the development of an ELISA based assay. Antibodies to this protein are present in approximately 80% of patients with pernicious anemia and a small percentage of general adult population. The latter percentage increases with age and may reflect the presence of atrophic gastritis. A negative test does not exclude a diagnosis of pernicious anemia. A test for intrinsic factor blocking antibody (IFab) may provide serological evidence in support of the diagnosis in some of these patients.

206 - ACETYLCHOLINE RECEPTOR BINDING ANTIBODY 4/27/2023 9:00 PM Reported:

Final Status: Lab Name: QUEST DIAGNOSTICS INCORPORATED,

NICHOLS INSTITUTE

Test Name Result

ACETYLCHOLINE RECEPTOR nmol/L < 0.30 **BINDING ANTIBODY**

Reference Range

Reference Range

Flags

Flags

Reference Ranges for Acetylcholine Receptor Binding Antibody:

Negative: < or =0.30 nmol/L Equivocal: 0.31-0.49 nmol/L Positive: > or =0.50 nmol/L

93881 - ACETYLCHOLINE RECEPTOR GANGLIONIC (ALPHA 3) AB

4/27/2023 9:00 PM Reported: Status: Final

Test Name

QUEST DIAGNOSTICS INCORPORATED, Lab Name: NICHOLS INSTITUTE

<55

Result

ACETYLCHOLINE RECEPTOR <55 pmol/L GANGLIONIC (ALPHA 3) AB

Acetylcholine Receptor Ganglionic Antibody: <55 pmol/L Negative:

Reference Ranges for

Borderline: 55-160 pmol/L Positive: >160 pmol/L This test was developed and its analytical performance

characteristics have been determined by Quest Diagnostics Nichols Institute San Juan Capistrano. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

Performing Laboratory Information QUEST DIAGNOSTICS Name:

MASSACHUSETTS LLC 200 FOREST STREET, 3RD FLOOR, SUITE A, MARLBOROUGH MA 01752-3023, SALIM E KABAWAT, MD TEST PERFORMED AT: QUEST Name: DIAGNOSTICS NICHOLS INSTITUTE -

CHANTILLY 14225 NEWBROOK DRIVE, PO BOX 10841, CHANTILLY, VA 20153 KENNETH

NICHOLS INSTITUTE

33608 ORTEGA HWY., SAN JUAN

CAPISTRANO, CA 92675

L. SISCO, MD QUEST DIAGNOSTICS INCORPORATED, Name: