

16020 Linden Ave North, Shoreline, Washington 98133, USA

## **Celiac Antibody Panel**

## **Complete Report**

Physician: Sample Report

Patient:

Received:

Completed:

Accession #: 2017000000 Completed:

Sex: Date of Birth:

Age: Sample Type: Serum

## Reference Range (chemiluminescent units, CU)

Analyte	Result	Indication	Negative	Weak Positive	Positive
Deamidated Gliadin Peptide IgA (DGP IgA)	24	Positive	<20	20 - 30	>30
Deamidated Gliadin Peptide IgG (DGP IgG)	42.1	Positive	<20	20 - 30	>30
Tissue Transglutaminase IgA (h-tTG IgA)	28.5	Positive	<20	20 - 30	>30
Tissue Transglutaminase IgG (h-tTG IgG)	4	Negative	<20	20 - 30	>30

<sup>&</sup>lt;: less than reportable range.

Commentary (semi-quantitative chemiluminescent immunoassay, CIA)

The results of this test were obtained with the FDA-approved INOVA QUANTA Flash® CIA immunoassay. Values obtained with different manufacturers' assay methods may not be used interchangeably.

Clinical sensitivity and specificity of h-tTG IgA QUANTA Flash® are reported at 94.0% and 98.1%, respectively.

Clinical sensitivity and specificity of DGP IgA QUANTA Flash® are reported at 71.4% and 100%, respectively.

Not all patients with celiac disease are positive for h-tTG IgA autoantibodies or DGP IgA antibodies. A negative result in an untreated suspect patient may be explained by selective IgA deficiency, a relatively frequent finding in this population. The presence of h-tTG IgG autoantibodies and DGP IgG antibodies can therefore aid in the patient assessment.

Clinical sensitivity of this method for h-tTG IgG autoantibodies has been shown to be 85.7% in a subset of selective IgA deficient patients.

Individuals on a gluten-free diet prior to testing may show low serological values.

Results of this assay should not be interpreted in the absence of a complete clinical history.

Confirmation of celiac disease requires small bowel biopsies demonstrating immune-mediated villous atrophy in addition to resolution of symptoms following the introduction and maintenance of a strict gluten-free diet.

This test is not intended to diagnose, treat, cure, or prevent any disease or replace the medical advice and/or treatment obtained from a qualified healthcare practitioner.

End of Report

CLIA: 50D0965661 COLA accredited

Director: Stephen Markus, MD ©2015 US BioTek Laboratories, Inc.

<sup>&</sup>gt;: greater than reportable range.