

16020 Linden Ave North, Shoreline, Washington 98133, USA

Celiac Panel, IgG&IgA/DGP + IgG&IgA/tTG

Complete Report

Provider: Collected:
Patient: Received:
Accession #: Completed:
Sex: Date of Birth:
Age: Sample Type:

| | | | Reference Range | | | |
|--|--------|---------------|-----------------|----------------------|-----------------|--|
| Analyte | Result | Indication | Negative | Weak Positive | Positive | |
| Deamidated Gliadin Peptide IgA (DGP IgA) | 6.7 | Negative / | <20 | 20 - 30 | >30 | |
| Deamidated Gliadin Peptide IgG (DGP IgG) | 7.5 | Negative | <20 | 20 - 30 | >30 | |
| Tissue Transglutaminase IgA (h-tTG IgA) | 149.1 | Positive | <20 | 20 - 30 | >30 | |
| Tissue Transglutaminase IgG (h-tTG IgG) | 20.8 | Weak Positive | <20 | 20 - 30 | >30 | |

<: 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: Jillian Harrington, PhD, HCLD (ABB)

>: greater than reportable range.