

Patient ID:  
Specimen ID:

DOB:  
Age:  
Sex:

# Patient Report

Ordering Physician:



Ordered Items: **Celiac Disease HLA DQ Assoc.; Venipuncture**

Date Collected:	Date Received:	Date Reported:	Fasting:
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## Celiac Disease HLA DQ Assoc.

Test	Current Result and Flag	Previous Result and Date	Units	Reference Interval
DQ2 (DQA1 0501/0505,DQB1 02XX) <sup>01</sup>	Negative			
DQ8 (DQA1 03XX, DQB1 0302) <sup>01</sup>	Negative			

**Final Results:**

DQA1\*01:DMNEZ, 01:DMWEC

DQB1\*05:DMPXJ, 05:DMZDX

**Code Translation:**

DMNEZ 01/04/05/12/18/22/26/27/29/34/35/37/49/53/55/56

DMPXJ 05:01/05:07/05:11/05:12/05:18/05:20/05:21/05:22/05:25/05:27/05:30/05:31/05:32/05:44/05:45/05:48/05:49/05:51/05:54/05:55/05:61/05:62/05:68/05:71/05:72/05:73/05:74/05:75/05:76/05:80/05:81/05:82/05:84/05:88/05:89/05:92/05:93/05:95/05:103/05:104/05:105/05:107/05:110N/05:111/05:112/05:114/05:115/05:120/05:122/05:124/05:126/05:128N/05:133/05:137/05:138/05:139/05:144/05:148/05:150/05:151/05:152/05:154/05:155/05:157/05:158/05:159/05:160/05:162/05:163/05:164/05:166/05:167/05:168/05:169/05:171/05:173/05:176/05:177/05:180/05:182/05:183/05:184/05:185N/05:188/05:190/05:193/05:194/05:195/05:197/05:215N/05:216/05:217/05:219/05:223/05:225/05:226/05:228/05:230/05:232/05:234/05:237/05:240/05:242/05:246/05:248/05:249/05:252/05:255/05:256/05:258/05:261/05:263/05:266/05:267/05:268/05:269/05:270/05:271/05:272

DMWEC 01/02/04/05/12/18/22/26/27/29/34/35/37/49/53/55/56

DMZDX 05:03/05:06/05:08/05:10/05:13/05:15/05:16/05:23/05:24/05:28/05:38/05:39/05:40/05:41N/05:42/05:43/05:50/05:56/05:60/05:66/05:67/05:78/05:85/05:91/05:96/05:98/05:108/05:109/05:121/05:130/05:134/05:140/05:147/05:149/05:161/05:191/05:200/05:201/05:202/05:203/05:204/05:205/05:206N/05:207/05:208N/05:209/05:211/05:212/05:213/05:214/05:218/05:220/05:221/05:224N/05:233/05:235N/05:236N/05:244/05:245/05:259/05:260/05:264/05:265N/05:273N

The patient is not positive for any of the HLA DQ risk alleles. Celiac Disease risk from the HLA DQA/DQB genotype is approximately 1:2518 (<0.04%). Allele interpretation for all loci based on IMGT/HLA

Patient ID:  
Specimen ID:

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## Patient Report

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### Celiac Disease HLA DQ Assoc. (Cont.)

database version 3.43.0

HLA Lab CLIA ID Number 34D0954530

Greater than 95% of celiac patients are positive for either DQ2 or DQ8 (Sollid and Thorsby, (1993) Gastroenterology 105:910-922). However these antigens may also be present in patients who do not have Celiac disease.

#### Comment:<sup>01</sup>

This test was performed using Polymerase Chain Reaction/(PCR)Sequence Specific Oligonucleotide Probes (SSOP) (Luminex) technique. Sequence Based Typing (SBT) and/or Sequence Specific Primers (SSP) may be used as supplemental methods when necessary. Please contact HLA Customer Service at 1-800-533-1037 if you have any questions.  
Director of HLA Laboratory  
Dr George C Maha, PhD

#### Additional Information:<sup>01</sup>

- 357:1731-1743.
- Megiorni F, Mora B, Bonamico M et al. HLA-DQ and risk gradient for celiac disease. Hum Immunol 2009; 70:55-59.
  - Pietzak MM, Schofield TC, McGinnis FM et al. Stratifying risk for celiac disease in a large at-risk United States population by using HLA alleles. Clin Gastroenterol Hepatol 2009; 7:966-971.
  - Sollid LM and Lie BA. (2005). Celiac Disease Genetics: Current Concepts and Practical Applications. Clin Gastroenterol and Hepat 3:843-851.
  - Snyder CL, Young DO, Green PHR, et al. Celiac Disease. In: Pagon RA, Bird TC, Dolan CR, Stephens K, editors. GeneReviews (Internet), University of Washington, Seattle, July 3, 2008:1-27. <http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=genepart=celiac> PMID 20301720 (PubMed)
  - Treem W. Emerging concepts in celiac disease. Curr Opin Pediatr 2004;16:552-559.

#### Disclaimer

The Previous Result is listed for the most recent test performed by Labcorp in the past 3 years where there is sufficient patient demographic data to match the result to the patient.

#### Icon Legend

▲ Out of reference range ■ Critical or Alert

#### Performing Labs

DOB:

## Patient Report



Patient ID:  
Specimen ID:

Age:  
Sex:

Ordering Physician:

### Patient Details

Phone:  
Date of Birth:  
Age:  
Sex:  
Patient ID:  
Alternate Patient ID:

### Physician Details

**Request A Test**  
**7027 Mill Road Suite 201, BRECKSVILLE, OH,**  
**44141**

Phone: **888-732-2348**  
Physician ID:  
NPI:

### Specimen Details

Specimen ID:  
Control ID:  
Alternate Control Number:  
Date Collected:  
Date Received:  
Date Entered:  
Date Reported:  
Rte:

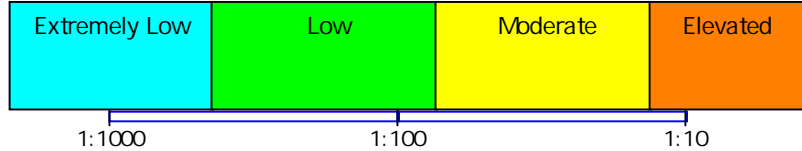
Specimen Number	BM Specimen #	Patient Name	Sex	Age	Date of Birth	Account #	Account Phone	Route
Control Number	Patient ID	Patient SSN	Physician Name		NPI	Physician ID		
Patient Address			Patient Phone		Account Address			
Additional Information				Party ID	NMDP/UNOS ID	Drive Number		
				Date Collected	Date Entered	Date Reported		

**Celiac Disease HLA DQA/DQB Association**

Result: NEGATIVE for celiac-associated allele(s)

Genetic Risk: Extremely Low

1:2518



HLA DQ alleles detected

DQA1\*01:DMNEZ, 01:DMMEC  
DQB1\*05:DMPXJ, 05:DMZDX

DQ2

DQA1\*05:01/05:05  
DQB1\*02:01/02:02

NEGATIVE for DQ2

DQ8

DQA1\*03:XX  
DQB1\*03:02

NEGATIVE for DQ8

HLA allele interpretation based on IMGT/HLA database version 3.43.0

The patient is not positive for any of the HLA DQ risk alleles. Celiac Disease risk from the HLA DQA/DQB genotype is approximately 1:2518 (< 0.04%).

Code/G Group Translation

DMNEZ 01/04/05/12/18/22/26/27/29/34/35/37/49/53/55/56

DMPXJ 05:01/05:07/05:11/05:12/05:18/05:20/05:21/05:22/05:25/05:27/05:30/05:31/05:32/05:44/05:45/05:48/05:49/05:51/05:54/05:55/05:61/05:62/05:68/05:71/05:72/05:73/05:74/05:75/05:76/05:80/05:81/05:82/05:84/05:88/05:89/05:92/05:93/05:95/05:103/05:104/05:105/05:107/05:110N/05:111/05:112/05:114/05:115/05:120/05:122/05:124/05:126/05:128N/05:133/05:137/05:138/05:139/05:144/05:148/05:150/05:151/05:152/05:154/05:155/05:157/05:158/05:159/05:160/05:162/05:163/05:164/05:166/05:167/05:168/05:169/05:171/05:173/05:176/05:177/05:180/05:182/05:183/05:184/05:185N/05:188/05:190/05:193/05:194/05:195/05:197/05:215N/05:216/05:217/05:219/05:223/05:225/05:226/05:228/05:230/05:232/05:234/05:237/05:240/05:242/05:246/05:248/05:249/05:252/05:255/05:256/05:258/05:261/05:263/05:266/05:267/05:268/05:269/05:270/05:271/05:272

DMMEC 01/02/04/05/12/18/22/26/27/29/34/35/37/49/53/55/56

Specimen Number	BM Specimen #	Patient Name	Sex	Age	Date of Birth	Account #	Account Phone	Route
Control Number	Patient ID	Patient SSN	Physician Name		NPI	Physician ID		
Patient Address		Patient Phone	Account Address					
Additional Information			Party ID	NMDP/UNOS ID	Drive Number			
			Date Collected	Date Entered	Date Reported			

**Code/G Group Translation**

DMZDX 05:03/05:06/05:08/05:10/05:13/05:15/05:16/05:23/05:24/05:28/05:38/05:39/05:40/05:41N/05:42/05:43/  
05:50/05:56/05:60/05:66/05:67/05:78/05:85/05:91/05:96/05:98/05:108/05:109/05:121/05:130/05:134/  
05:140/05:147/05:149/05:161/05:191/05:200/05:201/05:202/05:203/05:204/05:205/05:206N/05:207/  
05:208N/05:209/05:211/05:212/05:213/05:214/05:218/05:220/05:221/05:224N/05:233/05:235N/05:236N/  
05:244/05:245/05:259/05:260/05:264/05:265N/05:273N

The range of genetic risk for individuals with a celiac disease-associated genotype is 1:1842 (0.05%) to 1:7 (14.3%). See table "Genetic Risk from HLA-DQA/QB Genotypes" on page 2.

**The ACTUAL risk for this individual to have celiac disease may be significantly higher if there are symptoms of celiac disease, positive results from celiac antibody tests, positive intestinal biopsy, or family members with celiac disease.**

Greater than 90% of celiac patients are positive for DQ2, 5-10% carry DQ8, and the remaining carry half of the DQ molecules (Green and Cellier, 2007). However, the majority of individuals positive for celiac-associated HLA alleles do not develop celiac disease, and detection of these alleles alone is not sufficient for a diagnosis of celiac disease. Relatives of individuals positive for one or more celiac-associated HLA alleles are also at risk for being positive.

This test was performed using a Polymerase Chain Reaction/(PCR) Sequence Specific Oligonucleotide Probes (SSOP) technique on the Luminex platform. This test has been cleared by the U.S. Food and Drug Administration. Analytic sensitivity and specificity are > 99.9%. Sequence-based Typing (SBT) and/or Sequence Specific Primers (SSP) may be used as supplemental methods when necessary. This test evaluates HLA-DQA and DQB genotypes and cannot detect abnormalities elsewhere in the genome. It should be realized that there are many possible sources of diagnostic error including sample misidentification, rare technical errors, trace contamination of PCR reactions, and rare genetic variants that interfere with analysis.

This laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA-88) as qualified to perform high complexity clinical laboratory testing. Please contact HLA customer service at 1-800-533-1037 if you have any questions.

## INFORMATION ABOUT CELIAC DISEASE GENETICS

Celiac disease is a chronic immune-mediated inflammatory disorder with multi-systemic manifestations, both gastrointestinal and non-gastrointestinal. In genetically susceptible individuals, ingestion of gluten can cause inflammation and damage to the small intestine mucosa. Celiac disease has an incidence of 1:100 in the United States.

In order for celiac disease to develop, human leukocyte antigen (HLA) molecule DQ2 (encoded by alleles DQA1\*0501 or \*0505 plus DQB1\*0201 or \*0202), half of the DQ2 molecule, or DQ8 (encoded DQA\*03 plus DQB1\*0302) must be present. These molecules confer susceptibility to celiac disease by binding to gluten and interacting with intestinal T cells, leading to a pathologic immune response involving autoimmunity. The familial nature of susceptibility to celiac disease is shown by an 11-18% prevalence of this disorder in siblings of individuals with celiac disease and a 70% concordance rate between identical twins.

Among celiac disease patients, >90% carry DQ2, 5-10% carry DQ8, and the remaining carry half DQ2. The presence of DQ2, half DQ2, or DQ8 alone is not sufficient for a diagnosis of celiac disease. Clinical symptoms, positive test results for endomysial, tissue transglutaminase or deamidated gliadin peptide antibodies, or abnormal small bowel biopsy results all support a diagnosis of celiac disease. Most individuals with a positive genetic result do not develop celiac disease. The risk for developing celiac disease in individuals with a positive genetic result approaches 40% if there is a known first degree relative with celiac disease.

### Genetic Risk from HLA-DQA/DQB Genotypes

Genotype	Risk
DQ2 + DQ8	1:7 (14.3%)
DQ2 + DQ2 OR DQ2 Homozygous *02	1:10 (10%)
DQ8 + DQ8	1:12 (8.4%)
DQ8 + DQB1*02	1:24 (4.2%)
Homozygous DQB*02	1:26 (3.8%)
DQ2 alone	1:35 (2.9%)
DQ8 alone	1:89 (1.1%)
Population risk (genotype unknown)	1:100 (1%)
1/2 DQ2: DQB1*02	1:210 (0.5%)
1/2 DQ2: DQA1*05	1:1842 (0.05%)
No HLA-DQA/DQB susceptibility alleles	1:2518 (<0.04%)

From Megiorni et al. 2009 for all genotypes except DQ8+DQ8

DQ8+DQ8 risk is from Pietzak et al. 2009

### Other influences on risk for celiac disease

The overall risk for an individual to develop celiac disease is influenced not just by genetic risk from the HLA-DQA/DQB genotype, but by presence of symptoms of celiac disease, positive results for celiac antibody tests or intestinal biopsy, and having relatives with celiac disease. Celiac disease risk is also higher in individuals with IgA deficiency, Down syndrome, Turner syndrome, and the autoimmune disorders Type I diabetes mellitus, Sjogren syndrome, and thyroiditis. There are also additional genetic influences on the development of celiac disease in individuals predisposed to the disorder.

## REFERENCES

1. **Green PHR** and Cellier C. Celiac Disease. *N Eng J Med* 2007; 357:1731-1743.
2. **Megiorni F**, Mbra B, Bonamico M et al. HLA-DQ and risk gradient for celiac disease. *Hum Immunol* 2009; 70:55-59.
3. **Pietzak MM**, Schofield TC, McGinnis FM et al. Stratifying risk for celiac disease in a large at-risk United States population by using HLA alleles. *Clin Gastroenterol Hepatol* 2009; 7:966-971.
4. **Sollid LM and Lie BA**. (2005). Celiac Disease Genetics: Current Concepts and Practical Applications. *Clin Gastroenterol and Hepat* 3:843-851.
5. **Snyder CL**, Young DO, Green PHR, et al. Celiac Disease. In: Pagon RA, Bird TC, Dolan CR, Stephens K, editors. GeneReviews (Internet), University of Washington, Seattle, July 3, 2008:1-27.  
<http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=genepart=celiac> PMID 20301720 (PubMed)
6. **Treem W**. Emerging concepts in celiac disease. *Curr Opin Pediatr* 2004; 16:552-559.