HLA-DQA/DQB genotyping to aid in excluding the diagnosis of celiac disease and for risk assessment

Background
- Celiac disease is estimated to affect 1 out of 100 people, but fewer than 5% of patients are diagnosed as having the disease.¹ ³
- Early diagnosis and lifelong treatment with a gluten-free diet is critical to relieve symptoms and reduce the risk of complications such as secondary autoimmune disorders.¹ ³
- Diagnosis is challenging due to variable, nonspecific symptoms and varying age of onset.³ ⁴
- Tissue transglutaminase (tTG) and endomysial (EMA) IgA antibody testing are the most commonly used screening tests.¹ ³

Clinical Utility
- The HLA-DQ2 allele has been found in 90% to 95% of celiac cases, and HLA-DQ8 has been identified in most remaining cases.¹
- A negative HLA-DQA/DQB genotyping result essentially excludes celiac disease as the diagnosis.¹ ³ ⁵
- A positive result is not diagnostic but indicates a higher probability for celiac disease.³

Indications for Testing
- When diagnosis of celiac disease is unclear
  - Ambiguous antibody test result (tTG IgA or EMA IgA)³
  - Equivocal small bowel biopsy results³
  - Discrepancy between antibody and biopsy findings⁶
- When patient is on a gluten-free diet
  - Antibody testing is not helpful in this setting.³
  - HLA-DQA/DQB genotyping is unaffected by diet and is thus a useful alternative.³
  - If patient has maintained a gluten-free diet with no reduction in symptoms, HLA DQA/DQB genotyping can aid in excluding celiac as the source of symptoms.³ ⁶
- Evaluation of asymptomatic relatives
  - Presence or absence of celiac-associated HLA alleles can help assess celiac risk in first-degree relatives of affected patients.³
Scientific Excellence

- HLA-DQA/DQB genotyping provides detection of DQ2 (DQA1*0501, DQA1*0505, and DQB1*0201/*0202) and DQ8 (DQB1*0302)
- Report includes DQ2, DQ8, half DQ2, homozygosity for DQB1*02, and complete DQA and DQB genotypes
- Easy-to-interpret risk assessment
- Assay performed using FDA-cleared HLA test kits

Estimated Celiac Risk from Associated HLA-DQA/DQB Genotypes

<table>
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<tr>
<th>Genotype</th>
<th>Risk</th>
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<tr>
<td>DQ2+DQ8</td>
<td>1:7 (14.3%)</td>
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<tr>
<td>DQ2+DQ2 or DQ2 Homozygous DQB1*02</td>
<td>1:10 (10%)</td>
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<tr>
<td>DQ8+DQ8</td>
<td>1:12 (8.42%)</td>
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<tr>
<td>DQ8+DQB1*02</td>
<td>1:24 (4.2%)</td>
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<tr>
<td>Homozygous DQB1*02</td>
<td>1:26 (3.8%)</td>
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<tr>
<td>DQ2 alone</td>
<td>1:35 (2.9%)</td>
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<tr>
<td>DQ8 alone</td>
<td>1:89 (1.1%)</td>
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<tr>
<td>General population risk (genotype unknown)</td>
<td>1:100 (1%)</td>
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<tr>
<td>½ DQ2: DQB1*02</td>
<td>1:210 (0.5%)</td>
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<tr>
<td>½ DQ2: DQA1*05</td>
<td>1:1842 (0.05%)</td>
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<tr>
<td>No HLA-DQA/DQB susceptibility alleles</td>
<td>1:2518 (&lt;0.04%)</td>
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Note: Actual risk for celiac disease may be greater than shown above when there are symptoms of celiac disease, positive results for celiac antibody tests or small bowel biopsy, or if there is a family history of celiac disease.

Superior Service

- Comprehensive test menu for celiac disease, including antibody testing
- Consultative services from our scientific staff
- Broad network of managed care health plans
- Nationwide network of patient service centers

Visit the online Test Menu at www.LabCorp.com for full test information, including CPT codes and specimen collection requirements.

References

For more information or celiac genetic consultation, please call HLA customer service at 800-533-1037.