

Pathology Report

DAHLSTROM, VERA

For Surgery Use Urgent Ring Patient Make Appointment Note in Chart File

UR No.

Patient HANNON, PAUL

Patient Address 22 MULGRAVE RD YUNGABURRA QLD 4884

Sex M Age 74 years DOB 30/09/1950 Requested 16/01/2025

Report For DAHLSTROM, VERA

Collected 14/02/2025 09:00 AM

Ref. by/copy to DAHLSTROM, VERA

Reported 24/02/2025 02:29 PM

COELIAC DISEASE GENOTYPING RESULTS

SPECIMEN:

Blood

HLA-DQ2.5 alleles

HLA-DQA1*05:01 = **Heterozygous** (NG_032876.1:g.5702C>T)

HLA-DQB1*02:01 = Heterozygous

HLA-DQ2.2 alleles

HLA-DQA1*02:01 = Not Detected
HLA-DQB1*02:02 = Not Detected

HLA-DQ7 alleles

HLA-DQA1*05:05 = Not Detected
HLA-DQB1*03:01 = Not Detected

HLA-DQ8 alleles

HLA-DQA1*03:01 = Not Detected HLA-DQB1*03:02 = Not Detected

INTERPRETATION:

At-risk genotype detected: Heterozygous HLA-DQ2.5 (intermediate risk). This genotype is associated with genetic susceptibility for coeliac disease.

This result does not confirm a diagnosis of coeliac disease. Further clinical work up, including serology and small intestinal histology, is required for diagnosis (PMID: 25827511).

METHOD:

Testing for HLA haplotypes associated with genetic susceptibility to coeliac disease (DQ2.5, DQ2.2, DQ8 and DQ7.5) is performed with the LABType SSO HLA-DQA1/DQB1 Typing Kit (One Lambda, USA) utilising sequence-specific oligonucleotides probes on the LABScan3D platform (Luminex Corp., USA).

Typing and haplotype assignment is performed using the LabScan 3D software (Immucor, USA), and genetic susceptibility risk levels are

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assigned as per Horan et al, 2018 (PMID: 29764931). DQ haplotypes not associated with coeliac disease are not reported. Rare HLA-DQA1 and HLA-DQB1 alleles may not be detected. Additional loci that may contribute to the development of coeliac disease are not assessed by this test.

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Genomic Diagnostics