

HLA-B*58:01 Genotyping, Allopurinol Hypersensitivity

Client: Example Client ABC123 123 Test Drive Salt Lake City, UT 84108 UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB	9/18/1984
Sex:	Male
Patient Identifiers:	01234567890ABCD, 012345
Visit Number (FIN):	01234567890ABCD
Collection Date:	01/01/2017 12:34

HLA-B*58:01 Genotyping, Allopurinol Hype	Positive *
	Result: Positive for the HLA-B*58:01 allele
	Interpretation: The HLA-B*58:01 allele was detected in this patient. The presence of this allele increases risk for allopurinol-induced severe cutaneous adverse reactions (SCAR), including Stevens-Johnson syndrome (SJS) or toxic epidermal necrolysis (TEN). Allopurinol treatment is contraindicated. Therapy should be discontinued immediately if symptoms of SJS or TEN develop. Alternative medication should be used as first line therapy. Other genetic or non-genetic factors that may affect hypersensitivity to allopurinol are not identified. Rare and undocumented alleles may occur which could lead to false positive or false negative results. Performed By:
	BACKGROUND INFORMATION: HLA- B*58:01 Genotyping, Allopurinol Hypersensitivity CHARACTERISTICS: Allopurinol is the most commonly used drug for the treatment of hyperuricemia and gout. It inhibits xanthine oxidase, a key enzyme involved in uric acid formation. However, allopurinol is one of the most common causes of life-threatening severe cutaneous adverse reactions (SCAR), which include drug hypersensitivity syndrome, Stevens-Johnson syndrome (SJS) and toxic epidermal necrolysis (TEN). The presence of HLA-B*58:01 allele shows strong association with allopurinol-induced SCAR, including TEN and SJS. Although allopurinol-induced SCAR is rare with an estimated risk of 0.1-0.4 percent in allopurinol users, the severity can be high, with a mortality rate of up to 25 percent. Symptoms include rash, combined with eosinophilia, leukocytosis, fever, hepatitis and progressive kidney failure. Due to the severity of adverse reactions, it is recommended to test for the HLA-B*58:01 allele prior to initiation of the drug. INCIDENCE: HLA-B*58:01 allele frequency varies by ethnicity. In the US population, the highest incidence at 5.3 percent in Native Hawaiians or Pacific Islanders, 1.35 percent in Hispanics, 1.19 percent in African Americans or Alaska Natives and 0.8 percent in Caucasians. Frequencies may be higher in other countries, up to 20 percent in Singapore, Taiwan and among Han Chinese, 15.4 percent in India, 14.2 percent in Hong Kong, 12 percent in China and Korea, 11 percent in Indonesia. CAUSE: Allopurinol-induced SCAR, including SJS and TEN, is strongly associated with the presence of one or two copies of HLA-B*58:01 allele. The mechanism is immune mediated and involves direct interactions the allopuring metabolito

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

ARUP LABORATORIES | 800-522-2787 | aruplab.com 500 Chipeta Way, Salt Lake City, UT 84108-1221 Jonathan R. Genzen, MD, PhD, Laboratory Director

Patient: Patient, Example ARUP Accession: 22-119-136384 Patient Identifiers: 01234567890ABCD, 012345 Visit Number (FIN): 01234567890ABCD Page 1 of 2 | Printed: 12/21/2022 11:18:15 AM oxypurinol, and HLA-B*58:01, which may result in drug-induced changes in peptide presentation, allowing activation of self-reactive T lymphocytes. ALLELES TESTED: HLA-B*58:01 allele. CLINICAL SENSITIVITY AND SPECIFICITY: 71 percent sensitivity and 92 percent specificity, overall mean values from pooled populations (Yu KH et al, Int J Rheum Dis 2017). Higher in populations with increased HLA-B*58:01 allele frequency. METHODOLOGY: PCR followed by Sequence Specific Oligonucleotide Probe Hybridization of HLA-B locus. ANALYTICAL SENSITIVITY AND SPECIFICITY: Greater than 99 percent. LIMITATIONS: Copy number of HLA-B*58:01 will not be reported. Other genetic and non-genetic factors that influence allopurinol hypersensitivity are not evaluated. Other rare, or novel alleles may occur which may lead to false positive or false negative results.

Test systems were developed and their performance characteristics determined by the H&I laboratory at the University of Utah Health, under the accreditation guidelines from the American Society for Histocompatibility and Immunogenetics (ASHI).

VERIFIED/REPORTED DATES					
Procedure	Accession	Collected	Received	Verified/Reported	
HLA-B*58:01 Genotyping, Allopurinol Hype	22-119-136384	4/29/2022 3:09:00 PM	5/2/2022 7:44:57 AM	5/3/2022 1:49:00 PM	

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

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