

Report Summary

112

Total Variants

3

Categories

1

Pathway Figures

24

High Risk (+/+)

37

Moderate Risk (+/-)

51

Low Risk (-/-)

Table of Contents

Report Overview 1

Genetic Variants (3 categories) 2

HLA (56 variants) △ △ 2

Covid (29 variants) △ △ 4

Other Immune Factors (27 variants) △ △ 5

Important Disclaimer 6

Quick Navigation: [Overview](#) [Categories](#) [Disclaimer](#)

HLA

56 variants found 7 23 26
(See Figure 6)

Risk Summary: 7 high risk variants, 23 moderate risk variants - Consider discussing with healthcare provider

Quick Navigation: [Overview](#) [Categories](#) [Disclaimer](#)

SNP ID	SNP Name	Risk Allele	Your Allele	Result
rs2844846	HLA-F-AS1, MICE 4945T>A	T	TT	+/-
rs1632933	HLA-G 8178C>T	T	TC	+/-
rs3823339	HLA-A 7727C>G	G	GC	+/-
rs1061235	HLA-A 8057A>T	T	TA	+/-
rs3893538	HLA-W 915T>C	T	TT	+/-
rs2517701	HLA-W 2462A>T	T	AA	-/-
rs2001181	HLA-C 7911A>G	C	TT	-/-
rs1058026	HLA-B 8305T>G	C	AA	-/-
rs3819299	HLA-B His363Pro	G	TT	-/-
rs2523608	HLA-B 7431C>T	A	AG	+/-
rs3094228	HLA-X 441A>G	C	TT	-/-
rs3135394	HLA-DRA 5055G>A	G	AA	-/-
rs9268645	HLA-DRA 5085C>G	G	CC	-/-
rs3129878	HLA-DRA 5293C>A	A	AC	+/-
rs3129881	HLA-DRA 6041T>C	T	TC	+/-
rs3129882	HLA-DRA 6087A>G	G	AG	+/-
rs3129883	HLA-DRA 6694C>T	T	TC	+/-
rs9268658	HLA-DRA 7273G>A	A	GG	-/-
rs3135391	HLA-DRA 7544G>A	G	AG	+/-
rs8084	HLA-DRA 7592A>C	C	AA	-/-
rs2239804	HLA-DRA 8080T>C	C	TT	-/-
rs7192	HLA-DRA Leu242Val	T	TT	+/-
rs3129888	HLA-DRA 8283A>G	G	AG	+/-
rs2239803	HLA-DRA 8390C>T	T	CC	-/-
rs2239802	HLA-DRA 8403G>C	C	CG	+/-
rs3177928	HLA-DRA 8958G>A	A	GG	-/-
rs1041885	HLA-DRA 9332A>T	A	AT	+/-
rs9268831	HLA-DRB9 7239C>T	T	CC	-/-
rs2858331	HLA 4126892A>G	G	AA	-/-
rs2858331	HLA 4126892A>G	G	AA	-/-
rs9276431	HLA-DQA2 4158042T>C	T	TT	+/-
rs2239800	HLA-DQA2 4159063A>G	G	AA	-/-
rs7453920	HLA-DQB2 4175809A>G	G	AA	-/-
rs2051549	HLA-DQB2 4175883G>A	G	GG	+/-
rs1573649	HLA-DQB2 Met1Thr	G	GG	+/-
rs11244	HLA-DOB 9102C>T	A	GG	-/-
rs2856997	HLA-DOB 8050G>T	C	AC	+/-

SNP ID	SNP Name	Risk Allele	Your Allele	Result
rs2071473	HLA-DOB 7221G>A	C	TC	+/-
rs7383287	HLA-DOB 6740T>C	G	AA	-/-
rs2621326	HLA-DOB 5930C>T	G	AG	+/-
rs2071554	HLA-DOB Arg18Gln	T	CC	-/-
rs2071469	HLA-DOB 5043G>A	C	TC	+/-
rs23544	HLA-DMB 4348055C>T	T	CC	-/-
rs151719	HLA-DMB 4348318T>C	C	TC	+/-
rs3128935	HLA-DOA 9986A>G	T	TT	+/*
rs1044429	HLA-DOA 9748G>A	T	CC	-/-
rs376892	HLA-DOA 9503C>T	G	AA	-/-
rs416622	HLA-DOA 9109A>G	C	TT	-/-
rs9276977	HLA-DOA 8549C>T	A	AG	+/-
rs2581	HLA-DOA 7989C>A	T	TG	+/-
rs399604	HLA-DOA 7376A>G	C	TC	+/-
rs2284191	HLA-DOA 5736C>T	A	AG	+/-
rs86567	HLA-DPA1 20534T>C	G	TT	-/-
rs3077	HLA-DPA1 18960C>T	A	AG	+/-
rs2301226	HLA-DPA1 16159A>G	A	GG	-/-
rs9277535	HLA-DPB1 16159A>G	G	AG	+/-

This report is intended to translate your results into an easier to understand form. It is not intended to diagnose or treat. For diagnosis or treatment, please present this to your doctor (or find a doctor on MTHFRSupportTM website under Find a Practitioner). Additionally, genetic mutations are flags that something **could** be wrong and not a guarantee that you are having all or any of the associated issues. Other factors like environment, ethnic background, diet, age, personal history, etc all have a factor in whether a mutation starts to present itself or not and when. Copyright all rights reserved MTHFR SupportTM

Covid

29 variants found 10 10 9

Risk Summary: 10 high risk variants, 10 moderate risk variants - Consider discussing with healthcare provider

Quick Navigation: [Overview](#) [Categories](#) [Disclaimer](#)

SNP ID	SNP Name	Risk Allele	Your Allele	Result
rs2027432	NLRP3 variant	G	AG	+/-
rs3806265	NLRP3 variant	T	TC	+/-
rs10754557	NLRP3 variant	G	AG	+/-
rs1539019	NLRP3 variant	C	AC	+/-
rs10157379	NLRP3 variant	T	TC	+/-
rs1143634	IL1B variant	G	GG	+/*
rs3136558	IL1B variant	A	AA	+/*
rs1143627	IL1B variant	G	AA	-/-
rs2069827	IL6 variant	T	TG	+/-
rs1800797	IL6 variant	G	AG	+/-
rs2066992	IL6 variant	T	GG	-/-
rs2069849	IL6 variant	C	CC	+/*
rs3918226	NOS3 variant	T	CC	-/-
rs2853792	NOS3 variant	G	AA	-/-
rs3918227	NOS3 variant	C	CC	+/*
rs2853796	NOS3 variant	G	TT	-/-
rs743507	NOS3 variant	C	TT	-/-
rs1360485	HMGB1 variant	T	TT	+/*
rs1412125	HMGB1 variant	T	TT	+/*
rs4145277	HMGB1 variant	T	TT	+/*
rs4932178	FURIN variant	C	CC	+/*
rs17514846	FURIN variant	C	AC	+/-
rs11538758	PRNP V210I	A	CC	-/-
rs1799990	PRNP M129V	G	GG	+/*
rs16990018	PRNP N171S	G	AA	-/-
rs28933385	PRNP E200K	A	GG	-/-
rs2070788	TMPRSS2 variant	A	AG	+/-
rs12329760	TMPRSS2 variant	T	TC	+/-
rs2048683	ACE2 variant	G	GG	+/*

Other Immune Factors

27 variants found 7 4 16

Risk Summary: 7 high risk variants, 4 moderate risk variants - Consider discussing with healthcare provider

Quick Navigation: [Overview](#) [Categories](#) [Disclaimer](#)

SNP ID	SNP Name	Risk Allele	Your Allele	Result
rs10210302	ATG16L1 C234158839T	C	TC	+/-
rs6822844	4q27 Region 123509421G>T	G	GG	+/ ⁺
rs6822844	4q27 Region 123509421G>T	G	GG	+/ ⁺
rs479405	DMGDH G67591T	C	AA	-/-
rs479405	DMGDH G67591T	C	AA	-/-
rs479405	DMGDH G67591T	C	AA	-/-
rs479405	DMGDH G67591T	C	AA	-/-
rs2253262	DMGDH T372G	A	CC	-/-
rs2253262	DMGDH T372G	A	CC	-/-
rs2253262	DMGDH T372G	A	CC	-/-
rs2253262	DMGDH T372G	A	CC	-/-
rs2069812	IL5 A131879916G	G	AG	+/-
rs20541	IL-13 variant	A	AG	+/-
rs1800629	TNF -308 variant	A	GG	-/-
rs361525	TNF -238 variant	A	GG	-/-
rs660895	HLA-DRB1 variant	G	AA	-/-
rs660895	HLA-DRB1 variant	G	AA	-/-
rs28940879	TYR (MeFV) V726A variant	A	GG	-/-
rs28940578	MeFV M694I variant	T	CC	-/-
rs11466023	MeFV P369S variant	A	AG	+/-
rs1801275	IL4R Q576R variant	G	AA	-/-
rs7216389	GSDB variant	T	CC	-/-
rs17851582	GAMT C9110T	G	GG	+/ ⁺
rs17851582	GAMT C9110T	G	GG	+/ ⁺
rs17851582	GAMT C9110T	G	GG	+/ ⁺
rs17851582	GAMT C9110T	G	GG	+/ ⁺
rs17851582	GAMT C9110T	G	GG	+/ ⁺

⚠ Important Disclaimer

Medical Disclaimer: This report is intended to translate your results into an easier to understand form. It is **not intended to diagnose or treat** any medical condition.

Professional Consultation: For diagnosis or treatment, please present this report to your qualified healthcare provider or find a practitioner on the MTHFRSupport™ website under "Find a Practitioner".

Genetic Interpretation: Genetic mutations are indicators that something *could* be affected, but they are **not a guarantee** that you are experiencing all or any of the associated issues.

Contributing Factors: Many factors influence whether genetic variants manifest as health issues, including:

- Environmental factors
- Ethnic background
- Diet and lifestyle
- Age and personal history
- Other genetic interactions

© Copyright all rights reserved MTHFR Support™