

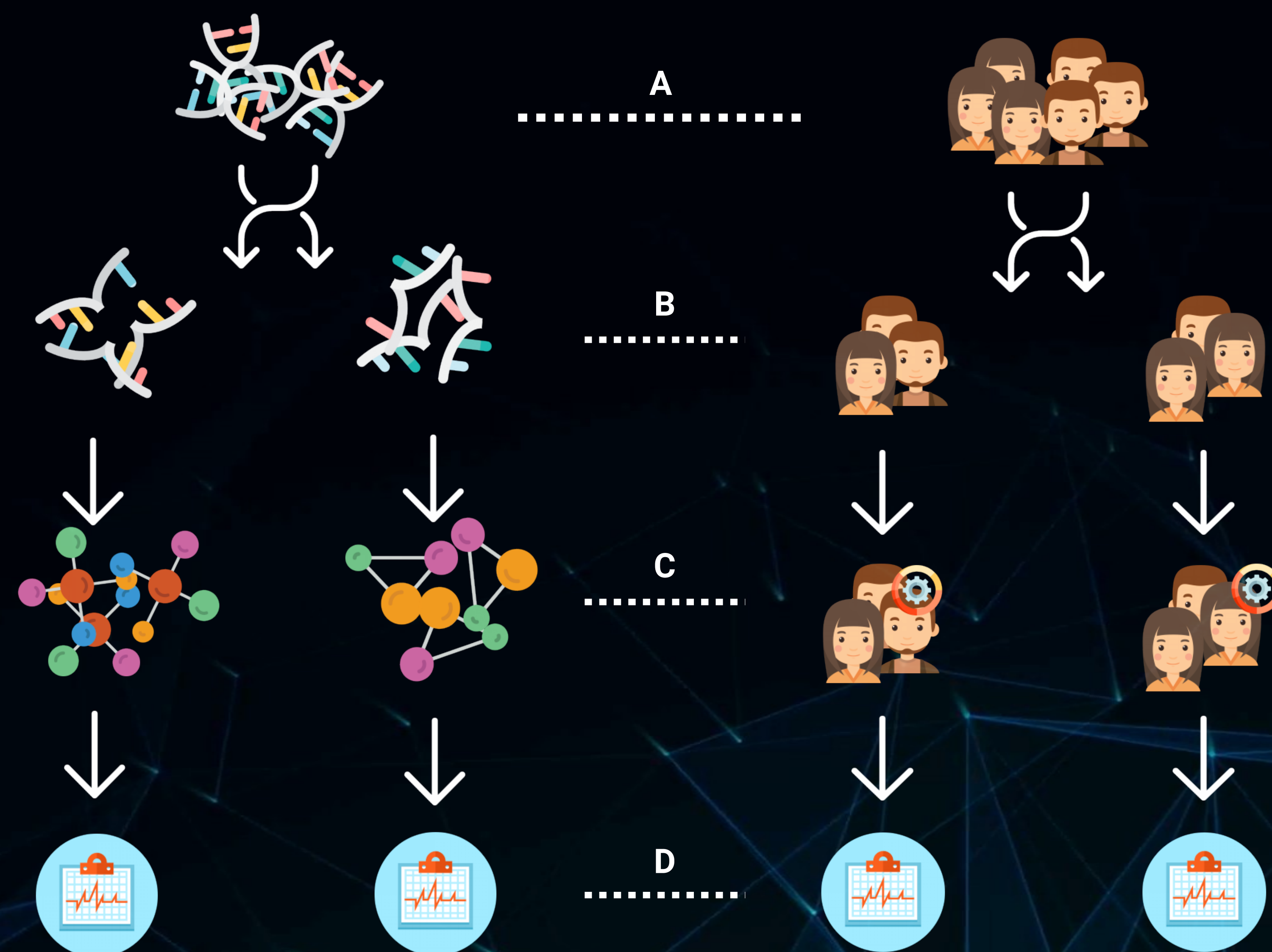
Celiac Disease Triggers

Using Mendelian Randomization to infer causality

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INTRODUCTION

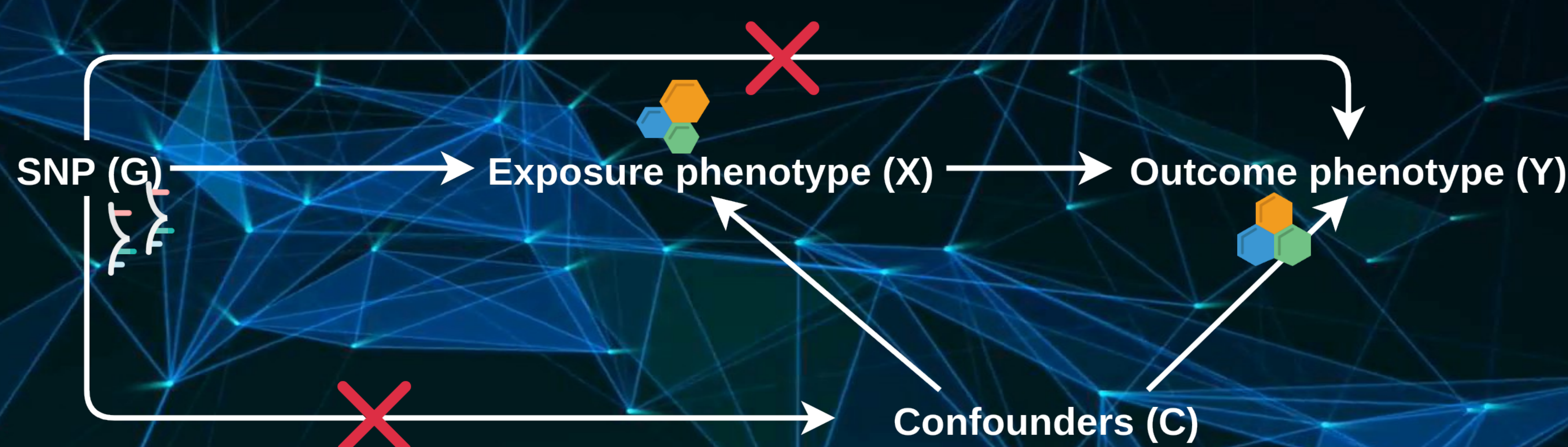
Celiac disorder affects up to two percent of the European population, and it is hereditary. People with a first-degree relative with celiac disease have 30 times higher risk of developing celiac disease. The inheritance model of celiac disease is oligogenic.^[1] In this model, HLA-DQ2 is a genetic variant in the major histocompatibility complex (MHC) region that accounts for 40% of the inheritable risk. But, this genetic variant alone is not sufficient enough to develop celiac disease. 30% of the Europeans are carriers of the variant, however 3% express celiac disease. We expect to find multiple causal or protective clinical parameters or molecular mechanisms that will be quantified to understand the impact of the causal or protective relationship, that can potentially explain what is triggering celiac disease in the 3% from the 30% of the Europeans that are carriers of HLA-DQ2.^[2]

MATERIALS AND METHODS

We investigated over 500.000 publicly available clinical parameters and molecular phenotypes, available from databases, and published genome-wide association studies (GWAS), with the Mendelian Randomization^[3] (MR) approach. We used two-samples MR methods: Inverse-Variance Weighted^[4] (IVW) method and MR-Egger method^[5] to infer causality links by combining summary statistics from GWAS.

RESULTS

We identified 1133 significant ($FDR < 0.05$) clinical parameters or molecular mechanisms that cause or protect for celiac disease. For example Type 1 Diabetes (T1D), which describes the co-occurrence of T1D and celiac disease.^[6] A shared association between celiac disease and crohn's disease.^[7] The effect of eosinophils is a risk factor for celiac disease, and that lymphocytes is associated as a protective factor for celiac disease.^[8] These findings are consistent with literature.



OTHER (NON-AUTOIMMUNE) DISEASE OF DIGESTION SYSTEM

HYPERTENSION

THROMBOCYTOSIS

THYROID ABNORMALITIES

PBMC NATURAL KILLER CELLS, CD4+

REDUCED BODY FAT/MASS

WHITE BLOODCELLS EOSINOPHILS, LYMPHOCYTES

NEUROLOGICAL DISORDERS

INFECTIONS, ALLERGIES

OTHER AUTO-IMMUNE DISEASES

ABNORMAL ERYTHROPOIESIS ANEMIA

BONE-RELATED DISORDERS

SKIN DISORDERS

CELIAC DISEASE

CONCLUSION & DISCUSSION

Mendelian Randomization approach gives the ability to predict causality links between a phenotype of interest and a medically relevant outcome by statistical methods that combine summary statistics from GWAS. Our results indicate that what is confirmed by other studies that celiac disease is a complex immune-mediated disease that needs both HLA-DQA1 and HLA-DQB1 alleles but are not sufficient to develop the disease.^[9] Here we present potential causal risk factors that may be involved in the development of celiac disease.

REFERENCES. ¹Robert Di Niro, et al. High abundance of plasma cells secreting transglutaminase 2-specific IgA autoantibodies with limited somatic hypermutation in celiac disease intestinal lesions; ²Gujral, N., Freeman, H. J & Thomson, A. B. Celiac disease: Prevalence, diagnosis, pathogenesis and treatment; ³David M. Evans and George Davey Smith. Mendelian Randomization: New Applications in the Coming Age of Hypothesis-Free Causality; ⁴Debbie A. Lawlor. Commentary: Two-sample Mendelian randomization: opportunities and challenges. International Journal of Epidemiology; ⁵Stephan Burgess, Simon G. Thompson. Interpreting findings from Mendelian randomization using the MR-Egger method; ⁶William Hagopian, Hye-Seung Lee, Edwin Liu, et al. Co-occurrence of Type 1 Diabetes and Celiac Disease Autoimmunity; ⁷Festen EAM, Goyette P, Green T, Boucher G, Beauchamp C, et al. A Meta-Analysis of Genome-Wide Association Scans Identifies IL18RAP, PTPN2, TAGAP, and PUS10 As Shared Risk Loci for Crohn's Disease and Celiac Disease. ⁸William J. Astle, Heather Elding, Tao Jiang, et al. Landscape of Human Blood Cell Trait Variation and Links to Common Complex Disease; ⁹Gosia Trynka, Karen A Hunt, Nicholas A Bockett. Dense genotyping identifies and localizes multiple common and rare variant association signals in celiac disease.