PGP-UK Genomics Report for uk9A677F

1 Summary

This is the genome report was produced using collaborative research tools, including SNPedia and GetEvidence. This section shows an overview of all the small variants which were found in the genome for this individual, when compared with a reference genome. These variants are summarised in Table 1 and the pie-charts in Figures 2, 3 and 4.

This report was generated automatically and is not clinically approved. It is provided for <u>personal and research purposes</u> only.

This document contains hyperlinks, shown in grey, that will take you to external websites where you can find more detailed explanations. Some of the technical terms are also explained in more detail in the Ensembl Glossary. We would welcome your feedback about this report, for example, if you would like more information about anything or if any of the links have become inactive. You can contact us on: pgp-uk@ucl.ac.uk.

This summary shows an overview of all the variants which were found in the genome for this individual. The "variants remaining after filtering" refers to any differences in the DNA identified when compared to the reference genome. Of these, the majority will have already been found in some other sequenced individual and put on a database (existing variants) while others have not yet been annotated (novel variants).

"Overlapped genes" refers to the number of times where a variant was found in a region of the genome containing a gene. The diagram in Figure 1 is a simplification of the usual gene structure. "Exon" refers to the part of the gene which goes on to form a protein, and variants in this part of the gene are more likely to cause changes in the shape of the protein. Upstream, downstream, intronic and intergenic variants are more likely to alter the regulation of that gene but will not change the protein itself.

A transcript for a protein-coding gene can include the exons, introns and other gene features that are transcribed and important for gene function but might not be translated into the final protein. Not all transcripts are for protein-coding genes, with many containing non-coding RNAs that can be overlapping other genes, in introns or in intergenic regions.

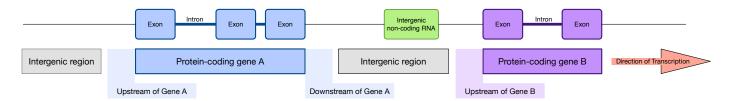


Figure 1: Diagram of gene structure indicating locations of potential variants

Feature	Count
Lines of input read	4907085
Variants filtered out	0
Novel / existing variants	$467525\ (9.5)\ /\ 4428203\ (90.5)$
Overlapped genes	56645
Overlapped transcripts	67435
Overlapped regulatory features	166463

Table 1: Variant calling summary

There are several different types of genomic variants. The most common change is when one single building block of the DNA (called a nucleotide) is changed, called a single nucleotide variants (SNV). Other variant types include insertions, where the DNA in the individual is longer than the reference sequence due to the insertion of one or more nucleotides; and deletions, where a few nucleotides are missing compared to the reference sequence.

Some of these changes will have no effect on the protein, while some changes may alter the protein function to varying degrees. The PolyPhen analysis software attempts to quantify the effect each mutation will have on the protein function. This ranges from "benign" where no change to the protein function is expected, to "probably damaging" where it is predicted that the mutation will affect protein function. It is nevertheless important to note that what is "damaging" for the protein is not necessarily damaging for the individual.

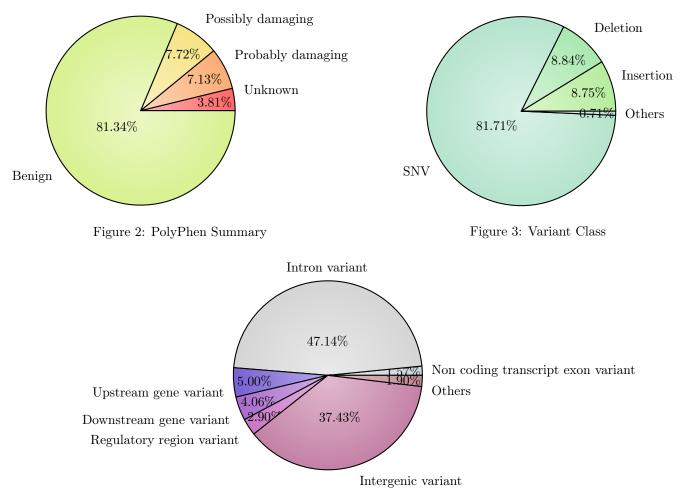


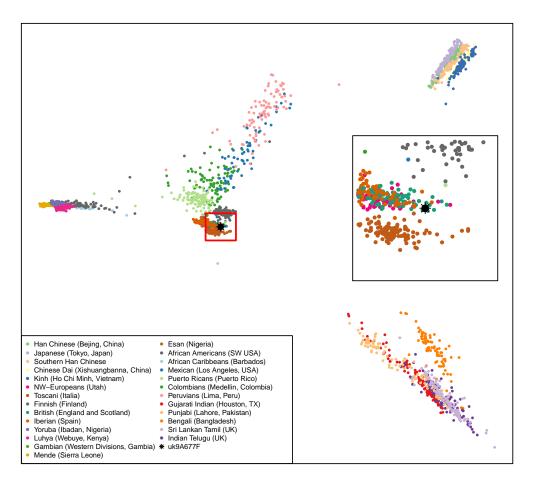
Figure 4: Consequence type

2 Ancestry

This plot shows the distribution of the genomes of different populations. Data from several studies which used whole genome sequencing was used to see the relationships between the genomes of the populations. It shows how closely related certain populations are genetically: Groups which cluster closely are more genetically similar than groups which are further apart. The black star symbol shows where this PGP-UK participant sits in relation to other populations, indicating their ancestry and their most closely related populations according to genetic sequence.

Based on the populations defined in the 1000 genomes project (1kGP), the ancestry composition for this individual is inferred to be 100.0 percent European [British in England and Scotland].

Please note that this analysis is limited by the populations available in the 1kGP data. If there are European subpopulations reported, and the ancestry of the participant does not correspond to any of the 1kGP populations, the closest 1kGP sampled subpopulation will be shown (even though it might be different from the participant's actual ancestry).



Ancestry uk9A677F

Figure 5: Ancestry Principal Component Analysis

3 Traits (based on SNPedia information)

Existing research has associated many variants with phenotypic traits, some of which can be perceived as beneficial while others appear to have a harmful effect. Some traits are complex and can be affected by several variants. It is likely that some of these would confer a higher risk while others a lower risk of trait manifestation. These can not be combined linearly to produce an actual risk of disease.

It is important to note that in most cases genomic data is probabilistic, not deterministic- i.e. having a genetic predisposition for a disease is not a diagnosis; rather, it shows an increased likelihood of developing that disease. Also, one person can have both potentially beneficial and harmful variants in the same gene, or associated with the same disease.

Some variants can also affect certain populations more, or will only affect a particular gender. For example, a variant for higher risk of endometriosis in the sequence of a male will not directly affect that person, but can be passed on to descendants.

While many traits are the result of a unique variant, many are the combination of several variants throughout the genome. In SNPedia, these are called genosets. These can integrate some of the information already present in the single variant tables, or be the combination of variants that have no phenotypic effect on their own, but contribute to a trait when together.

The variants in the following tables are sorted by magnitude. This is an subjective measure defined in SNPedia to highlight the perceived importance of the genotype described. At the moment this scale goes from 0 to 10. You can read more about it by visiting their explanatory webpage.

As our knowledge grows, the interpretation of the effect of certain variants might change. Clicking on the links in the genome report tables will take you to websites containing more information about each variant.

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
2.1	rs2511989	(A;G)	0.63x decreased age-related macular degeneratio		Link	0 / 0.2
2	rs1012053	(A;C)	0.625x reduced risk of Bipolar Disorder.		Link	
2	rs10468017	(C;T)	Associated with higher HDL cholesterol		Link	
2	rs1136410	(C;T)	0.80x reduced risk for glioblastoma	Link	Link	
2	rs1864163	(G;G)	Associated with higher HDL cholesterol		Link	
2	rs2060793	(A;A)	Lower serum levels of vitamin D			
2	rs2073963	(T;T)	Reduced risk of baldness			
2	rs2235015	(G;T)	Somewhat more likely to respond to certain anti	Link	Link	
2	rs2241423	(A;G)	0.79 decreased risk for obesity			
2	rs2542052	(C;C)	Better odds of living to 100			
2	rs2707466	(A;A)	Stronger bones	Link	Link	
2	rs2908004	(T;T)	Stronger bones	Link	Link	
2	rs3736309	(A;G)	0.44x decreased risk for chronic obstructive pu			
2	rs3750817	(C;T)	0.78x reduced risk for breast cancer			
2	rs3764261	(G;T)	Associated with higher HDL cholesterol		Link	Link
2	rs3782179	(C;T)	3x lower odds of testicular cancer risk for men			
2	rs3819331	(T;T)	Lower risk of autism	Link		
2	rs4149268	(A;G)	Associated with higher HDL cholesterol		Link	
2	rs6505162	(A;C)	0.58x decreased risk for esophageal cancer	Link		
2	rs6511720	(G;T)	Slightly lower odds of developing CHD.		Link	Link
2	rs6807362	(G;G)	Decreased autism risk	Link	Link	
2	rs6855911	(A;G)	0.62x decreased risk for gout		Link	
2	rs763110	(T;T)	$\sim 0.80 \text{x}$ reduced cancer risk			Link
2	rs801114	(T;T)	0.78x decreased Basal Cell Carcinoma risk.		Link	
1.8	rs187238	(C;G)	Hypertension not a risk factor for sudden cardi			
1.8	rs3814113	(C;C)	0.8x decreased risk for ovarian cancer		Link	
1.8	rs4714156	(C;C)	<0.61x risk for restless legs			

3.1 Possibly Beneficial Traits

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
1.5	rs1026732	(A;G)	0.70x risk for restless legs		Link	
1.5	rs1063192	(C;C)	0.71x reduced risk of myocardial infarction			
1.5	rs11136000	(C;T)	0.84x decreased risk for Alzheimer's disease		Link	
1.5	rs11212617	(A;C)	Somewhat increased likelihood of treatment succ			Link
1.5	rs11635424	(A;G)	0.70x risk for restless legs		Link	
1.5	rs12593813	(A;G)	0.71x risk for restless legs		Link	
1.5	rs309375	(G;G)	Smaller mosquito bites			
1.5	rs3784709	(C;T)	0.71x risk of developing restless legs syndrome		Link	
1.5	rs3790844	(C;T)	Slightly reduced risk (0.77x) for pancreatic ca			
1.5	rs3851179	(A;G)	0.85x decreased risk for Alzheimer's disease		Link	
1.5	rs4149274	(C;T)	Associated with higher HDL (good) cholesterol			
1.5	rs4489954	(G;T)	0.69x risk risk of developing restless legs syn		Link	
1.5	rs4939883	(C;C)	Associated with higher HDL cholesterol		Link	
1.5	rs5968255	(C;C)	Slower AIDS progression (8 years)			
1.5	rs9939609	(T;T)	Lower risk of obesity and Type-2 diabetes		Link	
1.4	rs1165205	(A;T)	0.85x decreased gout risk		Link	
1.4	rs6700125	(C;C)	0.7x decreased risk for ALS			
1.25	rs10088218	(A;G)	0.76x decreased risk for ovarian cancer			
1.2	rs4320932	(A;G)	0.87x decreased risk for ovarian cancer			
1.2	rs9306160	(C;T)	0.75x (reduced) risk for metastasis in LN-/ER+	Link	Link	
1.1	rs2293347	(G;G)	Among NSCLC patients: better Gefitinib response	Link		Link
1.1	rs4988235	(T;T)	Can digest milk			Link
1.1	rs7568369	(T;T)	0.90x reduced risk of obesity			
1	rs10248420	(A;G)	7x more likely to respond to certain antidepres		Link	
1	rs11983225	(C;T)	7x more likely to respond to certain antidepres		Link	
1	rs182549	(T;T)	Can digest milk.			Link
1	rs2235040	(A;G)	7x more likely to respond to certain antidepres	Link	Link	
1	rs2235067	(A;G)	7x more likely to respond to certain antidepres			
1	rs2952768	(C;T)	Slightly less drug dependence: decreased effect			Link
1	rs4148739	(A;G)	7x more likely to respond to certain antidepres		Link	
1	rs7850258	(A;G)	Typical odds of developing primary hypothyroidi			
1.0	rs11246226	(C;C)	Decreased risk of schizophrenia in limited stud		Link	
0.1	rs891512	(G;G)	Lower blood pressure than those with an A allel	Link		
0	rs1047781	(A;A)	ABH blood group "Secretor" status if Japanese	Link	Link	Link
0	rs1126809	(A;G)	Slight increase in skin cancer risk	Link	Link	Link
0	rs12252	(T;T)	More resistant to influenza	Link		Link
0	rs16990018	(A;A)	PrP Codon 171 Asn - Non-pathogenic variant	Link		Link
0	rs17244841	(A;A)	More responsive to statin treatment		Link	Link
0	rs1799782	(C;C)	Lower risk for skin cancer	Link	Link	
0	rs1800562	(G;G)	Not a C282Y hemochromatosis carrier.	Link	Link	Link
0	rs242941	(G;G)	Better response to inhaled corticosteroid in pa		Link	
0	rs28933385	(G;G)	Prion protein Codon 200 (E) - Non pathogenic va			Link
0	rs5065	(A;A)	1.12x risk on diuretic; if hypertensive: better	Link	Link	Link
0	rs74315403	(G;G)	$\Pr PrP$ codon 178 (D) - non pathogenic variant			Link
0	rs7495174	(A;A)	Blue/gray eyes more likely		Link	
0	rs9394492	(C;C)	<0.76x risk for restless legs			
0	rs9951307	(A;G)	0.10 decreased risk for brain edema after a str			

3.2 Possibly Harmful Traits

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
3.5	rs10490924	(T;T)	8.2x risk for age related macular degeneration	Link	Link	Link
3	rs10897346	(C;C)	If depressed: 2.6x more likely to not respond t			
3	rs1800460	(A;G)	(TPMT*3B) impaired drug metabolism	Link	Link	Link
3	rs2306402	(C;C)	1.18x increased risk for late-onset Alzheimer's			
3	rs2981582	(C;T)	1.3x higher risk of ER+ breast cancer		Link	
3	rs6920220	(A;G)	1.2x risk Rheumatoid Arthritis		Link	
3.0	rs1142345	(A;G)	TPMT*3C . impaired drug metabolism	Link	Link	Link
2.7	rs10830963	(C;G)	Increased type-2 diabetes risk; higher gestatio		Link	
2.6	rs110419	(A;A)	2.6x increased risk for neuroblastoma			
2.5	rs11190870	(T;T)	Possibly even more increased risk of scoliosis			
2.5	rs12803066	(A;G)	Increased risk of myopia			
2.5	rs13266634	(C;T)	Increased risk for type-2 diabetes	Link	Link	Link
2.5	rs3780374	(A;G)	Substantially increased odds of developing V617			
2.5	rs5888	(C;T)	3x higher risk for age-related macular degenera	Link		
2.5	rs613872	(G;T)	~5 fold higher risk for Fuchs' dystrophy: a cor			
2.5	rs660895	(G;G)	6x higher risk of rheumatoid arthritis		Link	
2.5	rs664143	(C;T)	Higher risk for number of cancers			
2.4	rs7966230	(G;G)	Slightly lower levels of plasma VWF			
2.2	rs2004640	(G;T)	1.4x increased risk for SLE		Link	Link
2.2	rs2231137	(G;G)	~1.5-3x increased risk for ischemic stroke	Link	Link	Link
2.2	rs944289	(T;T)	1.69x increased thyroid cancer risk		Link	
2.1	rs10811661	(T;T)	1.2x increased risk for type-2 diabetes		Link	
2.1	rs1329428	(G;G)	2x increased risk for macular degeneration			
2.1	rs1360780	(C,C) (T;T)	1.3x increased risk for depression		Link	
2.1	rs1695	(G;G)	3.5x asthma risk in certain populations	Link	Link	Link
2.1	rs17070145	(C;C)	Reduced memory abilities	LIIIK	Link	Link
2.1	rs2231142	(A;C)	1.74x increased gout risk; gefinitib takers 4x	Link	Link	Link
2.1	rs2254958	(C;T)	1.24x increased risk for Alzheimer's	Dilli	Dim	Dillin
2.1	rs2494732	(C;C)	Greater odds of cannabis-associated psychosis	Link	Link	
2.1	rs380390	(C;C)	Increased risk for ARMD	LIIIK	Link	
2.1	rs4402960	(C,C) $(\mathrm{T};\mathrm{T})$	1.2x increased risk for type-2 diabetes: 1.5x r		Link	Link
2.1	rs4693596	(1,1) (C;C)	2x odds of myopathy if taking statins		LIIIK	LIIIK
2.1	rs5186	(C,C) (A;C)	~1.4x increased risk of hypertension	Link	Link	Link
2.1	rs5751876	(T;T)	Significantly higher anxiety levels after moder	Link	LIIIK	LIIIK
2.1	rs6457617	(T,T) $(T;T)$	5.2x risk of rheumatoid arthritis	LIIIK	Link	
2.1	rs646776	$(\mathbf{A};\mathbf{A})$	1.2x risk of coronary artery disease		Link	
2.1	rs1024611	$(\mathbf{C};\mathbf{T})$	Increased risk of exercise induced ischemia		LIIIK	Link
2	rs1045642	(C,T) (C;T)	Slower metaboliser for some drugs	Link	Link	Link
2			Ű	LIIIK	LIIIK	LIIIK
2	rs10871777 rs10984447	(A;G)	Adults likely to be 0.22 BMI units higher >1.17x increased risk for multiple sclerosis		Link	
2		(A;A)	*		Link	
	rs11045585	(A;G)	63% chance (higher than average) of docetaxel-i			
2	rs11171739	(C;C)	1.75x risk of developing Type-1 diabetes		Link	
2	rs1160312	(A;G)	1.6x increased risk of Male Pattern Baldness.		Link	
2	rs1219648	(A;G)	1.20x risk for breast cancer		Link	
2	rs12696304	(C;G)	Prone to aging faster: at least in European pop			
2	rs1333048	(A;C)	1.3x increased coronary artery disease risk		T ' 1	
2	rs1537415	(G;G)	2x increased risk for periodontitis		Link	
2	rs1544410	(A;A)	Increased risk of low bone mineral density diso		Link	
2	rs1585215	(A;G)	2x increased risk for Hodgkin lymphoma			
2	rs16942	(A;G)	Very slightly increased breast cancer risk	Link	Link	Link
2	rs16944	(G;G)	Increased risk of mental disorders		Link	
2	rs17228212	(C;T)	1.26x increased risk for heart disease		Link	
2	rs1734791	(A;A)	1.4x increased risk for lupus			

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
2	rs17576	(A;G)	Higher risk for MI and lung cancer: and COPD in	Link	Link	
2	rs17696736	(A;G)	1.34x risk of type-1 diabetes		Link	
2	rs17782313	(C;T)	Adults likely to be 0.22 BMI units higher		Link	Link
2	rs2056116	(G;G)	1.41x risk for breast cancer			
2	rs2201841	(T;T)	2.4x increased risk for Graves' disease		Link	
2	rs2383206	(A;G)	1.4x increased risk for heart disease			
2	rs2383207	(A;G)	Increased risk for heart disease			
2	rs2420946	(C;T)	1.20x risk for breast cancer			
2	rs25487	(G;G)	2x higher risk for skin cancer; possibly other	Link	Link	Link
2	rs2736100	(T;T)	Higher risk of Interstitial lung disease: and t		Link	
2	rs326	(A;A)	Lower HDL cholesterol		Link	Link
2	rs358806	(C;C)	1.78x increased risk of developing Type-2 diabe		Link	
2	rs3738579	(C;C)	0.6x decreased risk for cervical cancer: but 1			
2	rs3746444	(C;T)	$\sim 1.2x$ increased risk for cancer	Link		
2	rs3793784	(C;G)	1.5x risk for ARMD		Link	Link
2	rs4420638	(A;G)	~3x increased Alzheimer's risk; 1.4x increased		Link	Link
2	rs4464148	(C;C)	1.35x increased risk for colorectal cancer			
2	rs4633	(T;T)	Higher risk for endometrial cancer	Link	Link	Link
2	rs4792311	(A;G)	Increased risk of prostate cancer	Link	Link	Link
2	rs4825476	(G;G)	1.9x higher risk of suicidal thoughts when taki		Link	
2	rs486907	(A;A)	2x increased prostate cancer risk	Link	Link	Link
2	rs493258	(A;G)	1.15x risk of Age Related Macular Degeneration			
2	rs4961	(G;T)	1.8x increased risk for high blood pressure	Link	Link	Link
2	rs5174	(A;G)	1.3x increased risk for heart disease	Link	Link	Link
2	rs520354	(A;G)	Increased risk in men for biliary conditions			
2	rs629242	(C;T)	Somewhat higher risk for prostate cancer			
2	rs6441286	(G;T)	1.54x chance of developing primary biliary cirr		Link	
2	rs6498169	(A;A)	>1.14x risk of multiple sclerosis		Link	
2	rs669	(G;G)	3.8x or higher increased risk for Alzheimers	Link	Link	Link
2	rs6897932	(C;C)	1.08x increased risk for multiple sclerosis	Link	Link	Link
2	rs6997709	(G;G)	1.5x higher risk for hypertension			
2	rs699	(C;T)	Increased risk of hypertension	Link	Link	Link
2	rs7216389	(T;T)	1.5x increased risk for Childhood Asthma.		Link	
2	rs7442295	(A;A)	~4x higher risk for hyperuracemia		Link	
2	rs744373	(C;T)	1.17x risk of Alzheimer's			
2	rs7639618	(C;T)	1.45x increased osteoarthritis risk	Link		
2	rs7776725	(C;C)	Weaker bones		Link	
2	rs7807268	(C;G)	1.3x risk for Crohn's disease		Link	
2	rs7961152	(A;C)	1.2x higher risk for hypertension			
2	rs800292	(C;C)	5% higher risk of Age related macular degenerat	Link	Link	Link
2	rs828907	(G;T)	Slightly increased risk of bladder cancer and 2			
2	rs854560	(A;T)	Higher risk for heart disease: diabetic retinop	Link	Link	Link
2	rs9652490	(A;A)	$\sim 2x$ increased risk for Parkinson's disease: and		Link	
2	rs965513	(A;G)	1.77x increased thyroid cancer risk		Link	
2	rs9954153	(G;G)	$\sim 5x$ higher risk for Fuchs' dystrophy: a corneal			
2.0	rs1434536	(A;A)	1.94x increased breast cancer risk			
2.0	rs2156921	(G;G)	1.29x increased risk for depression			
2.0	rs2305795	(A;A)	1.64x higher risk of narcolepsy compared to (G;			Link
2.0	rs4911414	(G;T)	2-4x higher risk of sun sensitivity if part of		Link	
1.9	rs7923837	(A;G)	1.6x risk for T2D			
1.8	rs1136287	(C;T)	$1.5 \mathrm{x}$ increased risk of wet ARMD in a Taiwanese	Link	Link	
1.8	rs2278206	(T;T)	1.16x increased risk for asthma	Link	Link	
1.8	rs4474514	(A;G)	3x increased testicular cancer risk for men		Link	
1.7	rs8055236	(G;T)	1.9x risk for heart disease		Link	
1.6	rs3764880	(A;A)	1.2 - 1.8x increased tuberculosis risk	Link	Link	

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
1.6	rs3775948	(C;G)	Slightly higher risk for gout			
1.5	rs10464059	(A;G)	Slightly increased risk of developing Parkinson			
1.5	rs10757272	(C;T)	1.30x increased risk for Coronary artery diseas			
1.5	rs10859871	(A;C)	Slight ($\sim 1.2x$) increase in endometriosis risk			
1.5	rs10883365	(A;G)	1.2x increased risk for developing Crohn's dise		Link	
1.5	rs10895068	(A;G)	2.5x increased odds of breast cancer among horm			
1.5	rs1169300	(A;G)	$\sim 1.5 x$ increased lung cancer risk			
1.5	rs12037606	(A;G)	1.22x risk of developing Crohn's disease			
1.5	rs1223271	(A;G)	Slightly increased risk of developing Parkinson		Link	
1.5	rs1375144	(C;C)	1.59x increased risk of developing bipolar diso			
1.5	rs140701	(A;G)	Increased risk for anxiety disorders			
1.5	rs144848	(G;T)	Very slightly increased breast cancer risk	Link	Link	Link
1.5	rs1571801	(A;A)	>1.36x risk for prostate cancer			
1.5	rs1801274	(C;T)	Complex; generally greater risk for cancer prog	Link	Link	Link
1.5	rs1867277	(A;G)	1.5x increased risk for thyroid cancer			
1.5	rs1975197	(C;T)	1.3x increased risk of developing restless legs		Link	
1.5	rs2076295	(G;G)	Slightly increased risk for pulmonary fibrosis			
1.5	rs2177369	(C;C)	1.5x increased risk for Alzheimer's disease			
1.5	rs2272127	(C;C)	Associated with herpes and schizophrenia			
1.5	rs2280714	(A;G)	1.4x increased risk of SLE			
1.5	rs2464196	(C;T)	$\sim 1.5 x$ increased lung cancer risk	Link	Link	Link
1.5	rs2697962	(A;G)	Slightly increased risk of developing Parkinson			
1.5	rs2736990	(C;T)	Slightly increased risk of developing Parkinson		Link	
1.5	rs27388	(A;G)	Slightly increased risk of developing schizophr			
1.5	rs2881766	(T;T)	Slightly increased risk for pregnancy-induced h			
1.5	rs3087243	(G;G)	Increased risk for autoimmune diseases		Link	
1.5	rs3212227	(A;A)	1.43x increased risk of developing psoriasis an			
1.5	rs3745516	(A;G)	Slightly increased risk of developing primary b			
1.5	rs393152	(A;A)	Increased risk of both PD and AD	Link	Link	
1.5	rs401681	(C;T)	\sim 1.2x increased risk for several types of cance		Link	
1.5	rs4027132	(A;G)	1.39x increased risk of developing bipolar diso			
1.5	rs4506565	(A;T)	1.4x increased risk for type-2 diabetes		Link	
1.5	rs464049	(C;T)	Increased risk of schizophrenia in limited stud			
1.5	rs4656461	(A;G)	1.5x increased risk for open angle glaucoma			
1.5	rs4982731	(C;C)	Possible higher risk of childhood acute lymphob			
1.5	rs5219	(C;T)	1.3x increased risk for type-2 diabetes	Link	Link	Link
1.5	rs5746059	(A;A)	Slightly higher fat mass			
1.5	rs6896702	(C;T)	Slightly increased risk of developing Parkinson			
1.5	rs6908425	(C;T)	1.63x increased risk of developing Crohn's dise		Link	
1.5	rs7341475	(G;G)	1.58x increased schizophrenia risk for women		Link	
1.5	rs7454108	(C;T)	Single HLA-DQ8 haplotype			
1.5	rs7774434	(C;T)	Slightly increased risk of developing primary b			
1.5	rs9561778	(G;T)	$\sim 2x$ increased risk of adverse drug reactions fr		Link	
1.5	rs9642880	(G;T)	1.2x increased bladder cancer risk		Link	
1.4	rs1126497	(T;T)	1.4x increased risk for breast cancer	Link	Link	Link
1.4	rs12770228	(A;G)	1.4x increased risk for meningioma			
1.4	rs2046210	(C;T)	1.4x increased breast cancer risk		Link	Link
1.4	rs2230201	(A;G)	1.4x risk of lupus	Link		
1.4	rs3131296	(G;G)	1.4x increased risk for schizophrenia		Link	
1.4	rs3184504	(C;T)	Slightly increased risk for celiac disease	Link	Link	
1.4	rs4977756	(G;G)	1.93x higher risk for glioma development		Link	
1.4	rs6010620	(G;G)	1.4x higher risk for glioma development; but th		Link	
1.3	rs1042713	(A;G)	1.3x increased risk that pediatric inhaler use	Link	Link	Link
1.3	rs10947262	(C;C)	1.3x increased risk for osteoarthritis			
1.3	rs1746048	(C;C)	1.03 increased risk for coronary heart disease		Link	

Mag.	Identifier	Genotype	Summary	ExAC	GetEvidence	ClinVar
1.3	rs2059693	(C;T)	1.3x increased risk for testicular cancer			
1.3	rs2295490	(A;G)	1.32x increased risk of early-onset type-2 diab	Link	Link	
1.3	rs356219	(A;G)	1.3x increased risk for Parkinson's disease			
1.25	rs748404	(T;T)	Slightly increased risk (1.25) for lung cancer		Link	
1.2	rs10865331	(A;G)	1.2x higher risk for ankylosing spondylitis			
1.2	rs11037909	(T;T)	1.47x type II diabetes risk	Link		
1.2	rs143383	(C;T)	1.1x increased risk for osteoarthritis		Link	Link
1.2	rs2814707	(A;G)	1.2x increased risk for ALS		Link	
1.2	rs3740878	(A;A)	1.46x type II diabetes risk; common	Link		Link
1.2	rs3849942	(A;G)	1.2x increased risk for ALS		Link	
1.2	rs4324715	(C;C)	>1.5x increased testicular cancer risk for men			
1.2	rs449647	(A;T)	Possibly lower levels of ApoE			
1.2	rs4795067	(A;G)	Slight increase in risk for psoriatic arthritis			
1.17	rs17465637	(A;C)	1.17x higher risk for myocardial infarction	Link	Link	
1.1	rs11110912	(C;G)	1.3x high blood pressure risk			
1.1	rs11650494	(A;G)	Slightly higher prostate cancer risk			
1.1	rs13387042	(A;G)	1.12x increased risk for breast cancer		Link	
1.1	rs1344706	(G;T)	1.1x increased risk for schizophrenia		Link	
1.1	rs2653349	(G;G)	2-6x increased risk for cluster headaches	Link	Link	
1.1	rs2828520	(G;G)	1.35x major depressive disorder risk			
1.1	rs34516635	(G;G)	Less longevity for Ashkenazi Jewish women.	Link		Link
1.1	rs688034	(C;T)	1.1x risk higher risk for coronary artery disea		Link	
1.1	rs7171755	(A;A)	Very slight descrease in cortical thickness and			
1.1	rs7412	(C;C)	More likely to gain weight if taking olanzapine	Link	Link	Link
1.1	rs925391	(C;C)	More likely to go bald; common			
1.05	rs2291834	(C;T)	Very slightly higher risk for myocardial infarc			
1	rs10504861	(G;G)	Major allele: normal risk of migraine			
1	rs10761659	(A;G)	1.2x risk of Crohn's disease		Link	
1	rs1143674	(A;A)	1.3x increased autism risk	Link		
1	rs2546890	(A;G)	Higher risk of multiple sclerosis			
1	rs3194051	(A;G)	1.12x risk of type-1 diabetes	Link	Link	Link
1	rs6166	(G;G)	Females slightly more likely to be sterile	Link	Link	Link
1	rs6932590	(T;T)	1.1x increased risk for schizophrenia		Link	
1	rs987525	(A;C)	2.5x increased risk for cleft lip		Link	
0.1	rs601338	(A;G)	Susceptible to Norovirus infections	Link	Link	Link
0	rs1004819	(C;C)	1.5x risk of Crohn's disease		Link	
0	rs1061646	(C;C)	1.16x increased risk for breast cancer	Link		Link
0	rs1495965	(A;A)	1.2x higher risk for spondylitis			
0	rs3813929	(C;C)	Possible weight gain if taking olanzapine		Link	Link
0	rs4293393	(T;T)	1.25x Increased Risk of CKD for T allele in			
0	rs4712653	(T;T)	2x increased risk for neuroblastoma			
0	rs6314	(C;C)	Higher risk for RA	Link	Link	

3.3 Genosets (Multi-variant Phenotypes)

Magnitude	Identifier	Summary
3.1	gs191	Problem metabolizing NSAIDs
3	gs127	Intermediate warfarin metabolizer
2.9	gs192	MTHFR polymorphisms affecting homocysteine
2.5	gs155	CYP3A5 non-expressor
2.5	gs161	CYP2C9 Intermediate Metabolizers
2.5	gs281	Part of the 88% of the population claimed not t
2	gs101	Probably able to digest milk
2	gs154	NAT2 Slow metabolizer
2	gs188	One copy of APOE4 is possible: but not certain
1.5	gs247	Parkinson's Disease Risk
1	gs182	CYP2D6*39
0	gs158	CYP1A2 normal metabolizer

4 Raw Data

The raw data used to create this report has been assigned the identifier ERS1176558 in the European Nucleotide Archive (ENA) hosted at the European Bioinformatics Institute (EBI).

These data will not be accessible unless the report is approved. This will happen by default one month after the report is issued, or if the report is approved for immediate release within the one month period. Participants can also withdraw from the study at any time in which case the report and the data will not be released and will be deleted.

If the data has already been released, it can be accessed at: http://www.ebi.ac.uk/ena/data/view/ERS1176558

5 Report Metadata

Resource	Version	Website
Genome	GRCh38	Link
BWA	0.7.12	Link
SAMtools	1.3	Link
GATK	3.4-46	Link
PLINK	v1.90b3.35	Link
VEP	88	Link
SNPedia	30-Jul-2017	Link
ExAC	v0.3.1	Link
GetEvidence	16-Dec-2016	Link
ClinVar	16-Dec-2016	Link

Table 5: Analysis Pipeline Versions

Report generated on August 2, 2017.