

Your Reports Summary

This is an overview of your 23andMe reports. It provides brief descriptions of your results but does not provide detailed information that may be important for understanding your results. 23andMe reports do not include all possible variants or account for other factors related to these conditions and traits.

Log into your 23andMe account for more details about each of your results. **If you have concerns about your results, talk to a healthcare professional.**

Your Reports Highlights

Certain results are highlighted because they may be unique or contain important information relevant to your health. The reports in your 23andMe account can provide more details about each of these results.

Genetic Health Risk Reports 1 highlighted of 7 reports
Age-Related Macular Degeneration **Slightly increased risk**

Ancestry Reports 1 highlighted of 5 reports
Ancestry Composition 92.5% Eastern European
10 populations

Carrier Status Reports 0 highlighted of 42 reports

Traits Reports 0 highlighted of 22 reports

Wellness Reports 3 highlighted of 8 reports
Genetic Weight Predisposed to weigh less than average
Lactose Intolerance Likely intolerant
Saturated Fat and Weight Likely similar weight



Genetic Health Risk Reports 1 highlighted report of 7 reports available

Learn whether you have specific genetic variants that can influence your risk for certain health conditions. **Consider talking to a healthcare professional if you have a personal or family history of one of these conditions or have concerns about your results.**

Our reports do not include all possible genetic variants that could affect these conditions. Other factors can also affect your risk of developing these conditions, including lifestyle, environment, and family history.

Age-Related Macular Degeneration

Slightly increased risk

Late-Onset Alzheimer's Disease

Variant not detected

Parkinson's Disease	Variants not detected
Alpha-1 Antitrypsin Deficiency	Variants not detected
Celiac Disease	Variants not detected
Hereditary Hemochromatosis (HFE-Related)	Variants not detected
Hereditary Thrombophilia	Variants not detected



Carrier Status Reports 0 highlighted reports of 42 reports available

Learn whether you have specific genetic variants that may not affect your health, but could affect your children's health. **Consider talking to a healthcare professional before making any major lifestyle changes or if you have any concerns about your results.**

If you see "Variant not detected" for a Carrier Status report, it means you're not a carrier of the tested variant (s). Keep in mind that while our Carrier Status reports cover many variants, they don't include all possible variants associated with each condition. **So it's still possible to be a carrier of a variant not included in our test.**

ARSACS	Variant not detected
Agenesis of the Corpus Callosum with Peripheral Neuropathy	Variant not detected
Autosomal Recessive Polycystic Kidney Disease	Variant not detected
Beta Thalassemia and Related Hemoglobinopathies	Variant not detected
Bloom Syndrome	Variant not detected
Canavan Disease	Variant not detected
Congenital Disorder of Glycosylation Type 1a (PMM2-CDG)	Variant not detected
Cystic Fibrosis	Variant not detected
D-Bifunctional Protein Deficiency	Variant not detected
Dihydrolipoamide Dehydrogenase Deficiency	Variant not detected
Familial Dysautonomia	Variant not detected
Fanconi Anemia Group C	Variant not detected
GRACILE Syndrome	Variant not detected
Gaucher Disease Type 1	Variant not detected
Glycogen Storage Disease Type Ia	Variant not detected
Glycogen Storage Disease Type Ib	Variant not detected
Hereditary Fructose Intolerance	Variant not detected
Herlitz Junctional Epidermolysis Bullosa (LAMB3-Related)	Variant not detected
Leigh Syndrome, French Canadian Type	Variant not detected
Limb-Girdle Muscular Dystrophy Type 2D	Variant not detected
Limb-Girdle Muscular Dystrophy Type 2E	Variant not detected
Limb-Girdle Muscular Dystrophy Type 2I	Variant not detected
MCAD Deficiency	Variant not detected
Maple Syrup Urine Disease Type 1B	Variant not detected
Mucopolidosis Type IV	Variant not detected

Neuronal Ceroid Lipofuscinosis (CLN5-Related)	Variant not detected
Neuronal Ceroid Lipofuscinosis (PPT1-Related)	Variant not detected
Niemann-Pick Disease Type A	Variant not detected
Nijmegen Breakage Syndrome	Variant not detected
Nonsyndromic Hearing Loss and Deafness, DFNB1 (GJB2-Related)	Variant not detected
Pendred Syndrome and DFNB4 Hearing Loss	Variant not detected
Phenylketonuria and Related Disorders	Variant not detected
Primary Hyperoxaluria Type 2	Variant not detected
Rhizomelic Chondrodysplasia Punctata Type 1	Variant not detected
Salla Disease	Variant not detected
Sickle Cell Anemia	Variant not detected
Sjögren-Larsson Syndrome	Variant not detected
Tay-Sachs Disease	Variant not detected
Tyrosinemia Type I	Variant not detected
Usher Syndrome Type 1F	Variant not detected
Usher Syndrome Type 3A	Variant not detected
Zellweger Syndrome Spectrum (PEX1-Related)	Variant not detected



Wellness Reports 3 highlighted reports of 8 reports available

Find out how your DNA may affect your body's response to diet, exercise, and sleep. **Consider talking to a healthcare professional before making any major lifestyle changes or if you have any concerns about your results.**

Alcohol Flush Reaction	Unlikely to flush
Caffeine Consumption	Likely to consume less
Deep Sleep	Less likely to be a deep sleeper
Genetic Weight	Predisposed to weigh less than average
Lactose Intolerance	Likely intolerant
Muscle Composition	Common in elite power athletes
Saturated Fat and Weight	Likely similar weight
Sleep Movement	Likely more than average movement



Ancestry Reports 1 highlighted report of 5 reports available

Discover the story of your ancient ancestors, your origins, and your ancestral background.

Ancestry Composition

European	99.5%
Eastern European	92.5%
Ashkenazi Jewish	3.0%
Southern European	1.5%
Balkan	0.8%
Broadly Southern European	0.7%
Northwestern European	0.7%
Finnish	0.2%
British & Irish	0.2%
Broadly Northwestern European	0.3%
Broadly European	1.9%
East Asian & Native American	0.5%
East Asian	0.5%
Yakut	0.5%
Broadly East Asian & Native American	Less than 0.1%
Maternal Haplogroup	H2
Neanderthal Ancestry	More Neanderthal variants than 57% of customers
Paternal Haplogroup	N-M46
Your DNA Family	1154 DNA Relatives



Traits Reports

0 highlighted reports of 22 reports available

Explore the genetics behind your appearance and senses.

Asparagus Odor Detection	Likely can smell
Back Hair	Likely little upper back hair
Bald Spot	Likely no bald spot
Bitter Taste	Likely can taste
Cheek Dimples	Likely no dimples
Cleft Chin	Likely no cleft chin
Earlobe Type	Likely detached earlobes
Early Hair Loss	Likely no hair loss
Earwax Type	Likely wet earwax
Eye Color	Likely brown or hazel eyes
Finger Length Ratio	Likely ring finger longer
Freckles	Likely little freckling
Hair Texture	Likely straight or wavy
Light or Dark Hair	Likely dark
Newborn Hair	Likely lots of baby hair
Photic Sneeze Reflex	Likely no photic sneeze reflex
Red Hair	Likely no red hair
Skin Pigmentation	Likely lighter skin
Sweet vs. Salty	Likely prefers salty
Toe Length Ratio	Likely big toe longer
Unibrow	Likely no unibrow
Widow's Peak	Likely no widow's peak

Roman Khmelevskiy's Reports Summary, printed on 2018-02-20 UTC



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