23andMe

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Reports Summary

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.**

Health Predisposition Reports **44+** reports available

Genetic factors that may influence your chances of developing 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.

Late-Onset Alzheimer's Disease	**Slightly increased risk**
Eczema (Atopic Dermatitis)	**Increased likelihood**
Hashimoto's Disease	**Increased likelihood**
Type 2 Diabetes	**Increased likelihood**
BRCA1/BRCA2 (Selected Variants)	**Variants not detected**
Hereditary Prostate Cancer (HOXB13-Related)	**Choose whether to receive this report**
MUTYH-Associated Polyposis	**Variants not detected**
Parkinson's Disease	**Variants not detected**
Age-Related Macular Degeneration	**Variants not detected**
Alpha-1 Antitrypsin Deficiency	**Variants not detected**
Anxiety	**Typical likelihood**
Asthma	**Typical likelihood**
Atrial Fibrillation	**Typical likelihood**
Attention-Deficit/Hyperactivity Disorder (ADHD)	**Typical likelihood**
Celiac Disease	**Variants not detected**
Chronic Kidney Disease (APOL1-Related)	**Variants not detected**
Coronary Artery Disease	**Typical likelihood**
Depression	**Typical likelihood**
Diverticulitis	**Typical likelihood**
Familial Hypercholesterolemia	**Variants not detected**

Fibromyalgia	**Typical likelihood**
G6PD Deficiency	**Variants not detected**
Gallstones	**Typical likelihood**
Glaucoma	**Typical likelihood**
Gout	**Typical likelihood**
HDL Cholesterol	**Typical likelihood**
Hereditary Amyloidosis (TTR-Related)	**Variants not detected**
Hereditary Hemochromatosis (HFE-Related)	**Variants not detected**
Hereditary Thrombophilia	**Variants not detected**
High Blood Pressure	**Typical likelihood**
Irritable Bowel Syndrome	**Typical likelihood**
Kidney Stones	**Typical likelihood**
LDL Cholesterol	**Typical likelihood**
Lupus	**Typical likelihood**
Migraine	**Typical likelihood**
Nonalcoholic Fatty Liver Disease	**Typical likelihood**
Obstructive Sleep Apnea	**Typical likelihood**
Psoriasis	**Typical likelihood**
Restless Legs Syndrome	**Typical likelihood**
Rosacea	**Typical likelihood**
Severe Acne	**Typical likelihood**
Skin Cancer (Basal and Squamous Cell Carcinomas)	**Typical likelihood**
Skin Cancer (Melanoma)	**Result not available**
Triglycerides	**Typical likelihood**

Pharmacogenetics Reports **3+** reports available

Learn whether you have specific genetic variants that may influence how your body processes certain medications. **Please talk to a healthcare professional if you are interested in learning more about how DNA variants may impact processing of some medications, or if you have concerns about your results.**

CYP2C19 Drug Metabolism	**Tutorial required before viewing**
DPYD Drug Metabolism	**Tutorial required before viewing**
SLCO1B1 Drug Transport	**Tutorial required before viewing**

Carrier Status Reports

46+ 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**
Familial Hyperinsulinism (ABCC8-Related)	**Variant not detected**
Familial Mediterranean Fever	**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**
Mucolipidosis 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 (SLC26A4-Related)	**Variant not detected**
Phenylketonuria and Related Disorders	**Variant not detected**
Pompe Disease	**Variant not detected**
Primary Hyperoxaluria Type 2	**Variant not detected**
Pyruvate Kinase Deficiency	**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 Spectrum Disorder (PEX1-Related)	**Variant not detected**

Wellness Reports **12+** 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**
Cat Allergy	**Increased likelihood**
Deep Sleep	**Less likely to be a deep sleeper**
Dog Allergy	**Typical likelihood**
Genetic Weight	**Predisposed to weigh about average**
Lactose Intolerance	**Likely intolerant**
Muscle Composition	**Common in elite power athletes**
Nearsightedness	**Typical likelihood**
Saturated Fat and Weight	**Likely similar weight**
Seasonal Allergies	**Typical likelihood**
Sleep Movement	**Likely more than average movement**

Ancestry Reports 6+ reports available

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

Ancestry Composition

Ancestry Composition

Sub-Saharan African	94.6%
West African	81.4%
Nigerian	36.8%
Ghanaian, Liberian & Sierra Leonean	31.5%
Senegambian & Guinean	4.1%
Broadly West African	9.0%
Congolese & Southern East African	12.5%
Angolan & Congolese	12.3%
Broadly Congolese & Southern East African	0.2%
African Hunter-Gatherer	0.4%
Broadly Sub-Saharan African	0.3%
European	4.9%
Northwestern European	4.6%
British & Irish	4.1%
Broadly Northwestern European	0.5%
Southern European	0.3%
Spanish & Portuguese	0.3%
East Asian	0.4%
Filipino & Austronesian	0.4%
Unassigned	0.1%

Maternal Haplogroup	L3e2b1a
Neanderthal Ancestry	Fewer Neanderthal variants than 94% of customers
Neanderthal Ancestry	Less Neanderthal variants than 100% of customers
Paternal Haplogroup	E-Z6018

Traits Reports

37+ reports available

Explore the genetics behind your appearance and senses.

Ability to Match Musical Pitch	**About a 50/50 chance of being able to match a musical pitch**
Asparagus Odor Detection	**Likely can't smell**
Back Hair	**Likely little upper back hair**
Bald Spot	**Likely no bald spot**
Bitter Taste	**Likely can't taste**
Bunions	**Less likely than average to have had a bunion**
Cheek Dimples	**Likely no dimples**
Cilantro Taste Aversion	**Odds of disliking cilantro not increased**
Cleft Chin	**Likely no cleft chin**
Dandruff	**About a 50/50 chance of getting dandruff**
Earlobe Type	**Likely detached earlobes**
Early Hair Loss	**Likely no hair loss**
Earwax Type	**Likely wet earwax**
Eye Color	**Likely brown or hazel eyes**
Fear of Heights	**Less likely than average to be afraid of heights**
Fear of Public Speaking	**Less likely to have a fear of public speaking**
Finger Length Ratio	**Likely ring finger longer**
Flat Feet	**More likely than average to have flat feet**
Freckles	**Likely little freckling**
Hair Photobleaching	**Less likely to experience hair photobleaching**
Hair Texture	**Likely straight or wavy**
Hair Thickness	**Less likely to have thick hair**
Ice Cream Flavor Preference	**More likely to prefer vanilla over chocolate ice cream**
Light or Dark Hair	**Likely dark**
Misophonia	**Less likely to hate chewing sounds**
Mosquito Bite Frequency	**Likely bitten as often as others**
Motion Sickness	**Less likely to experience motion sickness**
Newborn Hair	**Likely lots of baby hair**
Photic Sneeze Reflex	**Likely no photic sneeze reflex**
Red Hair	**Likely no red hair**
Skin Pigmentation	**Likely darker skin**
Stretch Marks	**Less likely to have stretch marks**

Sweet vs. Salty	**Likely prefers salty**
Toe Length Ratio	**Likely big toe longer**
Unibrow	**Likely at least a little unibrow**
Wake-Up Time	**Likely to wake up around 7:37 am**
Widow's Peak	**Likely no widow's peak**

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