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 **40** 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.

Age-Related Macular Degeneration	**Variant detected, not likely at increased risk**
Asthma	**Increased likelihood**
Celiac Disease	**Slightly increased risk**
Hashimoto's Disease	**Increased likelihood**
Hereditary Hemochromatosis (HFE-Related)	**Variant detected, not likely at increased risk**
Skin Cancer (Basal and Squamous Cell Carcinomas)	**Increased likelihood**
BRCA1/BRCA2 (Selected Variants)	**Variants not detected**
Late-Onset Alzheimer's Disease	**Variant not detected**
MUTYH-Associated Polyposis	**Variants not detected**
Parkinson's Disease	**Variants not detected**
Alpha-1 Antitrypsin Deficiency	**Variants not detected**
Anxiety	**Typical likelihood**
Atrial Fibrillation	**Typical likelihood**
Chronic Kidney Disease (APOL1-Related)	**Variants not detected**
Coronary Artery Disease	**Typical likelihood**
Diverticulitis	**Typical likelihood**
Eczema (Atopic Dermatitis)	**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 Thrombophilia	Variants not detected
High Blood Pressure	Typical likelihood
Irritable Bowel Syndrome	Typical likelihood
Kidney Stones	Typical likelihood
LDL Cholesterol	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 (Melanoma)	Typical likelihood
Triglycerides	Typical likelihood
Type 2 Diabetes	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	Predicted intermediate metabolizer
Citalopram (Celexa®)	Likely typical response
Clopidogrel (Plavix®)	Likely less effective
DPYD Drug Metabolism	Predicted normal metabolizer
SLCO1B1 Drug Transport	Predicted normal function

Carrier Status Reports **45** 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.**

Cystic Fibrosis	**Variant detected**
Rhizomelic Chondrodysplasia Punctata Type 1	**Variant detected**
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**
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
Primary Hyperoxaluria Type 2	Variant not detected
Pyruvate Kinase Deficiency	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 more
Cat Allergy	Increased likelihood
Deep Sleep	Less likely to be a deep sleeper
Dog Allergy	Increased likelihood
Genetic Weight	Predisposed to weigh less than average
Lactose Intolerance	Likely tolerant
Muscle Composition	Common in elite power athletes
Nearsightedness	Increased likelihood
Saturated Fat and Weight	Likely similar weight
Seasonal Allergies	Typical likelihood
Sleep Movement	Likely average or less movement

Ancestry Reports 6 reports available

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

Ancestry Composition

Ancestry Composition

European	99.5%
Northwestern European	98.2%
British & Irish	73.5%
French & German	23.2%
Broadly Northwestern European	1.5%
Ashkenazi Jewish	0.7%
Eastern European	0.6%
Western Asian & North African	0.4%
Arab, Egyptian & Levantine	0.2%
Levantine	0.2%
Northern West Asian	0.2%
Iranian, Caucasian & Mesopotamian	0.2%
Unassigned	0.1%
Maternal Haplogroup	T2a1a
Neanderthal Ancestry	More Neanderthal variants than 90% of customers
Neanderthal Ancestry	More Neanderthal variants than 76% of customers
Paternal Haplogroup	E-V13

Traits Reports

37 reports available

Explore the genetics behind your appearance and senses.

Ability to Match Musical Pitch	**Less likely to be able to match a musical pitch**
Asparagus Odor Detection	**Likely can 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	**Slightly higher odds of disliking cilantro**
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 blue or green eyes**
Fear of Heights	**Less likely than average to be afraid of heights**
Fear of Public Speaking	**More likely to have a fear of public speaking**
Finger Length Ratio	**Likely ring finger longer**
Flat Feet	**Less likely than average to have flat feet**
Freckles	**Likely a lot of freckles**
Hair Photobleaching	**About a 50/50 chance of experiencing hair photobleaching**
Hair Texture	**Likely straight or wavy**
Hair Thickness	**Less likely to have thick hair**
Ice Cream Flavor Preference	**About a 50/50 chance of preferring vanilla or chocolate ice cream**
Light or Dark Hair	**Likely dark**
Misophonia	**Less likely to hate chewing sounds**
Mosquito Bite Frequency	**Likely bitten more often than others**
Motion Sickness	**About a 50/50 chance of experiencing motion sickness**
Newborn Hair	**Likely little baby hair**
Photic Sneeze Reflex	**Likely photic sneeze reflex**
Red Hair	**Likely no red hair**
Skin Pigmentation	**Likely lighter skin**
Stretch Marks	**Less likely to have stretch marks**

Sweet vs. Salty	**Likely prefers salty**
Toe Length Ratio	**Likely second toe longer**
Unibrow	**Likely at least a little unibrow**
Wake-Up Time	**Likely to wake up around 8:16 am**
Widow's Peak	**Likely no widow's peak**

Ethan [redacted]s Reports Summary, printed on 2023-01-29 UTC

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