

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 Report	s 1 highlighted of 4 reports	Ancestry Reports 2 highli	ghted of 5 reports
 Late-Onset Alzheimer's Disease 	Slightly increased risk	 Ancestry Composition Your DNA Family 	24.4% British & Irish 12 populations
Carrier Status Reports 0 hig	ghlighted of 42 reports		populations
		Traits Reports 3 highlighte	ed of 19 reports
Wellness Reports 3 highlig	nted of 8 reports	 Asparagus Odor 	61% chance can smell,
 Genetic Weight Lactose Intolerance 	Predisposed to weigh more than average Likely tolerant	Detection Bitter Taste 	39% chance can't smell 51% chance can taste, 49% chance can't taste
 Saturated Fat and Weight 	Likely similar weight	Sweet Taste	50% chance prefers salty, 50% chance prefers sweet

Genetic Health Risk Reports 1 highlighted report of 4 reports available

These reports tell you about genetic variants that may increase your risk 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
Parkinson's Disease	Variants not detected
Alpha-1 Antitrypsin Deficiency	Variants not detected
Hereditary Thrombophilia	Variants not detected



These reports tell you about variants that may not affect your health, but could affect the health of your future family. For the conditions included in these reports, a person can be a carrier even if they don't have a personal or family history of the disease. **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 21	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
https://you.23andme.com/reports/print/	2/6

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

These reports help you understand how your DNA influences your body's response to environmental factors like diet or lifestyle. 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
Deep Sleep	Less likely to be a deep sleeper
Genetic Weight	Predisposed to weigh more than average
Lactose Intolerance	Likely tolerant
Muscle Composition	Common in elite power athletes
Saturated Fat and Weight	Likely similar weight
Sleep Movement	Likely average or less movement

Ancestry Reports 2 highlighted reports of 5 reports available

These reports let you explore what your DNA says about your origins and ancient ancestors.

Ancestry Composition

European	99.5%
Northwestern European	73.6%
British & Irish	24.4%
French & German	15.8%
Scandinavian	2.2%
Broadly Northwestern European	31.3%
Southern European	19.4%
Balkan	4.4%
Italian	2.1%
Broadly Southern European	12.9%
Ashkenazi Jewish	Less than 0.1%
Broadly European	6.4%
Sub-Saharan African	0.3%
West African	0.3%
Middle Eastern & North African	0.1%
Broadly Middle Eastern & North African	0.1%
East Asian & Native American	Less than 0.1%
East Asian	Less than 0.1%
Broadly East Asian	Less than 0.1%
Unassigned	Less than 0.1%
ternal Haplogroup	
Maternal	H1
aternal Haplogroup	{'haplogroup_id': 'H1'}
anderthal Ancestry	More Neanderthal variants than 53% of customers
ur DNA Family	1251 DNA Relatives
ar or a craining	

Traits Reports 3 highlighted reports of 19 reports available

These reports are a fun way to learn about how your DNA influences your physical appearance, preferences, and physical responses. The predictions are based on current knowledge of how genetic factors influence our traits.

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Asparagus Od	or Detection	61% chance can smell, 39% chance can't smell
Bitter Taste		51% chance can taste, 49% chance can't taste
Cheek Dimple	S	58% chance no dimples, 42% chance dimples
Cleft Chin		88% chance no cleft chin, 12% chance cleft chin
Earlobe Type		33% chance attached earlobes, 67% chance detached earlobes
Earwax Type		93% chance wet earwax, 7% chance dry earwax
Eye Color		35% chance lighter eyes, 65% chance darker eyes
Light	er eyes	35%
	Blue	3%
	Greenish blue	3%
	Green	9%
	Light hazel	20%
Dark	er eyes	65%
	Dark hazel	25%
	Light brown	13%
	Dark brown	27%
Finger Length	Ratio	26% chance index finger longer, 74% chance ring finger longer
Freckles		45% chance a lot of freckles, 55% chance little freckling
Hair Curliness		6% chance curly, 94% chance straight or wavy
Curly	1	6%
	Big curls	4%
	Small curls	2%
	Very tight curls	< 1%
Straig	ght or wavy	94%
	Wavy	13%
	Slightly wavy	41%
	Straight	40%

Light or Dark Hair	22% chance dark, 78% chance light
Dark	22%
Black	1%
Dark brown	21%
Light	78%
Light brown	33%
Dark blond	35%
Light blond	10%
Newborn Hair Amount	30% chance lots of baby hair, 70% chance little baby hair
Photic Sneeze Reflex	36% chance photic sneeze reflex, 64% chance no photic sneeze reflex
Red Hair	6% chance red hair, 94% chance no red hair
Skin Pigmentation	96% chance lighter skin, 4% chance darker skin
Lighter skin	96%
Very fair	39%
Moderately fair	32%
Light beige	25%
Darker skin	4%
Olive	3%
Light brown	1%
Dark brown	< 1%
Sweet Taste	50% chance prefers salty, 50% chance prefers sweet
Toe Length Ratio	58% chance big toe longer, 42% chance second toe longer
Unibrow	98% chance little or no unibrow, 2% chance moderate to thick unibrow
Little or no unibrow	98%
No unibrow	81%
Little unibrow	17%
Moderate to thick unibrow	2%
Moderate to thick unibrow	2%
Widow's Peak	83% chance no widow's peak, 17% chance widow's peak

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