





YOUR FITNESS REPORT

INTRODUCTION

This is your personal genomic fitness report, providing you with invaluable genetic insights into your personal physical fitness. Topics covered include exercise and heart-rate, strength, recovery, balance, mobility, reflexes, agility, precision, injury risk, BMI, fat, muscle, metabolism, blood pressure, cholesterol, insulin and glucose. On top of that, also shown in the report are personality traits, including optimism, leadership, conscientiousness, extraversion, learning, impulsivity, anger, and even combativeness.

Your fitness report, when presented to your doctor/healthcare professional/qualified personal trainer, could help you with your fitness goals. You may, for example, be interested in using your report to put together a fitness plan to become physically stronger, or to obtain a healthier BMI. Your report could help you to set and manage achievable fitness goals and even maximise the potential of your own personality traits day to day.

Do not modify diets/meal-plans or supplements recommended or prescribed by your doctor/qualified healthcare professional without first consulting them.

If you have any nutrition, diet, or healthcare related questions, please make sure that you promptly consult your doctor/qualified healthcare professional.

Genetic testing identifies and analyses your DNA and can help to rule out suspected genetic disorders and/or help determine your chances of developing or passing on a specific genetic disorder. Huge benefits from genetic testing also include:

1. Useful results for family planning

2. Guidance for your doctor/qualified healthcare professional in diagnosis and suitable treatments/interventions

3. Early interventions when it comes to medical conditions - which can even save a life.

As we all differ so much in their characteristics and traits, so do we in our genetic makeup. Our genetic differences are shown through variations in our DNA: the building blocks of life.

When a family member has a trait for a disease its wise to have genetic testing and screening to find out if you are a carrier or have the disease yourself.

Some genetic variations can increase our risk of developing certain health conditions, other variations can reduce them, whilst some people will develop a health condition without being at a higher genetic risk. Whichever way, it's wise to find out your own personal genetic composition and where possible, reduce your risk through lifestyle changes.

As a species, we humans are incredibly genetically diverse and complex. We find that there are some genetic variations more pronounced in one ethnicity over another, for example. Its in us identifying our own personal genetic complexities and variations that we can then work towards making constructive changes in our lives to better our health and wellbeing. Put simply, fully understanding your own genetic variations helps to provide a pathway to optimum health and a fruitful life.

Personal, appropriate genetic testing provides valuable insights but also has its limitations (see below).



LIMITATIONS AND OTHER IMPORTANT INFORMATION

Genetic testing provides you with personal information on your own genetic risk(s), based upon the assessment of specific genetic variants. However, it doesn't report on your entire genetic profile. It doesn't report on all of the genetic variants that relate to a specific disease or condition, and if a variant that is tested doesn't show up, that doesn't mean that there aren't any other genetic variants present that could be related. Indeed, other genetic risk tests could identify separate genetic variants that are related to the same disease/condition. In addition, your environment and lifestyle can be contributory risk factors affecting your risk of developing a disease or health condition.

Future versions of this report may include additional polymorphisms to provide a higher degree of accuracy than this SNP alone does.

Remember that testing is not a substitute for visits to your doctor/qualified healthcare professional. Any questions or concerns that you might have about your genetic test results or your current health status must be brought up in an appointment with them.

Lots of people find speaking with a genetic counselor or board-certified clinical geneticist very helpful in their genetic journey.

Genetic counselling can be valuable in helping you with worries and concerns that you might have about your genetic testing (both prior and post-results), and clinical geneticists can prove essential after you've received your results; to analyse them and potentially diagnose you with a condition/disease. You can access genetic counselors via the Genetic Counselor United Arab Emirates website: http://www.geneticcounselor.ae

The results of a genetic test do not provide you with a diagnosis or provide you with information about your current state of health/wellbeing and they mustn't be used by you to make any medical decisions, such as altering any medications that you are taking.

Our genetic testing reports haven't been evaluated by the FDA and by the UAE Ministry of Health & Prevention.Our product isn't intended to diagnose, treat, cure or prevent any medical condition/disease.

INFORMATION FOR HEALTH CARE PROFESSIONALS

Our genetic testing is not intended to diagnose a patient with a disease, to determine suitable medical treatments, or to inform the patient of anything regarding their current or future state of health. Instead, the test is intended to provide patients with personal genetic information to help influence their conversations with clinicians and to help them make constructive lifestyle decisions.

Any decisions regarding patient diagnosis or treatment should be based upon clinical diagnostic testing and/or other information that you consider to be appropriate.

This report is NOT intended for US persons and was not submitted for approval to the US Food and Drug Administration (FDA).

The above information provides an overview of predicted genetic risks to the patient. All information is based solely on genotype data and does not replace a consultation with a doctor/qualified healthcare professional.

Doctors/qualified healthcare professionals should also consider family history, symptoms presented, current medical prescriptions, and other factors prior to making any clinical or therapeutic decisions.

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KEY SUMMARY

	This colour means that you could have an enhanced beneficial reaction linked to the associated trait
•	This colour means that you could have an average reaction linked to the associated trait
	This solaur manys that you could have a reduced happing is reaction linked to the associated trait
•	This colour means that you could have a reduced beneficial reaction linked to the associated trait

QUICK SUMMARY

CARDIO		
CONDITION NAME	RESULTS	MAIN MESSAGE
Endurance Workout		People with your genetic profile are likely to receive enhanced health benefits from an endurance workout
Sprint	•	People with your genetic profile are likely to have an increased ability to sprint
Prepared for speed sports		People with your genetic profile are likely to have high propensity to benefit from speed sports
Predisposition to intense sports	•	People with your genetic profile are likely to have a high predisposition to benefit from intensive sports
Pace and variability of gait	•	People with your genetic profile are likely to walk at a higher pace
World-class athletic resistance	•	People with your genetic profile are likely to have high physical resistance
Respiratory capacity	•	People with your genetic profile are likely to have a regular respiratory capacity
Aerobic metabolism	•	People with your genetic profile are likely to have a normal capacity for oxygen consumption
STRENGTH		
CONDITION NAME	RESULTS	MAIN MESSAGE
Strength Workout	٠	People with your genetic profile are likely to receive more benefits from a strength workout
Muscle Strength	٠	People with your genetic profile are predisposed to have high muscle strength
Post Exercise Recovery	•	People with your genetic profile are likely to need longer recovery times post exercise
Hand Grip Strength	•	People with your genetic profile tend to have an increased grip ability following physical activity
Quadriceps strength	•	People with your genetic profile are likely to increase quadriceps strength easier during workouts
BALANCE AND FLEXIBILITY		
CONDITION NAME	RESULTS	MAIN MESSAGE



BALANCE AND FLEXIBILITY						
CONDITION NAME	RESULTS	MAIN MESSAGE				
Equilibrium	•	People with your genetic profile are likely to have altered equilibrium				
Joint mobility	•	People with your genetic profile have a lower range of joint motion and less joint laxity and flexibility				
OTHER KEY FACTORS						
CONDITION NAME	RESULTS	MAIN MESSAGE				
Reaction time (reflexes)	•	People with your genetic profile are likely to have a low reaction time				
Agility	•	People with your genetic profile are likely to have good agility				
Precision	•	People with your genetic profile are likely to have good eyesight precision				
BODY COMPOSITION - ADIP	OSE					
CONDITION NAME	RESULTS	MAIN MESSAGE				
Body mass index	•	People with your genetic profile are predisposed to have a tendency to gain weight easily				
Body mass index (non-smoking vs smokers interaction)	•	People with your genetic profile are likely to have an increase in body mass index in response to cigarette smoking				
Obesity in the absence of metabolic diseases	•	People with your genetic profile have a greater tendency to become obese				
Waist circumference adjusted for body mass index	•	People with your genetic profile tend to have a smaller waist circumference than average				
Waist-hip ratio		People with your genetic profile are likely to have a greater distribution of body fat				
Waist-hip ratio (energy interaction with the diet)		People with your genetic profile are likely to have a typical distribution of body fat in response to diet				
Subcutaneous adipose tissue	•	People with your genetic profile are likely to have regular accumulation of subcutaneous adipose tissue				
Ratio of visceral vs subcutaneous adipose tissue	•	People with your genetic profile are likely to have an unbalance between visceral adipose tissue/subcutaneous adipose tissue.				
Taking dietary macronutrients	•	People with your genetic profile tend to lose more weight by taking personalised dietary macronutrients				
Loss of Body Fat Response to Physical Activity	•	People with your genetic profile are likely to not have any enhanced benefits from exercise to lose body fat				
BODY COMPOSITION - MUS	CULATUR	E				
CONDITION NAME	RESULTS	MAIN MESSAGE				
Lean mass		People with your genetic profile are likely to have a regular lean mass				
Growth potential of lean mass	•	People with your genetic profile are likely to have a lower growth potential of lean mass				
Percentage of fibers that make up the muscle: white	•	People with your genetic profile are likely to have a low white muscle fiber percentage				
Percentage of fibers that make up the muscle: red	•	People with your genetic profile are likely to have an average red muscle fiber percentage				
BODY COMPOSITION - META	BOLISM					
CONDITION NAME	RESULTS	MAIN MESSAGE				
Body energy expenditure (24 hours)		People with your genetic profile are likely to have an enhanced metabolic rate				



BODY COMPOSITION - META	BOLISM			
CONDITION NAME	RESULTS	MAIN MESSAGE		
Metabolism at rest		People with your genetic profile are likely to have regular catalytic rate at rest		
Energy expenditure	•	People with your genetic profile are likely to have a higher consumption of energy		
EXERCISE AND HEALTH				
CONDITION NAME	RESULTS	MAIN MESSAGE		
Physical Activity in Weight Loss	•	People with your genetic profile are predisposed to being overweight		
RESPONSE TO PHYSICAL AC	CTIVITY			
CONDITION NAME	RESULTS	MAIN MESSAGE		
Blood Pressure Response to Physical Activity	•	People with your genetic profile have an increased likelihood of elevated blood pressure		
Interaction between LDL cholesterol levels and physical activity	•	People with your genetic profile are likely to have a stronger reduction of LDL levels in response to physical activity		
HDL (good) Cholesterol Response to Physical Activity	•	People with your genetic profile are likely to have an enhanced benefit in their HDL levels through exercise		
Insulin Sensitivity Response		People with your genetic profile are likely to have an enhanced insulin sensitivity in response to exercise		
Increase of glucose uptake in response to exercise	•	People with your genetic profile are likely to have a regular glucose uptake in muscle fiber in response to exercise		
Increased heart rate in response to exercise	•	People with your genetic profile are susceptible to an excessive increase in heart rate following physic activity		
Heart rate response to post- recovery exercise	•	People with your genetic profile are likely to have a regular heart rate reduction during recovery		
Athletic difficulties due to reduced heart rate	•	People with your genetic profile are likely to have a lower heart rate		
Cognitive Benefits (high motor coordination)	•	People with your genetic profile are likely to receive cognitive benefits for motor coordination from physical activity		
Predisposition to moderate to vigorous physical activity levels	•	People with your genetic profile are likely to have a low predisposition to benefit from vigorous physical activity levels		
Muscle response to resistance training	•	People with your genetic profile are likely to have low muscle response to resistance training		
Creatine kinase		People with your genetic profile are likely to have a costant creatine kinase level after racing		
INJURY RISK				
CONDITION NAME	RESULTS	MAIN MESSAGE		
Sport Injury Risk	•	People with your genetic profile are likely to have a lower tendency to incur sport injuries		
Tendinopathy	•	People with your genetic profile have an increased likelihood of getting tendonitis		
Predisposition to tendinopathies	•	People with your genetic profile are likely to have a high predisposition to tendonitis and other tendinopathies		
JOINTS				
CONDITION NAME	RESULTS	MAIN MESSAGE		
Joint fragility	•	People with your genetic profile are likely to have an increased chance of joint fragility		
Shoulder dislocation		People with your genetic profile are likely to have a lower tendency to dislocate their shoulder		



JOINTS		
CONDITION NAME	RESULTS	MAIN MESSAGE
Rotator cuff disease	•	People with your genetic profile are likely to have a higher susceptibility to rotator cuff disease
Cruciate ligament/anterior Cruciate ligament injuries	•	People with your genetic profile likely have a high risk factor for anterior cruciate ligament injuries
Ankle injury	•	People with your genetic profile are likely to not have an increased risk to develop Ankle injury
MUSCLES		
CONDITION NAME	RESULTS	MAIN MESSAGE
Muscle Cramps		People with your genetic profile are likely to have a decreased risk factor to develop muscle cramps after intense workouts
Propensity to the development of muscle pain	•	People with your genetic profile are likely to have a low propensity to feel muscle pain
Risk of suffering muscle damage		People with your genetic profile are likely to have a regular risk of suffering muscle damage
Exercise-induced myopathy	•	People with your genetic profile are likely to have a low tendency to experience myopathy
Slow muscle repair	•	People with your genetic profile are likely to have a good muscle repair ability
Predisposition to the development of inguinal stress hernias	•	People with your genetic profile are likely to have a low predisposition for inguinal stress hernias
STRESS FRACTURE		
CONDITION NAME	RESULTS	MAIN MESSAGE
Stress fracture period prevalence	•	People with your genetic profile are likely to have a lower tendency to develop stress fractures
Stress fracture	•	People with your genetic profile are likely to have a regular tendency to get stress fractures
PERSONALITY TRAITS		
CONDITION NAME	RESULTS	MAIN MESSAGE
Optimism		People with your genetic profile are predisposed to have a high sense of optimism
Leadership	•	People with your genetic profile are likely to have higher leadership abilities
Conscientiousness		People with your genetic profile are likely to have a less conscientious personality.
Extraversion	•	People with your genetic profile are likely to be more extraverted.
Insufficiency of experiential learning		People with your genetic profile are likely to have a good ability for experiential learning
Impulsivity		People with your genetic profile are likely to have a lower tendency to act impulsively
Response to Anger		People with your genetic profile are likely to have a good ability to handle feelings of anger
Combative personality		People with your genetic profile are likely to have a higher tendency to have a combative personality



ENDURANCE WORKOUT

RESULTS					
Endurance workouts are a type of intensive workout that helps improve stamina and build cardiovascular aerobic fitness. Examples include running, sprinting, swimming, and biking [1]. Rs2016520 is a specific genetic polymorphism that is linked to endurance in a person's athletic performance. The Rs2016520 polymorphism is a variation of the PPARD gene which can help to increase a person's endurance and strength [2]. People with your genetic profile are likely to receive enhanced health benefits from an endurance workout					
	_				
		SCIENTIFIC DE IAILS			
Gene		rsID	Genotype		
PPARD	PPARD rs2016520 CT				
REFERENCES					
[1] Morici, G., Gruttad'Auria, C. I., Baiamonte, P., Mazzuca, E., Castrogiovanni, A., & Bonsignore, M. R. (2016). Endurance training: is it bad for you?. Breathe (Sheffield, England), 12(2), 140–147.					

[2] Is there an interaction between PPARD T294C and PPARGC1A Gly482Ser polymorphisms and human endurance performance? Nir Eynon, Yoav Meckel, Alberto Jorge Alves, Chen Yamin, Michael Sagiv, Ehud Goldhammer, Moran Sagiv.

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SPRINT



[2] Voisin S, Cieszczyk P, et al. EPAS1 gene variants are associated with sprint/power athletic performance in two cohorts of European athletes. BMC Genomics. 2014; 15(1): 382. Published online 2014 May 18. doi: 10.1186/1471-2164-15-382.



PREPARED FOR SPEED SPORTS





PREDISPOSITION TO INTENSE SPORTS

RESULTS



Intense sports require high intensity and speed endurance training, each of which help to slow down fatigue and preserve energy and muscle oxidative capacity [1]. However, intense training induces oxidative stress and the inflammatory cytokine response. Based on scientific research, after an intense sport activity, genes (mainly the L17B, IL2RB, TXLNA and IL18R1 genes) became inflamed and regulated twice as much as other genes. A balanced pro-inflammatory and anti-inflammatory expression, and an immune response is vital in maintaining a hemostatic state in skeletal muscles, and in improving endurance training [2].

People with your genetic profile are likely to have a high predisposition to benefit from intensive sports





PACE AND VARIABILITY OF GAIT

RESULTS



Gait refers to the pattern of our limb movements, involving joint and muscle actions. This could be walking on the ground, or cycling; which involves swing movements [1]. Gait is key when it comes to fitness and health and gait speed is vital for high endurance athletes. Variations in genes that control neurological and cardiovascular functions affect a persons gait speed.

People with your genetic profile are likely to walk at a higher pace

SCIENTIFIC DETAILS				
Gene	rsID	Genotype		
PRKG1	rs10823991	AT		
REFERENCES				
[1] Hughes, J., & Jacobs, N. (1979). Normal huma	[1] Hughes, J., & Jacobs, N. (1979). Normal human locomotion. Prosthetics and Orthotics International, 3(1), 4–12.			



WORLD-CLASS ATHLETIC RESISTANCE

RESULTS



Resistance training is a type of physical exercise used to increase muscle mass, skeletal muscle capability, and muscle strength. Having stronger muscles can help professionals to improve their performance at a world-class level. Studies on resistance training have shown that it builds up an athlete's strength and plays a key part in enabling them to perform at an above-average level. Although resistance training might not improve sports ability, evidence indicates that it indirectly improves sport performance by reducing the likelihood of a person incurring an injury [1].

People with your genetic profile are likely to have high physical resistance





RESPIRATORY CAPACITY



[2] GSTCD and INTS12 Regulation and Expression in the Human Lung Maen Obeidat ,Suzanne Miller ,Kelly Probert,Charlotte K. Billington,Amanda P. Henry,Emily Hodge,Carl P. Nelson,Ceri E. Stewart,Caroline Swan,Louise V. Wain,María Soler Artigas,Erik Melén,Kevin Ushey,Ian P. Hall. Published: September 18, 2013https://doi.org/10.1371/journal.pone.0074630



AEROBIC METABOLISM



Review Miroslav Petr 1, Petr Stastny 2, Adam Zajac 3, James J Tufano 4, Agnieszka Maciejewska-Skrendo 5



STRENGTH WORKOUT





MUSCLE STRENGTH

	RESULTS			
(170)	Muscular strength and endurance are measured by looking at the maximum amount of force and number of contractions that a muscle group can exert within a certain time frame without leading to any muscular injury [1]. 30-85% of muscle strength and 50-80% of lean muscle mass are heritable traits, meaning that specific genes are associated with potential muscle strength. Studies suggest that a variation of the Gremlin1 gene is associated with lean mass and that the less frequently occurring variation in the CNTF gene (an A/A genotype) influences someone's muscle strength. Although the significance is not clear, some studies have suggested that variations of the ACTN3 gene affect skeletal muscle strength [2].			
	People with your genetic profile are predisposed to have high mus	scle strength		
	SCIENTIFIC DETA	AILS		
Gene	rsID	Genotype		
ACTN3	rs1815739	СС		
REFERENCES				
[1] Kell, R. T., Bell, G., & Quinney, A. (2001). Musculoskeletal fitness, health outcomes and quality of life. Sports medicine (Auckland, N.Z.), 31(12), 863–873. [2] Eynon N, Hanson ED, Lucia A, Houweling PJ, Garton F, et al. (2013) Genes for elite power and sprint performance: ACTN3 leads the way. Sports Med 4:				

803-817. 10.1007/s40279-013-0059-4



POST EXERCISE RECOVERY

RESULTS					
Post-exercise recovery time is the number of hours between when a person finishes exercising and their cardiovascular system returns to its "resting state" and it accounts for the physiological processes taking place in the body following exercise activity [1]. Muscle recovery is the vital mechanism that takes place during a post exercise recovery period, and certain genetic variations affect that process. People with your genetic profile are likely to need longer recovery times post exercise					
SCIENTIFIC DETAILS					
Gene	rsID	Genotype			
CRP	rs1205	СС			
REFERENCES					
[1] Romero, S. A., Minson, C. T., & Halliwill, J. R. (2017). The cardiovascular system after exercise. Journal of applied physiology (Bethesda, Md. : 1985), 122(4), 925–932.					



HAND GRIP STRENGTH



[2] Bohannon R. W. (2019). Grip Strength: An Indispensable Biomarker For Older Adults. Clinical interventions in aging, 14, 1681–1691.



QUADRICEPS STRENGTH

RESULTS					
Your quadriceps (often known as quads) are the big muscle group in your thighs. A deficit in quadricep strength is often the result of an anterior cruciate ligament injury and limits many sports people from performing at the levels at which they might have done in the past [1]. The Fokl polymorphism of the VDR gene alters how it is decoded, and the C allele of Fokl reduces a persons quadriceps strength. Studies show that Bsml (which is another VDR polymorphism) impacts the functions of skeletal muscles and the bb genotype of this gene variation might increase quadricep strength in healthy women [2]. People with your genetic profile are likely to increase quadriceps strength easier during workouts					
	SCIENTIFIC DETAILS				
Gene		rsID	Genotype		
VDR		rs2228570	AA		
REFERENCES					
[1] Palmieri-Smith, R. M., & Lepley, L. K. (2015). Quadriceps Strength Asymmetry After Anterior Cruciate Ligament Reconstruction Alters Knee Joint Biomechanics and Functional Performance at Time of Return to Activity. The American journal of sports medicine, 43(7), 1662–1669.					

[2] Hopkinson, N. S., Li, K. W., Kehoe, A., Humphries, S. E., Roughton, M., Moxham, J., Montgomery, H., & Polkey, M. I. (2008). Vitamin D receptor genotypes influence quadriceps strength in chronic obstructive pulmonary disease. The American journal of clinical nutrition, 87(2), 385–390.



EQUILIBRIUM

RESULTS				
During any athletic activity, cardio exercises, resistance/strength training it is vital to maintain the body's equilibrium and balance in order to try and avoid injury, muscle strain and suchlike. Maintaining a dynamic equilibrium during physical activities can aid sports people to perform better and for longer periods of time [1]. Variations in neurology genes can affect a person's mobility and balance. A variation in the Beta-III spectrin gene can cause Spinocerebellar ataxia (Lincoln ataxia) which impacts a person's cerebellum; which is responsible for regulating balance and equilibrium in the body [2].				
SCIENTIFIC DETAILS				
Gene	rsID	Genotype		
SLC7A2	rs2248010	AG		
REFERENCES				
[1] Hrysomallis C. (20 [2] Public Library of	 Hrysomallis C. (2011). Balance ability and athletic performance. Sports medicine (Auckland, N.Z.), 41(3), 221–232. Public Library of Science. (2012, December 6). New genetic disorder of balance and cognition discovered. ScienceDaily. Retrieved May 2, 2021 			



JOINT MOBILITY

		RESUL	٢S		
Joint mobility is defined as the degree of flexibility, articulation, and movement between two joints without any muscular restrictions. Flexibility and articulation are required in any form of sport activities and are determinants of overall physical fitness and performance. Muscle strength and joint mobility are strictly correlated [1]. Joint hypermobility syndrome due to mutation in the COL3A1, COL5A2 and COL6A3 causes loose and overly flexible joints which gives greater agility. Although a syndrome, this condition is known to benefit activities that require high flexibility such as acrobatics and gymnastics [2]. Other studies show that variation of the COL12A1 with T/T or C/T genotype reduces joint flexibility and movement and therefore protects sports people against severe ligament rupture [3].					
	People with your genetic p	profile have a lower range of joir	it motion and less joi	nt laxity and flexibility	
		SCIENTIFIC	DETAILS		
Gene		rsID		Genotype	
COL12A1	COL12A1 rs970547 TT				
REFERENCES					
 Micheo, W., Baer and rehabilitation, Grahame R. (19 O'Connell K, Ki Posthumus M, Cier 	ga, L., & Miranda, G. (2012). Ba: 4(11), 805–811. 99). Joint hypermobility and ge night H, Ficek K, Leonska-Dunie szczyk P, Collins M. Interactions	sic principles regarding strengt enetic collagen disorders: are th ec A, Maciejewska-Karlowska A, 5 between collagen gene variant	h, flexibility, and stabil ey related?. Archives Sawczuk M, Stepien- s and risk of anterior	iity exercises. PM & R : the journal of injury, function, of disease in childhood, 80(2), 188–191. Slodkowska M, O'Cuinneagain D, van der Merwe W, cruciate ligament rupture. Eur J Sport Sci.	

2015;15(4):341-50.



REACTION TIME (REFLEXES)

RESULTS				
A person's reaction time is the period of time that they require to respond to stimulus. The DRD4- 521 CT polymorphism (dopamine D4 receptor gene) can negatively affect a person's cognitive function, attention span, and reaction time, whilst the COMT Val158Met polymorphism involved in dopamine neurotransmission can also reduce a person's attention control. These impacts can lead to a person having a slower reflex [1]. The metalloproteinase 3 promoter is bound by a protein encoded by the TCF20 gene. The TCF20 regulates general cognitive functions and reaction times [2].				
SCIENTIFIC DETAILS				
Gene	rsID	Genotype		
TCF20	rs5758659	СТ		
REFERENCES				
 Szekely, A., Balota, D.A., Duchek, J.M., Nemoda, Z., Vereczkei, A. and Sasvari-Szekely, M. (2011), Genetic factors of reaction time performance: DRD4 7- repeat allele associated with slower responses. Genes, Brain and Behavior, 10: 129-136. 				

[2] Davies G, et al. Genetic contributions to variation in general cognitive function: a meta-analysis of genome-wide association studies in the CHARGE consortium (N=53949). Mol Psychiatry. 2015 Feb;20(2):183-92. doi: 10.1038/mp.2014.188. Epub 2015 Feb 3. PMID: 25644384; PMCID: PMC4356746.

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AGILITY



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PRECISION

RESULTS



The term precision is related to accuracy and validity, which in sport is needed in order to hit a target. High precision requires accuracy, coordination, agility, and often speed, and for this, eyesight and hearing are generally required. Having a good awareness of ones body, senses, functions, and suchlike is fundamental for the young; in order to develop controlled motor skills. Myopia is a condition which affects a persons eyesight and causes objects in the distance to appear blurry. Hepatocyte Growth Factor (HGF), Matrix Metalloproteinases (MMPs), and variations of the PAX6 and PPFIA2 genes seem to be associated with the development of Myopia, which can negatively affect athletes that require precision and accuracy in their sport [1,2]. Toll-Like Receptors (TLRs) play a significant role in the immune system and can be found on different types of membranes in the body, such as the retinal pigment: epithelium. However, genetic variations lead to TLRs not protecting the external retina or the choroid (the vascular layer of the eye) as as they should and this can negatively impact a persons levels of accuracy in sports such as archery [3].

People with your genetic profile are likely to have good eyesight precision

SCIENTIFIC DETAILS

Gene	rsID	Genotype	
TLR3	rs3775291	СС	
REFERENCES			

[1] Valle, M. S., Lombardo, L., Cioni, M., & Casabona, A. (2018). Relationship between accuracy and complexity when learning underarm precision throwing. European journal of sport science, 18(9), 1217–1225.

[2] Singh, M., & Tyagi, S. C. (2018). Genes and genetics in eye diseases: a genomic medicine approach for investigating hereditary and inflammatory ocular disorders. International journal of ophthalmology, 11(1), 117–134.

[3] Ma L, Tang F, et al. Association of toll-like receptor 3 polymorphism rs3775291 with age-related macular degeneration: a systematic review and metaanalysis. Sci Rep. 2016; 6:19718.



BODY MASS INDEX

DECULTO				
RESULIS				
Body Mass Index (BMI) is an indicator of body fat composition across the body which indicates a person's fitness. It is calculated by measuring the weight of an individual in kilograms and then dividing that number by the square of their height in meters. People with BMI values below 18.5 are considered to be underweight, those who have a BMI between 18.5 to 24.9 are considered to be in the healthy weight range, and those who have a BMI above 29.9 are considered to be in the obese range [1]. According to medical research, five main genes (MC4R, PCSK1, POMC, ADRB2 and BDNF) are associated with obesity and can increase a person's risk of developing obesity [2, 3]. People with your genetic profile are predisposed to have a tendency to gain weight easily				
	0			
	SCIENTIFIC DETAILS			
Gene	Gene rsID Genotype			
ADRB2	rs1042713	GA		
REFERENCES				
[1] Nuttall F. Q. (2015). Body Mass Index: Obesity, BMI, and Health: A Critical Review. Nutrition today, 50(3), 117–128.				
[2] Choquet, H., & Meyre, D. (2011). Genetics of Obesity: What have we Learned?. Current genomics, 12(3), 169–179.				
[2] Mitra CD, Tap, DV	Amini E Association of ADDD2 m10/2712 with Obesity and Obes	ity Delated Departures and Its Interaction with Distany Est in		

[3] Mitra SR, Tan PY, Amini F. Association of ADRB2 rs1042713 with Obesity and Obesity-Related Phenotypes and Its Interaction with Dietary Fat in Modulating Glycaemic Indices in Malaysian Adults. J Nutr Metab. 2019 Mar 17;2019:8718795. doi: 10.1155/2019/8718795. PMID: 31007954; PMCID: PMC6441509.



BODY MASS INDEX (NON-SMOKING VS SMOKERS INTERACTION)

RESULTS



Body Mass Index (BMI) is a calculation that can help people find out what their healthy weight is/would be. It provides indications of any medical problems that a person classified as having an unhealthy BMI might face. Research indicates that people who smoke often tend to have a lower body weight than non-smokers, people who stop smoking tend to gain weight [1]. This is because smoking can cause insulin resistance which can then lead to fat accumulation.

People with your genetic profile are likely to have an increase in body mass index in response to cigarette smoking



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
CFH	rs460184	TT	
REFERENCES			
[1] Abramowitz, M. K. et al. (2018). Muscle mass, BMI, and mortality among adults in the United States: A population-based cohort study. PLOS ONE, 13 (4), s. e0194697.			



OBESITY IN THE ABSENCE OF METABOLIC DISEASES

RESULTS		
Metabolic syndrome is a series of conditions that increase the risk of heart disease, stroke and type 2 diabetes. Metabolic syndrome is also closely linked to being overweight or obese [1]. However, obesity could be due to genetic factors in absence of metabolic disease and is often inherited. In obesity, the participation in the regulation of energy expenditure by the ADRB2, ADRB3, ADRB1 genes makes themselves excellent candidates for studying this condition. They belong to the B-adrenoceptor gene families [2]. People with your genetic profile have a greater tendency to become obese		
SCIENTIFIC DETAILS		
Gene	rsID	Genotype
ADRB1	rs1801253	CC
REFERENCES		
[1] Hunt, K. J., Heiss, G., Sholinsky, P. D., & Province, M. A. (2000). Familial history of metabolic disorders and the multiple metabolic syndrome: the NHLBI family heart study. Genetic epidemiology, 19(4), 395–409.		

[2] Plotnikov D, Williams C, Guggenheim JA. Association between birth weight and refractive error in adulthood: a Mendelian randomisation study. Br J Ophthalmol. 2020 Feb;104(2):214-219. doi: 10.1136/bjophthalmol-2018-313640. Epub 2019 May 16. PMID: 31097437.



WAIST CIRCUMFERENCE ADJUSTED FOR BODY MASS INDEX

RESULTS			
Waist circumference measurement (often used to assess the level of fat in the area), along with any adjustment for Body Mass Index (BMI), can be useful to help measure a person's obesity related health risks [1]. Experimental studies show that the MC4R gene is a key regulator of a person's energy balance; affecting their food intake and energy expenditure through changing hormonal neuronal pathways. If the MC4R gene changes its signal, it can affect a person's glucose utilization and insulin sensitivity, so impacting their waist circumference [2]. People with your genetic profile tend to have a smaller waist circumference than average			
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
MC4R		rs12970134	GG
REFERENCES			
[1] Berentzen, T. L. et al. (2012). Waist Circumference Adjusted for Body Mass Index and Intra-Abdominal Fat Mass. PLOS ONE, 7 (2), s. e32213. doi:10.1371/journal.pone.0032213			

[2] Kang J, Guan RC, Zhao Y, Chen Y. Obesity-related loci in TMEM18, CDKAL1 and FAIM2 are associated with obesity and type 2 diabetes in Chinese Han patients. BMC Med Genet. 2020 Mar 30;21(1):65. doi: 10.1186/s12881-020-00999-y. PMID: 32228543; PMCID: PMC7106578.



WAIST-HIP RATIO

RESULTS			
The Waist-Hip Ratio (WHR) compares a person's Body Mass Index (BMI) and waist circumference to help assess a their physical fitness and risk of developing Cardiovascular Disease (CVD), type 2 diabetes, hypertension (also known as high blood pressure) and elevated cholesterol levels [1]. The impact of genetics upon a person's fat distribution is complex with only about 48 Single-Nucleotide Polymorphisms (SNPs) (DNA sequence variations) identified in the study. However, according to a recent Genome-Wide Association Study (GWAS), the TBX15, HOXC13, RSP03 and CPEB4 genes were strongly associated with fat distribution and obesity risk factors such as insulin resistance, metabolic change and changes of the adipokine levels. Adipokine are a group of adipose tissue-derived bioactive molecules that are important in regulating diverse processes such as appetite, metabolism, fat distribution, insulin activity, and inflammation [2].			
	People with your genetic pl	othe are likely to have a greater distribution of body	/ fat
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
RSP03		rs9491696	CG
REFERENCES			
 [1] Emdin, C. A. et al. (2017). Genetic Association of Waist-to-Hip Ratio With Cardiometabolic Traits, Type 2 Diabetes, and Coronary Heart Disease. JAMA, 317 (6), s. 626. doi:10.1001/jama.2016.21042 			
[2] Schleinitz D. Bött	[2] Schleinitz, D., Böttcher, Y., Blüher, M., & Kovacs, P. (2014). The genetics of fat distribution. Diabetologia, 57(7), 1276–1286.		



WAIST-HIP RATIO (ENERGY INTERACTION WITH THE DIET)

RESULTS



Diet plays an essential role in energy consumption which then influences a person's Waist-Hip Ratio (WHR). Foods with a low Glycemic Index (GI) (do not produce high insulin release peaks) and low energy density are associated with lower levels abdominal fat and weight gain. Studies suggest that the TFAP2B gene is associated with a person's Body Mass Index (BMI) and waist circumference, which affects their WHR. TFAP2B is normally expressed in fat tissue and a variation of the gene can lead to an overexpression in the fat cells, increasing glucose uptake and insulin resistance [1,2]. ADAMTS9 T2D risk alleles are also associated with insulin resistance and differences in body fat distribution [3].

People with your genetic profile are likely to have a typical distribution of body fat in response to diet



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
ADAMTS9	rs6795735	СС	
REFERENCES			

[1] Romaguera, D., Ängquist, L., Du, H., Jakobsen, M. U., Forouhi, N. G., Halkjær, J., Feskens, E. J., van der A, D. L., Masala, G., Steffen, A., Palli, D., Wareham, N. J., Overvad, K., Tjønneland, A., Boeing, H., Riboli, E., & Sørensen, T. I. (2011). Food composition of the diet in relation to changes in waist circumference adjusted for body mass index. PloS one, 6(8), e23384.

[2] Stocks, T. et al. (2012). TFAP2B Influences the Effect of Dietary Fat on Weight Loss under Energy Restriction. PLOS ONE, 7 (8), s. e43212. doi:10.1371/journal.pone.0043212

[3] Heid IM, Jackson AU, Randall JC, et all. Meta-analysis identifies 13 new loci associated with waist-hip ratio and reveals sexual dimorphism in the genetic basis of fat distribution. Nat Genet. 2010 Nov;42(11):949-60. doi: 10.1038/ng.685. Epub 2010 Oct 10. Erratum in: Nat Genet. 2011 Nov;43(11):1164. PMID: 20935629; PMCID: PMC3000924.



SUBCUTANEOUS ADIPOSE TISSUE

RESULTS			
The tissue beneath the skin is called the Subcutaneous Adipose Tissue (SAT) or subcutaneous fat. Women mainly tend to store this fat in their gynoid regions, e.g. their hips, thighs, and gluteal region (buttocks). SAT is different from visceral fat tissue (fat stored within the abdominal cavity) as it is present in the lining of a persons internal organs [1]. A high accumulation of SAT leads to health risks such as metabolic syndrome, hypertension, and insulin resistance, whilst fat in the gynoid and gluteal regions lowers the risk of developing Cardiovascular Disease (CVD). Studies suggest that rs17782313 of MC4R and rs4846567 of LYPLAL1 are strongly associated with obesity and subcutaneous fat mass [2].			
	People with your genetic profile are likely to have regular accu	mulation of subcutaneous adipose tissue	
SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
PLIN1	rs2304795	АА	
REFERENCES			
 Mittal B. (2019). Subcutaneous adipose tissue & visceral adipose tissue. The Indian journal of medical research, 149(5), 571–573. Daily, J.W., Yang, H.J., Liu, M. et al. Subcutaneous fat mass is associated with genetic risk scores related to proinflammatory cytokine signaling and interact with physical activity in middle-aged obese adults. Nutr Metab (Lond) 16, 75 (2019). 			

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RATIO OF VISCERAL VS SUBCUTANEOUS ADIPOSE TISSUE

RESULTS



The ratios of visceral fat tissue (fat stored within the abdominal cavity) compared to Subcutaneous Adipose Tissue (SAT) (subcutaneous fat) is associated with cardiometabolic risk factors such as insulin resistance, blood pressure, and dyslipidemia. However, visceral fat storage is riskier to hold than SAT. According to a Genome-Wide Association Study (GWAS) study, genetic variants in the TFAP2B gene and near the MSRA gene significantly influence visceral fat gain in visceral fat tissues. The Pro12Ala variant of the PPARG gene increases a persons insulin sensitivity and reduces their risk of type 2 diabetes by reducing fat storage in their visceral and subcutaneous tissues [1].

People with your genetic profile are likely to have an unbalance between visceral adipose tissue/subcutaneous adipose tissue.



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
ROBO1	rs17377726	GG	
LGALS12	rs933186	СТ	
AL139390.1 - RREB1	rs2842895	СС	
AC116362.1	rs10060123	СС	
REFERENCES			
[1] Kaess, B. M. et al. (2012). The ratio of visceral to subcutaneous fat, a metric of body fat distribution, is a unique correlate of cardiometabolic risk.			

Diabetologia, 55 (10), s. 2622–2630. doi:10.1007/s00125-012-2639-5



TAKING DIETARY MACRONUTRIENTS



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t ID: GFX0435522

LOSS OF BODY FAT RESPONSE TO PHYSICAL ACTIVITY

RESULTS



Although physical endurance training increases a person's ability to process food (including fatty acids), carbohydrates are still the main energy source when exercising skeletal muscles. It has been suggested that the major effects of exercise on fat metabolism may occur after exercise, although this has never clearly been proven. Therefore, how exercise leads to a reduction in fat mass is still not completely clear. Since fat oxidation is increased during exercise, and since physical endurance exercise training increases the capacity to oxidize fat, by extension, it is assumed that 1) more fat is oxidized on a day that exercise is performed; and 2) people will oxidize more fat over a 24 hour period if they have exercise [1,2].

People with your genetic profile are likely to not have any enhanced benefits from exercise to lose body fat



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
LPL	rs328	СС	
REFERENCES			
 Booth, F. W., Chakravarthy, M. V. & Spangenburg, E. E. (2002). Exercise and gene expression: physiological regulation of the human genome through physical activity. The journal of physiology, 543 (2), s. 399–411. doi:10.1113/jphysiol.2002.019265 Hainer, V., Zamrazilová, H., Spálová, J., Hainerová, I., Kunesová, M., Aldhoon, B., & Bendlová, B. (2008). Role of hereditary factors in weight loss and its maintenance. Physiological research, 57 Suppl 1, S1–S15. 			

Kit ID: GFX0435522



LEAN MASS

RESULTS A person's weight is a combination of the body fat required for energy and the lean muscles, bones, organs, and suchlike. A lower lean mass is an important determinant of body health and the opposite can lead to disability and chronic diseases [1]. Studies show that the rs2276541 of the ACVR2B gene is strongly associated with lean body mass, particularly that of the muscles [2]. People with your genetic profile are likely to have a regular lean mass SCIENTIFIC DETAILS Gene rsID Genotype TRHR rs16892496 AA REFERENCES [1] Pomeroy, E., Macintosh, A., Wells, J. C. K., Cole, T. J. & Stock, J. T. (2018). Relationship between body mass, lean mass, fat mass, and limb bone crosssectional geometry: Implications for estimating body mass and physique from the skeleton. American journal of physical anthropology, 166 (1), s. 56-69. doi:10.1002/ajpa.23398

[2] Xiao-Gang Liu, Li-Jun Tan, Shu-Feng Lei, Yong-Jun Liu, Hui Shen, Liang Wang, Han Yan, Yan-Fang Guo, Dong-Hai Xiong, Xiang-Ding Chen, Feng Pan, Tie-Lin Yang, Yin-Ping Zhang, Yan Guo, Nelson L. Tang, Xue-Zhen Zhu, Hong-Yi Deng, Shawn Levy, Robert R. Recker, Christopher J. Papasian, Hong-Wen Deng, Genome-wide Association and Replication Studies Identified TRHR as an Important Gene for Lean Body Mass, The American Journal of Human Genetics, Volume 84, Issue 3, 2009, Pages 418-423, ISSN 0002-9297


GROWTH POTENTIAL OF LEAN MASS

RESULTS					
Strength training can increase a persons lean body mass [1]. Studies suggest that the rs1056513 variant in the PATJ gene, and the rs12439003 variant of the CAPN3 gene could help a person to increase their lean mass, to a degree [2,3]. People with your genetic profile are likely to have a lower growth potential of lean mass					
	SCIENTIFIC DI	ETAILS			
Gene	Gene rsID Genotype				
PATJ	PATJ rs1056513 GA				
REFERENCES					
 [1] Thomas, M. H., & Burns, S. P. (2016). Increasing Lean Mass and Strength: A Comparison of High Frequency Strength Training to Lower Frequency Strength Training. International journal of exercise science, 9(2), 159–167 [2] Klimentidis, Y. C., Bea, J. W., Thompson, P., Klimecki, W. T., Hu, C., Wu, G., Nicholas, J. S., Ryckman, K. K., & Chen, Z. (2016). Genetic Variant in ACVR2B Is Associated with Lean Mass. Medicine and science in sports and exercise, 48(7), 1270–1275. 					

[3] Novel Genetic Loci Identified for the Pathophysiology of Childhood Obesity in the Hispanic Population Anthony G. Comuzzie, Shelley A. Cole, Sandra L. Laston, V. Saroja Voruganti, Karin Haack, Richard A. Gibbs, Nancy F. Butte Published: December 14, 2012.



PERCENTAGE OF FIBERS THAT MAKE UP THE MUSCLE: WHITE

RESULTS				
W	White muscle fibers (also known as "fast-twitch" fibers), are composed of certain enzymes and proteins. Although a high resting energetic state is present within white muscle fibers, they fatigue quickly [1,2]. People with your genetic profile are likely to have a low white muscle fiber percentage			
SCIENTIFIC DETAILS				
Gene rsID Genotype				
ACVR2B	ACVR2B rs2268757 CT			
REFERENCES				
 Herbison, G. J., Jaweed, M. M., & Ditunno, J. F. (1982). Muscle fiber types. Archives of physical medicine and rehabilitation, 63(5), 227–230. Tajsharghi H. (2008). Thick and thin filament gene mutations in striated muscle diseases. International journal of molecular sciences, 9(7), 1259–1275. 				



PERCENTAGE OF FIBERS THAT MAKE UP THE MUSCLE: RED

RESULTS				
Red muscle fibers are "slow-twitch" muscle fibers and have a completely different composition and energy demand compared to white muscle fibers. Slow-paced, low-intensity exercise requires oxidative red muscle fibers. The red muscle has slow isoforms and its oxidative capacity is threefold higher than that of white muscle fibers [1]. An overexpression of the MSTN gene by the rs1805086 variant with the CC and TC genotype has a negative effect on muscle mass but leads to a higher red muscle fiber percentage [2,3].				
	People with your genetic profile are likely to have an average red muscle	fiber percentage		
Gene		Genotype		
MSTN	MSTN rs1805086 TT			
REFERENCES				
 [1] Glancy, B., & Balaban, R. S. (2011). Protein composition and function of red and white skeletal muscle mitochondria. American journal of physiology. Cell physiology, 300(6), C1280–C1290. [2] Fuku, N. et al. (2016). Muscle-Related Polymorphisms (MSTN rs1805086 and ACTN3 rs1815739) Are Not Associated with Exceptional Longevity in 				

Japanese Centenarians. PLOS ONE, 11 (11), s. e0166605. doi:10.1371/journal.pone.0166605 [3] Chen X, Guo Y, et al. (2018). Arginine promotes skeletal muscle fiber type transformation from fast-twitch to slow-twitch via Sirt1/AMPK pathway. The

Journal of Nutritional Biochemistry, 61, 155-162.



BODY ENERGY EXPENDITURE (24 HOURS)

RESULTS



A person's energy expenditure is determined by looking at their body size, its composition, their food consumption and their physical activity. The person's resting energy expenditure depends upon their body size, body composition (to upkeep more weight requires a higher energy expenditure), and food intake. Any form of physical activity causes energy expenditure [1].

People with your genetic profile are likely to have an enhanced metabolic rate

SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
UCP2	rs659366	СТ	
REFERENCES			
[1] Westerterp, K. R., & Plasqui, G. (2004). Physical activity and human energy expenditure. Current opinion in clinical nutrition and metabolic care, 7(6), 607–613.			



METABOLISM AT REST

	RESULT	S	
A person's Resting Metabolic Rate (RMR) is the total number of calories burned by their body when they are resting and which enables them to maintain a "physiological homeostasis and equilibrium". RMR is associated with blood circulation, breathing, and organ and neurological function [1,2]. The RMR is proportional to lean body mass and it decreases with fat percentage gain. Energy deficiency during a high RMR can cause growth impairment in people with SS genotype Sickle cell disease [3]. People with your genetic profile are likely to have regular catalytic rate at rest			
SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
CYP1A1	CYP1A1 rs1048943 TT		
REFERENCES			
[1] Blundell, J. E., Cau hunger and appetite [2] McMurray, R. G., S Medicine and science	dwell, P., Gibbons, C., Hopkins, M., Naslund, E., King, N., & Finlay control: a new formulation. Disease models & mechanisms, 55 joares, J., Caspersen, C. J., & McCurdy, T. (2014). Examining var e in sports and exercise, 46(7), 1352–1358.	rson, G. (2012). Role of resting metabolic rate and energy expenditure in (5), 608–613. iations of resting metabolic rate of adults: a public health perspective.	

[3] Singhal, A., Parker, S., Linsell, L. & Serjeant, G. (2002). Energy intake and resting metabolic rate in preschool Jamaican children with homozygous sickle cell disease. The american journal of clinical nutrition, 75 (6), s. 1093–1097. doi:10.1093/ajcn/75.6.1093



ENERGY EXPENDITURE

RESULTS					
	Energy expenditure refers to the amount of energy a person requires to maintain the functions of their body during physical activity. Total daily energy expenditure is the sum of a person's Basal Metabolic Rate (BMR): the heat that they produce after a meal, and the energy that they use up in physical activity. To avoid obesity and weight loss, a person's daily energy expenditure must equate to their consumption of calories. [1,2,3] People with your genetic profile are likely to have a higher consumption of energy				
	_				
SCIENTIFIC DETAILS					
Gene	Gene rsID Genotype				
LEP rs7799039		rs7799039	GG		
REFERENCES					
 [1] Heaney J. (2013) Energy: Expenditure, Intake, Lack of. In: Gellman M.D., Turner J.R. (eds) Encyclopedia of Behavioral Medicine. Springer, New York, NY. [2] Braud, S., Ciufolini, M., & Harosh, I. (2010). 'Energy expenditure genes' or 'energy absorption genes': a new target for the treatment of obesity and Type II diabetes. Future medicinal chemistry, 2(12), 1777–1783. 					

[3] Jiang, L., Penney, K. L., Giovannucci, E., Kraft, P. & Wilson, K. M. (2018). A genome-wide association study of energy intake and expenditure. PLOS ONE, 13 (8), s. e0201555. doi:10.1371/journal.pone.0201555



PHYSICAL ACTIVITY IN WEIGHT LOSS

RESULTS				
	Exercise plays a crucial role in a person's weight loss as it utilizes any excess calories that are stored as fat. During exercise, the body uses its fat stores to produce and then break down glucose to meet the body's demand for energy. Glucose breakdown is essential for weight loss. The GCK gene plays a vital role in breaking down glucose in the body, and variations in the gene could significantly affect the breakdown of glucose in the body and ultimately lead to weight gain [1,2]. People with your genetic profile are predisposed to being overweight			
SCIENTIFIC DETAILS				
Gene rsID Genotype				
FTO	rs1121980	GA		
REFERENCES				
 [1] Cox C. E. (2017). Role of Physical Activity for Weight Loss and Weight Maintenance. Diabetes spectrum : a publication of the American Diabetes Association, 30(3), 157–160. 				

[2] Matschinsky, F. M. & Wilson, D. F. (2019). The Central Role of Glucokinase in Glucose Homeostasis: A Perspective 50 Years After Demonstrating the Presence of the Enzyme in Islets of Langerhans. Frontiers in physiology, 10. doi:10.3389/fphys.2019.00148

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BLOOD PRESSURE RESPONSE TO PHYSICAL ACTIVITY

RESULTS			
<u>S</u>	Systolic Blood Pressure (BP) is the measurement of the pressure placed on a person's arterial walls with each heartbeat. This of course increases during physical activity where the heart rate is raised due to an increase in muscular demand for oxygen. However, when a person's systolic BP significantly increases during exercise, they can also experience a Hypertensive Response to Exercise (HRE). The 235T and 174M variants of the Angiotensinogen (AGT) gene can potentially lead to hypertension, and the EDN1 gene plays an important role in encoding Endothelin 1 (ET-1) (a 21-amino acid peptide which controls blood pressure) [1,2].		
	-		
SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
EDN1	rs5370	GT	
REFERENCES			
[1] Kim, D. & Ha, JW. (2016). Hypertensive response to exercise: mechanisms and clinical implication. Clinical hypertension, 22 (1). doi:10.1186/s40885-016-0052-y			

[2] Effect of Endothelin 1 Genotype on Blood Pressure Is Dependent on Physical Activity or Fitness Levels Tuomo Rankinen, Timothy Church, Treva Rice, Nathan Markward, Arthur S. Leon, Dabeeru C. Rao, James S. Skinner, Steven N. Blair, and Claude Bouchard. Originally published 15 Oct 2007https://doi.org/10.1161/HYPERTENSIONAHA.107.093609Hypertension. 2007;50:1120–1125



INTERACTION BETWEEN LDL CHOLESTEROL LEVELS AND PHYSICAL ACTIVITY

RESULTS



Exercise alone does not lower a persons "bad" cholesterol levels (formed of Low-Density Lipoproteins (LDLs)); also required is an improvement in their diet. Reducing LDLs through physical activity (endurance exercises are often the most beneficial [1]) and a managed diet can reduce a persons risk of developing Cardiovascular Disease (CVD). The LDL receptors present in the liver transport cholesterol out of the bloodstream, but mutations to the LDLR, APOB, PCSK9, LRP1B and LDLRAP1 genes in the LDL receptors can cause hypercholesterolemia, a dangerous condition associated with CVD [2,3,4].

People with your genetic profile are likely to have a stronger reduction of LDL levels in response to physical activity



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
LRP1B	rs7583934	TT	
REFERENCES			

[1] Albarrati, A. M., Alghamdi, M., Nazer, R. I., Alkorashy, M. M., Alshowier, N., & Gale, N. (2018). Effectiveness of Low to Moderate Physical Exercise Training on the Level of Low-Density Lipoproteins: A Systematic Review. BioMed research international, 2018, 5982980.

[2] Hobbs, H. H., Brown, M. S., & Goldstein, J. L. (1992). Molecular genetics of the LDL receptor gene in familial hypercholesterolemia. Human mutation, 1(6), 445–466.

[3] Paththinige, C., Sirisena, N. & Dissanayake, V. (2017). Genetic determinants of inherited susceptibility to hypercholesterolemia – a comprehensive literature review. Lipids in health and disease, 16 (1). doi:10.1186/s12944-017-0488-4.

[4] Chun-Xiang Liu, Yonghe Li, Lynn M. Obermoeller-McCormick, Alan L. Schwartz, Guojun Bu, The Putative Tumor Suppressor LRP1B, a Novel Member of the Low Density Lipoprotein (LDL) Receptor Family, Exhibits Both Overlapping and Distinct Properties with the LDL Receptor-related Protein^{*}, Journal of Biological Chemistry, Volume 276, Issue 31, 2001, Pages 28889-28896, ISSN 0021-9258,



HDL (GOOD) CHOLESTEROL RESPONSE TO PHYSICAL ACTIVITY

RESULTS



Exercise can decrease levels of the cholesterol known as Low-Density Lipoproteins (LDL) and increase the cholesterol known as High-Density Lipoproteins (HDL) (the former being of detriment to the body, the latter being of benefit to the body). The function of HDL is to transport any surplus bad cholesterol from the blood to the liver where it is then broken down and excreted from the body [1]. Moderate to intense exercise is associated with higher HDL. People with certain metabolic syndromes have a low HDL level which can lead to obesity, high blood pressure and type 2 diabetes. In the Heritage Family Study, people with the C/T and T/T genotypes for the rs2076167 of the PPARD gene variants were more likely to have enhanced HDL levels when exercising than those without the variants. People with the latter might be able to increase their levels through exercise but may not experience enhanced effects.

People with your genetic profile are likely to have an enhanced benefit in their HDL levels through exercise

SCIENTIFIC DETAILS				
Gene	rsID	Genotype		
PPARD	rs2076167	СТ		
REFERENCES				
[1] Ruiz-Ramie, J. J., Barber, J. L. & Sarzynski, M. A. (2019). Effects of exercise on HDL functionality. Current opinion in lipidology, 30 (1), s. 16–23. doi:10.1097/mol.000000000000568				



INSULIN SENSITIVITY RESPONSE

RESULTS				
	Insulin is a hormone produced in the pancreas that enables people to keep their glucose levels low. In healthy people and in those with type 2 diabetes, exercise increases their insulin-stimulated glucose uptake and regulates their lipid metabolism [1]. People with your genetic profile are likely to have an enhanced insulin sensitivity in response to exercise			
SCIENTIFIC DETAILS				
Gene	Gene rsID Genotype			
LIPC		rs1800588	СС	
REFERENCES				
[1] Venkatasamy, V. V., Pericherla, S., Manthuruthil, S., Mishra, S., & Hanno, R. (2013). Effect of Physical activity on Insulin Resistance, Inflammation and Oxidative Stress in Diabetes Mellitus, Journal of clinical and diagnostic research : JCDR, 7(8), 1764–1766.				



INCREASE OF GLUCOSE UPTAKE IN RESPONSE TO EXERCISE

RESULTS				
	The skeletal muscle contractions caused by exercise increase a persons glu uptake 50-fold; where the glucose is brought to the muscle, the glucose e People with your genetic profile are likely to have a regular glucose uptake	cose uptake. Exercise can increase a person's glucose nters the muscle cells, and then the cells process it [1,2]. in muscle fiber in response to exercise		
SCIENTIFIC DETAILS				
Gene rsID Genotype				
BDKRB2	BDKRB2 rs1799722 CC			
REFERENCES				
 [1] Richter, E. A., Derave, W. & Wojtaszewski, J. F. P. (2001). Glucose, exercise and insulin: emerging concepts. The journal of physiology, 535 (2), s. 313–322. doi:10.1111/j.1469-7793.2001.t01-2-00313.x, [2] Sylow L. Kleinert, M. Richter, F. et al. Exercise-stimulated glucose uptake — regulation and implications for glycaemic control. Nat Rev Endocrinol 13. 				

[2] Sylow, L., Kleinert, M., Richter, E. et al. Exercise-stimulated glucose uptake — regulation and implications for glycaemic control. Nat Rev Endocrinol 133–148 (2017).



INCREASED HEART RATE IN RESPONSE TO EXERCISE

RESULTS



It is a well-known fact that a person's heart rate increases significantly during exercise and remains higher than normal for a time afterward [1]. There is a strong association between an increased heart rate and variants of the YWHAQ gene; a member of the family of signalling proteins which take part in processes such as cellular suicide, cell division, and metabolism. Individuals with a certain YWHAQ gene variant can be at an increased vulnerability of developing an excessive heart rate during physical exercise [2].

People with your genetic profile are susceptible to an excessive increase in heart rate following physical activity



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
YWHAQ	rs12692388	СТ	
REFERENCES			

[1] Jagoda, A., Myers, J. N., Kaminsky, L. A., & Whaley, M. H. (2014). Heart rate response at the onset of exercise in an apparently healthy cohort. European journal of applied physiology, 114(7), 1367–1375.

[2] Rankinen T, Sung YJ, Sarzynski MA, Rice TK, Rao DC, Bouchard C. Heritability of submaximal exercise heart rate response to exercise training is accounted for by nine SNPs. J Appl Physiol (1985). 2012;112(5):892-897.





HEART RATE RESPONSE TO POST-RECOVERY EXERCISE

RESULTS			
	During intense exercise, a person will have a high heart rate, but within about 2 minutes, it will quickly return to normal [1, 2]. Post-exercise recovery time is the number of hours between when a person finishes exercising and their cardiovascular system returns to its "resting state" and it accounts for the physiological processes taking place in the body following exercise activity [1, 2]. The protein hormone called Insulin-like Growth Factor 2 (IGF-2) regulates different metabolic processes and affects heart rate during the cardiovascular recovery process. People with your genetic profile are likely to have a regular heart rate reduction during recovery		
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
IGF-II		rs680	ТТ
REFERENCES			
[1] Javorka, M., Zila, I., Balhárek, T. & Javorka, K. (2002). Heart rate recovery after exercise: relations to heart rate variability and complexity. Brazilian journal of medical and biological research, 35 (8), s. 991–1000. doi:10.1590/s0100-879x2002000800018.			

[2] Romero, S. A., Minson, C. T. & Halliwill, J. R. (2017). The cardiovascular system after exercise. Journal of applied physiology, 122 (4), s. 925–932. doi:10.1152/japplphysiol.00802.2016.



ATHLETIC DIFFICULTIES DUE TO REDUCED HEART RATE

RESULTS			
Athletic Heart Syndrome (AHS) is a condition in which a person's heart is enlarged, causing the resting heart rate to be lower than it would be, were it in a "normal" state. This condition can be had by athletes or people who do practice sport for more than one hour per week. Although the syndrome is not potentially harmful, it can cause issues such as dizziness and fainting [1]. Bradycardia (a slower than normal heart rate) can cause Atrial Fibrillation (AF) which can then increase a person's risk of heart failure or stroke.			
SCIENTIFIC DETAILS			
Gene	rsID Genotype		
CHRM2	CHRM2 rs324640 GA		GA
REFERENCES			
[1] Moses, S., (2008) Athletic Heart Syndrome. Family Practice Notebook.here online Archived 26 May 2008 at the Wayback Machine.			

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COGNITIVE BENEFITS (HIGH MOTOR COORDINATION)

RESULTS			
Physical activities that involve motor coordination can improve a persons motor skills and cognitive function [1]. Cognitive function is, however, largely influenced by genetics. One study suggests that the APOE gene and brain-derived neurotrophic factor (BDNF Val66Met) polymorphisms interact to affect a persons cognitive function and can reduce their episodic memory. These genetic variations could ultimately lead to cognitive function disorders such as dementia and Alzheimer's disease [2]. People with your genetic profile are likely to receive cognitive benefits for motor coordination from physical activity			
		_	
SCIENTIFIC DETAILS			
Gene rsID Genotype			
BDNF	BDNF rs6265 CC		
REFERENCES			
[1] Zeng, N., Ayyub, M., Sun, H., Wen, X., Xiang, P., & Gao, Z. (2017). Effects of Physical Activity on Motor Skills and Cognitive Development in Early Childhood: A Systematic Review. BioMed research international, 2017, 2760716.			

[2] Fan, J., Tao, W., Li, X., Li, H., Zhang, J., Wei, D., Chen, Y., & Zhang, Z. (2019). The Contribution of Genetic Factors to Cognitive Impairment and Dementia Apolipoprotein E Gene, Gene Interactions, and Polygenic Risk. International journal of molecular sciences, 20(5), 1177.



PREDISPOSITION TO MODERATE TO VIGOROUS PHYSICAL ACTIVITY LEVELS

RESULTS



Physical activity is key in both obtaining and retaining physical and mental health. However, certain genetic variants can bring about risk in moderate to vigorous physical activities. The GNPDA2 gene encodes glucosamine-6-phosphate deaminase, which is involved in the biosynthesis of glucosamine and is expressed by a small part of the brain called the hypothalamus. People with variations in the GNPDA2 gene are less likely to follow intense workouts and instead prefer more moderate physical exercise [1]

People with your genetic profile are likely to have a low predisposition to benefit from vigorous physical activity levels



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
GNPDA2	rs10938397	AG	
REFERENCES			
[1] Zhang, X., & Speakman, J. R. (2019). Genetic Factors Associated With Human Physical Activity: Are Your Genes Too Tight To Prevent You			

Exercising?. Endocrinology, 160(4), 840–852.



MUSCLE RESPONSE TO RESISTANCE TRAINING

RESULTS



Resistance training can improve a person's skeletal muscle function through causing the regeneration of muscular tissue. During resistance training, muscle fibers and motor neurons work together as a "motor pool" to coordinate the contractions of the muscles. Variations in the NR3C1 gene (which is responsible for binding to glucocorticoid hormones) can decrease a person's muscle strength and reduce their lean mass which then affects their muscles' response to resistance training [1].

People with your genetic profile are likely to have low muscle response to resistance training



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
NR3C1	rs4634384	СТ	
REFERENCES			
[1] Bandy W. D. Lovelace-Chandler V. & McKitrick-Bandy B. (1990). Adaptation of skeletal muscle to resistance training. The Journal of orthopaedic and			

 Bandy, W. D., Lovelace-Chandler, V., & McKitrick-Bandy, B. (1990). Adaptation of skeletal muscle to resist sports physical therapy, 12(6), 248–255.



CREATINE KINASE

RESULTS				
Creatine Kinase (CK) is an enzyme present in several types of tissue, including the heart, the brain, and with the highest concentration found in skeletal and cardiac muscles. The enzyme levels are elevated when physical activities cause muscle damage [1]. People with your genetic profile are likely to have a costant creatine kinase level after racing				
SCIENTIEIC DETAILS				
Gene	Gene rsID Genotype			
SOD2	SOD2 rs4880 GG			
REFERENCES				
[1] Bais, R., & Edward	s, J. B. (1982). Creatine kinase	e. Critical reviews in clinical lab	poratory sciences, 16(4), 291–335.	



SPORT INJURY RISK





TENDINOPATHY

RESULTS Image: Substance of the second system of the secon



PREDISPOSITION TO TENDINOPATHIES

RESULTS



People experiencing tendinopathies often have a genetic predisposition to it. Tendonitis involves an inflammation of the tendon itself and its'enthesis (where the tendon is attached to the bone). Hypoxia (a lack of oxygen supply) is a trigger for tendinopathies because rather than it promoting the synthesis of type 1 collagen (typical for tendons), it promotes the synthesis of type III collagen, which is less resistant than type I and stimulates proinflammatory molecules [1].

People with your genetic profile are likely to have a high predisposition to tendonitis and other tendinopathies



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
TIMP2	rs4789932	GG	
REFERENCES			
[1] Xu, Y., & Murrell, G. A. (2008). The basic science of tendinopathy. Clinical orthopaedics and related research, 466(7), 1528–1538.			



JOINT FRAGILITY

	RESULT	S	
The joints represent the junctions between two bones covered by a layer of cartilage. Under "normal conditions", the cartilage has a smooth surface. Prolonged engagement in sport can lead to weak and fragile joints (especially in the knees and ankles), which can then predispose a person to joint injuries. Joints can often be painful; continuous joint pain may be felt. Ultimately it can lead to post-traumatic osteoarthritis: characterized by articular cartilage degeneration that causes joint pain and dysfunction. To try and avoid joint injury, it is advisable to moderate a persons exercise [1].			
1 11	People with your genetic profile are likely to have an increase	ed chance of joint fragility	
	SCIENTIFIC [ETAILS	
Gene	Gene rsID Genotype		
IL1B	IL1B rs16944 GG		
REFERENCES			
[1] Buckwalter J. A. (2003). Sports, joint injury, and posttraumatic osteoarthritis. The Journal of orthopaedic and sports physical therapy, 33(10), 578– 588.			



SHOULDER DISLOCATION

RESULTS				
Shoulder dislocations commonly fall into one of two categories. The first are anterior dislocations which occur either when a persons arm is abducted and externally rotated, or when the posterior humerus falls out of the joint. The second category is that of a posterior dislocation, which happens with an injury to the anterior shoulder. With a posterior dislocation a persons shoulder joint is prone to dislocations as the shallow glenoid (part of the shoulder socket) moves around the humerus (the small head of the bone). Dislocations can lead to nerve damage, tissue tears, fractures, and a type of fracture. Strong force or extreme rotation can pop the shoulder humerus out of the pocket. Some immediate relocation techniques include Scapular manipulation, the external rotation technique, the Cunningham technique, and the Stimson technique [1].				
	· · · · · · · · · · · · · · · · · · ·			
	SCIENTIFIC DETAILS			
Gene	Gene rsID Genotype			
COL1A1 rs1800012 CA		СА		
REFERENCES				
[1] Abrams R, Akbarnia H. Shoulder Dislocations Overview. [Updated 2020 Nov 1]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2021 Jan				



ROTATOR CUFF DISEASE

RESULTS				
S S I	Rotator cuff disease is a type of shoulder tendon injury that results from tears. Intensive activities, shoulder injuries, and prolonged use of the related joints are a risk factor for developing rotator cuff disease. Partial and anterior tears can develop into cuff degeneration. People with this disease show common onset symptoms such as acute pain and when examined, tenderness to the area, and muscle atrophy. People with this disease find it difficult to engage in overhead sports, to carry heavy objects, and often, carry out daily activities. Studies show that genetics play an important role and that the rs1800972 C>G variant of the DEFB1 gene is seen more frequently in people with the disease [1,2].			
	_			
SCIENTIFIC DETAILS				
Gene	Gene rsID Genotype			
DEFB1	DEFB1 rs1800972 GG			
REFERENCES				
[1] May T, Garmel GM. Rotator Cuff Injury. [Updated 2020 Jul 2]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2021 Jan- [2] Dabija, D. I., Gao, C., Edwards, T. L., Kuhn, J. E., & Jain, N. B. (2017). Genetic and familial predisposition to rotator cuff disease: a systematic review. Journal of shoulder and elbow surgery, 26(6), 1103–1112.				



CRUCIATE LIGAMENT/ANTERIOR CRUCIATE LIGAMENT INJURIES

RESULTS The Anterior Cruciate Ligament (ACL) is one of the 2 ligaments responsible for supporting a persons knee joint. Injury to the ACL is very common during sports activities, due to rapid movements, sudden changes in direction, sudden stops, and jumping/landing motions. Some of the most common injuries to the ACL are sprains and tears; where people will often hear a snap/popping sound, feel severe pain, and notice the swelling of the joint [1]. People with your genetic profile likely have a high risk factor for anterior cruciate ligament injuries SCIENTIFIC DETAILS Gene rsID Genotype COL5A1 rs12722 СТ **REFERENCES** [1] Evans J, Nielson JI. Anterior Cruciate Ligament Knee Injuries. [Updated 2021 Feb 19]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2021 Jan-.



ANKLE INJURY

RESULTS			
Acute ankle injuries are the most common type of injury experienced by athletes and sports people. They affect soft tissue; causing sprains and dislocations, mainly in the Anterior Talofibular Ligament (ATFL) and the Calcaneofibular Ligament (CFL). Severe injuries or dislocations can even require surgical intervention [1]. People with your genetic profile are likely to not have an increased risk to develop Ankle injury			
	_		
SCIENTIFIC DETAILS			
Gene	Gene rsID Genotype		
MGAT4C	MGAT4C rs10506913 AA		
REFERENCES			
[1] Polzer, H., Kanz, K. G., Prall, W. C., Haasters, F., Ockert, B., Mutschler, W., & Grote, S. (2012). Diagnosis and treatment of acute ankle injuries: development of an evidence-based algorithm. Orthopedic reviews, 4(1), e5.			



MUSCLE CRAMPS

RESULTS			
Muscle cramps are the involuntary contractions of skeletal muscles which usually last for a short period of time but which can cause severe pain and immobility of the muscle in question. They often occur either during or after exercise. Causes of the cramps are uncertain; with some studies stating that they are caused by an imbalance in the body's levels of salt and water, some say they are due to unusual spinal reflect movements, whilst others claim that they are caused by fatigue or dehydration [1]. Also associated with muscle cramps are long periods of exercise or physical work (especially those in hot weather) and certain medications and illnesses [1].			
	People with your genetic profile are likely to have a decreased	risk factor to develop muscle cramps after intense workouts	
Gene	rsID	Genotype	
MCT1	rs1049434	АА	
REFERENCES			
 Maughan, R. J., & Shirreffs, S. M. (2019). Muscle Cramping During Exercise: Causes, Solutions, and Questions Remaining. Sports medicine (Auckland, N.Z.), 49(Suppl 2), 115–124. 			



PROPENSITY TO THE DEVELOPMENT OF MUSCLE PAIN

RESULTS			
Muscle pain can be experienced during physical activity, straight after physical activity, and later on as the result of muscle cramps. Muscle pain from exercise is due to lactic acid formation, ion imbalances, hormonal factors, and proteins [1]. Certain genetic factors are associated with it and some variations can predispose a person to muscle damage and soreness. Studies suggest that the R577X polymorphism of the ACTN3 gene and the TT genotype of the SLC30A8 gene are biomarkers for exercise induced muscle damage or pain [2].			
	People with your genetic profile are likely to have a	low propensity to feel muscle pain	
SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
SLC30A8	rs13266634	CT	
REFERENCES			
[1] Miles, M. P., & Clark 203–216.	[1] Miles, M. P., & Clarkson, P. M. (1994). Exercise-induced muscle pain, soreness, and cramps. The Journal of sports medicine and physical fitness, 34(3), 203–216.		

[2] Baumert, P., Lake, M. J., Stewart, C. E., Drust, B., & Erskine, R. M. (2016). Genetic variation and exercise-induced muscle damage: implications for athletic performance, injury and ageing. European journal of applied physiology, 116(9), 1595–1625.



RISK OF SUFFERING MUSCLE DAMAGE



Exercise can lead to the risk of athletes/sports people experiencing muscle damage. Repetitive, strenuous exercise can cause disruption to muscle fibers, causing them to become inflamed and possibly damaged [1]. Studies indicate that polymorphisms in the ACTN3, IGF2, TNFA, and IL6 genes can increase a persons risk of muscle damage following exercise. Gene variants that decrease CCL2 or CCR2 levels can also increase the risk of muscle damage and cause recovery to be more challenging [2].

People with your genetic profile are likely to have a regular risk of suffering muscle damage



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
CCL2	rs1860189	АА	
REFERENCES			

[1] Clarkson, P. M., & Hubal, M. J. (2002). Exercise-induced muscle damage in humans. American journal of physical medicine & rehabilitation, 81(11 Suppl), S52–S69.

[2] Hubal, M. J., Devaney, J. M., Hoffman, E. P., Zambraski, E. J., Gordish-Dressman, H., Kearns, A. K., Larkin, J. S., Adham, K., Patel, R. R., & Clarkson, P. M. (2010). CCL2 and CCR2 polymorphisms are associated with markers of exercise-induced skeletal muscle damage. Journal of applied physiology (Bethesda, Md.: 1985), 108(6), 1651–1658.



EXERCISE-INDUCED MYOPATHY

RESULTS



Myopathy is a disorder that affects people's muscular structure and metabolism, commonly causing them muscle pain, weakness, cramps and stiffness after/during exercise [1]. Several studies show genetic mutations associated with exercise-induced myalgia (muscle pain). Skeletal muscle cells require an energy supply to function and to move the body. The enzyme called Aadenosine Monophosphate Deaminase (AMD) comes from the AMPD1 gene. AMD is found in the muscles used for voluntary movements (skeletal muscles), which play a role in energy production. Alterations to AMD can, amongst other things, show up as exercise-induced muscle pain.

People with your genetic profile are likely to have a low tendency to experience myopathy



SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
AMPD1	rs17602729	GG	
REFERENCES			
[1] Nagy H, Veerapaneni KD. Myopathy. [Updated 2021 Jan 20]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2021 Jan			



SLOW MUSCLE REPAIR

RESULTS				
Bodies tend to naturally repair their injured muscles by making changes to the Extracellular Matrix (ECM) [1] - which includes things such as collagen and enzymes. The CD163 gene, along with anti-inflammatory white blood cells, are involved in facilitating the production growth factors (including the Insulin-like Growth Factor-1 (IGF-1)) which reduce muscle inflammation, prevent cell apoptosis, and promote muscle repair [2]. However, the INSIGF2 variation could increase the risk of muscle damage and increase the muscle recovery period [3]. People with your genetic profile are likely to have a good muscle repair ability				
	SCIENTIFIC DE	TAILS		
Gene	rsID	Genotype		
INS-IGF2	rs7924316	TT		
	REFERENCES			
 [1] Petrosino, J. M., Leask, A., & Accornero, F. (2019). Genetic manipulation of CCN2/CTGF unveils cell-specific ECM-remodeling effects in injured skeletal muscle. FASEB journal : official publication of the Federation of American Societies for Experimental Biology, 33(2), 2047–2057. [2] Peake, J. M., Neubauer, O., Della Gatta, P. A., & Nosaka, K. (2017). Muscle damage and inflammation during recovery from exercise. Journal of applied physiology (Bethesda, Md. : 1985), 122(3), 559–570. 				

[3] Baumert P, Lake MJ, Stewart CE, Drust B, Erskine RM. Genetic variation and exercise-induced muscle damage: implications for athletic performance, injury and ageing. Eur J Appl Physiol. 2016;116(9):1595-1625.



PREDISPOSITION TO THE DEVELOPMENT OF INGUINAL STRESS HERNIAS

RESULTS				
An inguinal hernia is when the contents of a person's abdominal peritoneum protrude out of a part of their groin (inguinal hernia) [1] and evidence suggests that this could be down to a lack of collagen regulation in the body [2]. People with indirect inguinal hernias seem to have lower collagen levels and have a reduced ratio of type I collagen to type III collagen. People with your genetic profile are likely to have a low predisposition for inguinal stress hernias				
SCIENTIFIC DETAILS				
Gene		rsID	Genotype	
EFEMP1		rs11899888	АА	
REFERENCES				
[1] Purkayastha, S., Chow, A., Athanasiou, T., Tekkis, P., & Darzi, A. (2008). Inguinal hernia. BMJ clinical evidence, 2008, 0412.,				

[2] Sezer, S., Şimşek, N., Celik, H. T., Erden, G., Ozturk, G., Düzgün, A. P., Çoşkun, F., & Demircan, K. (2014). Association of collagen type I alpha 1 gene polymorphism with inguinal hernia. Hernia : the journal of hernias and abdominal wall surgery, 18(4), 507–512.



STRESS FRACTURE PERIOD PREVALENCE

RESULTS			
Stress fractures often occur when people are involved in continuous physical activity; which is very common in athletes/sports people. According to research, the prevalence of stress fractures in elite athletes is between 14% and 21% and could be influenced by genetics. One study indicated that the presence of the Single Nucleotide Polymorphism (SNP) rs3018362 of RANK in the heterozygous and homozygous alleles is strongly associated with the prevalence of stress fracture periods in athletes/sportspeople [1].			
SCIENTIFIC DETAILS			
Gene	rsID	Genotype	
RANKL	rs1021188	TT	
REFERENCES			
[1] Varley, I., Hughes, D. C., Greeves, J. P., Stellingwerff, T., Ranson, C., Fraser, W. D., & Sale, C. (2015). RANK/RANKL/OPG pathway: genetic associations with			



STRESS FRACTURE

RESULTS				
	Stress fractures are bone f exercises such as running, time, making them more pr People with your genetic p	ractures occurring in healthy bones that have expensive sprinting, and jumping. The stresses trigger the ost one to stress fractures [1].	rienced stresses after repeated and prolonged eoclastic mechanism which weakens bones over ess fractures	
SCIENTIFIC DETAILS				
Gene		rsID	Genotype	
RPS6KA5		rs1286083	TT	
REFERENCES				
[1] Knapik, J. J., Reynolds, K., & Hoedebecke, K. L. (2017). Stress Fractures: Etiology, Epidemiology, Diagnosis, Treatment, and Prevention. Journal of special operations medicine : a peer reviewed journal for SOF medical professionals, 17(2), 120–130.,				

Biogenix Fitness Report

Kit ID: GFX0435522



OPTIMISM

RESULTS			
When a person is optimistic, they feel positive and hopeful for the future. People with high levels of optimism during challenging times often have effective coping mechanisms, and positive stimuli [1]. Optimism is a strong indicator of ones mental health which then positively affects their physical health. Recent studies suggest that levels of optimism could be inherited and that the genetic variation in the LINC01470 gene could be responsible. The OXTR gene that is linked to oxytocin production appeared to indirectly influence optimism and depression [2]. People with your genetic profile are predisposed to have a high sense of optimism			
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
LINC01470		rs4958581	ТС
REFERENCES			
[1] Carver, C. S., Scheier, M. F., & Segerstrom, S. C. (2010). Optimism. Clinical psychology review, 30(7), 879–889.			
[-]			

[2] Rana, B. K., Darst, B. F., Bloss, C., Shih, P. A., Depp, C., Nievergelt, C. M., Allison, M., Parsons, J. K., Schork, N., & Jeste, D. V. (2014). Candidate SNP associations of optimism and resilience in older adults: exploratory study of 935 community-dwelling adults. The American journal of geriatric psychiatry : official journal of the American Association for Geriatric Psychiatry, 22(10), 997–1006.e5.
Biogenix Fitness Report

Kit ID: GFX0435522



LEADERSHIP

RESULTS			
Â	Leadership is a position of effectively guide others. Lea relationship of the one who genetic components involv People with your genetic p	responsibility, requiring skill, consistency, and certair adership is a valued skill in team sports as well as in o occupies the highest position in an organised soci ed in the transmission of the neuronal signal [1]. rofile are likely to have higher leadership abilities	n behaviors and attitudes; enabling people to the workplace. By leadership, we refer to the al structure. Leadership may be determined by
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
HTR2A		rs6311	СТ
REFERENCES			
[1] Vender R. J. (2015	5). Leadership: an overview. Th	ne American journal of gastroenterology, 110(3), 362	2-367.



CONSCIENTIOUSNESS



[2] A.Zeman, Consciousness, Brain, Volume 124, Issue 7, July 2001, Pages 1263-1289



EXTRAVERSION



[1] McCabe, K. U., & Fleeson, W. (2012). What is extraversion for? Integrating trait and motivational perspectiv extraversion. Psychological science, 23(12), 1498–1505.



INSUFFICIENCY OF EXPERIENTIAL LEARNING

RESULTS				
- COR	Experiential learning is a way of learning through experience and reflection; suitable in many environments [1]. People affected by cognitive disabilities do, however, tend to find this kind of learning practice rather challenging. DISC1 is a gene implicated in the regulations of cells, such as cell duplication and differentiation (when a cell changes from one type to another). If a persons DISC1 gene is not regulated properly, cognitive deficits can result. People with your genetic profile are likely to have a good ability for experiential learning			
SCIENTIFIC DETAILS				
Gene		rsID	Genotype	
DISC1		rs821616	AA	
REFERENCES				
 [1] Green A. J. (1995). Experiential learning and teachinga critical evaluation of an enquiry which used phenomenological method. Nurse education today, 15(6) 				



IMPULSIVITY

	RESULTS	
	Impulsivity is a trait associated with sudden and sometimes unclearly calculated decisions and responses to circumstances; rather than calm, mindful decisions where all factors are considered. Psychologists define impulsivity as when a person gives their behavior little to no forethought, they have a tendency to act on a whim, and they do not consider the potential consequences of their actions. Contributing factors to impulsivity could be motor activation, inattentiveness, and a lack of planning. Impulsivity can be an indicator of some psychiatric and cognitive disorders [1]. Impulsive actions tend to negatively impact a person's long-term goals. In genetics, it has been reported that the rs1406946 mutation in the HTR1E gene results in low protein expression and is associated with an increase in impulsivity.	
	People with your genetic profile are likely to have a lower tend	ency to act impulsively
SCIENTIFIC DETAILS		
Gene	rsID	Genotype
HTR1E	rs1406946	СС
REFERENCES		
[1] Bakhshani N. M. (2	014). Impulsivity: a predisposition toward risky behaviors. Interi	national journal of high risk behaviors & addiction, 3(2), e20428.



RESPONSE TO ANGER

RESULTS		
	Anger is a natural emotion but one which can be a serious obs are however treatments/therapies to help manage anger, some modifying how one interprets information, and finding ways to People with your genetic profile are likely to have a good ability	stacle in sports, negatively impacting focus and energy levels. There e of which include making changes to how one interacts socially, o desensitize oneself [1]. I to handle feelings of anger
SCIENTIFIC DETAILS		
Gene	rsID	Genotype
NRTN	rs1379868	AA
REFERENCES		
[1] Steffgen G. (2017). Anger Management - Evaluation of a Cognitive-Behavioral Training Program for Table Tennis Players. Journal of human kinetics, 55, 65–73.		



COMBATIVE PERSONALITY

RESULTS			
- All	Combative personalities are often associated with aggressive, hostile, angry and violent behaviors in people. People with these traits can have dysfunctional neural circuits responsible for regulating emotional control. On a biochemical level, the combative personality is associated with several enzymes, including the one encoded by the COMT gene; which determines levels of neurotransmitters like Dopamine and Serotonin. These neurotransmitters play a key role in determining people's reactions and moods. Since Serotonin is responsible for inhibition in the brain's Prefrontal Cortex (PFC), a deficiency of Serotonin can enable aggressive behaviors [1].		
	People with your genetic profile are likely to have a higher tendency to have a combative personality		
SCIENTIFIC DETAILS			
Gene		rsID	Genotype
COMT		rs4680	GA
REFERENCES			
[1] Pavlov, K. A., Chistiakov, D. A., & Chekhonin, V. P. (2012). Genetic determinants of aggression and impulsivity in humans. Journal of applied genetics, 53(1), 61–82.			



GLOSSARY		
ALLELE	An allele is a variant form of a gene that is located at a specific position, or genetic locus, on a specific chromosome. Humans have two alleles at each genetic locus, with one allele inherited from each parent.	
CHROMOSOME	A chromosome is a condensed thread-like structure of DNA that carries hereditary information, or genes. Human cells have 22 chromosome pairs plus two sex chromosomes, giving a total of 46 per cell.	
GENOTYPE	The genetic makeup of an individual organism. It may also refer to just a particular gene or set of genes carried by an individual. The genotype determines the phenotype, or observable traits of the organism.	
SNP	Single nucleotide polymorphisms, frequently called SNPs, are the most common type of genetic variation among people. A SNP is a variation in a single nucleotide that occurs at a specific position in the genome.	
GENE	Genes are the basic units of inheritance. They contain DNA and are present in two forms called alleles.	
ZYGOSITY	Zygosity describes whether you inherited one copy of this variant from one of your parents (heterozygous), or you inherited two copies from both of your partents (homozygous). Typically for pathogenic variants homozygosity cause a more severe form of the condition. In many cases, heterozygous variants do not lead to the condition becoming apparent in the patient (also known as a recessive condition) but do mean that the next generation is at risk of inheriting the condition (or themselves becoming a carrier).	
POLYMORPHISM	A polymorphism consists of the different forms or variants in which a certain stretch of DNA can occur.	