

Genetic Testing: The Book of Me

Type: Book Review

Received: June 22, 2026

Published: July 21, 2026

Citation:

Christina Rahm. "Genetic Testing: The Book of Me". PriMera Scientific Surgical Research and Practice 8.1 (2026): 290-300.

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Abstract

Personalized medicine, especially genetic testing, is one of the innovations that has transformed how individuals understand their genetic makeup and how it influences their wellbeing. Knowledge about genetic predispositions can help avoid certain illnesses and make suitable decisions regarding nutrition, exercise, and other aspects of living. This approach is more useful than the general model where the same treatment regimen is applied to all the patients involved. The DRC-Gx product line exemplifies this innovation, offering six specialized programs: The products are DRC-GxNutrient, DRC-GxPerform, DRC-GxSlim, DRC-GxRenew, DRC-GxCBD and DRC-GxCannabis. All of them are designed to address different health issues and fulfill individual requirements based on the genetic information.

The information that could be used contained in the genetic data cannot be overemphasized. The DRC products provide a quantitative assessment of the vast area of health, such as nutrient metabolism, exercise performance, body composition, and gastrointestinal function. For instance, DRC-GxNutrient is created for enhancing micronutrient utilization while DRC-GxPerform incorporates exercise techniques for enhanced performance. In the same manner, DRC-GxSlim offers recommendations on how to lose weight based on genetic makeup and DRC-GxRenew offers aspects to do with aging. Indeed, such knowledge may contribute to the creation of measures that would have a positive impact on the state of health and quality of life. The integration of the genetic testing in the PHM is a major advance to achieving maximum health and fitness for every individual according to DNA constitution.

Introduction

Genetic health analysis is a powerful concept for rethinking the principles of individual health from the genetic perspective. This field includes a number of techniques for measuring genetic risks that might affect health, nutrition, and athletic ability. While genetic health analysis examines the individual's potential genetic susceptibilities, it explains how an individual may derive nutrients, the best exercise regime, and other lifestyle matters (Guest et al., 2019). The developments in genetic testing technologies enabled the acquisition of significant amounts of data that provide molecular and mechanistic information to guide interventions based on an individual's specific genetic profile.

The program reveals the need for customized healthcare solutions in today's society. Conventional approaches adopted over the years have been standardized and thus do not consider the genetic differences between people; hence, they can be off by quite a margin. Genetic-based consumer health solutions make it possible to develop individualized plans for dieting, exercising, and healthy living. One learns one's chances of getting diseases, which would have been very hard to avoid had one not done a DNA test, and hence, one should do a DNA test to avoid diseases (Mahmoud et al., 2022). This targeted strategy is beneficial now that the public is growing more aware of the disadvantages of receiving broadly targeted health information.

Understanding Genetic Testing

Genetic testing can be defined as all the procedures employed in examining an individual's DNA, which is the chemical blueprint that determines the existence, growth, and operation of any living thing. These tests can reveal abnormalities of chromosomes and genes or proteins and are often classified into diagnostic testing, carrier testing, prenatal testing, newborn screening, and predictive testing. Newborn screening is performed on asymptomatic newborns to identify those who may have a genetic disorder. At the same time, prenatal testing is done to detect the presence of a particular genetic disorder in a fetus (Tiffon, 2018). Prenatal diagnosis determines the health condition of a fetus, while newborn screening is the identification of genetic disorders on the baby's birth. Carrier testing determines the likelihood of developing a particular disease at some point in life, a valuable tool in the organization of preventive medicine.

Understanding the history of genetic testing must tackle the historical past of genetic testing, starting from the discovery of the DNA structure in the 1950s. At the start of genetic testing, the only diseases being tested were those that were relatively rare, and the tests used were labor-intensive. Nevertheless, due to technology, the area has been enhanced (Guest et al., 2019). Decoded in 2003, the Human Genome Project was launched to create a human genome map, forming the base for most of today's genetic tests. Nowadays, next-generation sequencing platforms for the analysis of large volumes of genetic data are affordable, making genetic testing available to the general populace. These developments have not only enhanced the reliability and speed of the tests but have also opened new horizons in the field of molecular diagnostics, enabling the tests to be used in the developing field of pharmacogenomics, where the therapy can be designed as per the genetic markers of the individual.

Ethics is key in genetics utilization; nevertheless, some concerns include privacy of genetic information, job discrimination, healthcare discrimination with particular emphasis on insurance companies and psychological effect of receiving the test results. This means that genetic tests must be taken because people must be fully aware of the consequences of these tests, as well as the fact that they are limited in some ways (Guest et al., 2019). In addition, there is a discussion on how to approach incidental findings that can indicate a risk of a disease different from the reason for the test. As genetic testing technology is applied and integrated into healthcare systems today and in the future, addressing some of these ethical issues becomes crucial in serving the interest of the individual and protecting and promoting the rights of the patient who needs personalized health solutions.

The Role of Genetics in Health

Genes are the means through which health is determined because they control several bodily functions. Every person has specific genes and alleles that may lead to different diseases affecting how a person metabolizes food, and how his or her body reacts to diets and training (Tiffon, 2018). For example, genetic differences may impact the body's ability to absorb nutrients, produce hormones, or regulate inflammation. Knowledge of such gene factors enables people to adopt the correct lifestyle that suits them to reduce cases of diseases.

Through an analysis of current research, considerable genes linked to nutrition, performance, and weight control have been revealed. For instance, the FTO gene is linked to obesity and adiposity, and the MTHFR gene influences folate metabolism and, therefore, cardiovascular disease (Guest et al., 2019). Also, genes such as BCMO1 determine the biosynthesis of vitamin A from beta carotene, thus an individual's nutritional status. ACTN3 gene for example, is associated with muscle fiber type and size distribution and therefore, they are genetically coded to do well in certain types of sport (Meroni et al., 2020). Examining these genes, specific preventative measures can be proposed to improve an individual's diet and exercise routine according to specific genetic variation.

Scientific Basis of Genetic Testing

Capitalizing on the enhanced molecular science and technology, genetic testing has transformed into an indispensable tool in mapping human health for risk profiling according to conventional set standards like blood group and to identify particular and individual genetic markers (Jamshed et al., 2019). As for the program of DRC-GxRenew, there are several genetic markers that help to explain health and aging. (Claussnitzer et al., 2020). These genetic markers include skin aging (such as TYR, IRF4), fat loss reaction to cardio (such as ADRB2, LPL), and exercise intrinsic motivation (e.g., BDNF) (Dar & Rasheed, 2023). A particular marker is connected with physiological characteristics or susceptibilities, facilitating suggestions concerning nourishment, physical activity, and alterations in everyday practice directed at successful aging and the enhancement of quality of life.

Studies have continued to establish a link between genes and people's wellbeing. For example, research suggests that there are systematic differences between people's ability to process dietary components or respond to the acts of exercise. This relationship also points to the benefits of having individualized prevention strategies supported by genetic susceptibility (Mullins et al., 2020). Even more, one can use genes linked to metabolism, nutrient absorption, and physical performance to design weight loss programs, as well as exercise and dietary plans. Using this genetic information, people can make decisions that are concord with their biology.

As appealing as the ideas proposed by genetic testing might be, there are drawbacks to relying on DNA recommendations, and the reliability of such tests cannot be questioned. One major weakness is that the study of genetic factors does not consider the contribution of the environment, even though the environment significantly impacts health (Claussnitzer et al., 2020). Consequently, the genetic profile analysis alone can omit one factor, which can distort the results and provide a completely wrong picture of a given person's health condition.

However, it is also true that the predictive strength of some of these markers may not be consistent. For instance, some markers might have significant links with particular disease and characteristics while others might not (Guest et al., 2021). Such variability can lead to contradictions as regards the recommendations that are given based on genetic tests. Moreover, genetic information is best understood by professionals; poor understanding of the results might lead to incorrect decisions concerning one's state of health or, on the contrary, cause excessive concern about one's condition.

DRC-Gx Nutrient Analysis

DRC-Gx Nutrient analysis is an innovative genetic test project that is aimed at determining the subjects' ability of their body to assimilate the necessary micronutrients from the food they consume. From this analysis, it is possible to learn how different genetic changes affect the bodies' ability to absorb and metabolize various vitamins and minerals, which is important in designing a personal nutrition program (Dar & Rasheed, 2023). People may take foods with vitamins and minerals, but due to genetics, the body will not allow these nutrients to penetrate the cells. According to certain people's heredity, they can regulate their meals about specific nutrients and finally get the most of them.

The DRC-Gx Nutrient report highlights some of the micronutrients as consisting of vitamins; Vitamin A, B6, C, D, E, and B12 and minerals; Calcium, Iron, Magnesium and Zinc. Each nutrient is then evaluated for absorption tendency according to a specific genetic type. For example, people may have low propensities to synthesize and absorb Vitamin A and Vitamin B6 because of the differences in genes known as BCMO1 and NBPFF3. On the other hand, some people may have above-normal tendencies in Vitamin E and omega levels because of genetic imperatives that can be linked to ZPR1 and FADS1 (Dar & Rasheed, 2023). While body nutrient assimilation capacity is established for each nutrient, this comprehensive nutrition overview also reveals any possible deficiencies or overloads that may affect health.

Nutrient absorption and health are areas that have many consequences. Negligible nutrient absorption contributes to many diseases, such as weakened immune system, chronic diseases vulnerability, and reduced brain performance. Such deficiencies will include low levels of Vitamin D, which hampers bone health and could lead to osteoporosis and inadequate Vitamin B12 absorption, which

causes anemia and neurological complications (Prudent, 2019). Genetic profiling of the particular nutrient assimilation patterns allows people to avoid such risks by modifying their diet or using special supplements.

Moreover, the assessment of the nutrient profile of the DRC-Gx underlines the need to consider genetic factors that affect one's response to dietary modification. For example, a person with a low aptitude for Vitamin A absorption will benefit from eating food products rich in easily absorbed Vitamin A or integrating himself with special supplements (Guest et al., 2021). Likewise, persons with below-average iron absorption might require more iron-containing food combined with vitamin C to increase iron bioavailability. This means that people are enabled to make sound decisions that will suit their genetic endowment.

Besides proposing individual macronutrient portions, the DRC-Gx Nutrient analysis also emphasizes the interactions of micronutrients with one another. For instance, the lack of one nutrient impacts the processing of the other, so it is important to understand how they are processed for effective nutrition strategy formulation (Claussnitzer et al., 2020) since genetic information provides a clear picture of the deficits within the body, the best diets can be formulated to address the shortfalls while considering the interaction between the nutrients.

Regarding genetic research, the DRC-Gx Nutrient analysis is a major step forward in individualized nutrition. It accomplishes this with the help of genetics and dieting facts that offer practical recommendations to help everyone achieve better health according to their genetic makeup (Jamshed et al., 2019). Significantly, everybody can improve their health and prevent diseases associated with improper digestion of their bodies' key vitamins and minerals.

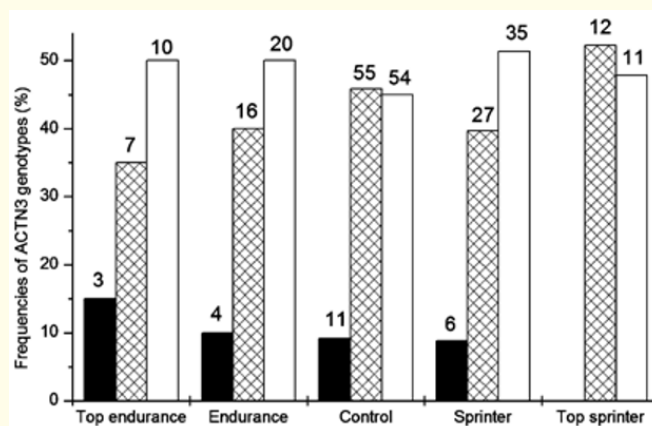
The DRC-Gx Nutrient analysis can be helpful to anybody interested in changing their nutrient status for the better because of understanding their genetics (Claussnitzer et al., 2020). This analysis reveals unique patterns concerning micronutrient assimilation. It thus provides specific direction regarding diet modifications or supplementation, which means that this analysis helps people to become active participants in the management of their health. Moving on to further revelations on the connection between genetics and diet, more tools such as the DRC-Gx Nutrient analysis will significantly be in the future management and optimization of health and wellbeing.

DRC-Gx Perform: Athletic Performance Insights

The DRC-Gx Perform program is the new face of DNA-based athletic performance that is aimed to help the athletes to develop health and performance management strategies based on their genes. The program is very informative as it employs DNA sequences to predict several training attributes, namely muscle mass, cardio, fat loss, and metabolism. This makes it easy for athletes to understand how their inherent traits can predict their training and performance (Guest et al., 2021). The program stresses that the genes are immutably fixed while changing the training and nutrition based on these findings will yield improved athletic outcomes.

Another factor that also contributes to the definition of an athlete in relation to muscularity and cardiovascular skillfulness is heredity. For instance, differences in the ACTN3 gene influence the muscle fiber type and an athlete's strength and power. Some types of people can possess more percent of fast twitch contractile muscle fibers, which is suitable for sprinting or weightlifting (Vesnina et al., 2020). In addition, variants of genes regulating cardiovascular adaptation to exercise training such as the ACE insert/deletion polymorphism and NOS3 are likely to affect oxygen consumption during endurance exercise. With this knowledge, athletes can make their schedules for training that involve the genetics they have including the weak ones.

The DRC-Gx Perform program also explains fat loss and nutrient intake based on the genetic make-up. For example, individuals with reduced fat utilization due to certain gene variants will have to correct the macronutrient ratios, or exert extra effort when aiming at achieving (Chen et al., 2021). However, those who have enhanced protein utilization can join well developed strength training programs that propel the muscles and aid in their repair. This means that athletes are able to gain the training goals set in the best way possible due to the unique plan on both diet and training.



Mitochondrial DNA and ACTN3 genotypes in Finnish elite endurance and sprint athletes

Aside from the somatic physical components, the DRC-Gx Perform program integrates the psychological factors like the motivation to exercise. BDNF biochemical measures, for example, tend to be higher in individuals with higher self-generated goals. This insight can be used by athletes when developing their training schedule or when creating personal goals because most athletes are more inclined to motivation. Knowledge of psychological profiles can improve participants' compliance with training initiatives and improve performance.

Recovery is another important goal area captured by the DRC-Gx Perform program. Predispositions to specific genes can determine the rate of an athlete's recovery from rigorous training or injuries (Guest et al., 2021). For instance, people's recessive genes determining the inflammatory response to injury might affect the duration it will take to recover. By identifying these factors, athletes can effectively adopt recovery protocols that are genetically based to include periods of rest or some form of treatment that would improve LTSP of athletes.

Additionally, the program provides advice on how to prevent injuries as informed by one's genetic makeup. Knowledge of which genes make athletes more susceptible to injury helps athletes avoid exacerbating the problem in training schedules (Guest et al., 2021). This may entail altering the frequency and type of exercise, adding preventive exercises or exercises that target parts of the body that may have been genetically predisposed to muscle injuries. The genetically personalized injury prevention approach enables athletes to minimize the risks of episodes that may slow their progress.

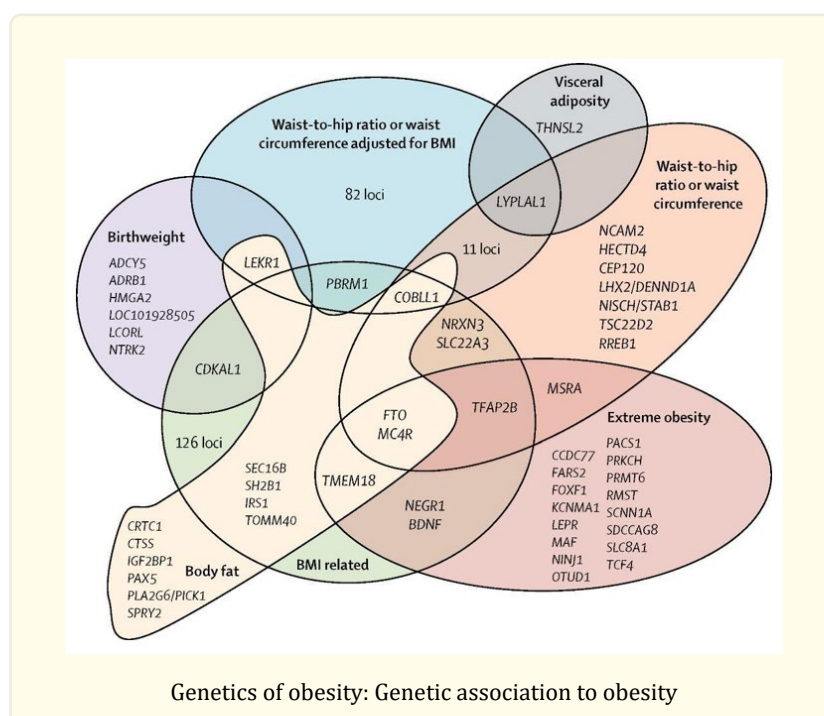
The DRC-Gx Perform program is quite advanced in adaptive athletic training, given its incorporation of genetic factors into performance enhancement plans. This program gives athletes the ability to understand muscle-building potential, cardiovascular fitness level, fat burn, recovery rate, and psychological factors based on one's genes to personalize their training programs (Prudent., 2019). As the research in genetics progresses within the field of sports science, programs such as the DRC-Gx Perform will be the foundation for the development and, potentially, the management of the future athlete as it relates to their health and selected sports.

DRC-Gx Slim: Weight Management Strategies

The DRC-Gx Slim is a plan for weight loss based on a person's genetic predisposition, and it offers a customized approach in terms of dieting and exercising. DRC-Gx Slim, as an example of a weight loss program that includes specific dietary advice based on an individual's genotype rather than relying on conventional health advice, trends or dieting. This approach makes sure that the strategies do not just work on a guesswork basis; it is actually founded on scientific knowledge on genetic aspects that can affect metabolism, appetite control and weight loss (Hibberd et al., 2019). The program's goal is to provide people with the only solution to diet and exercises they

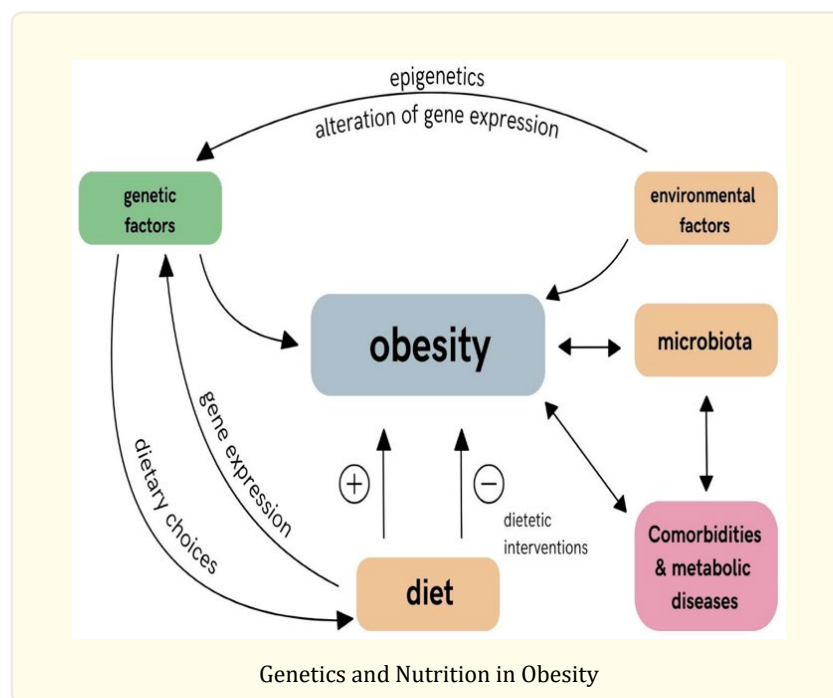
will definitely require throughout their life through DNA analysis.

Heritability of weight loss and body fat response is high and complex. Science has revealed different genes related to how people metabolize food changes or exercise programs. For example, the FTO and TCF7L2 genes are linked with body fat distribution and the ability to pack pounds. Some people have specific versions of these genes, and those with such versions can be less inclined or less likely to shed some extra pounds and sustain a healthy weight (Chen et al., 2021). Moreover, the LEPR gene, as well as CHRNA3, and CRY21 genes can influence how many calories are burned and how well metabolic pathways control hunger signals. Knowing such genetic inclinations of clients, the DRC-Gx Slim program can then make further appropriate recommendations to improve their weight loss endeavor.



The DRC-Gx Slim program offers individual diet plans based on people's genetic compatibility with specific nutrients and how those nutrients are processed in the body. For instance, people with a propensity for gluten sensitivity or lactose intolerance are likely to be advised to exclude gluten-containing products or dairy products as they may cause digestibility problems and interfere with the assimilation of nutrients. Additionally, the program measures the subject's preference for sweets and his/ her bitterness sensitivity threshold, which can help select foods that he/she would likely take with ease in adherence to the program's nutritional guidelines (Mahmoud et al., 2022). Thus, the program tries to make people follow the given dietary plans more consistently, as it addresses genetics, which might interfere with weight loss efforts.

Besides the eating plans, the DRC-Gx Slim program has been developed based on the importance of exercises according to genetic predispositions. The information refers to such aspects as an ability to determine genetic predisposition to physical activity in an individual, as well as to understand how this individual is going to react in a case if he/she begins to exercise, choosing any type of activity (Chen et al., 2021). For example, some people might have genes that mean they will burn fat more effectively during cardiovascular training. In contrast, others might gain more from weight training because of their genes related to muscle composition. In this way, the program offers exercise recommendations, which considers these genetic aspects to provide improved results.



Furthermore, the health impact assessment of the DRC-Gx Slim program also includes micronutrient status. Knowledge of how genetics influences micronutrient metabolism might identify a lack of important nutrients that could slow weight loss or impede overall wellbeing. Thus, low nutrient scores for some vitamins will require supplementation or a particular food plan that helps the body meet the nutritional requirements (Hibberd et al., 2019). This holistic strategy is appropriate for the weight issue and the overall health issue, the root cause of which may be nutritional problems.

The consequence of applying gene knowledge to weight loss is severe. These strategies, developed according to a person's genetic characteristics, bring better outcomes than regular, conventional methods could ever hope to achieve. This approach facilitates an understanding of a person's body and requirements, and assists in making lasting alterations in contrast to temporary fixes as are evident in diets (Vesnina et al., 2020). Through the DRC-Gx Slim program, people can manage their health by interacting with their genetic profiles highly scientifically.

This work is a step towards the positive evolution of the DRC-Gx Slim program as it introduces genetic analysis into the theoretical models of Weight Management Plans. This program has value since it incorporates a person's genetic profile in making recommendations about diet and exercise routines, something that most programs do not. As facts are still being discovered on how genetics is involved in the regulation of people's health and obesity, programs such as DRC-Gx Slim are expected to play a more significant role in the future as they provide long-term solutions for controlling obesity.

DRC-Gx Renew: Gut Health and Microbiome Support

Probiotics are more and more often considered as a key to maintaining good health, impacting not only the gastrointestinal tract but also other aspects of a person's health. The human stomach or gut contains trillions of microorganisms known as the microbiome responsible for regulating internal conditions. A healthy gut helps digestion and absorption of nutrients and can support elemental body immunity (Hibberd et al., 2019). On the other hand, an unhealthy bacteria profile will result in a myriad of diseases, for example, gastrointestinal disorders, autoimmune, and mental disorders such as anxiety and depression (Lee et al., 2020). Therefore, getting it right with the gut must be a priority as it determines one's overall health.

The microbiome plays a vital role in the absorption of nutrients and regulating inflammation in the body. It helps in the digestion of starches, the production of vitamins and the assimilation of minerals. For example, some of the gut microbiota that derive energy from fermented dietary fibers to generate SCFAs that have proline stimulating properties and immunomodulating effects and are critical for gut lining health. Any imbalance in the normal microbial flora interferes with these processes, thus causing malabsorption of nutrients and elevated systemic inflammation (Lee et al., 2020). Chronic inflammation that affects the body as a whole is associated with obesity, diabetes, and cardiovascular diseases. Hence, it is important to avoid the cause of inflammation in order to have a healthy microbiome.

The DRC-Gx Renew program, therefore stresses the need to appreciate the body's genetic structure when dealing with gut health. People also have different gene expressions that determine their reactions to foods, the vulnerability of their gut, and the nature of host microbiota. For example, some genes may point out that a person is gluten- or lactose intolerant, which affects the gut when not well regulated. By understanding these genetic elements, a person can get more information about what may be required to maintain the optimum conditions in the person's gut.

Some examples of the popular approaches to enhance gut health based on genetics are genetically personalized nutritional changes. For instance, a person with genetic traits that require them to avoid gluten will be healthy if they avoid foods containing gluten to avoid inflammation of the gut (Nichols & Davenport, 2021). Thus, people with lactose intolerance should avoid consuming dairy products or take products without lactose to prevent stomach upset. Furthermore, pre-and probiotic foods should be included in the diet to restore the microbiota's balance. Prebiotics are food elements that nourish good bacteria, whilst probiotics are live forms of bacteria that improve gut microorganisms.

Apart from modifications in diet, other aspects of people's lives help enhance gut health. Physical exercise is understood to positively affect the composition of the microbiota and the proper functioning of the gut (Lee et al., 2020). Other forms of relaxation, like guided breathing or even practice of yoga is also helpful since stress is said to have adverse effects on the gut by altering the microbiota and increasing permeability. The DRC-Gx Renew program encourages people to change their diets and lifestyles according to their genetic results.

Furthermore, the program also recognizes the importance of having periodic assessments of the actions to take in the process. It is important to understand that gut health is dynamic and may alter based on what a person eats, level of stress, and whether they are on certain medications or have been exposed to toxins (Hibberd et al., 2019). It is also recommended to perform the examination of gut health from time to time using genetics or any other method that may show how effective taken measures are. The principle of the cycle is that it is possible to make an endless number of tiny changes to one's diet and lifestyle until everything is ideal for their guts.

DRC-Gx CBD and Cannabis: Therapeutic Applications

The DRC-Gx CBD and Cannabis program provides a detailed report on the genetic makeup of an individual and how it determines his or her response to cannabis and CBD. With the legalization of cannabis for medical as well as recreational purposes steadily gaining traction across the globe, it will be important in the future to know what predisposes a body to specific outcomes regarding cannabis (Kaplan et al., 2022). This program offers information about several aspects concerning cannabinoid genetics - THC metabolism, memory, impulse behavior, pain and stress tolerance, and sleep quality (Almada et al., 2020). Such an approach will enable users to base their consumption of cannabis on their genetic makeup.

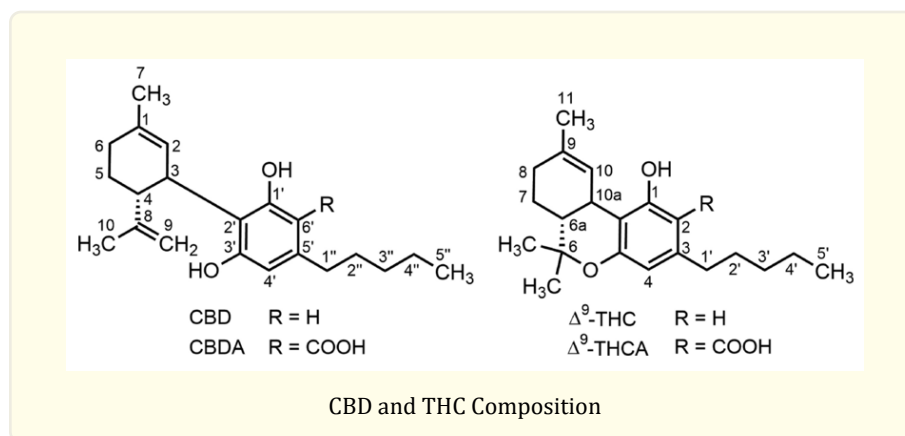
THC and CBD interact with the body's endocannabinoid system (ECS), a regulatory network that helps maintain balance across the nervous, immune, and digestive systems. THC binds directly to cannabinoid receptors, particularly CB1, influencing processes like mood, appetite, and pain perception, while CBD works indirectly by modulating receptor activity and supporting the body's own endocannabinoid signaling. Together, they influence how the body maintains physiological balance without acting as nutrients themselves. This complex biological system helps the body maintain homeostasis, or balance, in multiple physiological functions including the nervous, immune and endocrine systems, and the glycemic index. It shows that genetic makeup plays a huge role in determining how carcinogenic substances are metabolized and how effective they are in treating a given ailment (Leinen et al., 2023). For example,

different versions of the CYP2C9 gene can cause people to metabolize THC more quickly or more slowly, which will affect the length of time that the effects of the drug will last and the strength of the effects. Knowledge of such genetic factors can assist people in optimizing the dosing of cannabis to reap optimal therapeutic effects with minimal side effects.

There are many different potential benefits of CBD and cannabis, and they too may differ depending on an individual's genetic make-up. For instance, pain tolerance induced by some genetic imprint may find better relief in chronic pain conditions under cannabis products. Likewise, it will be wise for those with genetics that put them at risk of more severe cognitive decline to use THC in moderation to avoid worsening any memory or concentration problems (Leinen et al., 2023). Through such information about these tendencies in genetics, the DRC-Gx CBD and Cannabis program helps users choose products and concentrations that are appropriate for them.

Also, the program covers typical Wellness issues like stress and sleep. Some people genetically are better prepared to handle stress or respond to calming molecules such as cannabinoids found in CBD. For example, people with specific genotypes that affect stress response will receive more advantages from specific formulations of cannabinoids known to provide relaxation without a high (Almada et al., 2020). Furthermore, learning about one's heritability of sleep quality would enable people to make the right decision regarding cannabis varieties or any product that will enable them to have quality sleep.

With the increasing number of people using cannabis for medical reasons, it is critical to weigh the cost and advantage of the substance. The DRC-Gx CBD and Cannabis program, therefore recognizes that whilst genetics offer key information as to how an individual might react to cannabis, diet and exercise, and overall health also contribute significantly to the results (Kaplan et al., 2022). Thus, integrating genetic findings with other approaches to health improvement can help to increase the efficiency of using marijuana as a therapeutic resource.



Ethical and Legal Considerations

The issue of ethics and laws is vital in determining how genetic testing should be conducted and who should gain access to the results so that people's rights and privacy are protected, while at the same time, advancing the field of knowledge. This means that it is a requirement in genetic testing that the subjects know what will happen to them, the consequences, and the use of their genetic information. In this case, testing should be accompanied by information sharing about the pros and cons of such testing and the likelihood of coming across other unrelated but significant findings. Also, insisting on informed consent has a positive effect on the subject, healthcare personnel, and genetic testing services.

Another aspect of the ethical implications of genetic testing relates to data privacy. Genetic data is personal information that should not be disclosed or shared with anyone without the consent of the owner. Organizations planning to engage in genetic tests must ensure that they have appropriate mechanisms to protect this data. Moreover, individuals must have the ability to share or transfer the

rights to their Genomic data and also, the right to access; modify or delete information stored in Genomic data (Prudent, 2019). That is because data sharing, storage duration and third party access involves passing information to third parties and consequently clear policies regarding such areas can reduce risk and safeguard individuals against harm that may be occasioned by data breach or misuse.

The use of genetic tests has been faced with a major issue associated with genetic discrimination that should be tackled. This is the act of giving out a negative treatment on a person or refraining from offering the person services due to the person's genetic makeup as presented by the results of tests such as employment and insurance. In order to regulate these challenges, various legal acts have been adopted (Vesnina et al., 2020). For example, in the United States, there is the Genetic Information Non-Discrimination Act (GINA) that prohibits discrimination of people in issues regarding health insurance and employment on grounds of genetic information. However, there are still gaps in protection from discrimination in those areas such as life insurance or long-term care insurance and, therefore, it is necessary to have more sophisticated legal frameworks governing all these sectors.

Problems that arise in connection with the legal regulation of this sphere are constantly evolving, due to new technologies and greater understanding of genetics. The Food and Drug Administration (FDA) and other similar organizations have a significant role in monitoring the genetic tests sold directly to consumers to check the tests' validity and credibility. However, questions still arise regarding comparing the testing methods and results obtained (Nichols & Davenport, 2021). They pointed out that the absence of standard protocols increases the possibility of inconsistency in the development and implementation of the tests and the interpretation of the results across the various providers. Therefore, there is a need for continuous dialogue between stakeholders, such as policymakers, healthcare providers, and consumers, to develop clear standards to encourage proper ethics even as advanced genetic testing technologies are developed.

Future Directions in Genetic Health Analysis

New tendencies already at the forefront of genetic testing technologies are gradually transforming the sphere of health assessments and individualized medicine. The improved protocols in NGS to reduce both the costs and the time required to perform a comprehensive genomic profile have allowed more people to get broader information about them. Advanced technologies like CRISPR and whole-genome sequencing have opened up new avenues of precise genetic alteration and understanding of the multiple gene effects (Prudent, 2019). Furthermore, the use of AI in the assessment of genetic information is improving the influence of such findings by better predicting how specific genes affect an individual's health. These innovations are not only bringing genetic testing to a more accessible level but also providing more pertinent information to launch customized health interventions based on one's genetic makeup.

The possibility of using genetic testing in the framework of personalized medicine in the future is truly impressive. Since genetic information gives a broad view of an individual's susceptibility to certain diseases, healthcare providers can develop tailor-made treatment regimens for the patients (Nichols & Davenport, 2021). For instance, pharmacogenomics, which deals with how genes influence a particular individual's response to drugs, can help avoid drug prescriptions that result in side effects by recommending drugs that the patient would best respond to, given his or her genetic makeup. Additionally, as further studies are made to associate given genes with specific diseases, genetic tests would provide early identification and prevention measures for diseases like cancer, diabetes, and cardiovascular diseases. This approach could have significantly benefited patients based on their genetic profile from early intervention.

Nevertheless, there are several issues and possibilities in the further development of genetic health analysis, which can be considered insufficiently investigated and deserving further research. One of the main problems is the possible legal and ethical issues connected with the genetic databases, which refer to privacy issues, data protection, and genetic discrimination. Adequate protection of people's genetic data must be upheld to promote the public uptake of genetic testing services. Further, the concern for the quality of service to the patient requires standardization of the tests conducted and the interpretation of the results that different laboratory and service providers give. Opportunities: There is a constant trend in the preference for personalized health, implying a large market

for genetic testing services. With the continually growing knowledge about the genetic components of health, people are now looking for personalized preventive health programs that incorporate genetics into their daily routines.

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