VIOME



CHARLES WARDEN'S RESULTS

\'IOME

Dear Charles Warden,

The information on this report is for educational and informational use only. The information is not intended to be used by the customer for any diagnostic purpose and is not a substitute for professional medical advice. You should always seek the advice of your physician or other healthcare providers with any questions you may have regarding diagnosis, cure, treatment, mitigation, or prevention of any disease or other medical condition or impairment or the status of your health.



Test Name: Gut Intelligence Test

Authorized Order Person: Charles Warden

Customer Name: Charles Warden

DOB: 04/05/1985 **Gender:** Male

Customer Id: e16bdd01 **Sample Source:** Fecal

Date Collected: 03/11/2021
Date Received: 03/17/2021
Date Issued: 04/14/2021

Sample ID: 164CAFE3C491

Test Name: Human Gene Expression Test **Authorized Order Person:** Charles Warden

Customer Name: Charles Warden

DOB: 04/05/1985 **Gender:** Male

Customer Id: e16bdd01 **Sample Source:** Blood

Date Collected: Not Available

Date Received:

Date Issued: 04/14/2021

Sample ID: 23F498D42FE1



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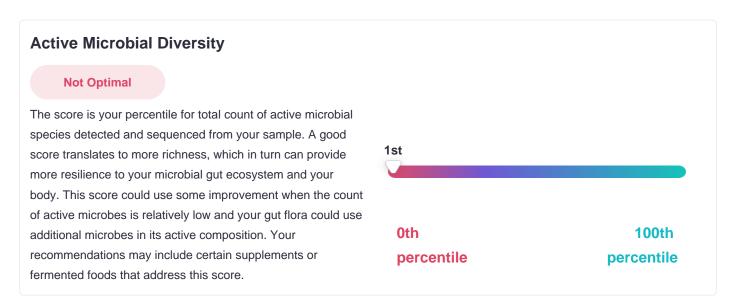
Lab Contact: https://support.viome.com (505) 672-5785

Customer Name: Charles Warden

DOB: 04/05/1985

All My Scores

Let's improve these.



Active Microbial Diversity Key

Reportable Range -13.6 to 8.53

Reference Ranges:

- Not Optimal -13.6 to -2.77 combined metric represents 0 to 5th percentile of the Viome population
- Average -2.76 to 2.44 combined metric represents 6th to 94th percentile of the Viome population
- Good 2.45 to 8.53 combined metric represents 95th to 100th percentile of the Viome population

Learn more by reading our references: https://viome.com/referenceresults



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Customer Name: Charles Warden

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Sulfide Gas Production Pathways

Not Optimal

This score assesses the levels of activity of all microbial pathways that result in the production of hydrogen sulfide gas. It can be made from some proteins that contain sulfur amino acids or from ingested sulfate or sulfite molecules found in foods like dried fruit, preserved meats, and some alcoholic beverages. This kind of activity, when high, contributes to proinflammatory patterns potentially harmful to the gut lining, as well as slowing of your motility (moving the food down your digestive tract). A good score means that the activity of sulfide production pathways is low.



Sulfide Gas Production Pathways Key

Reference Ranges:

- Not Optimal Represents 28% of the Viome population
- Average Represents 60% of the Viome population
- Good Represents 12% of the Viome population

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Putrescine Production Pathways Not Optimal This score assesses the levels of activity of all microbial pathways that lead to putrescine production. Putrescine is a molecular byproduct of protein fermentation - a microbial breakdown of protein. If the activities of putrescine production pathways are too high, it can be harmful to the gut environment and the intestinal barrier lining. It is also one of the signs that you may be eating too much protein that may

Putrescine Production Pathways Key

Reference Ranges:

not be digested properly.

- Not Optimal Represents 25% of the Viome population
- Average Represents 62% of the Viome population
- Good Represents 13% of the Viome population

Learn more by reading our references: https://viome.com/referenceresults

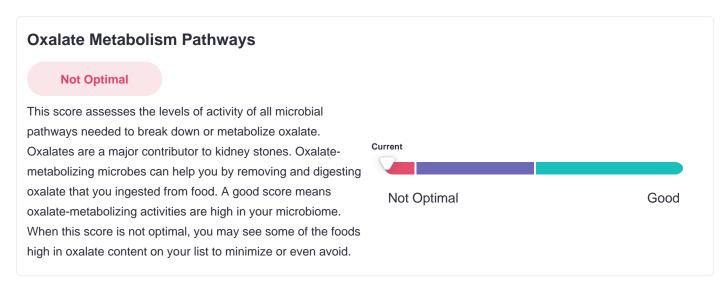


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Customer Name: Charles Warden

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Oxalate Metabolism Pathways Key

Reference Ranges:

- Not Optimal Represents 79% of the Viome population
- Average Represents 14% of the Viome population
- Good Represents 7% of the Viome population

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TMA Production Pathways Not Optimal This score assesses the levels of all activity of metabolic pathways that result in TMA production. TMA (trimethylamine) is a molecule that gets converted to TMAO (Trimethylamine Noxide) in the liver. TMAO is associated with unfavorable metabolic and cardiovascular effects. Since one of the substances used for microbial TMA production is choline, reducing high-choline-containing foods in the diet may be one of the options for improving this pattern. A good score means these TMA production pathway activity levels are low.

TMA Production Pathways Key

Reference Ranges:

- Not Optimal Represents 27% of the Viome population
- Average Represents 0% of the Viome population
- Good Represents 73% of the Viome population

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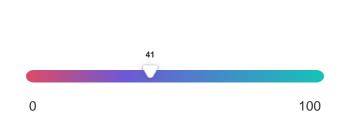
Customer Name: Charles Warden

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Gut Microbiome Health

Not Optimal

Your Gut Microbiome Health score integrates over 20 microbial functional scores. When this score is low it means that your gut microbiome may be producing chemicals that are causing inflammation (such as LPS, sulfide, or ammonia) or not producing enough nutrients that your body needs (such as butyrate, serotonin, and other vitamins). Our food and supplement recommendations are designed specifically for you to optimize your microbial functions and bring your gut microbiome into balance. Scroll down below to the section titled "How We Calculate This Score" to learn more. Did you know? In many ways, your gut bacteria are as vast and mysterious as the Milky Way. About 100 trillion bacteria, both good and bad, live inside your digestive system. Optimizing your microbial functions can help you achieve a healthy weight, boost energy, reduce stress, improve sleep, and strengthen your immunity.



Gut Microbiome Health Key

Reference Ranges:

- Not Optimal 0 to 43 which represents 17% of the Viome population
- Average 44 to 54 which represents 71% of the Viome population
- Good 55 to 100 which represents 12% of the Viome population

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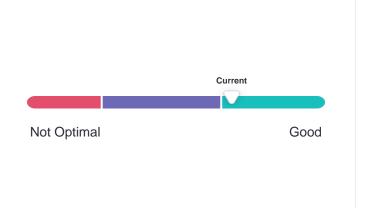
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Ammonia Production Pathways

Good

This score assesses the levels of activity of all microbial pathways that result in the production of ammonia. Ammonia gas can be made from amino acids as a byproduct of the breaking down of protein or from ingested nitrate or nitrite molecules found in things like food preservatives or additives, preserved meats, and dried fruit. This kind of activity, when high, contributes to pro-inflammatory patterns potentially harmful to the gut lining, as well as slowing of your motility (moving the food down your digestive tract), and is also one of the signs that your proteins may not be digested properly. A good score means that the activity of ammonia production pathways is low.



Ammonia Production Pathways Key

Reference Ranges:

- Not Optimal Represents 24% of the Viome population
- Average Represents 47% of the Viome population
- Good Represents 29% of the Viome population

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Good This score assesses the levels of activity of all microbial pathways that lead to the production of uric acid (or urate). Uric Acid is a normal byproduct that comes from the breakdown of compounds called purines, which can be found in beer, sugary sodas, seafood and shellfish, turkey, veal, bacon, and organ meats. Excessive amounts of uric acid can contribute to gout. A good score means that your uric acid

Uric Acid Production Pathways Key

production pathway levels are low.

Reference Ranges:

- Not Optimal Represents 29% of the Viome population
- Average Represents 52% of the Viome population
- Good Represents 19% of the Viome population

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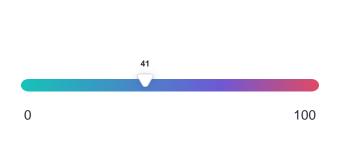
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Inflammatory Activity

Average

This score measures the activities of your microbes that can contribute to or reflect inflammation in your gut environment. Inflammation in your gut can be caused by harmful things your microbes produce when you are either inefficiently digesting your proteins, have excessive microbial gas production, or simply have a gut environment that your microbes perceive as threatening. A score in the red zone (not optimal) means that there are relatively more pro-inflammatory activities, as opposed to anti-inflammatory or protective ones. Everyone's pattern is unique, so if your score is in the red, some of your recommendations may focus on boosting more of the protective and healing anti-inflammatory functions, while others may focus more on controlling and balancing out the more harmful pro-inflammatory microbes and functions. Follow your recommendations to maintain a good range or improve this score.



Inflammatory Activity Key

Reference Ranges:

- Not Optimal 50 to 100 which represents 9% of the Viome population
- Average 36 to 49 which represents 69% of the Viome population
- Good 0 to 35 which represents 22% of the Viome population

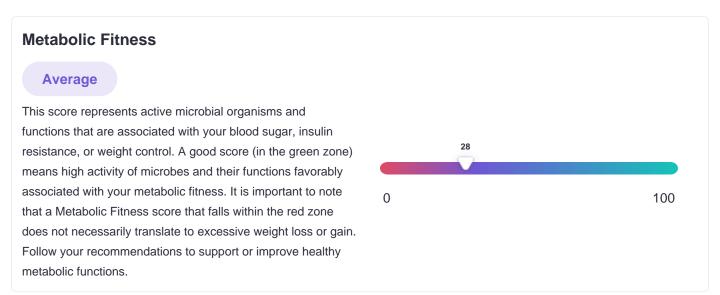
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Customer Name: Charles Warden

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Metabolic Fitness Key

Reference Ranges:

- Not Optimal 0 to 22 which represents 25% of the Viome population
- Average 23 to 30 which represents 58% of the Viome population
- Good 31 to 100 which represents 17% of the Viome population

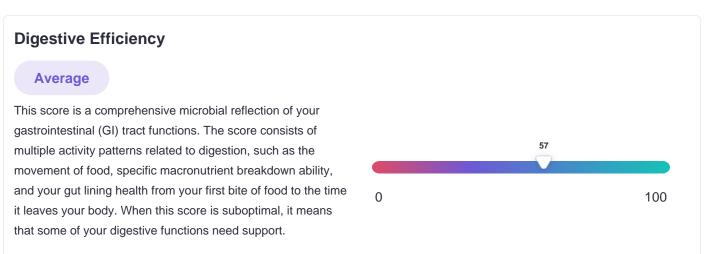
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Digestive Efficiency Key

Reference Ranges:

- Not Optimal 0 to 43 which represents 19% of the Viome population
- Average 44 to 64 which represents 63% of the Viome population
- Good 65 to 100 which represents 18% of the Viome population

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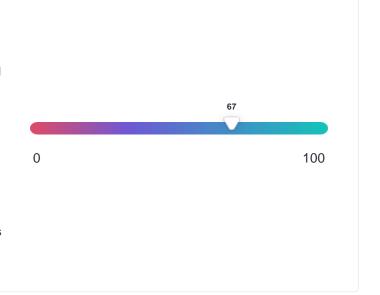
Customer Name: Charles Warden

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Gut Lining Health

Average

This score focuses on your gut lining (or intestinal barrier) and the health of the mucosal layer that protects it. When your gut lining is compromised, things from the outside environment, like toxins, medications, and harmful bacteria, can make their way into your bloodstream from your gut and negatively affect your immune system and overall wellbeing. A good score (in the green zone) means more optimal microbial functions that support your intestinal barrier and fewer disruptive or harmful functions are active in your gut. Follow your recommendations to address your specific pattern of microbial functions, and to prevent any intestinal permeability known as 'leaky gut'.



Gut Lining Health Key

Reference Ranges:

- Not Optimal 0 to 65 which represents 14% of the Viome population
- Average 66 to 77 which represents 65% of the Viome population
- Good 78 to 100 which represents 21% of the Viome population

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Protein Fermentation Average This score reflects whether or not you are digesting your proteins properly. Protein digestion begins when you first start chewing and continues down in your stomach. If the protein is not fully broken down through this process, your microbes will digest the excess protein available and may convert it into harmful byproducts. Overly high microbial protein fermentation translates into a score within the red zone, suggesting your

Protein Fermentation Key

protein digestion is suboptimal.

Reference Ranges:

- Not Optimal 65 to 100 which represents 26% of the Viome population
- Average 36 to 64 which represents 56% of the Viome population
- Good 0 to 35 which represents 18% of the Viome population

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Average This score is an assessment of your overall gas production activity by the microbes in your gut. Overall high microbial gas production has been associated with digestive difficulties, discomfort, and gut inflammation. A good score means that your microbes are not actively engaged in gas production functions.

Gas Production Key

Reference Ranges:

- Not Optimal 65 to 100 which represents 20% of the Viome population
- Average 36 to 64 which represents 65% of the Viome population
- Good 0 to 35 which represents 15% of the Viome population

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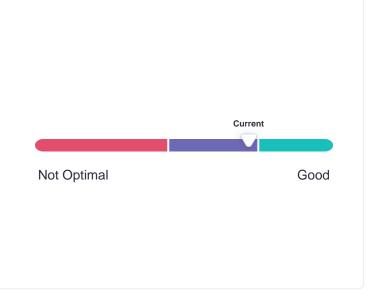
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Butyrate Production Pathways

Average

This score assesses the levels of activity of all microbial pathways that lead to the production of a beneficial nutrient -butyrate. Butyrate is a short-chain fatty acid known to beneficially affect many wellness areas from gut lining to insulin sensitivity and satiety (feeling full). A score that is not optimal means that your microbial butyrate production could really use a good boost! Individuals with low butyrate production activity would benefit from supplements or foods that either feed or add butyrate producing microbes into your gut ecosystem.



Butyrate Production Pathways Key

Reference Ranges:

- Not Optimal Represents 20% of the Viome population
- Average Represents 65% of the Viome population
- Good Represents 15% of the Viome population

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LPS Biosynthesis Pathways Average This score assesses the levels of activity of all microbial pathways leading to the production of LPS (lipopolysaccharides) in your gut. LPS is a pro-inflammatory molecule that gut microbes make, which can trigger your immune system response, especially if it passes to the bloodstream through the gut lining. This score is an important factor in assessing your inflammatory activity patterns.

LPS Biosynthesis Pathways Key

Reference Ranges:

- Not Optimal Represents 25% of the Viome population
- Average Represents 55% of the Viome population
- Good Represents 20% of the Viome population

Learn more by reading our references: https://viome.com/referenceresults

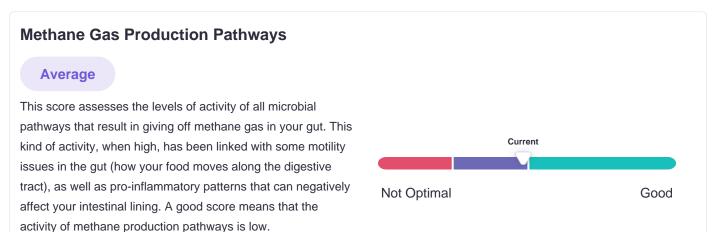


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Methane Gas Production Pathways Key

Reference Ranges:

- Not Optimal Represents 26% of the Viome population
- Average Represents 30% of the Viome population
- Good Represents 44% of the Viome population

Learn more by reading our references: https://viome.com/referenceresults



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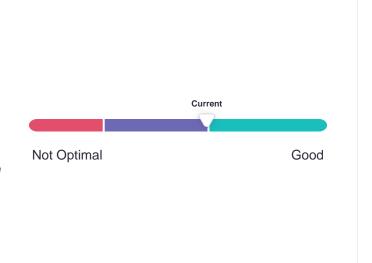
Customer Name: Charles Warden

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Flagellar Assembly Pathways

Average

This score assesses the levels of activity of all microbial pathways leading to the making of a structure called flagella. Flagellar structures serve as "fins" or "tails" for various microbes to help them move. A score that is not optimal suggests that these signaling pathway activities are high, indicating unrest in your microbiome as flagellar structures are helping beneficial organisms move away from a perceived threat. Higher than usual activity can also signal the presence of opportunistic organisms that are known to have these flagellar structures. This score is an important factor in assessing your inflammatory activity patterns.



Flagellar Assembly Pathways Key

Reference Ranges:

- Not Optimal Represents 24% of the Viome population
- Average Represents 55% of the Viome population
- Good Represents 21% of the Viome population

Learn more by reading our references: https://viome.com/referenceresults

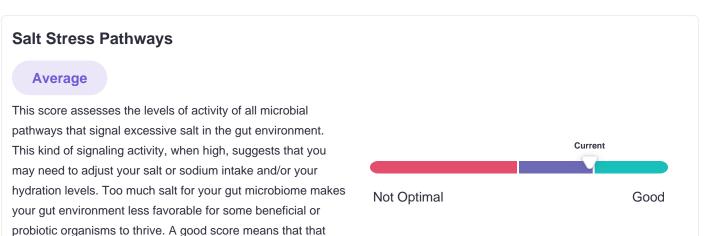


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Salt Stress Pathways Key

Reference Ranges:

Not Optimal Represents 11% of the Viome population

Average Represents 61% of the Viome population

pathway levels that signal microbial salt stress are low.

Good Represents 28% of the Viome population

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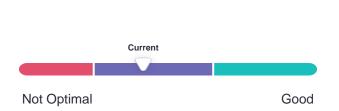
Customer Name: Charles Warden

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Biofilm, Chemotaxis, and Virulence Pathways

Average

This score assesses the levels of all activity of all metabolic pathways that suggest a pro-inflammatory or hostile environment in the gut. This includes virulence factors, biofilm formation, and chemotaxis signaling, which are all important parts of your overall inflammatory activity patterns. When this score is relatively high it means that there is some threat in the environment and your microbes are trying to either defend themselves, attack each other, or move. This type of a "microbial war zone" can negatively impact your gut environment, and some of the "bullets" secreted by the microbes may trigger an immune response. A good score means that these pathway activities are at low levels.



Biofilm, Chemotaxis, and Virulence Pathways Key

Reference Ranges:

- Not Optimal Represents 25% of the Viome population
- Average Represents 46% of the Viome population
- Good Represents 29% of the Viome population

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23 / 54

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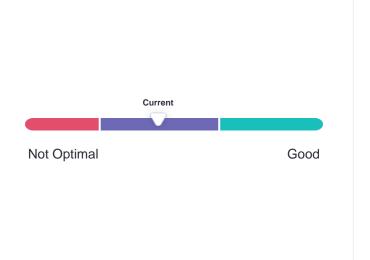
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Bile Acid Metabolism Pathways

Average

This score assesses the levels of activity of all metabolic pathways that include bile acids. Normally bile acids are made by the liver to help with fat digestion. Bile acids enter the colon in the form of bile salts. Your gut microbiota can change them back into bile acids, after which they can even be recycled back to the liver. If this activity is relatively high or excessive, it may be an indicator of your inability to break down fat or absorb nutrients properly, which can contribute to a proinflammatory environment or negative liver-related effects, as microbiome's bile acid pathways have been implicated in fatty deposits in the liver. A good score means these pathway activity levels are low in your sample.



Bile Acid Metabolism Pathways Key

Reference Ranges:

- Not Optimal Represents 31% of the Viome population
- Average Represents 49% of the Viome population
- Good Represents 20% of the Viome population

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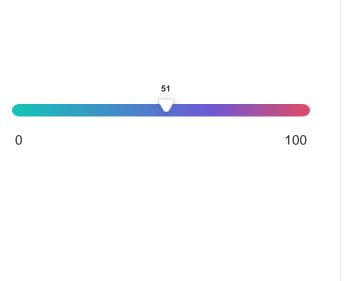
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Microbiome-Induced Stress Average

Your Microbiome-Induced Stress score offers insights about those microbial activities that can lead to stress or inflammatory response not only in your gut, but also in your body. Toxins and other molecules produced by the gut microbiome may enter the bloodstream and contribute to cellular stress and pro-inflammatory pathways throughout your body. If this score is not optimal, it may suggest that these microbial activities need to be mitigated by either suppressing them, balancing them out with beneficial and protective microbial activities, or by strengthening your gut lining to prevent them from crossing the gut lining and affecting the rest of your body.



Microbiome-Induced Stress Key

Reference Ranges:

- Not Optimal 61 to 100 which represents 20% of the Viome population
- Average 36 to 60 which represents 65% of the Viome population
- Good 0 to 35 which represents 15% of the Viome population

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Meet your probiotic microbes

These are microbes that are found in commercially available probiotic products that are also active in your sample. If there are no organisms listed, no probiotics were identified in your sample.

Bifidobacterium animalis subsp. lactis

P Probiotic

Lactobacillus delbrueckii subsp. bulgaricus

P Probiotic



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My Active Microbes





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Alistipes senegalensis JC50

B Bacterium

Alistipes shahii WAL 8301

B Bacterium

Alistipes sp. AL-1

B Bacterium

Alistipes sp. HGB5

B Bacterium

Anaerostipes caccae

B Bacterium

Anaerostipes hadrus strain BPB5

B Bacterium

Anaerostipes sp. 3_2_56FAA

B Bacterium

Anaerotruncus colihominis

B Bacterium

Anaerotruncus rubiinfantis

B Bacterium

Angelakisella massiliensis strain Marseille-P3217

B Bacterium

Atopobium

B Bacterium

Bacteroides acidifaciens

B Bacterium

Bacteroides barnesiae DSM 18169 = JCM 13652

B Bacterium



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Bacteroides caccae

B Bacterium

Bacteroides caecimuris strain 148

B Bacterium

Bacteroides cellulosilyticus strain WH2

B Bacterium

Bacteroides coprocola

B Bacterium

Bacteroides coprophilus

B Bacterium

Bacteroides dorei CL03T12C01

B Bacterium

Bacteroides eggerthii 1_2_48FAA

B Bacterium

Bacteroides faecichinchillae strain DSM

B Bacterium

Bacteroides faecis

B Bacterium

Bacteroides finegoldii

B Bacterium

Bacteroides fluxus

B Bacterium

Bacteroides fragilis

B Bacterium

Bacteroides fragilis str. 3-F-2 #6

B Bacterium



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Bacteroides helcogenes P 36-108

B Bacterium

Bacteroides massiliensis dnLKV3

B Bacterium

Bacteroides nordii

B Bacterium

Bacteroides ovatus strain ATCC

B Bacterium

Bacteroides plebeius

B Bacterium

Bacteroides sp. 1_1_30

B Bacterium

Bacteroides sp. 1_1_6

B Bacterium

Bacteroides sp. 2_1_22

B Bacterium

Bacteroides sp. 2_1_33B

B Bacterium

Bacteroides sp. 2_2_4

B Bacterium

Bacteroides sp. 3_1_13

B Bacterium

Bacteroides sp. 3_1_19

B Bacterium

Bacteroides sp. 3_1_23

B Bacterium



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Bacteroides sp. 3_1_33FAA

B Bacterium

Bacteroides sp. 3_1_40A

B Bacterium

Bacteroides sp. 4_1_36

B Bacterium

Bacteroides sp. 4_3_47FAA

B Bacterium

Bacteroides sp. 9_1_42FAA

B Bacterium

Bacteroides sp. D2

B Bacterium

Bacteroides sp. D20

B Bacterium

Bacteroides sp. D22

B Bacterium

Bacteroides sp. HMSC067B03

B Bacterium

Bacteroides sp. HMSC068A09

B Bacterium

Bacteroides sp. HMSC073E02

B Bacterium

Bacteroides sp. HPS0048

B Bacterium

Bacteroides sp. Marseille-P3108 sp. Marseille-P3108

B Bacterium



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Bacteroides sp. Marseille-P3208T strain Marseille-P3208

B Bacterium

Bacteroides stercoris ATCC 43183

B Bacterium

Bacteroides stercoris CC31F

B Bacterium

Bacteroides stercoris strain CL09T03C01

B Bacterium

Bacteroides stercoris strain DSM

B Bacterium

Bacteroides thetaiotaomicron VPI-5482

B Bacterium

Bacteroides thetaiotaomicron strain 7330

B Bacterium

Bacteroides uniformis

B Bacterium

Bacteroides vulgatus ATCC 8482

B Bacterium

Bacteroides xylanisolvens

B Bacterium

Bariatricus massiliensis strain AT12

B Bacterium

Bifidobacterium animalis strain A6

B Bacterium

Bifidobacterium animalis strain RH

B Bacterium



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Customer Name: Charles Warden

DOB: 04/05/1985

Bifidobacterium animalis subsp. lactis B Bacterium P Probiotic Bifidobacterium animalis subsp. lactis AD011 **B** Bacterium Bifidobacterium animalis subsp. lactis B420 **B** Bacterium Bifidobacterium animalis subsp. lactis BB-12 **B** Bacterium Bifidobacterium animalis subsp. lactis BLC1 **B** Bacterium Bifidobacterium animalis subsp. lactis Bi-07 **B** Bacterium Bifidobacterium animalis subsp. lactis BI-04 **B** Bacterium Bifidobacterium animalis subsp. lactis Bl12 **B** Bacterium Bifidobacterium animalis subsp. lactis CNCM I-2494 **B** Bacterium Bifidobacterium animalis subsp. lactis KLDS2.0603 **B** Bacterium Bifidobacterium animalis subsp. lactis V9 **B** Bacterium Bifidobacterium animalis subsp. lactis strain BF052 **B** Bacterium Bilophila sp. 4_1_30 **B** Bacterium



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Bilophila wadsworthia 3_1_6

B Bacterium

Bilophila wadsworthia ATCC 49260

B Bacterium

Blautia massiliensis sp. GD8

B Bacterium

Blautia obeum ATCC 29174

B Bacterium

Blautia obeum strain 2789STDY5608837

B Bacterium

Blautia obeum strain 2789STDY5608838

B Bacterium

Blautia obeum strain 2789STDY5834861

B Bacterium

Blautia obeum strain 2789STDY5834957

B Bacterium

Blautia sp. KLE 1732

B Bacterium

Blautia sp. Marseille-P2398

B Bacterium

Blautia sp. Marseille-P3087 sp. Marseille-P3087

B Bacterium

Blautia sp. Marseille-P3201T strain Marseille-P3201

B Bacterium

Blautia wexlerae

B Bacterium



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Burkholderiales bacterium 1_1_47

B Bacterium

Butyricicoccus desmolans ATCC 43058

B Bacterium

Butyricicoccus pullicaecorum

B Bacterium

Butyricimonas virosa

B Bacterium

Butyrivibrio crossotus DSM 2876

B Bacterium

Candidatus Stoquefichus sp. KLE1796

B Bacterium

Candidatus Stoquefichus sp. SB1

B Bacterium

Catabacter hongkongensis strain ABBA15k

B Bacterium

Christensenella timonensis

B Bacterium

Clostridia bacterium UC5.1-1D1

B Bacterium

Clostridia bacterium UC5.1-1D10

B Bacterium

Clostridia bacterium UC5.1-2F7

B Bacterium

Clostridia bacterium UC5.1-2H11

B Bacterium



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Clostridiaceae bacterium MS3

B Bacterium

Clostridiales bacterium

B Bacterium

Clostridiales bacterium KLE1615

B Bacterium

Clostridiales bacterium VE202-01

B Bacterium

Clostridiales bacterium VE202-03

B Bacterium

Clostridiales bacterium VE202-06

B Bacterium

Clostridiales bacterium VE202-07

B Bacterium

Clostridiales bacterium VE202-13

B Bacterium

Clostridiales bacterium VE202-15

B Bacterium

Clostridiales bacterium VE202-16

B Bacterium

Clostridiales bacterium VE202-21

B Bacterium

Clostridiales bacterium VE202-26

B Bacterium

Clostridiales bacterium VE202-27

B Bacterium



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Clostridiales bacterium VE202-28

B Bacterium

Clostridioides difficile P28

B Bacterium

Clostridium lentocellum DSM 5427

B Bacterium

Clostridium paraputrificum

B Bacterium

Clostridium phoceensis strain GD3

B Bacterium

Clostridium sp. AT4

B Bacterium

Clostridium sp. ATCC BAA-442

B Bacterium

Clostridium sp. DSM 4029

B Bacterium

Clostridium sp. HGF2

B Bacterium

Clostridium sp. KLE 1755

B Bacterium

Clostridium sp. L2-50

B Bacterium

Clostridium sp. M62/1

B Bacterium

Clostridium sp. Marseille-P3244 sp. Marseille-P3244

B Bacterium



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Coprobacillus sp. 3_3_56FAA

B Bacterium

Coprobacillus sp. 8_2_54BFAA

B Bacterium

Dialister invisus DSM 15470

B Bacterium

Dielma fastidiosa

B Bacterium

Dorea formicigenerans

B Bacterium

Dorea longicatena strain 2789STDY5608851

B Bacterium

Dorea longicatena strain 2789STDY5834914

B Bacterium

Dorea sp. 5-2

B Bacterium

Drancourtella massiliensis strain GD1

B Bacterium

Eggerthella lenta DSM 2243

B Bacterium

Eggerthella sp. 1_3_56FAA

B Bacterium

Eggerthella sp. HGA1

B Bacterium

Eisenbergiella tayi

B Bacterium



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Entamoeba

E Eukaryote

Enterobacter

B Bacterium

Enterococcus faecium NEF1

B Bacterium

Erysipelotrichaceae bacterium 5_2_54FAA

B Bacterium

Escherichia coli

B Bacterium

Eubacterium plexicaudatum ASF492

B Bacterium

Eubacterium ramulus

B Bacterium

Eubacterium sp. 3_1_31

B Bacterium

Eubacterium sp. SB2

B Bacterium

Eubacterium ventriosum ATCC 27560

B Bacterium

Faecalibacterium prausnitzii

B Bacterium

Flavonifractor plautii strain YL31

B Bacterium

Flintibacter

B Bacterium



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Fusicatenibacter saccharivorans

B Bacterium

Gemella sanguinis ATCC 700632

B Bacterium

Gokushovirus WZ-2015a

V Virus

Gordonibacter pamelaeae 7-10-1-b

B Bacterium

Gordonibacter urolithinfaciens

B Bacterium

Haemophilus sp. HMSC71H05

B Bacterium

Holdemania filiformis DSM 12042

B Bacterium

Holdemania massiliensis AP2

B Bacterium

Holdemania sp. Marseille-P2844 sp. Marseille-P2844

B Bacterium

Intestinimonas butyriciproducens strain AF211

B Bacterium

Intestinimonas massiliensis sp. GD2

B Bacterium

Klebsiella

B Bacterium

Lachnoclostridium sp. YL32 sp. YL32

B Bacterium



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Lachnospira pectinoschiza strain 2789STDY5834886

B Bacterium

Lachnospiraceae bacterium

B Bacterium

Lachnospiraceae bacterium 1_1_57FAA

B Bacterium

Lachnospiraceae bacterium 1_4_56FAA

B Bacterium

Lachnospiraceae bacterium 2_1_58FAA

B Bacterium

Lachnospiraceae bacterium 3_1_46FAA

B Bacterium

Lachnospiraceae bacterium 3_1_57FAA_CT1

B Bacterium

Lachnospiraceae bacterium 5_1_57FAA

B Bacterium

Lachnospiraceae bacterium 5_1_63FAA

B Bacterium

Lachnospiraceae bacterium 6_1_63FAA

B Bacterium

Lachnospiraceae bacterium 7_1_58FAA

B Bacterium

Lachnospiraceae bacterium M18-1

B Bacterium

Lactobacillus delbrueckii subsp. bulgaricus

B Bacterium P Probiotic





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Lactobacillus delbrueckii subsp. bulgaricus strain DSM

B Bacterium

Lactococcus

B Bacterium

Lactonifactor longoviformis DSM 17459

B Bacterium

Listeria

B Bacterium

Longibaculum

B Bacterium

Marvinbryantia formatexigens DSM 14469

B Bacterium

Marvinbryantia formatexigens strain I-52

B Bacterium

Massilioclostridium coli strain Marseille-P2976

B Bacterium

Mediterranea massiliensis strain Marseille-P2645

B Bacterium

Mogibacterium diversum

B Bacterium

Neglecta timonensis

B Bacterium

Odoribacter laneus YIT 12061

B Bacterium

Odoribacter splanchnicus DSM 20712

B Bacterium



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Oscillibacter sp. ER4

B Bacterium

Oscillibacter sp. KLE 1745

B Bacterium

Oscillospiraceae bacterium VE202-24

B Bacterium

Parabacteroides distasonis ATCC 8503

B Bacterium

Parabacteroides goldsteinii CL02T12C30

B Bacterium

Parabacteroides goldsteinii DSM 19448 = WAL 12034

B Bacterium

Parabacteroides gordonii DSM 23371

B Bacterium

Parabacteroides gordonii MS-1

B Bacterium

Parabacteroides johnsonii

B Bacterium

Parabacteroides merdae CL03T12C32

B Bacterium

Parabacteroides sp. 2_1_7

B Bacterium

Parabacteroides sp. D26

B Bacterium

Parabacteroides sp. HGS0025

B Bacterium



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Parabacteroides sp. Marseille-P3236 strain Marseille-P3236, sp. Marseille-P3136

B Bacterium

Parabacteroides sp. SN4 strain SN4, sp. SB4

B Bacterium

Paraprevotella xylaniphila YIT 11841

B Bacterium

Parasutterella excrementihominis YIT 11859

B Bacterium

Pepper mild mottle virus



Phocea massiliensis strain Marseille-P2769

B Bacterium

Prevotella bivia DNF00320

B Bacterium

Prevotella stercorea DSM 18206

B Bacterium

Pseudoflavonifractor capillosus ATCC 29799

B Bacterium

Pseudomonas aeruginosa strain W36662

B Bacterium

Ralstonia pickettii 12D

B Bacterium

Ralstonia pickettii 12J

B Bacterium

Romboutsia

B Bacterium



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Roseburia faecis

B Bacterium

Roseburia hominis A2-183

B Bacterium

Roseburia inulinivorans

B Bacterium

Roseburia sp. 499

B Bacterium

Roseburia sp. 831b

B Bacterium

Rothia mucilaginosa DY-18

B Bacterium

Ruminococcaceae bacterium D16

B Bacterium

Ruminococcaceae bacterium Marseille-P2963

B Bacterium

Ruminococcaceae bacterium cv2

B Bacterium

Ruminococcus bicirculans

B Bacterium

Ruminococcus champanellensis

B Bacterium

Ruminococcus flavefaciens strain Y1

B Bacterium

Ruminococcus gauvreauii

B Bacterium



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Ruminococcus lactaris

B Bacterium

Ruminococcus sp. 5_1_39BFAA

B Bacterium

Ruminococcus sp. DSM 100440

B Bacterium

Ruminococcus sp. JC304

B Bacterium

Ruminococcus torques ATCC 27756

B Bacterium

Ruthenibacterium lactatiformans strain 585-1

B Bacterium

Saccharomyces cerevisiae S288C

E Eukaryote

Saccharomyces sp. 'boulardii' strain unique28

E Eukaryote

Sellimonas intestinalis

B Bacterium

Streptococcus australis

B Bacterium

Streptococcus gallolyticus subsp. gallolyticus ATCC BAA-2069

B Bacterium

Streptococcus intermedius JTH08

B Bacterium

Streptococcus milleri

B Bacterium



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Streptococcus mitis SPAR10

B Bacterium

Streptococcus salivarius JIM8777

B Bacterium

Streptococcus salivarius strain HSISS4

B Bacterium

Streptococcus sp. 1171_SSPC

B Bacterium

Streptococcus sp. 263_SSPC

B Bacterium

Streptococcus sp. 343_SSPC

B Bacterium

Streptococcus sp. 400_SSPC

B Bacterium

Streptococcus sp. A12 sp. A12

B Bacterium

Streptococcus sp. FDAARGOS_192

B Bacterium

Streptococcus sp. HMSC072C09

B Bacterium

Streptococcus sp. HMSC078H12

B Bacterium

Streptococcus sp. I-G2 sp. I-G2

B Bacterium

Streptococcus sp. I-P16 sp. I-P16

B Bacterium



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Streptococcus thermophilus JIM 8232

B Bacterium

Streptococcus thermophilus strain KLDS

B Bacterium

Streptococcus thermophilus strain ND07

B Bacterium

Streptococcus thermophilus strain S9

B Bacterium

Streptococcus viridans

B Bacterium

Subdoligranulum sp. 4_3_54A2FAA

B Bacterium

Subdoligranulum variabile

B Bacterium

Tobacco mild green mosaic virus

V Virus

Tobacco mosaic virus

V Virus

Tomato brown rugose fruit virus

V Virus

Tomato mosaic virus

V Virus

Veillonella dispar

B Bacterium

Veillonella parvula DSM 2008

B Bacterium



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Veillonella parvula strain UTDB1-3

B Bacterium

Veillonella sp. 3_1_44

B Bacterium

Veillonella sp. 6_1_27

B Bacterium

[Bacteroides] pectinophilus

B Bacterium

[Clostridium] asparagiforme

B Bacterium

[Clostridium] bolteae

B Bacterium

[Clostridium] citroniae

B Bacterium

[Clostridium] clostridioforme 2_1_49FAA

B Bacterium

[Clostridium] clostridioforme 90A3

B Bacterium

[Clostridium] clostridioforme 90A4

B Bacterium

[Clostridium] clostridioforme 90A6

B Bacterium

[Clostridium] clostridioforme 90A7

B Bacterium

[Clostridium] clostridioforme 90A8

B Bacterium



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[Clostridium] clostridioforme 90B1

B Bacterium

[Clostridium] clostridioforme CM201

B Bacterium

[Clostridium] clostridioforme WAL-7855

B Bacterium

[Clostridium] clostridioforme strain 2789STDY5834865

B Bacterium

[Clostridium] clostridioforme strain NLAE-zl-G208

B Bacterium

[Clostridium] glycyrrhizinilyticum JCM 13369

B Bacterium

[Clostridium] innocuum

B Bacterium

[Clostridium] lactatifermentans

B Bacterium

[Clostridium] leptum DSM 753

B Bacterium

[Clostridium] scindens ATCC 35704

B Bacterium

[Clostridium] spiroforme

B Bacterium

[Clostridium] symbiosum WAL-14163

B Bacterium

[Clostridium] symbiosum WAL-14673

B Bacterium



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[Eubacterium rectale] ATCC 33656

B Bacterium

[Eubacterium] contortum strain 2789STDY5834876

B Bacterium

[Eubacterium] eligens ATCC 27750

B Bacterium

[Eubacterium] hallii DSM 3353

B Bacterium

[Eubacterium] rectale strain T1-815

B Bacterium

[Eubacterium] siraeum DSM 15702

B Bacterium

[Eubacterium] siraeum strain 2789STDY5834928

B Bacterium

[Ruminococcus] torques strain 2789STDY5834841

B Bacterium

bacterium LF-3

B Bacterium

https://www.viome.com/reportablerange



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Viome Methodology

Microbial total RNA is extracted, ribosomal RNA molecules are removed from total RNA, and the remaining RNA molecules are sequenced on Illumina NextSeq or NovaSeq. Proprietary bioinformatics algorithms are used to perform taxonomic classification and functional analysis of the sequencing data.

Whole blood total RNA is extracted, polyadenylated transcripts are captured from total RNA and sequenced on Illumina NextSeq or NovaSeq. Proprietary bioinformatics algorithms are used to perform quantitative gene expression analysis of the sequencing data. Results are reported to Viome customers in the context of integrative functional health themes communicated as scores derived largely from proprietary pathway content and analytics methodology. Each score is built to account for molecular pathway topology and strength of literature evidence manually curated by translational science experts in systems biology. Scoring results are CLIA-validated and are end-to-end automated in the production system, which uses each customer's gene expression data as input.

Method Limitation

Viome's results and recommendations are based on our ability to identify and quantify thousands of microbial taxa. Such vast diversity has not been captured in the genomic databases, so it is impossible to assess it comprehensively. There are microorganisms that thrive in the gut whose genomes have not been sequenced. Viome is unable to identify those specific organisms, but can identify their near neighbors, which have similar homology. There are also taxa that we cannot discriminate because of their sequence similarity, for example at the strain level. There are some RNA transcripts that may not always align and match to specific known organisms, which may be due to the fact that these sequences are poorly characterized, reliable consensus sequence may not be available for reference. Viome monitors the growth of public genomic databases and will update its own databases when there is sufficient new information to be worthy of incorporation.

Detection of a microorganism by this test does not imply having a disease. Similarly, not detecting a microorganism by this test does not exclude the presence of a disease-causing microorganism. Further, other organisms may be present that are not detected by this test. This test is not a substitute for established methods for identifying microorganisms or their antimicrobial susceptibility profile. Results are qualitative and identify the presence or absence of identified annotated organisms.

Viome's results and recommendations are based on our ability to identify and quantify thousands of human transcripts. While the test has been clinically validated and shows very high precision, it also has some limitations. As the presence of transcripts nears the limits of detection, the ability of the test to accurately detect them is diminished. This is simply due to the uneven distribution of molecules in liquid volumes, causing small random changes in the transcript concentrations. Scores rely on detection of expressed genes, as well as their levels of expression against the reference population cohort. Hence, certain sample results may be affected by any skewing or sampling biases of the reference cohort, as opposed to solely the biology of the given customer. Scores also are limited by our current understanding of actionable or biologically



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meaningful insights and literature coverage to date. As Viome's reference population expands and current knowledge grows, these limitations become more negligible.

The Gut Intelligence Test was developed by, and its performance characteristics determined by Viome Inc. It has not been cleared or approved by the US Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. This laboratory is registered under CLIA (32D2156145) to perform high complexity testing. Sequencing was performed at CLIA (). Contact Viome for any further questions.

The Human Gene Expression test was developed by, and its performance characteristics determined by Viome Inc. It has not been cleared or approved by the US Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. This laboratory is registered under CLIA 32D2156145 to perform high complexity testing. Sequencing was performed at Viome, Inc. CLIA 32D2156145. Contact Viome for any further questions.



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CHARLES WARDEN'S RESULTS

VERSION: 1.14.2