

Date/Time 12/30/2020 12/31/2020 01/01/2021 Specimens Collected 3

Verrucomicrobia

Tenericutes

Actinobacteria

Proteobacteria

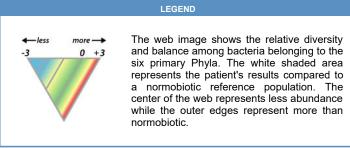
Bacteriodeles

Firmicutes

DI Score

Microbiome Abundance and Diversity Summary

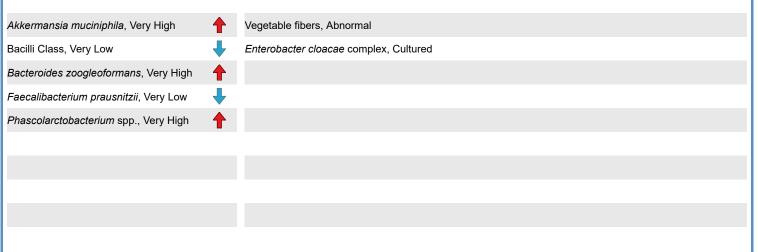
The abundance and diversity of gastrointestinal bacteria provide an indication of gastrointestinal health, and gut microbial imbalances can contribute to dysbiosis and other chronic disease states. The GI360™ Microbiome Profile is a gut microbiota DNA analysis tool that identifies and characterizes more than 45 targeted analytes across six Phyla using PCR and compares the patient results to a characterized normobiotic reference population. The web chart illustrates the degree to which an individual's microbiome profile deviates from normobiosis.

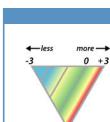


Dysbiosis Index

The Dysbiosis Index the (DI) is calculated strictly from the results of the Microbiome Profile, with scores from 1 to 5. A DI score above 2 indicates dysbiosis; a microbiota profile that differs from the defined normobiotic reference population. The higher the DI above 2, the more the sample deviates from the normobiotic profile. The dysbiosis test and DI does not include consideration of dysbiotic and pathogenic bacteria, yeast, parasites and viruses that may be reported in subsequent sections of the GI360[™] test.

Key Findings





Diagnostics



Order: SAMPLE REPORT Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Age	ient: e:56 k:Fem		le Pati			Dat Dat Dat	e Coll e Rec e Rep	Collectio lected eived oorted ns Colle	12/30/2020 12/31/2020 01/01/2021
-3 -2 -1 0		+1	+2	-	+3					rmobiotic population.
Very Low Low Within Reference	e Interval	High		Very Hig	gh	microbiota profile	in which	n microo	rganisms w	izes a composition of the ith potential health benefits r potentially harmful ones.
Actinobacteria	Result		-3	-2	-1	0	+1	+2	+3	Reference Interval
Actinobacteria	-1				\triangle					0
Actinomycetales	0									0
<i>Bifidobacterium</i> spp.	-1				\triangle					0
Bacteriodetes	Result		-3	-2	-1	0	+1	+2	+3	Reference Interval
Alistipes spp.	0									0
Alistipes onderdonkii	0									0
Bacteroides fragilis	0									0
Bacteroides spp. & Prevotella spp.	+1						Δ			0
Bacteroides stercoris	0									0
Bacteroides zoogleoformans	+2									0
Parabacteroides johnsonii	0									0
Parabacteroides spp.	0									0
Firmicutes	Result		-3	-2	-1	0	+1	+2	+3	Reference Interval
Firmicutes	0							_		0
Bacilli Class	-2									0
Catenibacterium mitsuokai	0									0
Clostridia Class	0									0
Clostridium L2-50	0									0

Notes:

The gray-shaded area of the bar graph represents reference values outside the reporting limits for this test. *This test was developed and its performance characteristics determined in a manner consistent with CLIA requirements. The U. S. Food and Drug Administration (FDA) has not approved or cleared this test; however, FDA clearance is not currently required for clinical use. The results are not intended to be used as a sole means for clinical diagnosis or patient management decisions.

Methodology: Multiplex PCR



Order: SAMPLE REPORT Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Age:	nt: Samp 56 Female	le Patie	ent		Dat Dat Dat	e Coll e Rec e Rep	Collectio lected eived lorted ns Colle	12/30/2020 12/31/2020 01/01/2021
Firmicutes	Result	-3	-2	-1	0	+1	+2	+3	Reference Interva
Dialister invisus	0								0
Dialister invisus & Megasphaera micronuciformis	0								0
<i>Dorea</i> spp.	0								0
Eubacterium biforme	0								0
Eubacterium hallii	0								0
Eubacterium rectale	0								0
Eubacterium siraeum	0								0
Faecalibacterium prausnitzii	-3								0
Lachnospiraceae	-1		4						0
Lactobacillus ruminis & Pediococcus acidilactici	0								0
Lactobacillus spp.	0								0
Phascolarctobacterium spp.	+3								0
Ruminococcus albus & R. bromii	0								0
Ruminococcus gnavus	0								0
Streptococcus agalactiae & Eubacterium rectale	0								0
Streptococcus salivarius ssp. thermophilus & S. sanguinis	0								0
<i>Streptococcus salivarius</i> ssp. <i>thermophilus</i>	-1		4	\land					0
Streptococcus spp.	0								0
<i>Veillonella</i> spp.	-1		4	\land					0

Notes:

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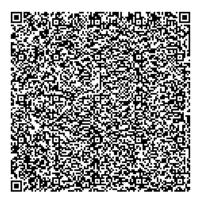


Order: SAMPLE REPORT Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Age:	nt: Samp 56 ⁻ emale	le Pat	ient		Dat Dat Dat	e Coll e Rec e Rep	eived		Date/Time 12/30/2020 12/31/2020 01/01/2021 3
Proteobacteria	Result	-3	-2	-1	0	+1	+2	+3	Refe	erence Interval
Proteobacteria	0								0	
Escherichia spp.	0								0	
Tenericutes	Result	-3	-2	-1	0	+1	+2	+3	Refe	erence Interval
Mycoplasma hominis	0								0	
Verrucomicrobia	Result	-3	-2	-1	0	+1	+2	+3	Refe	erence Interval
Akkermansia muciniphila	+2								0	



Microbiome Abundance Information:

• The GI360[™] Microbiome Profile is a focused gut microbiota DNA analysis tool that identifies more than 45 targeted analytes across six phyla using a CE-marked multiplex PCR system. Patient results are compared to a highly defined normobiotic reference population (n > 1,100). The white shadowed web plot within the hexagonal diagram illustrates the degree to which an individual's microbiome profile deviates from normobiosis. The center of the diagram represents less bacterial abundance while the outer edges represent greater than normobiosis. Deviation from a hexagon-shaped plot indicates variant diversity of the microbial community. Key findings for patient's microbiome profile are summarized in the table below the diagram, and detailed results for all of the analytes are presented on the next 3 pages of the report. Detailed results for the specific bacteria are reported as -3 to +3 standard deviations, as compared to the normobiotic reference population.



Notes:

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Order: SAMPLE REPORT Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Patient: Sample F Age: 56 Sex: Female	Patient	Sample Collection Date Collected Date Received Date Reported Specimens Collected	Date/Time 12/30/2020 12/31/2020 01/01/2021 3
Viruses		Result		
Adenovirus F40/41		Negative		
Norovirus GI/GII		Negative		
Rotavirus A		Negative		
Pathogenic Bacteria		Result		
Campylobacter (C. jejuni, C. coli and C. lari)		Negative		
Clostridioides difficile (Toxin A/B)		Negative		
Escherichia coli O157		Negative		
Enterotoxigenic Escherichia coli (ETEC) It/st		Negative		
Salmonella spp.		Negative		
Shiga-like toxin-producing Escherichia coli (S	TEC) stx1/stx2	Negative		
Shigella (S. boydii, S. sonnei, S. flexneri & S.	dysenteriae)	Negative		
Vibrio cholerae		Negative		
Parasites		Result		
Cryptosporidium (C. parvum and C. hominis)		Negative		
Entamoeba histolytica		Negative		
Giardia duodenalis (AKA intestinalis & lamblia	a)	Negative		





Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient Age: 56 Sex: Female

Sample CollectionDate/TimeDate Collected12/30/2020Date Received12/31/2020Date Reported01/01/2021Specimens Collected3

Protozoa	Result	
Balantidium coli	Not Detected	
<i>Blastocystis</i> spp.	Not Detected	
Chilomastix mesnili	Not Detected	
Dientamoeba fragilis	Not Detected	
Endolimax nana	Not Detected	
Entamoeba coli	Not Detected	
Entamoeba hartmanni	Not Detected	
Entamoeba histolytica/Entamoeba dispar	Not Detected	
Entamoeba polecki	Not Detected	
Enteromonas hominis	Not Detected	
Giardia duodenalis	Not Detected	
lodamoeba bütschlii	Not Detected	
Isospora belli	Not Detected	
Pentatrichomonas hominis	Not Detected	
Retortamonas intestinalis	Not Detected	
Cestodes - Tapeworms	Result	
Diphyllobothrium latum	Not Detected	
Dipylidium caninum	Not Detected	
Hymenolepis diminuta	Not Detected	
Hymenolepis nana	Not Detected	
Taenia	Not Detected	
Trematodes - Flukes	Result	
Clonorchis sinensis	Not Detected	
Fasciola hepatica/Fasciolopsis buski	Not Detected	
Heterophyes heterophyes	Not Detected	
Paragonimus westermani	Not Detected	
Nematodes - Roundworms	Result	
Ascaris lumbricoides	Not Detected	

Notes: Methodology: Microscopy Page: 6 of 17



Order: SAMPLE REPORT Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Patient: Sample Patient Age: 56 Sex: Female	Sample Collection Date Collected Date Received Date Reported Specimens Collected	Date/Time 12/30/2020 12/31/2020 01/01/2021 3
Nematodes - Roundworms	Result		
Capillaria hepatica	Not Detected		
Capillaria philippinensis	Not Detected		
Enterobius vermicularis	Not Detected		
Hookworm	Not Detected		
Strongyloides stercoralis	Not Detected		
Trichuris trichiura	Not Detected		
Other Markers	Result	Refer	ence Interval
Yeast	Few	Not D	etected – Rare
RBC	Not Detected	Not D	etected – Rare
WBC	Not Detected	Not D	etected – Rare
Muscle fibers	Not Detected	Not D	etected – Rare
Vegetable fibers	Moderate	Not D	etected – Few
Charcot-Leyden Crystals	Not Detected	Not D	etected
Pollen	Not Detected	Not D	etected
Macroscopic Appearance	Result	Refer	ence Interval
Color	Brown	Brown	า
Consistency	Soft	Soft	
Mucus	Negative	Nega	tive

Barasitology Information:

- This test is not designed to detect Cyclospora cayetanensis or Microsproridia spp.
- Intestinal parasites are abnormal inhabitants of the gastrointestinal tract that have the potential to cause damage to their host. The presence of any parasite within the intestine generally confirms that the patient has acquired the organism through fecal-oral contamination. Damage to the host includes parasitic burden, migration, blockage and pressure. Immunologic inflammation, hypersensitivity reactions and cytotoxicity also play a large role in the morbidity of these diseases. The infective dose often relates to severity of the disease and repeat encounters can be additive.
- There are two main classes of intestinal parasites, they include protozoa and helminths. The protozoa typically have two stages; the trophozoite stage that is the metabolically active, invasive stage and the cyst stage, which is the vegetative inactive form resistant to unfavorable environmental conditions outside the human host. Helminths are large, multicellular organisms. Like protozoa, helminths can be either free-living or parasitic in nature. In their adult form, helminths cannot multiply in humans.





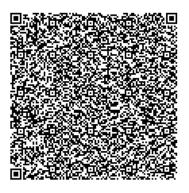
Precision Point Diagnostics

Patient: Sample Patient Age: 56 Sex: Female

Sample Collection Date/Time Date Collected 12/30/2020 Date Received 12/31/2020 **Date Reported** 01/01/2021 Specimens Collected 3

Parasitology Information:

- In general, acute manifestations of parasitic infection may involve diarrhea with or without mucus and or blood, fever, nausea, or abdominal pain. However these symptoms do not always occur. Consequently, parasitic infections may not be diagnosed or eradicated. If left untreated, chronic parasitic infections can cause damage to the intestinal lining and can be an unsuspected cause of illness and fatique. Chronic parasitic infections can also be associated with increased intestinal permeability, irritable bowel syndrome, irregular bowel movements, malabsorption, gastritis or indigestion, skin disorders, joint pain, allergic reactions. and decreased immune function.
- In some instances, parasites may enter the circulation and travel to various organs causing severe organ diseases such as liver abscesses and cysticercosis. In addition, some larval migration can cause pneumonia and in rare cases hyper infection syndrome with large numbers of larvae being produced and found in every tissue of the body.
- Red Blood Cells (RBC) in the stool may be associated with a parasitic or bacterial infection, or an inflammatory bowel condition such as ulcerative colitis. Colorectal cancer, anal fistulas, and hemorrhoids should also be ruled out.
- White Blood Cells (WBC) and Mucus in the stool can occur with bacterial and parasitic infections, with mucosal irritation, and inflammatory bowel diseases such as Crohn's disease or ulcerative colitis
- Muscle fibers in the stool are an indicator of incomplete digestion. Bloating, flatulence, feelings of "fullness" may be associated with increase in muscle fibers.
- **Vegetable fibers** in the stool may be indicative of inadequate chewing, or eating "on the run".





Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics	Age:	56 Female			Date C Date R Date R	collected acceived aported nens Coll	12/30/2020 12/31/2020 01/01/2021
Pathogenic Bacteria	Result	NG	1+	2+	3+	4+	Reference Interval
Aeromonas	NG						No Growth
Edwardsiella tarda	NG						No Growth
Plesiomonas shigelloides	NG						No Growth
Salmonella	NG						No Growth
Shigella spp.	NG						No Growth
Vibrio cholerae	NG						No Growth
Vibrio	NG						No Growth
Yersinia	NG						No Growth
Imbalance Bacteria	Result	NG	1+	2+	3+	4+	Reference Interval
Beta hemolytic strep, group B	2+			\triangle			No Growth
Citrobacter freundii complex	1+		Δ				No Growth
Comamonas jiangduensis	3+				\triangle		No Growth
Gamma hemolytic strep	2+			Δ			No Growth
Dysbiotic Bacteria	Result	NG	1+	2+	3+	4+	Reference Interval
Enterobacter cloacae complex	3+						No Growth
Yeast	Result	NG	1+	2+	3+	4+	Reference Interval
No yeast isolated	NG						

Patient: Sample Patient

Sample Collection

Date/Time

GI

Microbiology Information:

- Pathogenic bacteria consist of known pathogenic bacteria that can cause disease in the GI tract. They are present due to the consumption of contaminated food or water, exposure to animals, fish, or amphibians known to harbor the organism. These organisms can be detected by either Multiplex PCR or microbiology culture.
- Imbalanced bacteria are usually neither pathogenic nor beneficial to the host GI tract. Imbalances can occur when there are insufficient levels of beneficial bacteria and increased levels of commensal bacteria. Certain commensal bacteria are reported as dysbiotic at higher levels.
- **Dysbiotic bacteria** consist of those bacteria that have the potential to cause disease in the GI tract. They can be present due to a number of factors including: exposure to chemicals that are toxic to beneficial bacteria; the use of antibiotics, oral contraceptives or other medications; poor fiber intake and high stress levels.
- Yeast may normally be present in small quantities on the skin, in the mouth and intestine. While small quantities of yeast may be normal, yeast observed in higher quantities is considered abnormal.





Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient Age: 56 Sex: Female Sample CollectionDate/TimeDate Collected12/30/2020Date Received12/31/2020Date Reported01/01/2021Specimens Collected3





Order: SAMPLE REPOR Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics		Patient: Age: 56 Sex: Fer	Sample Pa nale	ıtient	Sample C Date Colle Date Rece Date Repo Specimen	ected12/30/2020bived12/31/2020
Digestion / Absorption	Result	Unit	L	WRI	Н	Reference Interval
Elastase	427	µg/mL				> 200
Fat Stain	Few					None – Few
Carbohydrates [†]	Negative					Negative
Inflammation	Result	Unit	L	WRI	н	Reference Interval
Lactoferrin	1.1	µg/mL				<7.3
Lysozyme*	117	ng/mL				≤ 500
Calprotectin	12	µg/g				≤50
Immunology	Result	Unit	L	WRI	Н	Reference Interval
Secretory IgA*	60.0	mg/dL				30-275
Short Chain Fatty Acids	Result	Unit	L	WRI	Н	Reference Interval
% Acetate [‡]	71					50-72
% Propionate [‡]	16					11 – 25
% Butyrate [‡]	12					11 – 32
% Valerate [‡]	1.3					0.8-5.0
Butyrate [‡]	1.3	mg/mL				0.8-4.0
Total SCFA's [‡]	11	mg/mL				5.0-16.0
Intestinal Health Markers	Result	Unit	L	WRI	Н	Reference Interval
рН	6.2					5.8-7.0
β-glucuronidase*	100	U/L				100 – 1200
Occult Blood	Negative					Negative

Chemistry Information:

• Elastase findings can be used for the diagnosis or the exclusion of exocrine pancreatic insufficiency. Correlations between low levels and chronic pancreatitis and cancer have been reported.

Notes:

RI= Reference Interval, L (blue) = Low (below RI), WRI (green) = Within RI (optimal), WRI (yellow) = Within RI (not optimal), H (red) = High (above RI) *This test was developed and its performance characteristics determined in a manner consistent with CLIA requirements. The U. S. Food and Drug Administration (FDA) has not approved or cleared this test; however, FDA clearance is not currently required for clinical use. The results are not intended to be used as a sole means for clinical diagnosis or patient management decisions.

[†]This test has been modified from the manufacturer's instructions and its performance characteristics determined in a manner consistent with CLIA requirements.

[‡]This test was developed and its performance characteristics determined in a manner consistent with CLIA requirements. The U.S. Food and Drug ^{Administration (FDA)} has not approved or cleared this test; however, FDA clearance is not currently required for clinical use.

Methodology: Elisa, Microscopy, Colormetric, Gas Chromotography, ph Electrode, Guaiac



Diagnostics



Patient: Sample Patient Age: 56 Sex: Female

Sample CollectionDate/TimeDate Collected12/30/2020Date Received12/31/2020Date Reported01/01/2021Specimens Collected3

Chemistry Information:

- Fat Stain: Microscopic determination of fecal fat using Sudan IV staining is a qualitative procedure utilized to assess fat absorption and to detect steatorrhea.
- Carbohydrates: The presence of reducing substances in stool specimens can indicate carbohydrate malabsorption.
- Lactoferrin and Calprotectin are reliable markers for differentiating organic inflammation (IBD) from function symptoms (IBS) and for management of IBD. Monitoring levels of fecal lactoferrin and calprotectin can play an essential role in determining the effectiveness of therapy, are good predictors of IBD remission, and can indicate a low risk of relapse.
- Lysozyme is an enzyme secreted at the site of inflammation in the GI tract and elevated levels have been identified in IBD patients.
- Secretory IgA (sIgA) is secreted by mucosal tissue and represents the first line of defense of the GI mucosa and is central to the normal function of the GI tract as an immune barrier. Elevated levels of sIgA have been associated with an upregulated immune response.
- Short chain fatty acids (SCFAs): SCFAs are the end product of the bacterial fermentation process of dietary fiber by beneficial flora in the gut and play an important role in the health of the GI as well as protecting against intestinal dysbiosis. Lactobacilli and bifidobacteria produce large amounts of short chain fatty acids, which decrease the pH of the intestines and therefore make the environment unsuitable for pathogens, including bacteria and yeast. Studies have shown that SCFAs have numerous implications in maintaining gut physiology. SCFAs decrease inflammation, stimulate healing, and contribute to normal cell metabolism and differentiation. Levels of **Butyrate** and **Total SCFA** in mg/mL are important for assessing overall SCFA production, and are reflective of beneficial flora levels and/or adequate fiber intake.
- pH: Fecal pH is largely dependent on the fermentation of fiber by the beneficial flora of the gut.
- Occult blood: A positive occult blood indicates the presence of free hemoglobin found in the stool, which is released when red blood cells are lysed.
- β-glucuronidase is an enzyme that breaks the tight bond between glucuronic acid and toxins in the intestines. The binding of toxins in the gut is protective by way of blocking their absorption and facilitating excretion.





Client #: 12345 Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient
Age: 56
Sex: Female

Sample CollectionDate/TimeDate Collected12/30/2020Date Received12/31/2020Date Reported01/01/2021Specimens Collected3

Enterobacter cloacae complex

Natural Agents	Low Sensitivity		High Sensitivity	
Berberine*				
Black Walnut*				
Caprylic Acid*				
Uva Ursi*				
Oregano*				
Grapefruit Seed Extract*				
Silver*				
Prescriptive Agents	Resistant	Intermediate	Susceptible	
Amoxicillin-Clavulanic Acid				
Ampicillin				
Cefazolin				
Ceftazidime				
Ciprofloxacin				
Sulfamethoxazole / Trimethoprim				



Susceptibility Information:

- **Natural antibacterial** agents may be useful for treatment of patients when organisms display in-vitro sensitivity to these agents. The test is performed by using standardized techniques and filter paper disks impregnated with the listed agent. Relative sensitivity is reported for each natural agent based upon the diameter of the zone of inhibition surrounding the disk. Data based on over 5000 individual observations were used to relate the zone size to the activity level of the agent. A scale of relative sensitivity is defined for the natural agents tested.
- **Susceptible** results imply that an infection due to the bacteria may be appropriately treated when the recommended dosage of the tested antimicrobial agent is used. **Intermediate** results imply that response rates may be lower than for susceptible bacteria when the tested antimicrobial agent is used. **Resistant** results imply that the bacteria will not be inhibited by normal dosage levels of the tested antimicrobial agent.



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Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient
Age: 56
Sex: Female

Sample CollectionDateDate Collected12Date Received12Date Reported01Specimens Collected3

Date/Time 12/30/2020 12/31/2020 01/01/2021

Introduction

This analysis of the stool specimen provides fundamental information about the overall gastrointestinal health of the patient. When abnormal microflora or significant aberrations in intestinal health markers are detected, specific commentaries are presented. If no significant abnormalities are found, commentaries are not presented.

Microbiome Abundance Information

Actinobacteria (phylum)

Actinobacteria is one of the largest bacterial phyla, comprised of Gram-positive bacteria. This phylum includes a wide range of species, with different morphological and physiological characteristics. Significant groups in the human colon include Actinomycetales and Bifidobacteriales. Actinomycetales were inversely associated with clinically significant depression in IBS patients, suggesting these bacteria may be depleted in depressed IBS patients. A strict vegetarian diet may increase the total count of *Actinomyces* spp. compared to following a Western diet.

👆 Bifidobacterium (genus)

Considered amongst the most beneficial commensal bacteria in the human gut, Bifidobacterium spp. are able to degrade monosaccharides, galacto-, manno-, and fructo-oligosaccharides, as well as some complex carbohydrates. Many of the nondigestible oligosaccharides, found as natural components in mother's milk, select for colonization of these species which dominate the infant gut shortly after birth. Bifidobacteria may provide health benefits directly through interactions with the host, and indirectly through interactions with other microorganisms. Bifidobacterium spp. take part in production and adsorption of vitamins, such as vitamins K and B12, biotin, folate, thiamine, riboflavin, and pyridoxine. They are also involved in lipid absorption and metabolism, glucose and energy homeostasis, and regulating intestinal barrier function. Although Bifidobacterium produce acetate over butyrate, healthy levels of Bifidobacterium spp. facilitate colonization of Faecalibacterium. prausnitzii. Polyphenols derived from chocolate, green tea, blackcurrant, red wine and grape seed extracts have been shown to increase Bifidobacterium species. The increased abundance of Bifidobacterium species has been associated with amelioration of inflammation. Multiple published studies have suggested that there is an association between obesity and a lower abundance of bifidobacteria. They may also be less abundant in elderly populations, patients with rheumatoid arthritis, and in individuals diagnosed with Alzheimer's disease. Patients with active inflammatory bowel disease (IBD) have a lower abundance of Bifidobacterium spp. than patients whose IBD is in remission. Taking a probiotic containing bifidobacteria, lactobacilli, and streptococci might help in controlling ulcerative colitis symptoms and preventing their recurrence. Some Bifidobacterium strains have been shown to have beneficial effects in irritable bowel syndrome (IBS). Bifidobacterium spp. abundance has been shown to be diminished with IBD and with long term use of macrolide antibiotics. Luminal bifidobacteria is reduced with restriction of fermentable carbohydrates, i.e. a low FODMAP diet. High fat dietary feeding is also associated with reduced abundance of bifidobacteria. Consumption of maize and barley-based whole grain products and red berries, which are comprised of anthocyans, are known to increase levels of bifidobacteria.

Bacteriodetes (phylum)

Bacteriodetes make up approximately 28% of the gut microbiota in healthy human adults. They are early colonizers of the infant gut and are amongst the most stable, at a species and strain level, in the host. A low preponderance of Bacteriodetes in relation to Firmicutes has been associated with obesity, though this can increase with weight loss and restricted calorie intake.

🔶 Prevotella (genus)

Prevotella-rich dysbiosis has been associated with insulin-resistance, obesity and hypertension. *Prevotella* have been shown to be significantly decreased in Crohn's disease and Parkinson's disease. High levels of fiber and carbohydrates from fruits and vegetables in a Mediterranean diet have been shown to increase the relative abundance of *Prevotella*.

Bacteroides (species)

Species in the genus *Bacteroides* carry out broad metabolic functions, including degradation of complex plant polysaccharides, proteolytic activities, de-conjugation of bile acids, mucosal barrier integrity, short chain fatty acid production, fatty acid storage and glucose metabolism. *Bacteroides* spp. are maintained at a higher abundance in breastfed individuals into adulthood. *Bacteroides fragilis* plays an important role in the prevention of intestinal inflammation. An energy-restricted diet has been shown to increase *B. fragilis* in overweight adolescents. An increase in *B. stercoris* has been associated with higher risk of colon cancer. Decreased levels of *Bacteroides* spp. have been reported in association with multiple sclerosis, rheumatoid arthritis and Parkinson's disease.





Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient
Age: 56
Sex: Female

Sample CollectionDateDate Collected12Date Received12Date Reported01Specimens Collected3

Date/Time 12/30/2020 12/31/2020 01/01/2021 3

Microbiome Abundance Information continued...

Firmicutes (phylum)

The phylum Firmicutes constitutes the most diverse and abundant group of gastrointestinal microbiota which are grouped into four classes, Bacilli, Clostridia, Erysipelotrichia, and Negativicutes. They constitute about 39% of gut bacteria in healthy adults, but may increase to as high as 80% in an imbalanced microbial community.

Faecalibacterium prausnitzii (species)

Faecalibacterium prausnitzii is one of the most abundant butyrate producing bacteria in a healthy gastrointestinal tract. As such, *F. prausnitzii* is a protective factor for the intestinal mucosa and supports very important intestinal barrier functions. *F. prausnitzii* exerts anti-inflammatory effects via metabolites such as short-chain fatty acids. *F. prausnitzii* is reduced in inflammatory bowel disease, irritable bowel syndrome, celiac disease and gastrointestinal inflammation in general. It is reduced in patients diagnosed with Parkinson's disease, bipolar disorder, colorectal cancer, diabetes and chronic idiopathic diarrhea. Diminished levels of *F. prausnitzii* were found in patients with major depressive disorder. The abundance of *F. prausnitzii* together with *E. coli* has been proposed as a discrimination tool between ulcerative colitis and Crohn's disease. *F. prausnitzii* has been correlated with pediatric obesity in instances of high consumption of foods that are rich in unabsorbed carbohydrate (banana, maize, rice). The prebiotic inulin has been shown to increase the proportion of *F. prausnitzii* in the human intestinal microbiota. Low FODMAP diets are associated with diminished *F. prausnitzii* and butyrate production.

Lachnospiraceae (family)

The Lachnospiraceae family is a diverse group of butyric acid producers, which have been associated with beneficial microbial and epithelial cell growth. Consumption of a Mediterranean diet decreased levels of species belonging to Lachnospiraceae. Lachnospiraceae are known to increase with intake of cruciferous vegetables and wheat bran, and decrease with a resistant starch diet.

Phascolarctobacterium (genus)

Phascolarctobacterium are in the Firmicutes phylum. *Phascolarctobacterium* can produce short chain fatty acids, including acetate and propionate, and may be associated with metabolic effects and mental state of the host. Patients diagnosed with major depressive disorder had increased levels of these species. Decreased levels of *Phascolarctobacterium* were found to be associated with Crohn's disease, ulcerative colitis and Alzheimer's disease. Consumption of cruciferous vegetables, such as broccoli, increases the abundance of *Phascolarctobacterium* in the gut.

Streptococcus (genus)

Higher abundance of *S. salivarius* and *S. thermophilus* (Firmicutes phylum) have been associated with a moderate to severe disease course in newly diagnosed ulcerative colitis (UC) patients. These findings are in accordance with a study that showed that UC patients have significantly increased *Streptococcus* spp. and depletion of *Bifidobacterium* spp. Higher levels of *Streptococcus* spp. were also observed in patients with colorectal cancer compared to healthy controls. Administration of *S. salivarius* together with *Bifidobacterium* bifidum was shown to reduce the incidence of acute diarrhea and rotavirus shedding in infants. *S. salivarius* and *S. thermophilus* are also widely used in dairy products like yogurt and cheese.

Veillonella (genus)

Veillonella (Firmicutes phylum) are known for their ability to ferment lactate, producing the short chain fatty acids propionate and acetate. *Veillonella* spp. were shown to be significantly increased in patients with Crohn's disease, type 1 diabetes, and patients diagnosed with liver cirrhosis. Increased amounts of *Veillonella* have been found in patients with constipation dominant irritable bowel syndrome (IBS-C). It is hypothesized that the relationship between *Veillonella* strains and IBS stems from its robust production of organic acids (propionate and acetate) which contribute to bloating, anxiety and abdominal pain. Higher levels of *Veillonella* were found in formula-fed infants compared to breast-fed infants.

Proteobacteria (phylum)

Proteobacteria include a wide variety of pathogens, including species within the *Escherichia*, *Shigella Salmonella*, *Vibrio*, and *Helicobacter* genera. The phylum includes a number of species that are permanent residents of the microbiota and capable of inducing nonspecific inflammation and diarrhea when their presence is increased. Proteobacteria make up approximately 2% of the gut microbiota in healthy adults.





Doctor: Sample Doctor Precision Point Diagnostics Patient: Sample Patient
Age: 56
Sex: Female

Sample CollectionDateDate Collected12/3Date Received12/3Date Reported01/0Specimens Collected3

Date/Time 12/30/2020 12/31/2020 01/01/2021 3

Microbiome Abundance Information continued...

Tenericutes (phylum)

Tenericutes are cell wall-less bacteria that do not synthesize precursors of peptidoglycan. Tenericutes consist of four main clades designated as the *Acholeplasma, Spiroplasma, Pneumoniae* and *Hominis* clusters. Tenericutes are typically parasites or commensals of eukaryotic hosts.

Verrucomicrobia (phylum)

Verrucomicrobia is a less common phylum in the human gut microbiota, but one with increasing recognition with regards to health. Verrucomicrobia includes *Akkermansia muciniphila*. The obligate anaerobe *A. muciniphila* constitutes 3-5% of total bacteria in a healthy microbiome, and has a protective or anti-inflammatory role in the intestinal mucosa.

Akkermansia muciniphila (genus)

Higher abundance of *Akkermansia muciniphila* has been associated with a milder disease course in newly discovered ulcerative colitis patients. Archaea and *Akkermansia* were significantly more prevalent after weight reduction. A Low FODMAP diet has been shown to decrease the abundance of *A. muciniphila* leading to recommendations against long-term use of such a diet. *A. muciniphila* is a mucolytic specialist that has potent anti-inflammatory effects in part associated with a specific surface coat protein (Amuc- 1100).

GI Pathogens

Introduction

The GI Pathogen profile is performed using an FDA-cleared multiplex PCR system. It should be noted that PCR testing is much more sensitive than traditional techniques and allows for the detection of extremely low numbers of pathogens. PCR testing does not differentiate between viable and non-viable pathogens and should not be repeated until 21 days after completion of treatment or resolution to prevent false positives due to lingering traces of DNA. PCR testing can detect multiple pathogens in the patient's stool but does not differentiate the causative pathogen. All decisions regarding the need for treatment should take the patient's complete clinical history and presentation into account.

Parasitology

Microscopic yeast

Microscopic examination has revealed more yeast in this sample than normal. While small quantities of yeast (reported as rare) may be normal, yeast observed in higher amounts (moderate to many) is considered abnormal. Yeast does not appear to be dispersed uniformly throughout the stool. Yeast may therefore be observed microscopically, but not grow out on culture even when collected from the same bowel movement. Further, some yeast may not survive transit through the intestines rendering it unviable for culturing. Therefore, both microscopic examination and culture are helpful in determining if abnormally high levels of yeast are present. If significant yeast are reported by microscopy, but not by culture, consider the presentation of patient symptoms.

Vegetable fibers

Excessive amounts of vegetable fibers were found in this stool specimen. The presence of vegetable fibers must be considered in conjunction with other parameters such as muscle fibers and Elastase for a proper assessment of maldigestion. Elevated levels of vegetable fibers may be indicative of inadequate chewing or rapid transit time.

Microbiology

Pathogenic/Dysbiotic Flora

In a healthy balanced state of intestinal flora, the beneficial bacteria make up a significant proportion of the total microflora. However, in many individuals there is an imbalance or deficiency of beneficial flora (insufficiency dysbiosis) and an overgrowth of non-beneficial (imbalance) or even pathogenic microorganisms. This can be due to a number of factors including: consumption of contaminated water or food; daily exposure of chemicals that are toxic to beneficial bacteria; the use of antibiotics, oral contraceptives or other medications; poor fiber intake and high stress levels.





Patient: Sample Patient
Age: 56
Sex: Female

Sample Collection Date Collected Date Received Date Reported Specimens Collected Speci

Date/Time 12/30/2020 12/31/2020 01/01/2021 3

Microbiology continued...

A number of toxic substances can be produced by the dysbiotic bacteria including amines, ammonia, hydrogen sulfide, phenols, and secondary bile acids which may cause inflammation or damage to the brush border of the intestinal lining. If left unchecked, long-term damage to the intestinal lining may result in leaky gut syndrome, allergies, autoimmune disease (e.g. rheumatoid arthritis), irritable bowel syndrome, fatigue, chronic headaches, and sensitivities to a variety of foods. In addition, pathogenic bacteria can cause acute symptoms such as abdominal pain, nausea, diarrhea, vomiting, and fever in cases of food poisoning.

Bacterial sensitivities to a variety of prescriptive and natural agents have been provided for the pathogenic bacteria that were cultured from this patient's specimen. This provides the practitioner with useful information to help plan an appropriate treatment regimen. Supplementation with probiotics or consumption of foods (yogurt, kefir, miso, tempeh, tamari sauce) containing strains of lactobacilli, bifidobacteria, and enterococci may help restore healthy flora levels. Soluble fiber and polyphenols derived from chocolate, green tea, blackcurrant, red wine and grape seed extracts have been found to increase the numbers of beneficial bacteria. Hypochlorhydria may also predispose an individual to bacterial overgrowth, particularly in the small intestine. Nutritional anti-inflammatories can aid in reversing irritation to the GI lining. These include quercetin, vitamin C, curcumin, gamma-linoleic acid, omega-3 fatty acids (EPA, DHA), and aloe vera. Other nutrients such as zinc, beta-carotene, pantothenic acid, and L-glutamine provide support for regeneration of the GI mucosa. A comprehensive program may be helpful in individuals in whom a dysbiotic condition has caused extensive GI damage.

Enterobacter cloacae complex

Enterobacter cloacae complex is part of the *Enterobacteriaceae* family. *E cloacae* complex is a group of six closely related species with similar resistance patterns: *E. cloacae, E. asburiae, E. hormaechei, E. kobei, E. ludwigii,* and *E. nimipressuralis.* This gram-negative bacterium is considered dysbiotic at levels of 3+ or greater. *E. cloacae* complex is considered an opportunistic pathogen associated with diarrhea in children. A Shiga-like toxin-producing *E. cloacae* was isolated from the feces of an infant with hemolytic-uremic syndrome. However, *E. cloacae* complex is most often involved in extraintestinal infections including the urinary tract, respiratory tract, and cutaneous wounds.

Widely distributed in the environment, *Enterobacter* spp. is commonly isolated from both human and animal feces. Environmental strains of *Enterobacter* spp. are capable of growth in foods at refrigeration temperature.

E. cloacae complex is known to possess inducible ß-lactamases. Isolates may become resistant to all cephalosporins after initiation of therapy. Avoid ß-lactam-inhibitor drugs such as amoxicillin/ clavulanate, ampicillin/sulbactam, and piperacillin/tazobactam.

Antibiotics may be indicated in systemic infections if symptoms are prolonged. Refer to the antimicrobial susceptibilities for treatment.

Imbalanced Flora

Imbalanced flora are those bacteria that reside in the host gastrointestinal tract and neither injure nor benefit the host. Certain dysbiotic bacteria may appear under the imbalanced category if found at low levels because they are not likely pathogenic at the levels detected. Imbalanced bacteria are commonly more abundant in association with insufficiency dysbiosis, and/or a fecal pH more towards the alkaline end of the reference range (6 - 7.8). Treatment with antimicrobial agents is unnecessary unless bacteria appear under the dysbiotic category.