Accession Number:

Reference Number:

Patient:

Age: 57 Sex: Male

Date of Birth:

 Date Collected:
 10/19/11

 Date Received:
 10/24/11

 Report Date:
 11/7/11

Telephone:
Fax:
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Comment:

Ordering Physician:

Institute of Alternative Medicine Brian Hardy DC,LAc,CCN,DACBN 301 North 200 East Suite 1 - C

St. George, UT 84770

Gastrointestinal Function Profile

Methodology: DNA Analysis, GC/MS, Microscopic, Colorimetric, Automated Chemistry, ELISA

Consistency = Formed/Normal



(E+007)E+007 **Predominant Bacteria Obligate anaerobes** 1.6 6.7 5.2 Bacteroides sp. >= 1.3 1.5 6.2 4.0 >= 1.0 Clostridia sp. 1.6 6.2 Prevotella sp. 2.1 >= 1.1 1.6 7.4 Fusobacteria sp. 3.4 >= 1.1 1.6 5.8 Streptomyces sp. 2.6 1.7 6.2 Mycoplasma sp. 2.4 **Facultative anaerobes** 1.8 7.8 Lactobacillus sp. 3.3 2.3 Bifidobacter sp. 2.7 **Obligate aerobes** Escherichia coli 4.7

Units and Reference Ranges

Organisms are detected by DNA analysis. One colony forming unit (CFU) is equivalent to one bacterium. Each genome detected represents one cell, or one CFU. Results are expressed in scientific notation, so an organism reported as 2.5 E7 CFU/gram is read as 25 million colony forming units per gram of feces. The cutoff for significance of Opportunistic Bacteria has been set at 1.0E+ 005 (100,000). These are levels above which clinically significant growth may be present. Rather than reporting semi-quantitative +1 to +4 levels, the new methodology provides full quantitative analysis.

Predominant Bacteria play major roles in health. They provide colonization resistance against potentially pathogenic organisms, aid in digestion and absorption, produce vitamins and SCFA's, and stimulate the GI immune system. DNA probes allow detection of multiple species (sp.) within a genus, so the genera that are reported cover many species.

Opportunistic Bacteria may cause symptoms and be associated with disease. They can affect digestion and absorption, nutrient production, pH and immune state. Antibiotic sensitivity tests will be performed on all opportunistic bacteria found, although clinical history is usually considered to determine treatment since the organisms are not generally considered to be pathogens.

Opportunistic Bacteria

No clinically significant amounts.

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D. (1 D. (1)		95% Reference
Pathogenic Bacteria		Range
Helicobacter pylori	<0.01	<=1.0E+005
E.H.E. coli	<0.01	<=1.0E+005
Clostridium difficile	<0.01	<=1.0E+005
Campylobacter sp.	<0.01	<=1.0E+005
V 1/5		Expected
Yeast/Fungi		Value

No clinically significant amounts.

Yeast/Fungi

Yeast overgrowth has been linked to many chronic conditions, in part because of antigenic responses in some patients to even low rates of yeast growth. Potential symptoms include diarrhea, headache, bloating, atopic dermatitis and fatigue. Positives are reported as +1, +2, +3 or +4 indicating >100, >1000, >10000 or >100000 pg DNA/g.

Parasites Expected Value Parasite present; taxonomy unavailable. Positive Neg

A taxonomy unavailable finding likely indicates an ingested protozoan and not a human parasite. It does not indicate treatment unless patient symptoms and other inflammatory markers are consistent with parasite infection.

Parasites

Parasite infections are a major cause of non-viral diarrhea. Symptoms may include constipation, gas, bloating, increased allergy response, colitis, nausea and distention.

Adiposity Index

Firmicutes 63

Bacteroidetes 37

Firmicutes 53

Firmicutes 54

Firmicutes 54

Firmicutes 55

Firmicutes

The **Adiposity Index** is derived by using DNA probes that detect multiple genera of the phyla Firmicutes and Bacteroidetes. Abnormalities of these phyla may be associated with increased caloric extraction from food.

Drug Resistance Genes aacA, aphD Neg gyrB, ParE Neg mecA Neg PBP1a, 2B Neg vanA, B, and C Neg

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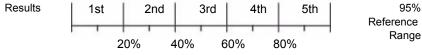
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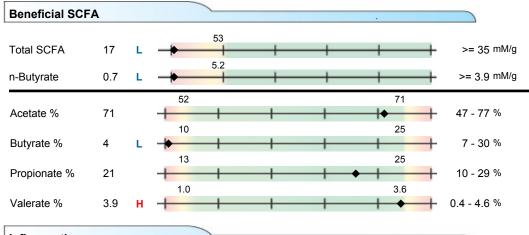
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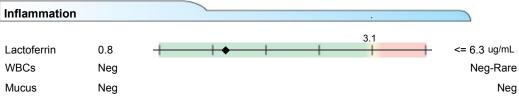
Gastrointestinal Function Profile

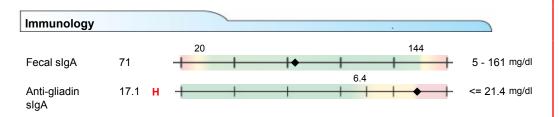
Methodology: DNA Analysis, GC/MS, Microscopic, Colorimetric, Automated Chemistry, ELISA

Percentile Ranking by Quintile 1st 2nd 3rd 4th 5th









Beneficial SCFA

Short chain fatty acids (SCFA) are produced by bacterial fermentation of dietary polysaccharides and fiber. The product, N-butyrate, is taken up and used to sustain the normal activity of colonic epithielial cells. Butyrate has been shown to lower the risk of colitis and colorectal cancer. A healthy balance of GI microbes depends on production of SCFA by one specie to allow the normal growth of another one in a complex cross-feeding network

Inflammation

Lactoferrin, an iron-binding glycoprotein, is released in IBD but not in non-inflammatory IBS. High levels are found in Crohn's, UC or infection. WBC's are elevated in general inflammation/infection. Mucus is often visualized in acute GI inflammation.

Immunology

High fecal slgA indicates immune system reactions to the presence of antigens from bacteria, yeast or other microbes. Low slgA can result from stress or malnutrition. Anti-gliadin slgA is a screening marker for gluten sensitivity.

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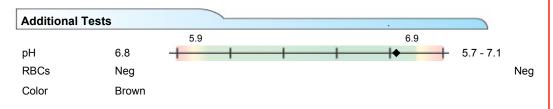
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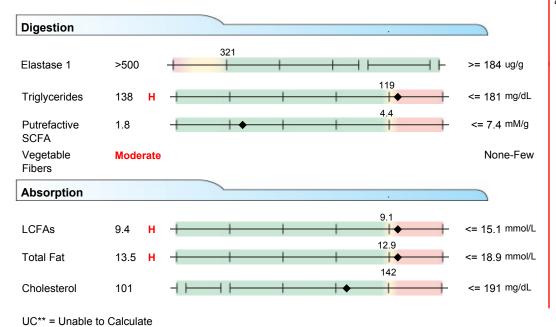
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Additional Tests

pH is influenced by numerous factors, but it is strongly related to the bacterial release of pH-lowering organic acids and pH-raising ammonia. Positive RBCs can signify GI tract bleeding. Color (other than brown) abnormalities can be due to upper GI bleeding, or bile duct blockage, steatorrhea or antibiotic use.

Digestion

Pancreatic elastase 1 levels below the reference limits are strongly correlated with pancreatic insufficiency. High triglycerides signify fat maldigestion. Putrefactive SCFA are a result of bacterial fermentation of undigested protein. High numbers of vegetable fibers indicate maldigestion.

Absorption

High **LCFA** indicates fat malabsorption due to pancreatic or biliary insufficiency, or acute bacterial infection that produces intestinal cell destruction. High total fat usually signals malabsorption, as does elevated fecal cholesterol.

Decisions involving diagnosis and treatment are the responsibility of the clinician.