

Comprehensive Stool Analysis / Parasitology x2

BACTERIOLOGY CULTURE

Expected/Beneficial flora

- 4+ *Bacteroides fragilis* group
- 1+ *Bifidobacterium* spp.
- 3+ *Escherichia coli*
- NG *Lactobacillus* spp.
- NG *Enterococcus* spp.
- 4+ *Clostridium* spp.

Commensal (Imbalanced) flora

- 3+ Alpha hemolytic strep
- 1+ Beta hemolytic strep, not group A or B
- 3+ Gamma hemolytic strep
- 1+ *Pseudomonas chlororaphis* group

Dysbiotic flora

NG = No Growth

BACTERIA INFORMATION

Expected / Beneficial bacteria make up a significant portion of the total microflora in a healthy & balanced GI tract. These beneficial bacteria have many health-protecting effects in the GI tract including manufacturing vitamins, fermenting fibers, digesting proteins and carbohydrates, and propagating anti-tumor and anti-inflammatory factors.

Clostridia are prevalent flora in a healthy intestine. *Clostridium* spp. should be considered in the context of balance with other expected/beneficial flora. Absence of clostridia or over abundance relative to other expected/beneficial flora indicates bacterial imbalance. If *C. difficile* associated disease is suspected, a Comprehensive *Clostridium* culture or toxigenic *C. difficile* DNA test is recommended.

Commensal (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 known pathogenic bacteria and those that have the potential to cause disease in the GI tract. They can be present due to a number of factors including: consumption of contaminated water or food, 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 CULTURE

Normal flora

- 1+ *Candida albicans*
- 1+ *Geotrichum* spp.

Dysbiotic flora

MICROSCOPIC YEAST

RESULT

None

EXPECTED

None – Rare

Yeast in stool is expected at a level of none-rare. A microscopic finding of yeast in stool of few, moderate, or many may be helpful in identifying potential yeast overgrowth, or non-viable or dietary yeast.

YEAST INFORMATION

Yeast may normally be present in small quantities in the skin, mouth, and intestine. When investigating the presence of yeast, disparity may exist between culturing and microscopic examination. Yeast are not uniformly dispersed throughout the stool and this may lead to undetectable or low levels of yeast identified by microscopy, despite culture and identified yeast species. Conversely, microscopic examination may reveal a significant amount of yeast present but no viable yeast cultured. Yeast may not always survive transit through the intestines. Nonviable diet-derived yeast may also be detected microscopically. Consideration of clinical intervention for yeast detected microscopically should be made in the context of other findings and presentation of symptoms.

SPECIMEN DATA

Date Collected: 10/24/2020

Date Received: 10/25/2020

Date Reported: 11/07/2020

Methodology: Culture and identification by MALDI-TOF and conventional biochemicals



**Aeromonas*, *Campylobacter*, *Plesiomonas*, *Salmonella*, *Shigella*, *Vibrio*, *Yersinia*, & *Edwardsiella tarda* have been specifically tested for and found absent unless reported.

Comprehensive Stool Analysis / Parasitology x2

Protozoa	PX1	PX2	
<i>Balantidium coli</i>	Not Detected	Not Detected	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.
<i>Blastocystis spp.</i>	Moderate	Moderate	
<i>Chilomastix mesnili</i>	Not Detected	Not Detected	
<i>Dientamoeba fragilis</i>	Not Detected	Not Detected	
<i>Endolimax nana</i>	Rare Cysts/Trophs	Few Cysts/Trophs	
<i>Entamoeba coli</i>	Not Detected	Not Detected	
<i>Entamoeba hartmanni</i>	Not Detected	Not Detected	
<i>Entamoeba histolytica/Entamoeba dispar</i>	Not Detected	Not Detected	
<i>Entamoeba polecki</i>	Not Detected	Not Detected	
<i>Enteromonas hominis</i>	Not Detected	Not Detected	
<i>Giardia duodenalis</i>	Not Detected	Not Detected	
<i>Iodamoeba bütschlii</i>	Not Detected	Not Detected	
<i>Isospora belli</i>	Not Detected	Not Detected	
<i>Pentatrichomonas hominis</i>	Not Detected	Not Detected	
<i>Retortamonas intestinalis</i>	Not Detected	Not Detected	
Nematodes - Roundworms			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 fatigue. 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.
<i>Ascaris lumbricoides</i>	Not Detected	Not Detected	
<i>Capillaria hepatica</i>	Not Detected	Not Detected	
<i>Capillaria philippinensis</i>	Not Detected	Not Detected	
<i>Enterobius vermicularis</i>	Not Detected	Not Detected	
<i>Strongyloides stercoralis</i>	Not Detected	Not Detected	
<i>Trichuris trichiura</i>	Not Detected	Not Detected	
Hookworm	Not Detected	Not Detected	
Cestodes - Tapeworms			
<i>Diphyllobothrium latum</i>	Not Detected	Not Detected	
<i>Dipylidium caninum</i>	Not Detected	Not Detected	
<i>Hymenolepis diminuta</i>	Not Detected	Not Detected	
<i>Hymenolepis nana</i>	Not Detected	Not Detected	
<i>Taenia</i>	Not Detected	Not Detected	
Trematodes - Flukes			
<i>Clonorchis sinensis</i>	Not Detected	Not Detected	
<i>Fasciola hepatica/Fasciolopsis buski</i>	Not Detected	Not Detected	
<i>Heterophyes heterophyes</i>	Not Detected	Not Detected	
<i>Paragonimus westermani</i>	Not Detected	Not Detected	
Other Markers			One negative parasitology x1 specimen does not rule out the possibility of parasitic disease, parasitology x3 is recommended. This test is not intended to detect <i>Cyclospora cayentanensis</i> or <i>Microsporidium spp.</i>
Yeast	Not Detected	Not Detected	
RBC	Not Detected	Rare	
WBC	Not Detected	Not Detected	
Charcot-Leyden Crystals	Not Detected	Not Detected	
Pollen	Not Detected	Not Detected	
Immunoassay			
<i>Giardia duodenalis</i>	RESULT: Negative	REFERENCE INTERVAL: Negative	
<i>Cryptosporidium</i>	Negative	Negative	

Sample Report

SPECIMEN DATA

Date Collected: 10/24/2020
Date Received: 10/25/2020
Date Reported: 11/07/2020
Methodology: Microscopy, Enzyme Immunoassay

Comprehensive Stool Analysis / Parasitology x2

DIGESTION / ABSORPTION			
	WITHIN	OUTSIDE	REFERENCE INTERVAL
Elastase	233		> 200 µg/mL
Fat Stain	None		None – Few
Muscle fibers	None		None – Rare
Vegetable fibers	Rare		None – Few
Carbohydrates [†]	Negative		Negative

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.

Fat Stain: Microscopic determination of fecal fat using Sudan IV staining is a qualitative procedure utilized to assess fat absorption and to detect steatorrhea.

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”.

Carbohydrates: The presence of reducing substances in stool specimens can indicate carbohydrate malabsorption.

INFLAMMATION			
	WITHIN	OUTSIDE	REFERENCE INTERVAL
Lactoferrin	<0.5		≤ 7 µg/mL
Calprotectin	<10		≤ 50 µg/g
Lysozyme*	147		≤ 500 ng/mL
White Blood Cells	None		None – Rare
Mucus	Negative		Negative

Lactoferrin and **Calprotectin** are reliable markers for differentiating organic inflammation (IBD) from functional 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.

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.

IMMUNOLOGY			
	WITHIN	OUTSIDE	REFERENCE INTERVAL
Secretory IgA*		10.1	30 – 275 mg/dL

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.

SPECIMEN DATA	
Date Collected:	10/24/2020
Date Received:	10/25/2020
Date Reported:	11/07/2020
Methodology:	Elisa, Microscopy, Colormetric, Macroscopic Observation

*This test was developed and its performance characteristics determined by Doctor's Data Laboratories 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 by Doctor's Data Laboratories in a manner consistent with CLIA requirements.

Comprehensive Stool Analysis / Parasitology x2

SHORT CHAIN FATTY ACIDS			
	WITHIN	OUTSIDE	REFERENCE INTERVAL
% Acetate [‡]	52		50 – 72 %
% Propionate [‡]	23		11 – 25 %
% Butyrate [‡]	20		11 – 32 %
% Valerate [‡]	4.4		0.8 – 5.0 %
Butyrate [‡]	0.85		0.8 – 4.0 mg/mL
Total SCFA's [‡]		4.2	5.0 – 16.0 mg/mL

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.

INTESTINAL HEALTH MARKERS			
	WITHIN	OUTSIDE	REFERENCE INTERVAL
Red Blood Cells	Rare		None – Rare
pH	6.7		5.8 – 7.0
Occult Blood	Negative		Negative

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.

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.

MACROSCOPIC APPEARANCE			
	WITHIN	OUTSIDE	EXPECTED
Color	Brown		Brown
Consistency		Loose/Watery	Soft


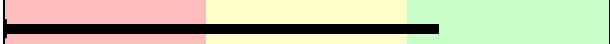
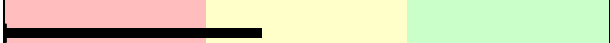


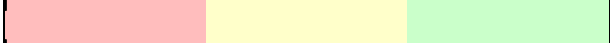
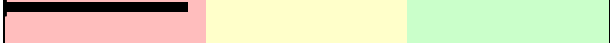
Color: Stool is normally brown because of pigments formed by bacteria acting on bile introduced into the digestive system from the liver. While certain conditions can cause changes in stool color, many changes are harmless and are caused by pigments in foods or dietary supplements.

Consistency: Stool normally contains about 75% water and ideally should be formed and soft. Stool consistency can vary based upon transit time and water absorption.

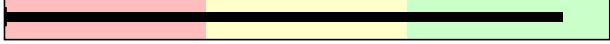
SPECIMEN DATA	
Date Collected:	10/24/2020
Date Received:	10/25/2020
Date Reported:	11/07/2020
Methodology:	Gas Chromotography, pH Electrode, Guaiac, Macroscopic Observation

[‡]This test was developed and its performance characteristics determined by Doctor's Data Laboratories 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.

Yeast Susceptibilities; *Candida albicans*

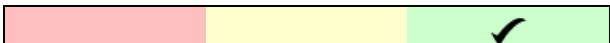
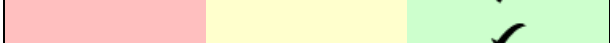
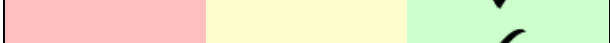
NATURAL ANTIFUNGALS		
	LOW SENSITIVITY	HIGH SENSITIVITY
Berberine*		
Caprylic Acid*		
Uva Ursi*		
Plant Tannins*		
Oregano*		
Undecylenic Acid*		
Grapefruit Seed Extract*		

Natural antifungal agents may be useful for treatment of patients when organisms display in-vitro susceptibility to these agents. The test is performed by using standardized techniques and filter paper disks impregnated with the listed agent. Relative activity is reported for each natural agent based upon the diameter of the zone of inhibition or no growth zone 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 activity is defined for the natural agents tested.

NON-ABSORBED ANTIFUNGALS		
	LOW SENSITIVITY	HIGH SENSITIVITY
Nystatin		

Non-absorbed antifungals may be useful for treatment of patients when organisms display in-vitro susceptibility to these agents. The test is performed using standardized commercially prepared disks impregnated with Nystatin. Relative activity is reported based upon the diameter of the zone of inhibition or no growth zone surrounding the disk.

Sample Report

AZOLE ANTIFUNGALS			
	RESISTANT	S-DD	SUSCEPTIBLE
Fluconazole			
Itraconazole			
Ketoconazole			


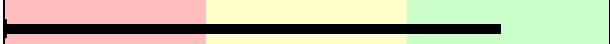



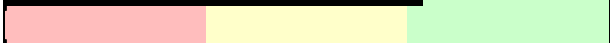
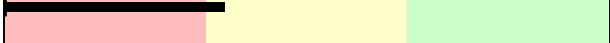
Susceptible results imply that an infection due to the fungus may be appropriately treated when the recommended dosage of the tested antifungal agent is used. **Susceptible - Dose Dependent (S-DD)** results imply that an infection due to the fungus may be treated when the highest recommended dosage of the tested antifungal agent is used. **Resistant** results imply that the fungus will not be inhibited by normal dosage levels of the tested antifungal agent.

Standardized test interpretive categories established for *Candida* spp. are used for all yeast isolates.

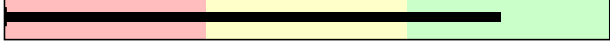
SPECIMEN DATA	
Date Collected:	10/24/2020
Date Received:	10/25/2020
Date Reported:	11/07/2020
Methodology:	Disk Diffusion

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Yeast Susceptibilities; Geotrichum spp.

NATURAL ANTIFUNGALS		
	LOW SENSITIVITY	HIGH SENSITIVITY
Berberine*		
Caprylic Acid*		
Uva Ursi*		
Plant Tannins*		
Oregano*		
Undecylenic Acid*		
Grapefruit Seed Extract*		

Natural antifungal agents may be useful for treatment of patients when organisms display in-vitro susceptibility to these agents. The test is performed by using standardized techniques and filter paper disks impregnated with the listed agent. Relative activity is reported for each natural agent based upon the diameter of the zone of inhibition or no growth zone 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 activity is defined for the natural agents tested.

NON-ABSORBED ANTIFUNGALS		
	LOW SENSITIVITY	HIGH SENSITIVITY
Nystatin		

Non-absorbed antifungals may be useful for treatment of patients when organisms display in-vitro susceptibility to these agents. The test is performed using standardized commercially prepared disks impregnated with Nystatin. Relative activity is reported based upon the diameter of the zone of inhibition or no growth zone surrounding the disk.

Sample Report

SPECIMEN DATA
<p>Date Collected: 10/24/2020 Date Received: 10/25/2020 Date Reported: 11/07/2020 Methodology: Disk Diffusion</p>

*This test was developed and its performance characteristics determined by Doctor's Data Laboratories 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.

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.

Microbiology

Beneficial Flora

One or more of the expected or beneficial bacteria are low in this specimen. Normally abundant bacteria include *Lactobacillus* spp, *Bifidobacteria* spp, *Clostridium* spp, *Bacteroides fragilis* group, *Enterococcus* spp, and *Escherichia coli*. The beneficial flora have many health-protecting effects in the gut, and as a consequence, are crucial to the health of the whole organism. Some of the roles of the beneficial flora include digestion of proteins and carbohydrates, manufacture of vitamins and essential fatty acids, increase in the number of immune system cells, break down of bacterial toxins and the conversion of flavonoids into anti-tumor and anti-inflammatory factors. *Lactobacilli*, *bifidobacteria*, *clostridia*, and *enterococci* secrete lactic acid as well as other acids including acetate, propionate, butyrate, and valerate. This secretion causes a subsequent decrease in intestinal pH, which is crucial in preventing an enteric proliferation of microbial pathogens, including bacteria and yeast. Many GI pathogens thrive in alkaline environments. *Lactobacilli* also secrete the antifungal and antimicrobial agents lactocidin, lactobacillin, acidolin, and hydrogen peroxide. The beneficial flora of the GI tract have thus been found useful in the inhibition of microbial pathogens, prevention and treatment of antibiotic associated diarrhea, prevention of traveler's diarrhea, enhancement of immune function, and inhibition of the proliferation of yeast.

In a healthy balanced state of intestinal flora, the beneficial bacteria make up a significant proportion of the total microflora. Healthy levels of each of the beneficial bacteria are indicated by either a 2+, 3+ or 4+ (0 to 4 scale). However, in some individuals there is an imbalance or deficiency of beneficial flora and an overgrowth of non-beneficial (imbalance) or even pathogenic microorganisms (dysbiosis). 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.

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, 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.

Antibacterial and antifungal susceptibility testing to a variety of prescriptive and natural agents may be provided for the pathogenic organisms that are cultured from this patient's specimen. This testing is intended to provide the practitioner with useful information to help plan an appropriate treatment regimen. A comprehensive program may be helpful in individuals in whom a dysbiotic condition has caused extensive GI damage.

Note: Not all genera or species can be tested for susceptibilities in the laboratory due to their specific growth requirements. In addition, the Centers for Disease Control and Prevention recommend not testing certain organisms such as those associated with food poisoning. If a practitioner has specific questions, please contact customer service.

Clostridium spp

Clostridia are expected inhabitants of the human intestine. Although most *clostridia* in the intestine are not virulent, certain species have been associated with disease. *Clostridium perfringens* is a major cause of food poisoning and is also one cause of antibiotic-associated diarrhea. *Clostridioides difficile* is a causative agent in antibiotic-associated diarrhea and pseudomembranous colitis. Other species reported to be prevalent in high amounts in patients with Autistic Spectrum Disorder include *Clostridium histolyticum* group, *Clostridium* cluster I, *Clostridium boltea*, and *Clostridium tetani*.

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.

Cultured Yeast

Small amounts of yeast (+1) may be present in a healthy GI tract. However higher levels of yeast (> +1) are considered to be dysbiotic. A positive yeast culture and sensitivity to prescriptive and natural agents may help guide decisions regarding potential therapeutic intervention for yeast overgrowth. When investigating the presence of yeast, disparity may exist between culturing and microscopic examination. Yeast grows in colonies and is typically not uniformly dispersed throughout the stool. Further, some yeast may not survive transit through the intestines rendering it unviable for culturing. This may lead to undetectable or low levels of yeast identified by culture, despite a significant amount of yeast visualized microscopically. Therefore, both microscopic examination and culture are helpful in determining if abnormally high levels of yeast are present.

Sample Report

Parasitology

Parasites

Parasites were detected by microscopic examination in this stool specimen. Intestinal parasites are abnormal inhabitants of the GI tract that live off and have the potential to cause damage to their host. Factors such as contaminated food and water supplies, day care centers, increased international travel, pets, carriers such as mosquitoes and fleas, and sexual transmission have contributed to an increased prevalence of intestinal parasites.

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 and eradicated. If left untreated, chronic parasitic infections can cause damage to the intestinal lining and can be an unsuspected cause of illness and fatigue. 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, decreased immune function, and fatigue.

Blastocystis spp

Blastocystis spp was identified in this specimen. *Blastocystis* is a common protozoan found throughout the world. *Blastocystis* is transmitted via the fecal-oral route or from contaminated food or water. Whether *Blastocystis* infection can cause symptoms is still considered controversial. Symptoms may be compounded by concomitant infection with other parasitic organisms, bacteria, or viruses. Often, *Blastocystis* is found along with other such organisms. Nausea, diarrhea, abdominal pain, anal itching, weight loss, and excess gas have been reported in some persons with *Blastocystis* infection.

Metronidazole has been traditionally considered the most effective drug (recommended adult dosage varies from 250 mg bid for 5-7 days to 750 mg tid x 10 days). Iodoquinol is also an effective medication (650 mg tid x 20 days). Recommended therapy can also eliminate *G. lamblia*, *E. histolytica* and *D. fragilis*. Various herbs may be effective, including oil of oregano. Limit refined carbohydrates in diet.

Endolimax nana

Endolimax nana, an amoeba, was identified in this specimen. *E. nana* is generally considered nonpathogenic or commensal. It lives in the large intestine of humans, mainly at the level of the cecum and feeds on bacteria. Infection occurs via fecal-oral route, and indicates increased risk of exposure to potential pathogens. Some research indicates that infection with *Endolimax nana* may be associated with diarrhea, urticaria, or reactive arthritis, possibly due to prolonged antigenic stimulation with formation of circulating antigen antibody complexes.

As *E. nana* is generally considered nonpathogenic, there is no treatment suggested in the Sanford Guide or Medical Letter. Natural agents include oil of oregano and quassia.

Sample Report

Stool Chemistries

Secretory IgA (sIgA) Low

The concentration of sIgA is abnormally low in this fecal specimen. Secretory IgA represents the first line of defense of the gastrointestinal (GI) mucosa and is central to the normal function of the GI tract as an immune barrier. Immunological activity in the gastrointestinal tract can be accessed via fecal sIgA levels in a formed stool sample. However, sIgA may be artefactually low due to fluid dilution effects in a watery or loose/watery stool sample.

Chronic mental and physical stress as well as inadequate nutrition have been associated with low fecal sIgA concentrations. This includes dietary restrictions, excessive alcohol intake, body mass loss, negative moods, and anxiety. One study found decreased levels of sIgA in malnourished children, particularly protein malnourishment, which responded well to nutritional rehabilitation with a significant increase in sIgA. A possible explanation for this may be the synthesis and expression of sIgA requires adequate intake of the amino acid L-glutamine. An increase of dietary L-glutamine may restore GI immune function by protection of cells that synthesize sIgA. *Saccharomyces boulardii* is a nonpathogenic yeast that has been used for the treatment of acute infectious enteritis and antibiotic-associated diarrhea. Restored levels of sIgA and subsequent enhanced host immune response have been found following *S. boulardii* administration (animal models). With low sIgA one might consider a salivary cortisol test.

Short Chain Fatty Acids (SCFAs)

The total concentration and/or percentage distribution of the primary short chain fatty acids (SCFAs) are abnormal in this specimen. Beneficial bacteria that ferment non-digestible soluble fiber produce SCFAs that are pivotal in the regulation of intestinal health and function. Restoration of microbial abundance and diversity, and adequate daily consumption of soluble fiber and polyphenols can improve SCFA status.

The primary SCFAs butyrate, propionate and acetate are produced by predominant commensal bacteria via fermentation of soluble dietary fiber and intestinal mucus glycans. Key producers of SCFAs include *Faecalibacterium prausnitzii*, *Akkermansia muciniphila*, *Bacteroides fragilis*, *Bifidobacterium*, *Clostridium* and *Lactobacillus* spp. The SCFAs provide energy for intestinal cells, and regulate the actions of specialized mucosal cells that produce anti-inflammatory and antimicrobial factors, mucins that constitute the mucus barriers, and gut active peptides that facilitate appetite regulation and euglycemia. The SCFAs also contribute to a more acidic and anaerobic microenvironment that disfavors dysbiotic bacteria and yeast. Abnormal SCFAs may be associated with dysbiosis (including insufficiency dysbiosis), compromised intestinal barrier function (intestinal permeability) and inappropriate immune and inflammatory conditions.

Stool Chemistries continued...

“Seeding” with supplemental probiotics may contribute to improved production and status of SCFAs, but it is imperative to “feed” the beneficial microbes. Sources of soluble fiber that are available to the microbes include chick peas, beans, lentils, oat and rice bran, fructo- and galacto- oligosaccharides, and inulin.

Sample Report