Small bowel diseases causing chronic diarrhea
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CHAPTER 1

DISEASES OF THE SMALL BOWEL IN CHRONIC DIARRHEA
ABSTRACT

The incidence of chronic diarrhea in (big hospitals at Asian countries) is between 0.8 - 1.0%. The diseases and abnormalities according to the location, which can cause chronic diarrhea, are divided into three locations: the small bowel, the large bowel and extraintestinal. The small bowel disorders include infectious and non-infectious diseases. The infectious diseases are bacterial infections, parasitic infections etc. and the non-infectious diseases include Crohn’s disease, Celiac sprue, Non Steroid Antiinflammatory Drugs (NSAID) enteropathy, lactose intolerance, benign tumor, carcinoid tumor, carcinoma, post surgery complications, laxative etc. The approach of diagnosis include a good medical history, careful physical examination, and laboratory tests. More specialized supporting examinations are X-ray of the colon, esophagastroduodenum, follow through, enteroclysis, ileo-colonoscopy and endoscopy of the upper portion of the digestive tract including small intestine with biopsy for histopathology examinations.

Keywords: Small bowel, chronic diarrhea, classification, etiology, approach.
INTRODUCTION

Chronic diarrhea is a condition that usually causes problems both for patients and clinicians. A morbidity study in North Jakarta(1) revealed that the number of patients with chronic diarrhea admitted to the hospital was around 1.0% of all patients with adults diarrhea, whereas the rate in Bangladesh was 0.8%.(2) Sometimes the diagnosis of this disease is difficult and treatment is not always successful. The quality of life of the patients with chronic diarrhea is frequently impaired. The Case Fatality Rate (CFR) of chronic diarrhea in children in Jakarta is about 20%. (3)

The etiologies of chronic diarrhea vary from one location to another. Some experts categorize the etiology into osmotic, secretory, abnormal motility, abnormal permeability, etc while others group of them into functional and organic or into infective and non-infective. (1,2,3,4) The diseases and abnormalities according to the location can be divided into three: the small bowel, the large bowel, and extraintestinal. Small bowel diseases, which play an important role in chronic diarrhea, are highly variable too. They include infectious (bacterial, parasitic etc.) and non-infectious (Crohn's disease, Celiac Disease, benign tumor, carcinoid tumor, carcinoma etc.).(4,5,6,7,8,9,10,11) Usually, watery and non-bloody non-steatorrheic stools are caused more by small bowel diseases and bloody stools are caused mainly by large bowel diseases. (4,5,6,7)

Patients who suffer from chronic diarrhea usually undergo unnecessary and costly examinations and treatments. We will discuss the definition, etiology, and diagnostic approach of chronic diarrhea especially in cases of small bowel abnormalities/diseases.

NORMAL SMALL BOWEL

The small bowel (small intestine) extends from the pylorus to the ileocecal valve and, depending on the tone of its muscles, measures from 3.5 to 6.5 meters in length. (10,11,12) It is divided into three regions, namely, duodenum (the first 25 cm), jejunum (the proximal 40% of the remainder of the small intestine), and ileum (the distal 60%).

A number of recent publications confirmed the important role of enteroscopy (small bowel endoscopy) in clinical practice, especially in chronic diarrhea cases. (11,13,14,15,16) The endoscopical examination of the small bowel is characteristic. Normally, the inner lining of the duodenal bulb is free of folds, but the other portions demonstrate characteristic folds of Kerckring. The papilla of Vater, located along the medial wall of the descending duodenum 3 to 6 cm distal to the apex of the bulb, corresponds to the main papilla which drains both the bile duct and the pancreatic duct. An accessory papilla (papilla of Santorini) may be located 2 to 4 cm proximal to the papilla of Vater along the medial duodenal wall. (10,12,17)

In normal histologic appearance, the villi are not all perfectly upright, finger-like structures standing in a row perpendicular to the lumen. (18) Rather, many villi tend to bend in different directions and to vary in their structure, ranging from slender, index finger-like structures to plumper, thumb-like structures with corrugated edges. Furthermore, not all villi are cylindrical with circular cross sections; many are normally leaf shaped, with elliptical cross sections.

In the first and the second portions of the duodenum, the underlying Brunner glands may be covered by broadened or shortened villi. Treitz ligament is located near the duodenojejunal junction where Brunner glands are normally rare. Villi on top of or adjacent to normal lymphoid nodules may be shortened, broadened, or absent. If one were to examine only those altered sections adjacent to a large lymphoid nodule without knowing that it was there, misinterpretation would be possible. (19)

The predominant cell covering the jejunal villi is the columnar absorptive cell (enterocyte). It has a refractile brush border. Occasional dark staining inter-epithelial lymphocytes are seen. Mitoses are confined to the crypt region and Paneth cells are located only at the crypt bases. The most common round cell of the normal lamina propria is the plasma cell, and it tends to concentrate around the bases of the crypts. Lymphocytes are fairly frequent and an occasional lymphoid nodule is normal. Isolated eosinophilic leukocytes are usually found. Occasionally
pigmented, iron-containing macrophages are seen near the tips of the villi. Polymorphonuclear leukocytes are not usually seen outside blood vessels within the lamina propria of the normal small intestine. In the normal submucosa there are blood vessels, nerve ganglion cells, lymphoid cells and lymphatics.(19)

Normally, the terminal ileum mucosa is smooth, and blood vessels may be seen, especially with air insufflation. Kerckring folds are less marked in the terminal ileum than in the duodenum and jejunum. The appearance of the mucosa is not glistening as in the colon but more villous and dull as in the duodenum.

DEFINITION OF DIARRHEA

Diarrhea is defined as the passage of stool of more than 200g per day or the passage of soft or watery stools more than 3 times per day with or without blood and/or mucous.(7,8,10,20,21)

The duration of chronic diarrhea can vary substantially. Some authors suggest that diarrhea is chronic if the duration exceeds 15 days.(4,5,8) Others suggest that the term chronic can only be used for diarrhea lasting at least 30 days or more.(7,10,11,20,21) Another authors suggest that diarrhea is categorized as chronic if it lasts for at least 3 to 6 weeks.(22)

Several experts use the term “persistent diarrhea” to refer to a diarrheal episode that begins acutely and continues for more than 2 weeks. This term does not include patients with chronic diarrhea due to non-infectious causes, such as gluten sensitive enteropathy, granulomatous disease or tumor producing gastro-intestinal hormones.

The term infective or infectious diarrhea will be used if infection is found to be the cause. The case will be diagnosed as a non-infective or non-infectious diarrhea if infection has been ruled out as being the cause.(8,9,23)

CLASSIFICATION

There are many classifications for diarrhea, such as infective or non-infective, organic or functional, stool characteristics etc.(23,24,25) Powel(24) groups the malabsorptive, secretory and inflammatory types of diarrhea into what he calls “enterocyte damage and death with inflammatory diarrhea”. He suggests a classification of chronic diarrhea into:

1. Steatorrhea (fat malabsorption).
2. Watery diarrheas that respond to fasting;
3. Watery diarrheas that may or may not respond to fasting (diarrheas of mixed or uncertain pathophysiology).
4. Watery diarrheas that do not respond to fasting (true secretory diarrheas).
5. Inflammatory diarrheas

ETIOLOGY

The abnormalities or diseases occurring in the small bowel, which are responsible for chronic diarrhea, include:(2,3,5,6,7,24,25,26)

A. Infective or infectious:

1. Bacterial infection and bacterial overgrowth: Pathogenic *Escherichia coli*, *Shigella* (dysenteriae, flexneri, etc.), *Staphylococcus aureus*, *Clostridium perfringens*, *Yersinia enterocolitica*, *Mycobacterium tuberculosis*, *Mycobacterium avium*, etc.
2. Protozoa: *Giardia lamblia*, *Cryptosporidium*, *Isospora*, *Microsporidia* etc.
3. Helminths: *Ascaris lumbricoides*, *Strongyloides*.
4. Fungus: *Candida albicans*.
5. Tropical sprue.
B. Non-infective or non-infectious:

1. Drugs or medications:
   - Laxatives such as magnesium sulfate, dioctil sodium sulfosuccinate.
   - Antacids containing magnesium hydroxide.
   - Sodium anion
   - Antibiotics: erythromycin and neomycin.
   - Antihypertensive drugs.
   - Antiarrhythmic agents.
   - Antineoplastic agents.
   - Sweeteners (sorbitol, fructose).
   - Ethanol.
   - Caffeine.

2. Hormones and neurotransmitters: Vasoactive Intestinal Polypeptide (Vipoma), Secretin, Prostaglandin E, Cholecystokinin, Cholinergic (acetylcholine), Serotonin, Calcitonine, Gastric Inhibitory Polypeptide, Glucagon and P substance.

3. Metabolic disturbances or abnormal motility of the small bowel: Diabetes Mellitus, Scleroderma, Amyloidosis, Thyrotoxicosis or hyperthyroidism, adrenal insufficiency, post gastric resection or vagotomy, diverticular disease.

4. Disturbance of electrolyte transport in the enterocyte: Congenital Chloride Diarrhea

5. Malabsorption:
   - Fat malabsorption:
     a. Intraluminal: Somatostatinoma
     b. Mucosal:
        - Drugs: colchicine, neomycin, p-aminosalicylic acid (PAS) and non-steroidal antinflammatory drugs (NSAID)
        - Immune system diseases: systemic mastocytosis, eosinophilic gastroenteritis
        - Celiac sprue
        - Dermatitis herpetiformis
        - Whipple's disease
        - Abetalipoproteinemia
     c. Post mucosal: Intestinal lymphangiectasia
   - Carbohydrate malabsorption:
     - Sorbitol and fructose diarrhea
     - Rapid intestinal transit
     - Glucose-galactose malabsorption
     - Lactase deficiency

6. Post Surgery:
   a. Bile acid diarrhea: resection or bypass of the distal ileum in Crohn's disease or postoperative adhesions
   b. Short bowel syndrome
   c. Gastrectomy
   d. Vagotomy
   e. Cholecystectomy

7. Ischemic bowel diseases

8. Radiation enteritis

9. Functional (idiopathic)

10. Inflammatory:
    a. Inflammatory Bowel Disease: Crohn’s disease
    b. Eosinophilic gastroenteritis
    c. Milk and Soy protein allergy
    d. Protein-losing enteropathy
    e. Miscellaneous: chronic mesenteric vascular ischemia, Behcet’s syndrome or Churg-
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Strauss syndrome, neutropenic enterocolitis, Cronkhite-Canada syndrome.

11. Tumor:
   - Benign: Polyp
   - Malignant: Adenocarcinoma, Carcinoid tumor, other tumor

12. Causes of steatorrhea include Crohn’s disease, radiation enteritis, celiac disease, intestinal lymphoma, bacterial overgrowth states in the small intestine and post gastrectomy syndrome.

13. Causes of non-bloody and non-steatorrhea diarrhea include exogenous agents such as laxative abuse and recognized food/drug causes, motility disturbance (diabetic autonomic neuropathy, thyrotoxicosis) and diverticular disease.

14. Others: Dihydroxy bile acid, Hydroxy fatty acid

Powell (24) describes the etiologies of chronic diarrhea as follows:

1. Steatorrhea (fat malabsorption):
   a. Intraluminal maldigestion: Cirrhosis and bile duct obstruction, bacterial overgrowth, pancreatic exocrine insufficiency, cystic fibrosis and somatostatinoma
   b. Mucosal malabsorption: drugs [colchicine, cholestyramine, neomycin, paraaminosalicylic acid (PAS) and non steroidal anti inflammatory drugs (NSAID)], infectious diseases (parasites like Giardia, Cryptosporidium, isospora, helminth such as strongyloides; mycobacterium avium; AIDS), immune system diseases (systemic mastocytosis, eosinophilic gastroenteritis), tropical sprue, Celiac sprue, dermatitis herpetiformis, Whipple’s disease, abetalipoproteinemia.
   c. Post mucosal malabsorption: intestinal lymphangiectasia
   d. Mixed causes of steatorrhea: short bowel syndrome, metabolic diseases (thyrotoxicosis adrenal insufficiency), protein calorie malnutrition, liver disease.

2. Watery Diarrheas that respond to fasting:
   a. Ingestion of non-absorbable solutes: magnesium-induced diarrhea and sodium anion diarrheas.
   b. Carbohydrate malabsorption: sorbitol and fructose diarrhea, lactose diarrhea, rapid intestinal transit and glucose-galactose malabsorption.
   c. Bile Acid Diarrhea:
      - Severe disease, resection, or bypass of the distal ileum: In Crohn’s disease or postoperative adhesions.
      - Primary bile acid malabsorption: congenital or acquired.
      - Bile acid malabsorption following upper abdominal surgery, either truncal vagotony or cholecystectomy
   d. Postvagotomy diarrhea

3. Watery diarrheas that may or may not respond to fasting (diarrheas of mixed or uncertain pathophysiology): Irritable bowel syndrome, food allergy, microscopic colitis (collagenous and microscopic colitis, pericrypt eosinophilic enterocolitis)

4. Watery diarrheas that do not respond to fasting (true secretory diarrheas): carcinoid syndrome, gastrinoma, vipoma or watery diarrhea-hypokalemia achlorhydria (WDHA) syndrome, medullary carcinoma of the thyroid, glucagonoma, villous adenomas, systemic mastocytosis, factitious diarrhea, chronic idiopathic diarrhea and pseudopancreatic cholera syndrome, diabetic diarrhea, alcoholic diarrhea, congenital secretory diarrhea.

5. Inflammatory diarrheas:
   a. Inflammatory bowel disease: Crohn’s of the small or large intestine or ulcerative colitis.
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b. Eosinophilic gastroenteritis
c. Milk and soy protein allergy
d. Protein-losing enteropathy
e. Chronic radiation enterocolitis
f. Miscellaneous diseases: Acute mesenteric arterial or venous thrombosis, chronic mesenteric vascular ischemia, gastrointestinal tuberculosis and histoplasmosis, Behcet's syndrome or Churg-Strauss syndrome, acute graft-versus-host disease after allogeneic bone marrow transplantation, Neutropenic enterocolitis, Cronkhite-Canada syndrome.

MEDICAL HISTORY AND CLINICAL SYMPTOMS

Steatorrhea is a type of diarrhea where the stool has a high fat content.(26) It is a rather watery stool with a light color, a strong smell, a tendency to float and it is difficult to wash off. Sometimes we can see a layer of oil on the surface of the water. Steatorrhea indicates a maldigestion or malabsorption of fat. In most cases, the clinical symptoms are not specific and are related to the effect of malabsorption of nutrients and lack of vitamins and electrolytes. Although the three main nutrients-fat, carbohydrate and protein-can all be affected by malabsorption, normally the clinical symptoms are caused by malabsorption of carbohydrate or fat.(24) Malabsorption of protein or amino acid (azotorrhea) can occur, but will be difficult to detect unless the condition is already so advanced that it has caused malnutrition—the exception are cases where defects in the transport of specific amino acids have resulted in a systemic congenital illness.

Malabsorption of electrolyte and water is part of the pathophysiology of malabsorptive diarrhea. Diarrhea with blood indicates a problem in the rectum or the left part of the colon. Diarrhea with blood also indicates mucosa ulceration. Clinical symptoms depend on etiology.

The clinical symptoms of bloodless diarrhea and non-steatorrheic also depend on the etiology. A patient with irritable bowel syndrome (IBS) usually shows a generally good condition, making his complaints disproportional to his physical condition. His diarrhea rarely occurs at night, intermittent with constipation and accompanied by abdominal pain and rumbling. Functional dyspepsia and other non-specific complaints can result from this disease. Frequently the patient can establish a connection between diarrhea and stress or pressure. The diarrhea will disappear or abate whenever he is relaxed or on holiday. Cases of thyrotoxicosis may also cause bloodless diarrhea and non-steatorrhea. In these cases, the patient usually has fever, perspires, palpitates and suffers weight loss. Laxative abuse is sometimes difficult to diagnose, and therefore a good medical history is crucial for the diagnosis.

Microscopic colitis, lymphocytic and collagenous colitis are defined as an inflammation of the colorectal mucosa that is diffuse at histologic examination. These conditions do not show any abnormality during endoscopic or X-ray barium tests. The symptom consists of a chronic watery diarrhea with no specific cause.(27,28) The connection between an immunologic disease and microscopic colitis indicates a possibility of an immunologic process. Drugs such as non-steroid anti-inflammation drug (NSAID) and vascular tonic should be considered as the cause. Microscopic colitis predominantly occurs in female patients aged 50-60, but certain cases can occur in all age groups including children. Bloodless, non-steatorrheic diarrhea can occur continuously or intermittently, with spontaneous or treatment-caused remission or relapses. Sometimes the patient complains of abdominal pain, nausea or vomiting. The patient is generally in a good condition and laboratory results are usually normal.

A 72-hour oral/enteral fast with intravenous hydration can occasionally determine the cause of the diarrhea. If it completely stops within 24 hours, it will be unnecessary to continue the fasting, which is usually due to osmotic diarrhea. Secretorific diarrhea usually ceases significantly with fasting, but may continue with more than 200 gram of stool every 24 hours. Diarrhea caused by malabsorption of food will stop with fasting.
PHYSICAL EXAMINATION

Conditions observed during the physical examination are usually caused by nutrient, vitamin and electrolyte deficiencies. Patients with chronic diarrhea may suffer weight loss down to a state of malnutrition, edema, muscle hypotrophy, dry skin, anemia, glossitis, dermatitis, paresthesia, peripheral neuropathy, increased susceptibility to bleeding and bruising, night blindness, weakness, tetany, bone pain, loss of hair, symptoms of dehydration and shock, pain when the abdomen is pressed; in the case of colon cancer, a lump may be palpable, etc. (26)

Loss of weight down to the state of malnutrition is usually caused by shortage of fat, protein and calories. Edema and muscle hypotrophy are the results of protein deficiency. Dry and snake-like skin is caused by deficiency of essential fatty acids and zinc. Anemia is caused by deficiency of iron, folic acid and vitamin B12. Glossitis and dermatitis are usually caused by deficiency in nicotinic acid. Paresthesia and peripheral neuropathy are usually the results of deficiencies of vitamins B6 and B12. Deficiency of vitamin K causes the patient to bleed or bruise easily. Night blindness is caused by deficiency of Vitamin A. Weakness is caused by deficiencies in Potassium, Sodium and Magnesium. Tetany and bone pain are usually caused by deficiency in Calcium, while loss of hair is usually the result of deficiencies in zinc and protein.

Other irregularities observed during physical exam are usually determined by the original disease or the etiology of the chronic diarrhea.

In patients with pancreatitis or pancreatic cancer, there will be abdominal pain with thrombosis.

In patients with Coeliac disease, the symptoms can be a small stature, delayed menarche, mouth ulcers, eruption of itchy skin, or evidence of dermatitis herpetiformis.

In the case of Whipple's disease, there will be polyarthritis with pigmentation

Patients with amyloidosis or rheumatoid arthritis will present with polyarthritis.

In patients with mesenteric ischemia and Crohn's disease, we will see abdominal angina, mouth ulcer, perianal fistel, sub-acute intestinal obstruction, abdominal masses, lymphadenopathy, lymphoma.

In biliary cirrhosis and sclerosing cholangitis, icterus may be noticeable.

In post-gastrectomy (Bill roth II), we will see scars on the abdomen, and blind loop formation is possible.

SUPPORTING EXAMINATIONS

Quite a few supporting examinations will be required, including three lots of stool tests that consist of routine tests, occult blood test, analysis of digestion, quantitative and qualitative tests of fat in stool, test for parasites in stool, tests for microbiology, routine blood test, blood chemistry, with liver functions, and kidney functions, blood glucose, test for antibody against amoebas, test for typhoid and virus infections, tests for HIV when suspected. Other supporting examinations will also be required such as ultrasonography of the abdomen, CT-scan of the abdomen, Ileocolonoscopic examination with biopsy for histopathology, gastroduodenojejunoscopic test with biopsy for histopathology, photography of the intestine (esophagogastro-duodenum follow-through or enteroclysis), endoscopic retrograde cholangiopancreatography (ERCP), magnetic resonance cholangio pancreatography (MRCP), H2 breath testing, etc. In selecting the supporting tests, especially for developing countries, attention should be given to the aspects of cost effectiveness and cost-benefit.

In the case of malabsorption, a special type of laboratory supporting test will be required to establish the clinical symptoms and certain pathophysiological mechanisms. If the patient suffers weight loss, a test of fat and serum protein will be required. (24) If the stool contains oil, it will be necessary to find out the amount of fat in the stool. If the fat in the stool exceeds 6 grams/24 hours, the condition is viewed as steatorrhea. In the case of anemia, it will be necessary to perform blood tests to find out if there are deficiencies of iron, pyridoxine, folic acid and vitamin B12. In the case of hemorrhagic diathesis, a prothrombin test is required. If
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the patient complains of bone pain, tests have to be conducted to determine the amounts of calcium, magnesium, phosphate and alkaline phosphatase. In the case of dermatitis and fish-like skin, it is necessary to check the contents of zinc serum in the urine.

DIAGNOSTIC APPROACHES

There are several different approaches to diagnosing chronic diarrhea, depending on the location, population, the purpose of the diagnosis and the specific consideration of each gastroenterology expert. Despite the variety, these procedures share the same principle, namely, it should begin with a medical history, followed by a physical examination, supporting laboratory tests, more specialized supporting examinations (including X-ray of the colon, esophagogastroduodenum follow-through, enterolysis, ileo-colonoscopy and endoscopy on the upper portion of the digestive tract including the small intestine (using duodeno-jejunoscopy) with biopsy for histopathology examinations (see Figure 1) (7,24,29,30)

It is fairly difficult to distinguish between problems in the small intestine, the colon or other organs. It often happens that problems in both the intestine and the colon occur at the same time. Usually, a diarrhea with blood is caused by a problem in the colon, although in rare cases it can be caused by a problem in the intestine. To determine whether the problem is in the intestine, we have to start by examining the colon or the other organs including the pancreas.

What is important is all the history data that support the probability of the case being an organic or a functional diarrhea. By organic diarrhea we mean a case of diarrhea that is caused by a disorder of the intestine that has been confirmed based on histological or biochemical tests. On the other hand, by functional diarrhea we mean idiopathic diarrhea, a diarrhea caused by food or a diarrhea caused by a problem in the intestine’s motility. Medical history, physical examination and laboratory screening test can determine 30 % of etiology of chronic diarrhea. A complete stool test, a proctosigmoidoscopy with a biopsy can determine a higher rate of the etiology of a chronic diarrhea—approximately 50%. With external evaluation, including collection of stool for 24 hours and fasting, an etiological diagnosis of 70-80% is attainable.

The diagnosis criteria to an organic disorder are the following: 1. The diarrhea lasts for a shorter period of time—usually less than three months; 2. The diarrhea is more dominant at night (nocturnal); 3. It occurs every day, not intermittently; 4. The onset was sudden; 5. It was followed by a weight loss exceeding five kilograms; 6. There is a history of dehydration or hypokalemia; 7. The erythrocyte sedimentation rate is high; 8. The level of haemoglobin is low; 9. The average volume of stool per day exceeds 400 g (some experts use 225 g as the borderline value). (22,23,24,29) The occurrence of at least three of these criteria is highly suspicion for an organic disorder, and the specificity of organic problems can reach 90 % or more. On the other hand, age, the presence of an abdominal pain, the number of times the patient has watery stool per day, and the description of the stool cannot serve to distinguish between an organic and a functional diarrhea. Incontinencia can occur in both of these diarrhea types.

For the test of laxative drugs use, see Table 1.

Table 1. Laboratory Evaluation for Laxative Abuse

<table>
<thead>
<tr>
<th>Test Description</th>
<th>Protocol</th>
</tr>
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<tbody>
<tr>
<td>Barium enema to test for cathartic colon (ahuastral right colon)</td>
<td>Specified by Donowitz(29)</td>
</tr>
<tr>
<td>Sigmoidoscopy for gross presence of pseudo-melanosis coli</td>
<td></td>
</tr>
<tr>
<td>Alkalination assay of stool: Phenolphthalein, some anthraquinones, and rhubarb</td>
<td></td>
</tr>
<tr>
<td>bisacodyl turns purple-blue</td>
<td></td>
</tr>
<tr>
<td>Spectrophotometry* or thin-layer chromatography of urine or stool water:</td>
<td></td>
</tr>
<tr>
<td>Detects anthraquinones, bisacodyl, phenolphthalein; can detect anthraquinones</td>
<td></td>
</tr>
<tr>
<td>&gt;32 hr after one dose</td>
<td></td>
</tr>
<tr>
<td>Measurement of stool osmollality: only useful with an osmolality of &lt;250 mOsm per</td>
<td></td>
</tr>
<tr>
<td>kilogram (implying dilution of stool with water or urine)</td>
<td></td>
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<tr>
<td>Stool osmotic gap: if &gt; 50 mOsm per kilogram is seen, measure stool magnesium</td>
<td></td>
</tr>
<tr>
<td>(normally &lt; 45 mmol per liter or &lt; 30 meq per day)</td>
<td></td>
</tr>
<tr>
<td>Measurement of stool sulfate and phosphate</td>
<td></td>
</tr>
</tbody>
</table>

cited from Donowitz(29)
Figure 1. Diagnostic Approach to the Etiology of Chronic Diarrhea
Table 2. Classification of Types of Chronic Diarrhea on the Basis of Responsiveness to Fasting

<table>
<thead>
<tr>
<th>Types (or causes) responsive to fasting</th>
</tr>
</thead>
<tbody>
<tr>
<td>Incontinence</td>
</tr>
<tr>
<td>Bile acid diarrhea</td>
</tr>
<tr>
<td>After cholecystectomy</td>
</tr>
<tr>
<td>After ileal resection</td>
</tr>
<tr>
<td>Steatorrhea</td>
</tr>
<tr>
<td>Osmotic diarrhea</td>
</tr>
<tr>
<td>Carbohydrate malabsorption</td>
</tr>
<tr>
<td>Excessive carbohydrate ingestion</td>
</tr>
<tr>
<td>Laxative (containing poorly absorbable anions: sodium sulfate, sodium phosphate, or sodium citrate, magnesium)</td>
</tr>
<tr>
<td>Food allergy</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Types (or causes) not responsive or only partly responsive to fasting</th>
</tr>
</thead>
<tbody>
<tr>
<td>Laxative or diuretic abuse</td>
</tr>
<tr>
<td>Inflammatory bowel diseases</td>
</tr>
<tr>
<td>Coeliac sprue</td>
</tr>
<tr>
<td>Intestinal lymphoma</td>
</tr>
<tr>
<td>Neuroendocrine tumors</td>
</tr>
<tr>
<td>Zollinger-Ellison syndrome</td>
</tr>
<tr>
<td>Pancreatic cholera</td>
</tr>
<tr>
<td>Carcinoid tumor</td>
</tr>
<tr>
<td>Medullary carcinoma of the thyroid</td>
</tr>
<tr>
<td>Systemic mastocytosis</td>
</tr>
<tr>
<td>Villous adenoma of the rectosigmoid</td>
</tr>
<tr>
<td>Chronic infection(e.g. Mycobacterium tuberculosis, Giardia, amebic infection)</td>
</tr>
<tr>
<td>Hyperthyroidism</td>
</tr>
<tr>
<td>Congenital diarrhea</td>
</tr>
<tr>
<td>Chloride-bicarbonate exchange deficiency</td>
</tr>
<tr>
<td>Sodium-hydrogen exchange deficiency</td>
</tr>
<tr>
<td>Microvillus inclusion disease</td>
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<tr>
<td>Bacterial overgrowth</td>
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</tbody>
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Cited from Donowitz(29)

Etiology of chronic diarrhea that may or may not respond to fasting is detailed in Table 2. Tests used for determining the causes of steatorrhea, bloody as well as non-bloody non-steatorrhea:(26,29)

1. Steatorrhea. This type of diarrhea can be caused by:
   a. Pancreatic disease, which requires other tests that will support the diagnosis, including Glucose Tolerance Test (GTT) examination, ultrasonography (USG) or computed tomography scan (CT-scan) of the abdomen as well as an endoscopical retrograde cholangiopancreatography (ERCP) test.
   b. Small intestine mucosal disease, which requires other tests to support the diagnosis, including GTT examination, test on xylose absorption, duodenum/jejunum biopsy, H₂ breath test with lactulose, barium meal follow-through/Enteroclysis examination and Schilling test.
   c. Liver problem, which requires additional tests including USG and CT-scan of the abdomen and ERCP.
   d. Post-gastrectomy symptoms, which require a H₂ breath test with lactulose, a follow-through/enteroclysis barium meal test.

2. Bloody diarrhea. In addition to medical history and physical examination, this type of diarrhea requires barium enema of the colon-loop and sigmoidoscopy/colonoscopy with biopsy. Stool analysis to detect presence of parasites are also necessary.

3. Non-bloody, non-steatorrheic diarrhea. In addition to clinical symptoms and physical examinations, this type of diarrhea requires other tests including tests for stool parasites.
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and colonoscopy with biopsy. In the case of irritable bowel syndrome (IBS), an exclusion of pathologic disorder of the bowel is required. Pseudomelanosis colon, a black coloration of the colon mucosa, indicative for laxative abuse, can be detected during colonoscopy.

If an HIV infection is suspected as being the cause of the disease, a blood screening for HIV infection is required as well as a special stool parasite tests such as Cryptosporidium, Isospora belli and Microsporidia. Thyroid functions test is required if thyrotoxicosis is suspected as the cause of the diarrhea.

CONCLUSION

Chronic diarrhea can be caused by a wide variety of small bowel diseases. A reliable diagnosis will require a systematic approach.

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