



Short Bowel Syndrome

Short bowel syndrome (SBS) occurs when the small intestine ceases to function properly due to trauma, disease activity, or when too much of it has been removed. SBS results in inadequate nutrient and fluid absorption as well as chronic and/or frequent diarrhea. It can lead to dehydration, malnutrition, fatigue, weight loss, and a variety of other health problems. Management might include extreme dietary modifications, medications, and further surgeries.

Intestinal Anatomy

The upper part of the gastrointestinal (GI) tract consists of the esophagus, stomach, and duodenum. The lower GI tract consists of the small intestine (about 6 metres in length) and the large intestine or colon (2 metres).

The upper 40% of the small intestine is the jejunum, which comes from the Latin word for empty. The lower 60% is the ileum, which, in Greek, means to roll or twist. These portions of the small intestine differ significantly in nature and function. The jejunum has a thicker wall and wider inside area (lumen) than the ileum. In addition, intestinal muscle contraction (peristalsis) is more forceful and rapid in the jejunum than in the ileum. Therefore, passage of material is usually faster through the upper section of the small intestine and slower as it goes farther along.

The principal function of the small intestine is to digest and absorb dietary nutrients, including proteins, carbohydrates, fats, vitamins, and minerals. The digestive tract works with support from colonic bacteria and other organs (the liver and pancreas), to break down complex foods and to extract the nutrients. Water and salt are absorbed into the body via the large intestine.

Causes

The need for surgical resection can occur for a number of

reasons, including blocked or restricted blood flow to the bowel, a strangulated intestinal hernia, Crohn's disease, gastrointestinal cancer, radiation therapy, a perforated bowel, a twisting or tangling of the small intestine (volvulus), congenital defects, and some other rare diseases. In infants and small children, the most common causes are congenital intestinal anomalies or a disease that causes bowel tissue death in premature babies (necrotizing enterocolitis).

The small intestine is quite adaptive; in fact, even with removal of up to 40% of it, appropriate digestion is still possible. However, removing more than this, or even removal of certain parts of the small intestine can have adverse consequences. For example, those who retain their duodenum, jejunum, and ileum are generally able to avoid severe complications because of the adaptive nature of the small intestine. Conversely, removing just 25% of the distal end of the small intestine that intersects with the large intestine (terminal ileum) can cause ongoing diarrhea and significant nutrient malabsorption.

Symptoms

The symptoms and severity of SBS vary according to the part of the intestine that is affected. For example, the duodenum absorbs iron, calcium, and magnesium. The jejunum absorbs fatty acids, amino acids, monosaccharides, and water-soluble vitamins. The ileum absorbs fat-soluble vitamins (A, D, E, K), vitamin B12, and bile acids. Therefore, loss of function in any one of these parts has its own implications as to what the body will end up having difficulty absorbing from the food an individual consumes.

Symptoms of SBS typically include chronic diarrhea, fatigue, bloating, cramping, weight loss, dehydration, and malnutrition. Many of the most devastating complications occur specifically because of this malnutrition.

Gastric acid hypersecretion can occur in individuals with SBS, especially in the immediate aftermath of a surgical removal of part of the intestine (resection). High levels of stomach acid can enter the altered intestine and interfere with absorption of nutrients.

Vitamin and mineral deficiencies can cause many additional aggravating symptoms, including visual disturbances, dryness of the eyes, prickling or tingling feelings on the skin, muscle spasms, easy bruising, blood clotting problems, and difficulty breathing on exertion.

Electrolyte imbalances can result when the small intestine is unable to absorb minerals such as potassium, sodium, and magnesium adequately, resulting in weakness, nausea, headaches, and irregular heartbeats.

Acidosis is an abnormally high level of lactic acid in the bloodstream resulting from the shortened bowel's reduced ability to digest carbohydrates. Undigested carbohydrates create lactic acid, which can build up due to a reduced ability to use and effectively dispose of it. Acidosis can result in confusion, blurry vision, and slurred speech.

Steatorrhea is dietary fat remaining in the feces after passing through the small intestine unabsorbed. In a normal small intestine, bile salts, which are essential for fat absorption, enter the gastrointestinal tract from the gallbladder. These bind to fat, which together are absorbed at the terminal ileum. When this part is missing or damaged, then an excess amount of fat passes in the stool. Steatorrhea causes stool that floats, is grey in colour, and is exceptionally foul smelling.

Kidney stones can also result when unabsorbed fat in the colon binds with calcium, preventing the normal combination of calcium and oxalate, a compound found in plant foods. When the colon absorbs oxalate that is unbound to calcium, the body passes it in higher than normal amounts in the urine, resulting in oxalate kidney stones.

Diagnosis

While having 50% or more of the small intestine removed is a strong indicator of SBS, physicians use these tools to make a diagnosis: blood tests, physical examination, and stool examination. Blood tests can reveal vitamin, mineral, and electrolyte deficiencies. Physical examination can help a physician observe loss of muscle mass, inability to maintain weight, and dermatological conditions related to vitamin deficiency, such as rashes or scaly skin. Stool examination is useful for identifying whether an individual is absorbing a healthy amount of fat and carbohydrates.

Management/Treatment

There are a variety of options for the management and treatment of short bowel syndrome, covering the spectrum of

nutrition, medication, and surgery.

Dietary Modifications

In the aftermath of intestinal resection, patients initially require the delivery of fluids, electrolytes, and liquid nutrients into the bloodstream through a tube placed in the vein (intravenous, or IV), called total parenteral nutrition (TPN) or parenteral nutrition (PN). This phase may last from 3-4 months. However, the major focus within 2-4 days after a surgical resection is on long-term intestinal adaptation. Depending on the extent of the resection, some individuals are able to resume a modified oral diet within a few weeks of surgery. Adaptation for others might take up to a year. Remarkably, during adaptation, the remaining intestinal villi will grow in length and thickness to compensate for the loss of intestine, thus increasing the remaining surface area for nutrient absorption.

Enteral and oral nutrition are two mechanisms that assist in intestinal adaptation. Enteral nutrition involves the delivery of a special liquid food mixture to the stomach or small intestine through a feeding tube. This mechanism stimulates adaptation of the intestinal villi. Patients must have a partially functioning GI tract for this to be effective.

Once out of the adaptation phase and able to eat orally, diet and eating habits might be able to help prevent malnutrition and dehydration. Dietitians will customize diet plans, which vary according to which parts of the remaining intestine are functioning.

General dietary guidelines include:

- eat small, more frequent meals (5-7 per day),
- drink liquids between meals instead of with them,
- consume high protein foods,
- eat complex carbohydrates, such as pasta, rice, potatoes, breads, and cereals,
- eat a low-fat diet, especially for those experiencing steatorrhea or who are missing their terminal ileum,
- limit alcohol and caffeinated beverages,
- consider limiting foods that are high in oxalate (for those without an ileum, but with retained colonic function),
- eat foods that help control diarrhea, including bananas, oatmeal, rice, tapioca, applesauce, and yogurt.

While oral nutrition is preferred, some individuals might be unable to acquire adequate nutrition through diet alone. In serious cases, even enteral nutrition might not work well and it could be necessary to use TPN long-term. However, this is complex therapy with risk of serious complications including central line infections, bone disease, and liver disease.

Medications

In addition to nutrition, a physician might prescribe medications, whether prescription or over-the-counter, to

relieve some symptoms and to address the underlying condition.

Anti-diarrheal medications slow down the time it takes food to pass through the small intestine by decreasing the rate of contraction and relaxation (peristalsis) of the intestinal muscles. Drugs in this category include diphenoxylate (Lomotil®) and loperamide (Imodium®). Codeine might also be helpful.

Gastric acid reducers come in two main types: histamine-2 receptor blockers (H2RAs) and proton pump inhibitors (PPIs), which can inhibit or reduce the release of stomach acid. Examples of H2RAs include cimetidine (Tagamet®), ranitidine (Zantac®), famotidine (Pepcid®), and nizatidine (Axid®). PPIs include omeprazole (Losec®), lansoprazole (Prevacid®), pantoprazole sodium (Pantoloc®), esomeprazole (Nexium®), rabeprazole (Pariet®), and pantoprazole magnesium (Tecta®), as well as dual delayed release PPI capsules in the form of dexlansoprazole (Dexilant®). Excessive acid passing from the stomach into the intestines can hinder intestinal adaptation and can cause pain and discomfort, especially right after surgery.

Bile acid sequestrant, cholestyramine (Olestyr®), reduces the incidence of bile-salt diarrhea by increasing the removal of bile acids from the body. As the body loses bile acids, it replaces them by converting cholesterol from the blood to bile acids, which also causes blood cholesterol levels to decrease.

Glutamine is the most abundant amino acid in the body and is involved in more metabolic processes than any other amino acid. It helps promote gut integrity by acting as an energy source and preventing the passage of intestinal bacteria through the gut mucosa into other bodily tissues and organs (bacterial translocation). Bacterial translocation can lead to buildup of harmful bacteria and their toxins (sepsis), typically through infection of a wound, and can be fatal. Glutamine acts as a major fuel source for the white blood cells of the immune system. It improves nitrogen balance, thereby helping those with SBS avoid the occurrence of nutritional stress in which the body uses more protein than it takes in. Glutamine might also reduce diarrhea because it aids in the reabsorption of sodium and water.

You can find oral glutamine in some health food stores, pharmacies, and online. Products include NutreStore® and GlutaSolve®. Discuss with your physician before using it for short bowel syndrome.

Teduglutide (Revestive®) was the first medication approved by Health Canada to treat adults and children with SBS who need extra nutrition or fluids from parenteral feeding. Teduglutide is a glucagon-like peptide (GLP-2), which helps the intestines absorb nutrients and fluid. It works by regenerating cells in the intestinal lining, improving intestinal absorption of fluids and nutrients, promoting growth in the surface area of the small intestine lining, and possibly restoring gastric emptying and secretion. Patients can inject it just under the skin (subcutaneously) once a day in the stomach area, upper legs, or upper arms.

Surgery

Regrettably, surgery might still be necessary when other therapeutic options are not working. These focus on increasing the intestines' absorptive properties and are either non-transplant or transplant surgeries.

Non-transplant surgeries involve manipulating the intestine to lengthen it. In the Bianchi procedure, a surgeon cuts a portion of the bowel lengthwise into two pieces and joins them together end to end. This results in a narrower, but longer segment of the bowel. Surgeons typically reserve the serial transverse enteroplasty (STEP) procedure for children who have enough small intestine remaining to lengthen and restore function. In the STEP procedure, a surgeon makes a series of cuts in the small intestine and staples it back together in a 'zig-zag' pattern, resulting in a narrower, longer bowel. Because of the intestine's new shape, food takes more time to move through the bowel, thus increasing its time in contact with the bowel lining and the chances of nutrient absorption.

In the case of small bowel transplantation, a surgeon removes the diseased small bowel and replaces it with a donor intestine. However, this procedure comes with potential complications, including organ rejection, infections, and the excess production of white blood cell lymphocytes that results in lymphoproliferative disease. To lessen the incidence of rejection, individuals are required to use immunosuppressive drugs. Nonetheless, surgical techniques and immunosuppressive drugs continue to improve, increasing the chances of successful transplants.

Outlook

For individuals with short bowel syndrome, especially those who have had a large part of their intestines surgically removed, the goal is to enhance intestinal adaptation, enhance nutrition, and eliminate the need for total parenteral nutrition. The future of treatments for SBS revolves around improving ways to maximize bowel adaptation and refining techniques for intestinal transplantation. For now, though, a variety of options exist for helping individuals manage symptoms.

A strong team for effective management will generally include gastroenterologists, surgeons, dietitians, nursing specialists, and pharmacists. Maintaining a strong relationship with these individuals is essential in order for those with SBS to find a treatment plan that is right for them. While there is no single diet or cure for short bowel syndrome, education and open communication with trusted medical professionals might help individuals avoid the harmful health consequences of malnutrition, dehydration, and chronic diarrhea associated with short bowel syndrome.

