Functional food for functional disorders
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Unexplained hypersensitivity to food is a major health problem affecting about 10% of the population, and is often associated with disabling symptoms, such as irritable bowel syndrome (IBS), musculoskeletal pain (fibromyalgia) and fatigue (chronic fatigue syndrome/myalgic encephalomyelitis; CFS/ME). Recent metabolomic studies suggest a hypometabolic state with excessive oxidative stress, impaired oxidative phosphorylation, and an energy production dependent on the metabolism of fat and amino acids.[1] 'Missing microbes' [2] following the use of antibiotics at an early age may be an important cause, possibly assisted by a Western diet high in simple carbohydrates.[3]

The present preliminary report is the result of a systematic survey of 438 consecutive patients examined by one of us (AB). Validated diagnostic criteria and scoring questionnaires for symptom severity were applied.[4] Briefly, the results are as follows. Age of the patients is on average 37 years; 2/3 are women. Virtually all patients have irritable bowel syndrome (IBS) with symptoms related to incomplete evacuation and altered intestinal fermentation. A selection of the patients who have undergone ultrasonography or MRI scans are found to have increased amount of fluid persisting in bowel loops one hour after the ingestion of lactulose, consistent with intestinal hypomotility.[5,6] Although perceived food intolerance was the 'entrance ticket' for medical investigation, IBS was rarely attributable to food allergy. In addition to IBS, about 70% of the patients had musculoskeletal pain and 85% chronic fatigue. About half of those with fatigue met the international criteria for an ME diagnosis, and this diagnosis correlated well with being placed on long-term sick leave. The history of illness averaged 26 years. The first symptoms were almost always related to abdominal discomfort, and very often (in about 90% of the cases) the symptoms appeared after prolonged antibiotic treatment at an early stage of life. Subsequently, the problems deteriorated, often after bouts of gastroenteritis and/or further courses of antibiotic treatment. Our results therefore suggest that both the perceived food hypersensitivity and IBS, acquired following antibiotics at early life, and the subsequently acquired musculoskeletal pain (including temporomandibular dysfunction), chronic fatigue and ME are aspects of one disease, not several different diseases.

Seminal studies show loss of lactic acid bacteria and increased growth of yeast (Candida albicans) in the bowel immediately following the use of antibiotics.[7,8] Previously, it was believed that this altered bowel flora would normalise in the course of a week. However, recent studies show that this is not always the case; sometimes the bowel flora never fully recover.[2,9] 'Missing microbes' following the use of antibiotics may lead to the growth of unfavourable flora, frequently in the form of a biofilm with yeast and facultatively anaerobic microbes.[10] This biofilm elicits little local damage, but stimulates innate immunity and the release of interferon gamma, which in turn activates indoleamine dioxygenase (IDO) and the metabolism of tryptophan along the kynurenine pathway.[11] The result is an altered immune response with increased immune tolerance, protecting the pathological flora.[12] However, tryptophan is also the precursor of important hormones such as serotonin and melatonin – and increased metabolism of tryptophan to kynurenines may lead to serotonin and melatonin deficiency.[13,14] Shortage of serotonin may explain poor intestinal peristalsis, sadness and 'brain fog', while melatonin deficiency may explain poor sleep and the feeling of never being completely refreshed. Increased physical and cognitive fatigue may largely be due to oxidative stress, possibly because a deficiency of lactic acid bacteria may result in excess oxygen levels within the bowel. By producing hydrogen peroxide, these microbes pick up remaining oxygen and thus help to maintain low oxygen tension. If the redox potential is too high, the metabolism of the host switches to an oxidative shielding response [15] with aerobic glycolysis and deficient mitochondrial energy (ATP) production (Crabtree effect).[16]

Some important aspects of the treatment are: dietary carbohydrate restriction may reduce IBS symptoms,[17] while commercial probiotic lactobacilli may...
do more harm than good.[18] Various forms of fecal microbiota transplantsations are currently being investigated.[19,20] Because tryptophan appears to play such important microbial and immunological roles in functional disorders,[21] we find it an interesting option to provide tryptophan as tryptophan-rich peptides.[22,23] Simultaneously, we are very concerned about the wide use of the herbicide glyphosate (Roundup), which blocks the synthesis of tryptophan in plants and microbes [24] and thus impoverishes our natural resources of the essential tryptophan.

**Disclosure statement**

No potential conflict of interest was reported by the authors.

**References**

[1] Naviaux RK, Naviaux JC, Li K, et al. Metabolic features of chronic fatigue syndrome. Proc Natl Acad Sci U S A. 2016;113(37):E5472–E5480.

[2] Blaser MJ. Missing microbes: how the overuse of antibiotics is fueling our modern plagues. New York (NY): Henry Holt; 2014.

[3] Sonnenburg ED, Smits SA, Tikhonov M, et al. Diet-induced extinctions in the gut microbiota compound over generations. Nature. 2016;529(7585):212–215.

[4] Berstad A, Undseth R, Lind R, et al. Functional bowel symptoms, fibromyalgia and fatigue: a food-induced triad? Scand J Gastroenterol. 2012;47:914–919.

[5] Undseth R, Berstad A, Klow NE, et al. Abnormal accumulation of intestinal fluid following ingestion of an unabsorbable carbohydrate in patients with irritable bowel syndrome: an MRI study. Neurogastroenterol Motil. 2014;26(12):1686–1693.

[6] Arslan G, Gilja OH, Lind R, et al. Response to intestinal provocation monitored by transabdominal ultrasound in patients with food hypersensitivity. Scand J Gastroenterol. 2005;40(4):386–394.

[7] Morotomi M, Watanabe T, Suegara N, et al. Distribution of indigenous bacteria in the digestive tract of conventional and gnotobiotic rats. Infect Immun. 1975;11(5):962–968.

[8] Savage DC. Microbial interference between indigenous yeast and lactobacilli in the rodent stomach. J Bacteriol. 1969;98(3):1278–1283.

[9] Sjölund M, Wreiber K, Andersson DI, et al. Long-term persistence of resistant Enterococcus species after antibiotics to eradicate Helicobacter pylori. Ann Intern Med. 2003;139(6):483–487.

[10] Hertzberger R, Arents J, Dekker HL, et al. H$_2$O$_2$ production in species of the Lactobacillus acidophilus group: a central role for a novel NADH-dependent flavin reductase. Appl Environ Microbiol. 2014;80(7):2229–2239.

[11] Fitzgerald P, Cassidy EM, Clarke G, et al. Tryptophan catabolism in females with irritable bowel syndrome: relationship to interferon-gamma, severity of symptoms and psychiatric co-morbidity. Neurogastroenterol Motil. 2008;20(12):1291–1297.

[12] Zelante T, Iannitti RG, Cunha C, et al. Tryptophan catabolites from microbiota engage aryl hydrocarbon receptor and balance mucosal reactivity via interleukin-22. Immunity. 2013;39(2):372–385.

[13] Barbaro MR, Di SA, Cremon G, et al. Interferon-gamma is increased in the gut of patients with irritable bowel syndrome and modulates serotonin metabolism. Am J Physiol Gastrointest Liver Physiol. 2016;310:G439–G447. apjgi.

[14] Bessede A, Gargaro M, Pallotta MT, et al. Aryl hydrocarbon receptor control of a disease tolerance defence pathway. Nature. 2014;511(7508):184–190.

[15] Naviaux RK. Oxidative shielding or oxidative stress? J Pharmacol Exp Ther. 2012;342(3):608–618.

[16] Molavian HR, Kohandel M, Sivaloganathan S. High concentrations of H$_2$O$_2$ make aerobic glycolysis energetically more favorable for cellular respiration. Front Physiol. 2016;7:362.

[17] Gibson PR, Shepherd SJ. Evidence-based dietary management of functional gastrointestinal symptoms: the FODMAP approach. J Gastroenterol Hepatol. 2010;25(2):252–258.

[18] Berstad A, Raa J, Midtvedt T, et al. Probiotic lactic acid bacteria - the fledgling cuckoos of the gut? Microb Ecol Health Dis. 2016;27:31557.

[19] Benno P, Befrits R, Norin E, et al. Is irritable bowel syndrome a dysbiotic bowel syndrome? Microb Ecol Health Dis. 2015;26:27637.

[20] Berstad A, Raa J, Valeur J. Indole - the scent of a healthy ‘inner soil’. Microb Ecol Health Dis. 2015;26:27997.

[21] Berstad A, Raa J, Valeur J. Tryptophan: ‘essential’ for the pathogenesis of irritable bowel syndrome? Scand J Gastroenterol. 2014;49(12):1493–1498.

[22] Nongonierma AB, FitzGerald RJ. Milk proteins as a source of tryptophan-containing bioactive peptides. Food Funct. 2015;6(7):2115–2127.

[23] Shimizu M. Interaction between food substances and the intestinal epithelium. Biosci Biotechnol Biochem. 2010;74(2):232–241.

[24] Samsel A, Seneff S. Glyphosate, pathways to modern diseases II: celiac sprue and gluten intolerance. Interdiscip Toxicol. 2013;6(4):159–184.