flora index = 6

intermediate dysbiosis

pronounced dysbiosis

mild dysbiosis

1 - 5: 6 - 12:

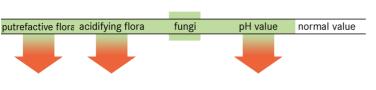
> 12:



## Intestinal health check

Summarised evaluation of the flora

# toxic load

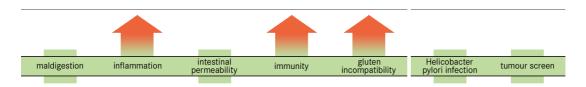


fermentation process

The analysis of the stool flora shows a reduction in the acidifying flora. The antagonistic putrefactive flora is, due to a reduction in Escherichia coli, also reduced. The decreased stool pH may be attributable to an accelerated gastrointestinal passage, e.g., due to a carbohydrate intolerance (lactose, fructose or sugar substitutes).

Decreased bacteroid and enterococcus counts are generally of no pathological relevance.

#### Summarised evaluation of the biochemical parameters



#### **Maldigestion**

• exclusion of maldigestion

#### **Inflammatory parameters**

• evidence of inflammatory changes in the mucous membrane

#### **Intestinal permeability**

• no evidence of an increased intestinal permeability

#### **Intestinal immunology**

- decreased level of activity of the mucosal immune system together with reduced secretory lgA
- suspected microbial infection of the intestinal mucosa together with increased B-defensin-2 levels

# Gluten incompatibility

suspected gluten incompatibility

#### Helicobacter pylori infection

• Exclusion of a Helicobacter pylori infection



# Early detection of colorectal carcinomasNo evidence of occult faecal blood.

Summarized evaluation of parasites and diarrhoea-associated pathogens

Diarrhoea-associated pathogens

• no evidence of clostridium difficile in stool

#### **Detailed results**

#### stool flora

	result in CFU/ g stool	assessment	norm
putrefactive flor	ra aerobic		
E.coli	< 10^5		10^6 - 10^7
E.coli-Biovare	< 10^4	T T	< 10^4
Proteus species	< 10^4		< 10^4
Klebsiella species	< 10^4		< 10^4
Pseudomonas specie	es < 10^4		< 10^4
Citrobacter species	< 10^4		< 10^4
Enterobacter sp.	< 10^4		< 10^4
Serratia species	< 10^4		< 10^4
Hafnia alvei	< 10^4		< 10^4
Morganella morganii	< 10^4		< 10^4
Kluyvera species	< 10^4		< 10^4
Providencia species	< 10^4	Ţ.	< 10^4
putrefactive flo	ra anaerobic		
Clostridium sp.	< 10^5		< 10^5
acidifying flora	anaerobic		
Bifidobacterium sp.	< 10^8	1	10^9 - 10^11
Lactobacillus sp.	< 10^5	Ţ	10^5 - 10^7
acidifying flora	aerobic		
Enterococcus sp.	< 10^5		10^6 - 10^7
neutral flora			
Bacteroides sp.	< 10^8		10^9 - 10^11
quantitative fun	gi detection		
Candida albicans	< 10^3		< 10^3
Candida sp.	< 10^3		< 10^3
Geotrichum sp.	< 10^3		< 10^3
Mould fungi	negative		negative
general investig	gations		
pH value	5.5		6 - 6.5
stool consistency	soft & unformed		soft & unforme



Putrefactive bacteria and acidifying bacteria behave antagonistically in the intestine. E. coli is a physiological intestinal bacterium, whereas all other putrefactive bacteria can be transiently found and are facultative pathogenic. Lactobacilli, bifidobacteria and enterococci belong to the physiological acidifying flora and, due to their metabolic activity, contribute to a physiological intestinal mucous membrane barrier.

The **Escherichia coli** species is a physiological component of the intestinal flora. An enduring decrease in Escherichia coli can lead to an inadequate stimulation of the mucosal immune system (MIS). The level of activity and readiness for defence are reduced.

**Enterococci** belong to the obligate flora of the intestinal wall. As a result of their resistance to acids and bile, enterococci can also be found in the small intestine. Enterococci inhibit the growth of pathogenic germs by acidifying the intestinal environment and by the production of bacteriostatic or bacteriocidal substances.

**Bifidobacteria** belong to the anaerobic acidifying flora. With a count of up to 10"11 CFU/g faeces they make up a considerable portion of the obligate intestinal flora. Bifidobacteria are purely saccharolytic micro-organisms, i.e., they exclusively metabolise carbohydrates. Degradation products of the carbohydrate metabolism are short-chain fatty acids, which by means of their acidifying actions and antagonistic effects on diverse putrefactive bacteria, play an important role in the resistance to colonisation.

**Lactobacilli** are the functionally most important component of a physiological small intestinal flora. Lactobacilli are purely saccharolytic micro-organisms i.e. they exclusively utilize non-cleavable carbohydrate bonds as well as constituents of the intestinal mucous. This action leads, in the first place, to the production of lactic acid. Lactobacilli are responsible for an acidification of the intestinal environment. Various metabolic products have a direct inhibitory effect on foreign germs and putrefactive bacteria such as Clostridium spp. and enterobacteria such as Proteus spp., etc.

**Bacteroides sp.** forms the numerically greatest proportion of the obligate colonic flora. Bacteroides predominantly metabolise proteins, although in comparison to other aerobic micro-organisms, they are less metabolically active.

#### Maldigestion

	result	unit	assessment	norm
Fat in stool	2.6	g/100g stool		< 3.5
1 at 111 51001	2.0	g/100g 31301		< 3.5
Protein in stool	1.1	g/100g stool		< 1
Sugar in stool	2	g/100g stool		< 2.5
Starch in stool	7.1	g/100g stool		9 - 13
Water in stool	81	g/100g stool		75 - 85
Pancreatic elastase	>500.0	μg/g		> 200
Bile acids	<1.7			< 1.7

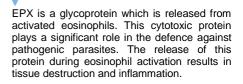
Borderline increased fat and protein residues are inconspicuous.

#### Inflammatory parameters

	result	unit	assessment	norm
Calprotectin	<7.8	mg/l		< 50
alpha-1-antitrypsin	<4.1	U/mI		< 27.5
EPX	1894.1	ng/ml		< 360
Lysozyme	279.2	ng/ml		< 600



The cause of an **elevated EPX (eosinophil protein X) value** can be a food allergy (lgE-mediated), parasitosis or a chronic inflammatory intestinal mucous membrane disease. In such chronic inflammatory intestinal illnesses, EPX can serve as a marker of the disease activity and the course of the illness.





**Secretory immunoglobin A (sIgA)** inhibits the invasion and colonisation of potentially pathogenic bacteria, viruses or fungi through the intestinal mucous membrane and neutralises antigens and toxins. The reduced concentration of slgA in the faeces suggests a decreased degree of activity in the mucosal immune system. A lasting reduction in the slgA value may be associated with an increased susceptibility to infections.

As a component of the inherited immune system, .8-defensin 2 contributes, by means of its antimicrobial effect, to the barrier function of the intestinal epithelium. An increased production of B-defensin-2 can be induced by proinflammatory cytokines and by micro-organisms such as fungi (e.g., Candida), viruses or bacteria. An elevated value may also accompany a Helicobactor pylori infection. Raised values can additionally be found with ulcerative colitis and other chronic inflammatory intestinal diseases.

#### Gluten incompatibility

	result	unit	assessment	norm
Transglutaminase antibody	213.2	U/I		< 100
Anti-gliadin antibody	<8.3	mU/g	<b>1</b>	< 100

An elevated faecal concentration of antibodies against transglutaminase was found. The high antibody concentration is indicative of gluten hypersensitivity.

#### Helicobacter pylori

	result	assessment	norm
Helicobacter pylori antigen	negative		negative

Up to 10% of patients suffering from a glutensensitive enteropathy have an undetected IgA deficiency. Because of this, in the case of a negative transglutaminase-IgA-antibody test in faeces and serum, together with persistent clinical symptoms, a determination of IgA and the IgA subgroups should be carried out.

In case of continuing gastric symptoms after exclusion of a Helicobacter pylori infection, a symptomatic therapy is recommended. An antibiotic eradication is not necessary. In high-risk patients (over 45 years old, long-term therapy with non-steroidal anti-rheumatics), a biopsy should nevertheless be taken to exclude the presence of gastric or duodenal ulcers of other origins.

#### Early detection of colorectal carcinomas



Besides the assessment of the haemoglobin-haptoglobin-complex in faeces, further sensitive parameters for the determination of colorectal carcinomas and adenomatous polyps are, above all, calprotectin, the proliferation marker M2-PK and haemoglobin. Through a combination of these parameters, the sensitivity and specificity for the detection of colorectal carcinomas can be distinctly increased.

#### Diarrhoea-associated pathogens



#### Therapy recommendation

#### Diet therapy

Raised values of a marker for inflammation are indicative of inflammatory mucous membrane alterations, which are, as a rule, associated with foodstuff incompatibilities. We recommend a mild and balanced diet.

In the case of a suspected **gluten incompatibility**, the clinical relevance should be clarified. For this, we recommend that a gluten-free diet should be adhered to temporarily, since in 90% of cases of gluten incompatibility or coeliac disease a considerable alleviation of symptoms can be achieved.

#### General recommendations for the prevention of colorectal tumours

Diet plays a decisive role in the development of colorectal carcinomas. A high-fat, low-fibre diet increases the risk of development. Increased meat consumption is also associated with an elevated risk of colonic cancer. However, there are many foodstuffs which can have a protective effect, as for example, secondary vegetable matter. Antioxidants such as vitamins, mineral nutrients and secondary vegetable matter should be ingested with the diet in sufficient amounts. The use of a dietary supplement to substitute a certain amount of these substances is recommended.

It is important to take preventive measures to avoid constipation in order to reduce the time over which carcinogenic substances remain in the intestine. For this, a high-fibre diet, adequate amounts of fluids and exercise are beneficial.



The replacement of lost fluids and electrolytes is of the highest priority during **acute diarrhoea**. Until improvement is seen, solids should be avoided. Thereafter the diet should be gradually normalized.



With a life-long, gluten-free diet, the effects of malabsorption can be prevented (osteoporosis, vitamin-deficiency syndrome), as well as the long-term consequences which include the frequent occurrence of malignancies.

Body weight also plays a further important role. Being overweight leads to a considerable increase in the cancer risk.

#### Anti-inflammatory therapy

As an antiphlogistic measure following the measurement of raised inflammatory markers we recommend the following preparations.

These preparations are also suitable for the treatment of fermentative dyspepsia, since this, in 60% of cases, is accompanied by an inflammatory mucous membrane irritation.

Colibiogen® Once daily, in the morning, in serious cases up to 3 times daily, 5 ml (= 1 teaspoon), 30 min before

meals

Myrrhinil intest® 3 times 4 dragees per day with fluids before meals

In children with Candida infections

3 times 3 dragees per day

MucoZink® 1 Tbsp. in 200 ml water perday

Duetocomponents such as L-glutamine and dexpanthenol, Muco Zink® promotes

wound healing and mucous membrane reconstruction.

CAUTION: MucoZink® contains fructose. In case of a known fructose incompatibility, not to be recommended.

#### Milieu-stabilising measures

For the regeneration of a balanced intestinal flora and/or in the case of a high yeast count, milieu-stabilising measures may be meaningful. Through the administration of Bifidobacteria and/ or Lactobacilli preparations, an acidification of the intestinal milieu can be obtained and favourable conditions for the restitution of the body's own acidifying flora achieved.

Due to the spectrum of intestinal micro-organisms found (L. acidophilus, L. lactis, L. casei, B. bifidum) we recommend

**proBiotik® pur** 1sachet(2.0g)perdaymixed with water and dietary supplement taken with a main meal.

# Immunomodulatory measures

In the case of decreased immunoglobulin A (slgA) together with high yeast counts, **glucans** can additionally be employed. They can contribute to a stabilisation of the immunological mucous membrane barrier due to their stimulating effects on macrophages.

nutriGlucan® 3 times 1 tablet per day (nutrimmun GmbH, Munster)

The immunomodulatory effect of live colibacteria by use of **AutoVACC-Oral E.c.**® can be employed with the following indications: atopical diseases, recurrent infections especially in the respiratory or urogenital tract. Instructions for administration can be obtained from GANZIMMUN AG.

**Mutaflor**® can support the intestinal flora and also have an immunomodulatory effect. Through defined metabolic products, the energy supply to the cells of the colonic mucous membrane can be improved, while at the same time, the physiological effects of the intestinal flora can be supported. Mutaflor® inhibits the infiltration of pathogenic germs into the intestinal cells. Additionally, the body-s own immune system is trained and the production of secretory

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The recommended doses are for adults. In children, they should be reduced corresponding to the child's age, or alternative preparations should be used.

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immunoglobin A stimulated.

Mutaflor® mite	1capsuleperday for 8 days, thereafter 2 capsules
	per day.
	In the case of persistent constipation
Mutaflor® 100 mg	vcan subsequently be prescribed. A dosage
G	increase of up to 4 capsules may be indicated, depending on the symptoms.
	depending on the symptoms.

# **Further diagnostics**

Further diagnostic measures should be anamnesis-orientated. Depending on the clinic, we recommend the following further diagnostic measures:

Allergy / Foodstuff incompatibility	- Prescreen Allergy
Detection of biogenic amines	· Histamine in urine
Carbohydrate intolerance	<ul> <li>bacterial cleavage of fructose, sorbitol, xylitol in faeces</li> </ul>
	<ul> <li>hydrogen breath test fructose, lactose, sorbitol</li> <li>genetic test fructose (ALDOB gene)/ lactose (LCT-gene) in EDTA blood</li> </ul>
Parasitesinfaeces	- Parasites, worms, wormeggs

Many thanks for your investigatory assignment.

Yours sincerely,

Dr. med. Ralf Kirkamm