ESPEN practical guideline: Clinical Nutrition in inflammatory bowel disease

Stephan C. Bischoff, Johanna Escher, Xavier Hébuterne, Stanisław Kłęk, Zeljko Krznaric, Stéphane Schneider, Raanan Shamir, Kalina Stardelova, Nicolette Wierdsma, Anthony E. Wiskin, Alastair Forbes

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1 ESPEN practical guideline: Clinical Nutrition in inflammatory bowel disease

Stephan C. Bischoff<sup>a\*</sup>, Johanna Escher<sup>b</sup>, Xavier Hébuterne<sup>c</sup>, Stanisław Kłęk<sup>d</sup>, Zeljko
Krznaric<sup>e</sup>, Stéphane Schneider<sup>c</sup>, Raanan Shamir<sup>f</sup>, Kalina Stardelova<sup>g</sup>, Nicolette Wierdsma<sup>h</sup>, Anthony E Wiskin<sup>j</sup>, Alastair Forbes<sup>k</sup>

- 5
- 6 Based on
- 7 ESPEN guideline: Clinical Nutrition in inflammatory bowel disease

8 Alastair Forbes, Johanna Escher, Xavier Hébuterne, Stanisław Kłęk, Zeljko Krznaric, Sté9 phane Schneider, Raanan Shamir, Kalina Stardelova, Nicolette Wierdsma, Anthony E.
10 Wiskin, Stephan C. Bischoff

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- 12
- 13 <sup>a</sup> University of Hohenheim, Institute of Nutritional Medicine, Stuttgart, Germany
- 14 <sup>b</sup> Erasmus Medical Center Sophia Children's Hospital, Rotterdam, The Netherlands
- 15 <sup>c</sup> Gastroentérologie et Nutrition Clinique, CHU de Nice, Université Côte d'Azur, Nice,
- 16 France
- 17 <sup>d</sup> General and Oncology Surgery Unit, Stanley Dudrick's Memorial Hospital, Krakow, Po-
- 18 land
- 19 <sup>e</sup> Clinical Hospital Centre Zagreb, University of Zagreb, Zagreb, Croatia
- 20 <sup>f</sup> Tel-Aviv University, Schneider Children's Medical Center of Israel, Petach-Tikva, Israel
- <sup>g</sup> University Clinic for Gasrtroenterohepatology, Clinal Centre "Mother Therese", Skopje,
- 22 Republic of North Macedonia
- 23 h Amsterdam university medical centers, Amsterdam, The Netherlands
- 24 <sup>j</sup> Paediatric Gastroenterology & Nutrition Unit, Bristol Royal Hospital for Children, Bris-
- 25 tol, United Kingdom
- 26 <sup>k</sup> Norwich Medical School, University of East Anglia, Norwich, United Kingdom
- 27
- 28 \*Institute of Nutritional Medicine, University of Hohenheim, Fruwirthstr. 12, 70593
- 29 Stuttgart, Germany E-Mail: bischoff.stephan@uni-hohenheim.de
- 30
- Keywords: Crohn's disease, ulcerative colitis, enteral nutrition, parenteral nutrition,
   inflammatory bowel disease, nutritional therapy
- 33 Abbreviations: CD, Crohn's disease; EN, enteral nutrition; IBD, inflammatory bowel dis-
- ease; ONS, oral nutritional supplements; PN, parenteral nutrition; UC, ulcerative colitis
- 35

#### 36 Introduction

37 Inflammatory bowel disease (IBD), predominantly ulcerative colitis (UC) and Crohn's disease (CD), is now common in the entire developed world. Malnutrition can occur as 38 39 well in UC and CD, but is a considerably greater problem in CD given its capacity to affect 40 any part of the gastrointestinal tract, unlike UC, which is restricted to the colon and has 41 few direct malabsorptive effects. As in adults, malnutrition is prevalent in paediatric IBD, mainly in active disease and more in CD than in UC. Since patients with IBD constitute a 42 high-risk population for malnutrition, they need screening for malnutrition, with its 43 44 subsequent assessment and management. Nutritional care is clearly important in the 45 treatment of patients with IBD and includes prevention of malnutrition and micronutri-46 ent deficiencies, prevention of osteoporosis, and, in children promotion of optimal growth and development. 47

48

#### 49 Methodology

50 The present practical guideline consists of 40 recommendations and is based on the ES-51 PEN Guideline: Clinical Nutrition in inflammatory bowel disease (1). The original guideline was shortened by restricting the commentaries to the gathered evidence and litera-52 ture on which the recommendations are based on. The recommendations were not 53 changed (except "artificial nutrition" was replaced by "medical nutrition" and language 54 55 was adapted to American English), but the presentation of the content was transformed 56 into a graphical presentation consisting of decision-making flow charts wherever possi-57 ble. The original guideline was developed according to the standard operating proce-58 dure (SOP) for ESPEN guidelines (2). This SOP is oriented on the methodology of the 59 Scottish Intercollegiate Guidelines Network (SIGN). Literature was searched and graded 60 into 1-4 according to evidence, and recommendations were created and graded into four 61 classes (A/B/0/GPP). All recommendations were not only based on evidence, but also 62 underwent a consensus process, which resulted in a percentage of agreement (%). 63 Whenever possible, representatives from different professions (physicians, dieticians, nurses, others) as well as patient representatives were involved. The guideline process 64 65 was funded exclusively by the ESPEN society. The guideline shortage and dissemination was funded in part by the UEG society, and also by the ESPEN society. For further details 66 67 on methodology, see the full version of the ESPEN guideline (1) and the ESPEN SOP (2). 68 The ESPEN practical guideline "Clinical Nutrition in inflammatory bowel disease" has

- 69 been structured according to a flow chart covering all nutritional aspects of IBD (Figure
- 70 1).
- 71

	Journal Pre-proof
72	Results
73	
74	I. Prevention of IBD (Figure 2)
75	
76	Recommendation 1
77 78 79	A diet rich in fruit and vegetables, rich in n-3 fatty acids, and low in n-6 fatty acids is associated with a decreased risk of developing CD or UC and is therefore recommended.
80	Grade of recommendation 0 – strong consensus (90 % agreement)

81

#### 82 **Commentary**

83 Smoking, antibiotic use, and diet are potentially reversible risk factors for IBD. Many studies have evaluated the effect of diet on the risk of developing IBD. However most of 84 them are retrospective case-control studies. In 2011 Hou et al. published the first sys-85 tematic review entitled "Dietary Intake and Risk of Developing IBD" (3). They used 86 87 guideline-recommended methodology to evaluate the association between pre-illness intake of nutrients (fats, carbohydrates, protein) and food groups (fruits, vegetables, 88 89 meats) and the risk of subsequent IBD diagnosis. Nineteen studies were included, en-90 compassing 2,609 IBD patients (1,269 with CD and 1,340 with UC), and over 4,000 con-91 trols. The main results are: (i) increased risk of developing UC and CD with high intake 92 of PUFAs, n-6 fatty acids, and meats, (ii) decreased risk of CD, but not UC, with high in-93 take of dietary fiber (>22 g/d) and fruits.

Fiber, fruit and vegetables (4): Compared to women with the lowest energy-adjusted
fiber intake, intake of fiber in the highest quintile (median 24 g/d) was associated with a
significant reduction in risk of CD [HR 0.59, 95% CI 0.39 – 0.90] but not UC.

97 In a meta-analysis including a total of 14 case-control studies (5), consumption of vege-98 tables was negatively associated with the risk of UC (OR=0.71), but not with CD 99 (OR=0.66). Higher consumption of fruit was negatively associated with the risk of UC 100 (OR=0.69) and CD (OR=0.57).

- **Dietary fat** (6): Cumulative energy-adjusted intake of total fat, saturated fats, unsaturated fats, n-6 and n-3 PUFA were not associated with risk of CD or UC. However, greater intake of long-chain n-3 PUFA was associated with a trend towards lower risk of UC (HR 0.72). In contrast, high long-term intake of trans-unsaturated fatty acids was associated with a trend towards on a special data of the trans-unsaturated fatty acids was associated with a trend towards was associated with a trend towards lower risk of UC (HR 0.72). In contrast, high long-term intake of trans-unsaturated fatty acids was associated with a trend towards was ass
- 105 with a trend towards an increased incidence of UC (HR 1.34).
- 106 In the EPIC study, 229,702 participants were recruited from nine European centers be-107 tween 1991 and 1998 (7). At recruitment, dietary intakes of DHA and fatty acids were
- 108 measured using validated food frequency questionnaires. In a nested case-control anal-109 ysis, each participant who developed incident UC (n=126) was matched with four con-
- 109 ysis, each participant who developed incident UC (n=126) was matched with four con-110 trols. The highest quartile of intake of linoleic acid was associated with an increased risk
- of UC (OR 2.49) with a significant trend across quartiles (OR 1.32 per quartile increase).
- 112

#### 113 **<u>Recommendation 2</u>**

- 114 Breastfeeding can be recommended, because it is the optimal food for infants and
- 115 **it reduces the risk of IBD.**
- 116 Grade of recommendation B strong consensus (93 % agreement)
- 117

#### 118 **Commentary**

Systematic reviews from 2004 and 2009 concluded strongly in favor of breastfeeding (8, 119 120 9) and subsequent studies have reinforced this interpretation. A case-control study from 121 New Zealand reported that breastfeeding was protective against IBD (CD OR 0.55 95%CI 122 0.41-0.74, UC OR 0.71 95%CI 0.52-0.96) with a duration-response effect (10). Compara-123 ble data were reported from a Danish cohort study, in which breastfeeding for more 124 than six months decreased the odds of IBD (OR 0.50, 95%CI 0.23-1.11) (11). Two further 125 publications confirmed this relationship, one from the US and another from Asia-Pacific (12,13). Breastfeeding for around six months or longer is desirable in all infants (14). 126

- 127
- 128 II. General aspects (Figure 3)

#### 129 **Recommendation 3A**

- 130 Patients with IBD are at risk and therefore should be screened for malnutrition at
- 131 the time of diagnosis and thereafter on a regular basis.
- 132 Grade of recommendation GPP strong consensus (96 % agreement)
- 133
- 134 **Recommendation 3B**
- 135 **Documented malnutrition in patients with IBD should be treated appropriately,**
- 136 because it worsens the prognosis, complication rates, mortality and quality of life.
- 137 Grade of recommendation GPP strong consensus (96 % agreement)
- 138
- 139Commentary for A/B

Adults with IBD are at increased risk of malnutrition, with deficits more common in patients with CD than UC (15). Obese patients may have covert deficits in lean mass which may be unmasked by tools such as skinfold thickness measurement. Patients with active IBD, particularly those whose disease is poorly responsive to medical therapy, are at highest risk of poor nutrition. In adults, risk of malnutrition can be assessed with vali-

- 145 dated screening tools (16).
- 146 Malnourished patients with IBD are more likely to be hospitalized following emergency
- 147 department attendance (17) and are more likely to be admitted to hospital due to infec-
- tion (18). In hospitalized patients, malnutrition is an independent risk factor for venous
- thromboembolism (19), non-elective surgery (20), longer admission (15, 20) and in-
- 150 creased mortality (15).

151 Malnutrition in children: Malnutrition in childhood CD is common at diagnosis and 152 may persist despite disease treatment (21). Children with UC are also at risk of poor nutrition, but nutritional deficits may not be immediately obvious on assessment of just 153 154 height and weight (22). Although a variety of screening tools exists, the tools have poor 155 ability to discern different levels of nutrition risk for children with IBD (23). Poor nutri-156 tion in childhood IBD contributes to disrupted pubertal development and impaired 157 growth velocity which may lead to short stature in adulthood. Of particularly im-158 portance in pediatric IBD is growth failure, which is the result of a combination of in-159 flammation and chronic malnutrition (24).

160

#### 161 **Recommendation 4**

# In general, the energy requirements of patients with IBD are similar to those of the healthy population; provision should be in line with this.

#### 164 **Grade of recommendation GPP – strong consensus (93 % agreement)**

165

#### 166 **Commentary**

For clarity this question can be formulated in two ways; firstly, do patients with IBD have an altered energy requirement compared to healthy individuals, and secondly do

- 169 energy requirements vary with disease activity.
- 170 There are relatively few studies examining energy expenditure in patients with UC and
- all studies are of only small numbers of patients. There may be an increase in metabolic
- activity at times of acute severe UC compared to remission in adults (25, 26) which is
- understandable considering that systemic disturbance (fever and tachycardia) is com-
- mon. However, an increase in resting energy expenditure is likely to be offset by reduc tion of physical activity. Significant reduction in dietary intake is common in acute UC
- 175 up physical activity. Significant reduction in dietary intake is common in a 176 and may result in negative energy halance (27)
- and may result in negative energy balance (27).
- 177 One single study has measured total energy expenditure in adults with CD and recorded 178 normal values (28). Measured resting energy expenditure per kilogram in adult patients 179 has been found to be higher than (29) or the same as (30) that measured in healthy con-180 trols. However, this could be due to inadequate consideration of body size and the rela-181 tive proportions of tissues of differing metabolic activity. No consistent association be-
- 182 tween CD activity and resting energy expenditure in adults has been demonstrated. In
- 183 children with CD, measured resting energy expenditure has not been demonstrated to
- 184 be significantly different. Measurement of resting energy expenditure by indirect calo-
- 185 rimetry could be used in troublesome cases.
- 186

## 187 **Recommendation 5A**

## 188 Protein requirement are increased in active IBD, and intake should be increased

189 (to 1.2-1.5 g/kg/d in adults) relative to that recommended in the general popula-

- 190 **tion.**
- 191 Grade of recommendation GPP strong consensus (96 % agreement)
- 192

#### 193 **Recommendation 5B**

- 194 The protein requirements in remission are generally not elevated and provision
- 195 should be similar (about 1g/kg/d in adults) to that recommended for the general 196 population.
- **Grade of recommendation GPP strong consensus (96 % agreement)**
- 198

#### 199Commentary for A/B

Patients with IBD develop a relative reduction in lean mass and increase in obesity over time. This may occur due to chronically poor dietary intake, increased rates of protein turnover and gut loss of nutrients during phases of active disease or from the effect of disease treatments. Corticosteroids increase net loss of protein in children (31) and adults (32) with CD. In contrast administration of elemental or polymeric feed as treatment of CD or as adjunctive nutrition support results in reduction of proteolysis and acquisition of lean tissue in children and adults (33-35).

207 Monitoring of anthropometry provides insight into which patients develop relative defi-208 cits in lean mass and therefore would benefit from nutritional supplementation. There is

no good evidence that the daily protein needs of IBD patients differ from those of healthy controls, but as discussed elsewhere poor appetite and restricted dietary intake

is commonplace. In patients receiving steroids and gut rest, enteral nutrition (EN) may

212 provide beneficial effects on protein turnover without deleterious consequences on dis-

- 213 ease activity.
- 214 There is no good evidence that the daily protein needs of IBD patients in remission differ
- 215 from those of healthy controls. Provision of 1g protein for each kilogram of body weight
- 216 is therefore reasonable. However, in active inflammation the proteolytic, catabolic re-
- sponse justifies an increase in provision to 1.2 to 1.5 g/kg bodyweight (36, 37).
- 218

#### 219 **<u>Recommendation 6</u>**

# Patients with IBD should be checked for micronutrient deficiencies on a regular basis and specific deficits should be appropriately corrected.

#### 222 Grade of recommendation GPP – strong consensus (100 % agreement)

223

#### 224 Commentary

Patients with IBD are vulnerable to micronutrient deficits due to gut loss from diarrhea
and inadequate dietary intake from anorexia accompanying disease activity. At times
when nutrition support is offered then multivitamin and micronutrient supplements
should also be offered to ensure an appropriately balanced nutritional intake.

- When interpreting blood results of micronutrients and trace elements it is important to consider that many serum values, or markers of status, are positive or negative acute phase reactants. Serum levels rise or fall, as part of the inflammatory response, for example ferritin, and copper increase but folate, selenium and zinc decrease in inflammation (38). In light of this, some authors have examined micronutrient status in patients
- in clinical disease remission and found deficits of a variety of micronutrients (39, 40).

- Furthermore, deficits may be present even in apparently well-nourished individuals (41). These observations highlight the need for routine monitoring (perhaps annually) to screen for deficiency. A daily multivitamin supplement may correct most deficiencies but is no guarantee of adequacy, even over the long term; iron, zinc and vitamin D are likely to require specific replacement regimens (42). Poor compliance, particularly in adolescents, is common with multivitamin supplements and patient education about the
- 241 rationale behind their use is important (43).
- 242 Consequences of deranged micronutrient status include anemia, impaired linear growth
- and poor bone health. Recent research has focused on vitamin D; it and its receptor may
- have some immunomodulatory properties, which further highlights the need for specific
- attention to micronutrient status in patients with IBD (Recommendation 11).
- 246

#### 247 **Recommendation 7A**

Iron supplementation is recommended in all IBD patients when iron deficiency anemia is present. The goal of iron supplementation is to normalize hemoglobin

- 249 anemia is present. The250 levels and iron stores.
- 251 Grade of recommendation A strong consensus (100 % agreement)
- 252

#### 253 **Recommendation 7B**

254 **Oral iron should be considered as first-line treatment in patients with mild ane-**

255 mia, whose disease is clinically inactive, and who have not been previously intol-256 erant to oral iron.

- 257 Grade of recommendation A strong consensus (100 % agreement)
- 258
- 259 **Recommendation 7C**

Intravenous iron should be considered as first-line treatment in patients with clinically active IBD, those with previous intolerance to oral iron, those with he-

moglobin below 100 g/L, and in patients who need erythropoiesis-stimulating

- 263 agents.
- 264 **Grade of recommendation A strong consensus (93 % agreement)**
- 265

#### 266 **Commentary for A/B/C**

267 Anemia is considered the most frequent extraintestinal manifestation of IBD, usually 268 complicating the course both in UC and CD. All patients with IBD regardless of their age 269 should be assessed for the presence of anemia (44). The major forms of anemia in IBD 270 are iron deficiency anemia, anemia of chronic disease and anemia of mixed origin [ECCO 271 Anemia Statement 1A] (44). Diagnostic criteria for iron deficiency depend on the level of 272 inflammation. For laboratory screening, complete blood count, serum ferritin, and Creactive protein should be used [ECCO Anemia Statement 1B]. For patients in remission 273 274 or mild disease, measurements should be performed every six to twelve months. In out-275 patients with active disease such measurements should be performed at least every 276 three months [ECCO Anemia Statement 1B]. In patients without clinical, endoscopic, or

277 biochemical evidence of active disease, serum ferritin  $<30 \mu g/L$  is an appropriate crite-278 rion for the diagnosis of iron deficiency anemia. In the presence of inflammation, a serum ferritin up to 100 µg/L may still be consistent with iron deficiency [ECCO Anemia 279 Statement 1D]. In the presence of biochemical or clinical evidence of inflammation, the 280 281 diagnostic criteria for anemia of chronic disease are a serum ferritin >100 µg/L and 282 transferrin saturation <20%. If the serum ferritin level is between 30 and 100  $\mu$ g/L, a 283 combination of true iron deficiency and anemia of chronic disease is likely [ECCO Ane-284 mia Statement 1E].

285 Iron supplementation is recommended in all IBD patients, whatever their age, when 286 iron-deficiency anemia is present [ECCO Anemia Statement 2A]. Quality of life improves 287 with correction of anemia, and this improvement is independent of clinical activity (45). 288 The European Crohn's and Colitis Organization (ECCO) guidelines (44) conclude that "IV 289 iron is more effective, shows a faster response, and is better tolerated than oral iron" 290 and state that "IV iron should be considered as first line treatment in patients with clini-291 cally active IBD, with previous intolerance to oral iron, with hemoglobin below 100 g/L, 292 and in patients who need erythropoiesis-stimulating agents; while oral iron may be used 293 in patients with mild anemia, whose disease is clinically inactive, and who have not been 294 previously intolerant to oral iron (44). The estimation of iron need is usually based on 295 baseline hemoglobin and body weight (Table 1) (46).

- 296
- 297

298 Table 1: Simple scheme for estimation of total iron need (46)

Hemoglobin g/L	Body weight <70 kg	Body weight ≥70 kg	
100-120 (women)	1000 mg	1500 mg	
100-130 (men)	1000 mg	1500 mg	
70-100	1500 mg	2000 mg	

299

After successful treatment of iron deficiency anemia with intravenous iron, re-treatment
 with intravenous iron should be initiated as soon as serum ferritin drops below 100
 µg/L or hemoglobin below 12 or 13 g/dL according to gender [ECCO Anemia Statement
 3E].

304

#### 305 III. Dietetic recommendations in active disease (Figures 4 and 5)

306

#### 307 Recommendation 8

There is no "IBD diet" that can be generally recommended to promote remission
 in IBD patients with active disease.

#### 310 Grade of recommendation GPP – strong consensus (96 % agreement)

311

#### 312 **Commentary**

- 313 RCT data regarding the effects of experimental diets such as specific carbohydrate,
- 314 paleolithic, gluten-free, low fermentable oligo-, di- and monosaccharides and polyols
- 315 (FODMAP), or  $\omega$ -3 PUFA enriched diets on intestinal inflammation or on inducing re-

mission are still lacking at this time. An adequately powered RCT of fructooligosaccharides showed no clinical benefit in patients with active CD (47). See also Recommendation 31. Therefore, no "oral IBD diet" can be generally recommended to promote remission in IBD patients with active disease. This recommendation does not prelude the needs of all IBD patients to receive an individual (nutritional) approach based on their specific personal situation, preferably with the active input of a dedicated dietician or nutritionist as part of the multidisciplinary approach.

323

#### 324 Recommendation 9A

325 IBD patients with severe diarrhea or a high output jejunostomy or ileostomy 326 should have fluid output and urine sodium monitored, and fluid input adapted 327 accordingly (decrease hypotonic fluid and increase saline solutions), with consid-328 eration of food intolerances that may enhance fluid output.

- 329 Grade of recommendation 0 strong consensus (93 % agreement)
- 330
- 331 **Recommendation 9B**
- Parenteral infusions (fluid and electrolytes) can be needed in the case of on-going
  high output stomas.
- 334 **Grade of recommendation 0 strong consensus (96 % agreement)**
- 335

#### 336 **Commentary for A/B**

337 Ongoing and severe diarrhea or increased/high output stoma can result in intestinal 338 insufficiency (48) with malabsorption, unintentional weight loss, malnutrition, nutritional deficiencies and/or dehydration. Malabsorption is an important contributing fac-339 340 tor to malnutrition in IBD (49). The retrospective study of Baker in 687 stoma patients 341 (50), showed that early high output (within three weeks) from an ileostomy is common 342 and although 49% resolved spontaneously, 51% needed ongoing medical treatment, 343 usually because of a short small-bowel remnant. 71% patients were treated with oral 344 hypotonic fluid restriction, glucose-saline solution and anti-diarrheal medication to 345 wean from parenteral infusions and 8% had to continue parenteral or subcutaneous 346 saline in home-setting. Satisfactory home management with oral fluid restriction and 347 monitoring of urine sodium content was demonstrated more than 35 years ago (51). In a 348 study in 13 adult (ileal) increased/high output stoma patients, oral rehydration solu-349 tions containing rice maltodextrins supplementation improved the sodium and potassi-350 um balance. The association of increased body weight with decreased serum renin con-351 centrations suggests that a positive water balance also occurred (52). In another study, 352 three different saline and/or glucose solutions were tested in six patients with jejunos-353 tomies. Based on this small group, a sipped glucose electrolyte solution seemed to be the 354 optimal mode of sodium replacement in patients with increased/high output stoma (53). 355 No RCTs are available on nutritional treatment of IBD related diarrhea or in-356 creased/high output stoma. Only case studies on treatment of CD with increased/high output stoma have been published, which show successful treatment with restriction of 357 358 hypotonic fluids, sodium enriched diets, exclusive enteral nutrition and/or parenteral 359 sodium-containing infusions.

360

#### 361 **Recommendation 10**

In CD patients with intestinal strictures or stenosis in combination with obstruc tive symptoms, a diet with adapted texture, or distal (post-stenosis) EN can be
 recommended.

#### 365 **Grade of recommendation GPP – strong consensus (95 % agreement)**

366

#### 367 Commentary

368 Depending on the severity (degree of obstruction) and site of intestinal strictures, nutri-369 tional support may become necessary while the effects of treatment are awaited. Such 370 treatment may be medical (with drugs) where the narrowing is mainly the result of in-371 flammation, or mechanical (by balloon dilatation or surgery) when there is fibrotic scar-372 ring. In patients with radiologically identified but asymptomatic stenosis of the intestine 373 it is conventional to recommend a modified diet which is low in insoluble fiber, but there 374 are no robust data to support this apparently logical approach. When symptoms are 375 present it may be necessary to adapt the diet to one of soft consistency, perhaps pre-376 dominantly of nutritious fluids.

377 Intestinal fibrosis is a common feature of CD and may appear as a stricture, stenosis, or 378 intestinal obstruction. Stenosing CD leads to a significantly impaired quality of life in 379 affected patients and constitutes a challenging treatment situation. A recent Chinese 380 prospective observational study in 59 adult CD patients with inflammatory bowel stric-381 tures showed that twelve weeks exclusive EN can effectively relieve inflammatory bowel 382 strictures; (81.4%) achieved symptomatic remission, 35 patients (53.8%) achieved ra-383 diologic remission, and 42 patients (64.6%) achieved clinical remission (54). Although it 384 is common practice to recommend a modified diet with adapted consistency perhaps 385 predominantly of nutritious fluids, at least in patients with radiologically identified ste-386 nosis of the (proximal) intestine and obstructive symptoms, or to feed distally by EN 387 whenever this is possible, there are no robust data to support these apparently logical 388 approaches.

389

## 390 **Recommendation 11**

In IBD patients (adults and children) with active disease and those who are steroid-treated, serum calcium and 25(OH) vitamin D should be monitored and supplemented if required to help prevent low bone mineral density. Osteopenia and osteoporosis should be managed according to current osteoporosis guidelines.

#### **Grade of recommendation B – strong consensus (96 % agreement)**

396

#### 397 **Commentary**

398 Significant risk factors for low bone mineral density studied in adult IBD populations

399 (n=116 and n=205) prove to be low serum vitamin D, male gender, Asian ethnicity, CD,

- 400 low BMI and corticosteroid use, whereas no consensus on role of age, or age at diagnosis
- 401 was found (55, 56). In children and adolescents with IBD risk factors associated with

402 low bone mineral density are cumulative corticosteroid dose, height-for-age Z-score,403 and BMI Z-score (57).

404 There is no overall consensus on the vitamin D status and necessary actions in children 405 and adolescents with IBD. An RCT of 132 adult osteopenic CD patients showed improved 406 bone mineral density at lumbar spine after two years of once weekly treatment course 407 with risedronate 35 mg, concomitant with calcium and vitamin D supplementation (58). 408 An earlier RCT showed no significant benefit of calcium supplementation (1 g/day) 409 alone on the bone mineral density at one year in corticosteroid-using IBD patients with 410 osteoporosis (59). Evaluation for vitamin D deficiency is recommended in IBD and en-411 suring always an adequate supply of calcium and vitamin D, especially in steroid-treated 412 IBD patients. Limitation of corticosteroid use helps to prevent low bone mineral density. 413

414 **<u>Recommendation 12A</u>** 

415 **CD** patients treated with sequestrants such as cholestyramine have minimal addi-

- tional risk of fat malabsorption, and therefore do not need differences in nutrition therapy compared to other patients with CD.
- 418 **Grade of recommendation GPP consensus (86 % agreement)**
- 419
- 420 **Recommendation 12B**

421 IBD patients with hyperoxaluria often also have fat malabsorption and these pa-

422 tients should be counselled regarding fat malabsorption.

423 Grade of recommendation GPP – consensus (88 % agreement)

424

#### 425 **Commentary for A/B**

426 The common causes of bile acid malabsorption in CD are ileal resection and inflamma-427 tion of the terminal ileum. Decreased reabsorption of conjugated gall bile acids leads to 428 excess transmission to the colon, where deconjugation by bacteria occurs. Osmotic diar-429 rhea and (in severe bile acid malabsorption) fat malabsorption might be a consequence 430 (60). If mild, bile acid diarrhea can be controlled by a sequestrant such as cholestyra-431 mine (61, 62). In a double-blind cross-over study in 14 CD patients who had undergone 432 ileal resection, no negative effect of cholestyramine treatment on jejunal fat absorption 433 was reported. In severe cases of bile acid malabsorption however, steatorrhea may 434 worsen as a result of cholestyramine treatment (63).

435 Enteric (secondary) hyperoxaluria (with increased risk of kidney stones) occurs in se-436 vere small bowel CD associated with fat malabsorption and a consecutive elevation of 437 intestinal oxalate absorption. Enteric hyperoxaluria may occur after ileal resection. 438 Presence of the colon is an important factor, as oxalate remains available for colonic ab-439 sorption because of concomitant fat malabsorption and its binding of calcium (64). Uri-440 nary oxalate excretion correlates with fat excretion, as was shown in one study in CD 441 patients undergoing intestinal resection. Increasing the dietary fat intake in these pa-442 tients further increased urinary oxalate excretion (65). Significantly lower mean values 443 of urinary oxalate excretion were found in pediatric than in adult CD patients (66). A reason for this may be the shorter history of CD, which usually also implies fewer bowel 444

- resections. This implies that a diet low in fat and oxalate and high in calcium should be
- 446 recommended in patients with hyperoxaluria. Restriction of dietary oxalate (teas and
- fruits mainly) seems warranted only in those with recurring urinary tract stones.
- 448

### 449 **Recommendation 13**

- 450 **Exclusion diets cannot be recommended to achieve remission in active CD, even if** 451 **the patient suffers from individual intolerances.**
- 452 **Grade of recommendation GPP strong consensus (96 % agreement)**
- 453

## 454 **Commentary**

455 The systematic enquiry revealed insufficient evidence to make firm recommendations

- 456 for exclusion diets as induction therapy. Exclusion diets have been described to alleviate457 symptoms (67), but only few uncontrolled studies report induction of remission (68, 69).
- 458 In an RCT, longer maintenance of remission (after successful induction of remission us-
- 459 ing elemental formula) was seen in patients using a stepwise dietary introduction pro-
- 460 gram excluding foods that worsened symptoms, compared to patients receiving cortico-
- 461 steroids on a tapering schedule while eating a normal diet (70). Similar results on
- 462 maintenance of remission were reported in an open label study by the same group using
- 463 a personal food exclusion diet (71). Another study reported maintenance of clinical re-
- 464 mission using an IgG4 guided exclusion diet in adult CD patients (72).
- Exclusion diets are labor-intensive for staff, and complex, challenging and often unpleasant for patients. The systematic enquiry revealed no evidence that exclusion diets are hazardous when applied under medical supervision. Evidence was not forthcoming to indicate that they contribute to nutritional deficiencies. Nonetheless it is good practice to monitor carefully for deficiencies that might be predicted from any particular set
- 470 of exclusions.
- 471

## 472 **Recommendation 14A**

- 473 Probiotic therapy using Lactobacillus reuteri or "VSL#3"\*, but not necessarily oth-
- 474 er probiotics, can be considered for use in patients with mild to moderate UC for 475 the induction of remission
- 475 **the induction of remission**.
- 476 **Grade of recommendation 0 strong consensus (92 % agreement)**
- 477
- 478 **<u>Recommendation 14B</u>**
- 479 **Probiotics should not be used for treatment of active CD.**
- 480 Grade of recommendation B strong consensus (95 % agreement)
- 481
- 482 **Commentary for A/B**
- 483 Two clinical trials in pediatric UC patients show a moderate effect of rectal enemas con-
- taining *Lactobacillus reuteri* in mild distal UC (73) and of an oral preparation of "VSL#3"

in active UC (74). The systematic enquiry indicated that probiotics were, in general, inef-

\*"VSL#3" refers only to the probiotic product used in the cited literature and equivalent

492 Recommendation 15A

fective in active CD.

493 Oral Nutrition Supplements (ONS) are the first step when medical nutrition is in494 dicated in IBD, but generally are a minor supportive therapy used in addition to
495 normal food.

496 **Grade of recommendation 0 – strong consensus (92 % agreement)** 

products independent from the present product labeling.

IV. Medical nutrition in active IBD (Figures 6 and 7)

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485

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491

#### 498 **Recommendation 15B**

499 If oral feeding is not sufficient then EN should be considered as supportive thera-

500 py. EN using formulas or liquids should always take preference over PN, unless it 501 is completely contraindicated.

- 502 Grade of recommendation A strong consensus (100 % agreement)
- 503

#### 504 Recommendation 15C

**PN is indicated in IBD (i) when oral nutrition or EN is not sufficiently possible, (e.g. when the GI tract is dysfunctional or in CD patients with short bowel), (ii) when there is an obstructed bowel where there is no possibility of placement of a feed**-**ing tube beyond the obstruction or where this has failed, or (iii) when other com**-**plications occur such as an anastomotic leak or a high output intestinal fistula.** 

- 510 Grade of recommendation B strong consensus (96 % agreement)
- 511

#### 512 **Commentary for A/B/C**

513 The decision on the optimal route of medical nutrition in IBD can be complex and in-514 volve several aspects, including the ability of the patient to eat, the absorptive capacity 515 of the GI tract, the nutritional status of the patient, and the therapeutic goals. Oral Nutri-516 tion Supplements (ONS) are the first step but generally are a minor supportive therapy 517 used in addition to normal food. By using ONS, a supplementary intake of up to 600 518 kcal/day can be achieved without compromising normal food intake in adults. If oral 519 feeding is not possible, feeding the patient through a nasogastric or nasoenteric tube 520 should be considered. EN should be considered in patients with a functional gastrointes-521 tinal tract but who are unable to swallow safely (75, 76). In situations when the gut can-522 not absorb all nutritional needs, EN should nonetheless be attempted with supplemen-523 tary PN (41, 77, 78). PN is indicated when there is an obstructed bowel where there is 524 no possibility of placement of a feeding tube beyond the obstruction or where this has 525 failed. It is required in patients with short bowel resulting in severe malabsorption of

- nutrients and/or fluid and electrolyte loss which cannot be managed enterally. PN is also indicated in surgical cases as above, and in any patient, who is intolerant of EN or in whom nutrition cannot be maintained by the enteral route (79). However, it must be recognized that these patients in need of PN are those with the most complicated disease (80).
- 531

#### 532 **Recommendation 16**

Exclusive EN is effective and is recommended as the first line of treatment to in duce remission in children and adolescents with acute active CD.

#### 535 Grade of recommendation B - strong consensus (92 % agreement)

536

#### 537 **Commentary**

538 Primary nutritional therapy in the form of exclusive EN should be considered in all pa-539 tients with acute active CD. This is a first choice in patients at high risk from alternative 540 therapy such as steroids. Old meta-analyses demonstrated that corticosteroids are bet-541 ter than exclusive EN in induction of remission in adults. The argument in favor of exclu-542 sive EN is stronger in pediatric practice and will normally be the first choice in many 543 centers. Firstly, this is because of the deleterious effects of undernutrition on growth. 544 Secondly, since growth is so essential in children, this increases the possibility of avoid-545 ing the use of steroids or delaying their introduction, which is of paramount importance. 546 Third, and most importantly, is the observed effect on induction of remission in pediat-547 ric studies demonstrating similar efficacy of steroids and exclusive EN (81), and that in 548 some settings (i. e. cconcomitant immunomodulatory treatment) exclusive EN might 549 even be superior to corticosteroids in children (82). However, these studies suffer from 550 methodological limitations. Recommendations in children are made only for exclusive 551 EN as limited data suggest that partial EN may be less effective (60), though one RCT 552 showed similar efficacy (83). The data are weaker for adult practice, and most centers 553 will continue to use steroids (or biologicals) as first-line therapy unless these agents are 554 actively contra-indicated. However, patient and disease characteristics also contribute 555 to the rapeutic management decisions and these may make EN therapy a first-line option 556 also in selected cases of adults with acute CD (84). EN is preferred, because PN has not been shown to offer any advantage in CD and should be used only to improve nutritional 557 558 status for surgery and when other modes of nutrition are not possible (85).

- 559
- 560 **Recommendation 17A**
- 561 **For EN in IBD, nasal tubes or percutaneous access can be used.**
- 562 Grade of recommendation B strong consensus (96 % agreement)
- 563
- 564 **Recommendation 17B**
- 565 **EN in CD should be administered via an enteral feeding pump.**
- 566 **Grade of recommendation B strong consensus (92 % agreement)**
- 567

#### 568 **Commentary for A/B**

- 569 EN can be safely delivered by nasogastric tube, or percutaneous endoscopic gastrostomy
- 570 (86-88). Continuous EN administered via an enteral feeding pump and increased slowly
- 571 to the full prescribed volume appears to have lower complication rates than bolus deliv-
- 572 ery (86-89). The most frequent complications of EN are mechanical (tube-related), then
- 573 metabolic and infectious, but these are not notably different from those seen in other
- 574 chronic conditions (88, 89).
- 575 Few patients with UC will need EN or PN other than during the most severe exacerba-
- tions and in the peri-operative phase. EN is most appropriate and associated with signif-
- 577 icantly fewer complications than PN in acute UC. Bowel rest through intravenous nutri-
- 578 tion does not alter the outcome, but nonetheless, there are no specific contraindications
- 579 for the use of PN in UC.
- In CD nutritional support is more often needed. There is no specific contraindication to the use of PN in patients with CD in comparison to other diseases, and a central or peripheral route may be selected according to its expected duration. There are not enough data to dictate the use of specific substrates in the composition of PN in CD. PN must however be adjusted to fulfil the needs of the individual patient. PN, especially at home, should be viewed as complementary non-exclusive nutrition, which can be tapered to a
- 586 minimal level when body composition has been sufficiently restored.
- 587

#### 588 Recommendation 18A

- 589 Standard EN (polymeric, moderate fat content, no particular supplements) can be
   590 employed for primary and supportive nutritional therapy in active IBD.
- 591 **Grade of recommendation 0 strong consensus (96 % agreement)**
- 592

#### 593 **Recommendation 18B**

- 594 **Specific formulations or substrates (e.g. glutamine, n-3-fatty acids) are not rec-**595 **ommended in use of EN or PN in IBD patients.**
- 596 **Grade of recommendation B strong consensus (96 % agreement)**
- 597

#### 598 **Commentary for A/B**

Several studies have compared the efficacies of different types (elemental, semi-599 600 elemental, oligomeric or polymeric diets) of enteral formulas in the management of ac-601 tive CD. A Cochrane meta-analysis of ten trials showed no statistically significant differ-602 ence between patients treated with elemental (n=188), and non-elemental diet (semi-603 elemental or polymeric diet; n=146) (90). The protein composition did not appear to 604 influence the therapeutic potential of EN. The present systematic enquiry reveals insuf-605 ficient evidence to make firm recommendations (90, 91). It is therefore advised that standard feeds are employed if primary nutritional therapy is being employed. 606

The use of feeds supplemented with growth factors, ones with lower levels of emulsifying data, or oligomeric feeds, as alternatives to standard feeds, is not supported by relia-

- 609 ble data. Equally there is no evidence that any of these alternatives is inferior to the use 610 of standard polymeric feeds (92).
- 611 There are not enough data to dictate the use of specific substrates in the composition of
- 612 PN in CD. PN must however be adjusted to fulfil the needs of the individual patient.
- 613

#### 614 **Recommendation 19**

# In CD patients every effort should be made to avoid dehydration to minimize the risk of thromboembolism.

- 617 Grade of recommendation GPP strong consensus (100 % agreement)
- 618

#### 619 **Commentary**

- 620 Although there are insufficient data to mandate routine anticoagulation, this should be 621 considered in all IBD patients and especially those on PN, with every effort made to
- 622 avoid dehydration (93-97).
- 623

#### 624 **Recommendation 20A**

- 625 **CD** patients with a distal (low ileal or colonic) fistula and low output can usually 626 receive all nutritional support via the enteral route (generally as food).
- 627 Grade of recommendation 0 strong consensus (100 % agreement)
- 628

#### 629 **Recommendation 20B**

- 630 **CD** patients with a proximal fistula and/or a very high output should receive nu-631 tritional support by partial or exclusive PN.
- 632 **Grade of recommendation B strong consensus (96 % agreement)**
- 633

#### 634 **Commentary for A/B**

Patients with CD are prone to fistulae formation between two intestinal sites or from intestine to another organ (especially skin, bladder and vagina). Most occur postoperatively. It is demonstrated that in surgical patients, early nutritional support, independently of the route of administration, decreases the occurrence and severity of fistulae (84, 98, 99). Malnutrition with BMI <20 appears as an independent risk factor (100).

640 Treatment of intestinal fistulae is usually complex, depending on the location, scale and 641 the nature of the symptoms, and warrants the input of a multidisciplinary team includ-642 ing gastroenterologist, surgeon and dietician (99). In patients with a distal (low ileal or colonic) fistula it may be possible to provide all necessary nutritional support via the 643 644 enteral route (101-103). In the patient with a proximal fistula and/or a very high output 645 it may be preferable to manage the situation with a rested gut and full PN (104, 105), 646 but even then, the psychological benefit of eating may warrant its inclusion in the nutri-647 tional regimen despite minimal expectations of useful nutrient absorption (102). Surgi-

- 648 cal correction is more likely to be successful if nutritional status has been optimized pre-649 operatively (106).
- 650

#### 651 **Recommendation 21**

In CD patients in whom nutritional deprivation has extended over many days,
 standard precautions and interventions to prevent refeeding syndrome are man datory, particularly with respect to phosphate and thiamine.

- 655 Grade of recommendation B strong consensus (100 % agreement)
- 656

#### 657 Commentary

658 Refeeding syndrome should not be a problem in the well-managed patient with IBD but 659 nonetheless it is not unusual to encounter patients in whom nutritional deprivation has 660 extended over many days and in whom this hot issue is pertinent. Standard precautions 661 and interventions are mandatory in these high-risk patients particularly in respect of

- 662 phosphate and thiamine (107-109).
- 663

#### 664 **Recommendation 22A**

- 665 **EN appears safe and can be recommended as supportive therapy according to** 666 **standard nutritional practice in patients with severe UC.**
- 667 Grade of recommendation GPP strong consensus (100 % agreement)
- 668
- 669 Recommendation 22B
- 670 **PN should not be used in UC unless intestinal failure occurs.**
- 671 Grade of recommendation 0 consensus (88 % agreement)
- 672

#### 673 Commentary for A/B

EN has not been adequately evaluated in active UC. However, it appears safe and can be
nutritionally adequate in patients with severe disease (110). Its efficacy needs to be
tested by additional studies in larger cohorts of patients.

PN is recommended in malnourished patients with UC and in those with severe disease,
only when they not able to tolerate EN, or cannot be fed effectively by either mouth or
enteric tube (110-112).

- 680
- 681 V. Surgical aspects of nutrition in IBD (Figures 8 and 9)
- 682

#### 683 Recommendation 23A

- 684 In most elective surgery cases, pre-operative fasting from midnight should not be
- 685 performed instead, an enhanced recovery (ERAS) protocol can be used.

#### 686 **Grade of recommendation B, see ESPEN Surgery guideline (113) – strong consen**-687 **sus (100 % agreement)**

688

#### 689 **Commentary**

ESPEN has produced guidance on nutrition in the surgical patient (113) and most of the
principles apply equally to the IBD patient undergoing surgical intervention. The subsequent guidance should be followed during the perioperative period. From a metabolic
and nutritional point of view, the key aspects of perioperative care include:

- avoidance of long periods of pre- operative fasting
- re-establishment of oral feeding as early as possible after surgery
- integration of nutrition into the overall management of the patient
- 697 metabolic control e. g. of blood glucose
- reduction of factors exacerbating stress related catabolism or impair GI function
- early mobilization to facilitate protein synthesis and muscle function.
- 700

#### 701 Recommendation 23B

In emergency surgery patients, medical nutrition (EN, PN) should be initiated if
the patient is malnourished at the time of surgery or if oral diet cannot be recommenced within 7 days after surgery.

- Grade of recommendation B, see ESPEN Surgery guideline (113) consensus
   (88 % agreement)
- 707

#### 708 **Commentary**

Nutritional support is indicated in patients with malnutrition and even in patients without significant malnutrition, if it is anticipated that the patient will be unable to eat for more than seven days perioperatively. It is also indicated in patients who cannot maintain oral intake above 60-75% of recommended intake for more than ten days. In these situations, it is recommended to initiate nutritional support (preferably by the enteral

- 714 route) without delay.
- 715

#### 716 **<u>Recommendation 24A</u>**

#### 717 Patients who do not meet their energy and/or protein needs from normal food

718 should be encouraged to take oral nutritional supplements (ONS) during the peri-719 operative period.

- 719 operative period.
- 720 Grade of recommendation B strong consensus (100 % agreement)
- 721

#### 722 **Commentary**

- 723 Insufficient preoperative intake is an indication for dietary counselling or ONS, because
- as Kuppinger et al. (114) showed for patients undergoing abdominal surgery, lower food
- intake before hospital admission is an independent risk factor for postoperative compli-
- 726 cations. Twenty-four trials on the use of ONS and EN have reported significant ad-

- 727 vantages from EN with particular regard to the reduction of infectious complications, 728 length of hospital stay and costs. In six RCTs postoperative and post-hospital admin-729 istration of ONS has been investigated (115-119). The available data do not show with 730 certainty that routine administration improves outcome, but they do show benefit in 731 terms of nutritional status, rate of minor complications, well-being and quality of life in
- 732 patients who cannot meet their nutritional requirements at home from normal food.
- 733

#### 734 **Recommendation 24B**

#### 735 Patients who do not meet their energy and/or protein needs from normal food 736 plus ONS should receive EN during the perioperative period.

- 737 Grade of recommendation B - strong consensus (100 % agreement)
- 738

#### 739 **Commentary**

- 740 As stated above, insufficient preoperative intake affects complication rates. Therefore, if 741 the oral intake is inadequate, regardless of the intervention (oral food or ONS), EN should be initiated (113). Postoperatively, EN should be continued/started as many 742
- 743 studies have shown the benefits and feasibility of feeding via a tube either inserted dis-
- tal to the anastomosis, e.g. needle catheter jejunostomy, or inserted via the nose with its 744
- 745 tip passed distally at the time of operation (nasojejunal tube) (120-125).
- 746

#### 747 **Recommendation 24C**

- 748 If malnutrition is diagnosed, then IBD surgery should be delayed for 7-14 days 749 whenever possible, and that time should be used for intensive medical nutrition.
- 750 Grade of recommendation A, see ESPEN Surgery guideline (113) - strong consen-751 sus (96 % agreement)
- 752

#### 753 **Commentary**

754 Undernutrition has a negative impact on the clinical course, the rate of postoperative 755 complications and on mortality (126-131). Therefore, patients with severe nutritional 756 risk will benefit from nutritional therapy prior to major surgery even if surgery has to be 757 delayed. "Severe" nutritional risk has been defined by an ESPEN working group (2006) 758 as the presence of at least one of the following criteria:

- 759 • Weight loss > 10-15% within six months
- 760 • BMI < 18.5 kg/m2
- 761 Serum albumin < 30g/l (with no evidence of hepatic or renal dysfunction) •
- 762

#### 763 **Recommendation 25A**

764 EN should always be preferred over the parenteral route, but combinations of EN

765 and PN should be considered in patients in whom there is an indication for nutri-

766 tional support and in whom >60% of energy needs cannot be met via the enteral 767 route.

- 768 Grade of recommendation A, see ESPEN Surgery Guideline (113) strong consen-
- 769 **sus (100 % agreement)**
- 770
- 771 Recommendation 25B
- PN in the perioperative period in IBD patients should be usually used as supple-mentary to EN.
- 774 **Grade of recommendation B strong consensus (96 % agreement)**
- 775
- 776 **Recommendation 25C**
- PN shall be used as the only intervention if EN is impossible (absence of access,
   severe vomiting or diarrhea) or contraindicated (intestinal obstructions or ileus,
- 779 severe shock, intestinal ischemia).
- 780 **Grade of recommendation A strong consensus (96 % agreement)**
- 781
- 782 **Commentary for A/B/C**
- The enteral route should always be preferred except when one or more of the followingcontraindications:
- Intestinal obstructions or ileus,
- Severe shock
- Intestinal ischemia
- High output fistula
- Severe intestinal hemorrhage
- 790 In those cases, PN may be needed for a period of days or weeks until the function of gas-
- trointestinal tract returns. For further details, see the ESPEN guideline on Clinical Nutrition in Surgery (113).
- 793

#### 794 **<u>Recommendation 26A</u>**

#### 795 Surgical patients with CD should obtain early nutritional support, because, inde-

- 796 pendently of the route of administration, it decreases the risk of postoperative 797 complications.
- 798 **Grade of recommendation B strong consensus (100 % agreement)**
- 799
- 800 **Commentary**
- 801 The advantages of early EN within 24 hours of surgery versus later commencement have
- been shown in two meta-analyses (one Cochrane systematic review) (132, 133).
- 803

#### 804 **Recommendation 26B**

805 In CD patients with prolonged gastrointestinal failure (such as patients in whom 806 resection has created a short bowel) PN is mandatory and life-saving at least in 807 the early stages of intestinal failure.

- 808 **Grade of recommendation B, see ESPEN surgery guideline strong consensus** 809 **(92 % agreement)**
- 810
- 811 **Commentary for A/B**

Although EN has proven to be the most beneficial in almost all patient populations, it is
relatively rare that it is sufficient in acute intestinal failure/ enterocutaneous fistulae
individuals because of the compromised integrity of the gastrointestinal tract. Therefore,
PN often represents the main option, alone or in association with EN (supplemental PN)
(72).

817

#### 818 **Recommendation 27A**

Normal food intake or EN can be commenced early after surgery in most IBD patients in the postoperative phase.

- 821 Grade of recommendation 0, see ESPEN surgery guideline strong consensus
  822 (100 % agreement)
- 823

#### 824 Recommendation 27 B

In the early phase after proctocolectomy or colectomy, water and electrolytes
 shall be administered to assure hemodynamic stability.

- 827 Grade of recommendation A, see ESPEN surgery guideline strong consensus
  828 (96 % agreement)
- 829

#### 830 **Commentary for A/B**

831 As stated in the Surgical Guidelines (113), early normal food or EN, including clear liq-832 uids on the first or second postoperative day, does not cause impairment of healing of 833 anastomoses in the colon or rectum and leads to significantly shortened hospital length 834 of stay. This has been emphasized by a Cochrane Systematic Review (129). Recent meta-835 analyses (133-135) showed significant benefits with regard to postoperative recovery 836 and infection rate. Early postoperative nutrition is associated with significant reductions 837 in total complications compared with traditional postoperative feeding practices and 838 does not negatively affect outcome such as mortality: anastomotic dehiscence, resump-839 tion of bowel function, or hospital length of stay (135).

840

#### 841 V. Dietetic recommendations during remission (Figures 10 and 11)

#### 842 Recommendation 28

843 All IBD patients in remission should undergo counselling by a dietician as part of

- 844 the multidisciplinary approach to improve nutritional therapy and to avoid mal-845 nutrition and nutrition-related disorders.
- 846 **Grade of recommendation GPP strong consensus (100 % agreement)**
- 847

#### 848 **Commentary**

849 There are very limited original data in this area, but at least nine papers include state-850 ments indicating that the input of a dietician is likely to be helpful in IBD management in 851 adults and children; the evidence base is poor. Nutritional deficiencies are self-evidently 852 more likely in patients with CD affecting the small bowel than in those with isolated co-853 lonic disease or UC, but the latter groups can be afflicted also (102). Nutritional screen-854 ing has been adopted as a mandatory component of gastrointestinal management in 855 many European countries, and it is further recommended that all IBD patients have ac-856 cess to a dietician with a special expertise in IBD

- 856 cess to a dietician with a special expertise in IBD.
- 857

#### 858 **Recommendation 29**

- 859 No specific diet needs to be followed during remission phases of IBD.
- 860 **Grade of recommendation 0 strong consensus (96 % agreement)**
- 861

#### 862 Commentary

863 In general, no specific diet needs to be followed during remission phases. None of the 864 alternative diets or semi-exclusive diets seems effective in obtaining remission. Howev-865 er, individual food intolerances are frequently seen in IBD patients, lactose and dairy 866 products, spices, herbs, fried, gas-generating and fiber rich products are often poorly 867 tolerated (136-139).

868 Patients with CD typically select a diet low in fiber and vegetables, and often one which 869 is hypocaloric and associated with multiple micronutrient deficiencies (40). Acquired 870 lactase deficiency is particularly prevalent in patients with proximal CD and will warrant 871 a lactose-restricted diet. Specific exclusion diets have been considered to have good effects by their protagonists, but for best results it is proposed that the diets should be 872 873 customized to avoid the patients' individual food intolerances. This strategy then makes 874 it difficult to generalize and there are no recent trials of exclusion diets. Limited con-875 trolled data support the elimination of lactose, dairy products in general, spices, herbs, 876 fried foods, gas-generating and fiber-rich products, but only when they are poorly tolerated. Their removal is then probably helpful in prolonging remission (140). Other stud-877 ies of reasonable quality have also included dietary manipulations, but alongside the use 878 879 of nutritional supplements; these studies are addressed in later sections. The use of an 880 exclusive EN regimen is clearly an extreme form of dietary exclusion.

881 EN has been thought to have a role in preventing relapse in children with inactive CD

- 882 (77, 90, 141, 142) and the effect has also been observed in a Japanese study of adult CD
- patient (143-145). Esaki et al. (146) considered from their trial of 145 patients with CD (mostly induced into remission with total PN) that, under maintenance with ele-

- Journal Pre-proof
- 885 mental/polymeric nutrition, the risk of recurrence was lower in those with small bowel
- rather than large bowel involvement. However, the present systematic enquiry has indi-
- cated that overall the use of elemental EN is ineffective in maintaining remission in CD.
- 888 This is therefore due for a verdict of not recommended. The panel considers this a con-
- 889 troversial conclusion, especially in view of a previous Cochrane evaluation which con-
- sidered that ongoing EN may help maintenance of remission and reduce use of cortico-
- steroids in CD (86, 146). No recommendation is therefore made.
- 892

### 893 **Recommendation 30**

#### 894 **Supplementation with n-3 fatty acids should not be advised to support mainte-**895 **nance of remission in patients with IBD.**

- 896 **Grade of recommendation B strong consensus (100 % agreement)**
- 897

## 898 **Commentary**

899 Systematic reviews have reached the conclusion that supplementing the diet with n-3 900 fats is ineffective in the maintenance of remission of patients with UC (147, 148). This is

- 901 therefore not advised. The above data were obtained in adults. It appears reasonable to
- 902 extrapolate the conclusions into pediatric practice. The latest Cochrane review (149) has
- 903 concluded that n-3 fatty acids are probably ineffective for maintenance of remission in904 CD.
- 905

## 906 **<u>Recommendation 31</u>**

# 907 Non-specific high fiber diets should not normally be recommended for mainte908 nance of remission in IBD.

## 909 Grade of recommendation 0 - strong consensus (96 % agreement)

910

## 911 **Commentary**

912 Much of the recent literature relates to the effects of specific agents chosen as prebiotics 913 and these are not considered here, but it is recognized that many forms of fiber will have 914 an important effect on the gut microbiota and thus possibly on the maintenance of re-915 mission in IBD. It is generally agreed that dietary fiber is unwise in patients known to 916 have intestinal stricturing (GPP), but the evolving literature suggests that prebiotic fi-917 bers may be useful in maintenance of remission in some patients with UC. Several small 918 controlled studies have shown apparent benefit from the addition of fiber to the diet of 919 patients with UC (150-152). Given that the effects in maintaining remission were similar 920 for germinated barley, ispaghula husk and *Plantago ovata* seeds it may be reasonable to

- 921 conclude that this is a generic effect of increased dietary fiber.
- 922 Fiber is more often relatively contra-indicated in CD because of the presence of stric-
- 923 tures, and fiber in the form of the prebiotic fructo-oligosaccharide is apparently ineffec-
- 924 tive in CD (47). However, in a loosely controlled study of wheat fiber supplementation 925 the supplemented patients did better in respect of quality of life and had no apparent
- adverse events (153). There is another recent study of fiber supplementation that also

- 927 claims benefit, and this was through the uncontrolled use of an ovo-vegetarian diet with
- 928 over 30 g of fiber for every 2000 kcal. Maintenance of remission to one year was a re-
- markable 92% (154). See also recommendation 8.
- 930

#### 931 Recommendation 32A

- 932 **Probiotic therapy should be considered for the maintenance of remission in UC.**
- 933 Grade of recommendation B strong consensus (96 % agreement)
- 934

#### 935 **Recommendation 32B**

- 936 **Probiotic therapy should not be used for maintenance of remission in CD.**
- 937 Grade of recommendation 0 strong consensus (100 % agreement)
- 938

#### 939 **Commentary for A/B**

940 The E. coli Nissle 1917 strain and the multispecies formulation "VSL#3" have benefit, 941 supported by meta-analysis (155) in the maintenance of remission in patients – includ-942 ing children - with mild to moderate UC, in comparison to 5-aminosalicylate compounds 943 (74, 156, 157). Other probiotic preparations have been studied but although they have usually been well tolerated with trends toward benefit, significant effectiveness has not 944 945 been demonstrated (158, 159). A cautionary note exists for Lactobacillus rhamnosus GG; 946 case reports in both children and adults describe bacteremia with the administered pro-947 biotic in patients with acute severe UC (160, 161).

Probiotics are probably ineffective in preventing disease recurrence for patients with CD
(157). Although some positive claims are made no unequivocal benefit can be discerned
(162-167). Probiotics are not currently recommended.

- 951 "VSL#3" refers only to the probiotic product used in the cited literature and equivalent
- 952 products independent from the present product labeling.
- 953

## 954 **Recommendation 33A**

#### 955 **Colectomized patients with a pouch and pouchitis should be treated with a probi** 956 **otic mixture ("VSL#3"), if antibiotic treatment has failed.**

- 957 **Grade of recommendation B strong consensus (96 % agreement)**
- 958
- 959 **Recommendation 33B**
- 960 The probiotic mixture "VSL#3" may be used for primary and secondary preven-
- tion of pouchitis in patients with UC who have undergone colectomy and pouch anal anastomosis.
- 963 Grade of recommendation B strong consensus (100 % agreement)
- 964
- 965 **Commentary for A/B**

Antibiotics (ciprofloxacin, metronidazole) are the treatment of reference of acute 966 967 pouchitis (168). Two double-blind placebo-controlled trials performed in adults showed effectiveness of a particular probiotic mixture ("VSL#3" containing 450 billion colony 968 969 forming units of eight lactic acid bacteria: B. breve, B. longum, B. infantis, L. acidophilus, L. 970 casei, L. delbrueckii, L. plantarum and Streptococcus salivarius subsp. thermophilus) in 971 maintaining remission in patients with chronic pouchitis (169, 170). A pooled analysis 972 of these two studies (76 participants) suggests that this bacteriotherapy may be more 973 effective than placebo for maintenance of remission. Eighty-five per cent (34/40) of 974 verum patients maintained remission at nine to twelve months compared to 3% (1/36) 975 of placebo patients (RR 20.24). A GRADE analysis indicated that the quality of evidence 976 supporting this outcome was low due to very sparse data (35 events) (171). In another 977 study (168) effects of this bacteriotherapy were evaluated as an adjunctive to a standard 978 therapy. The decrease in UC disease activity index (UCDAI) scores of 50% or more was 979 higher in the verum group than in the placebo group (63.1 vs. 40.8; per protocol 980 P=0.010). Remission was higher in the verum group than in the placebo group (47.7% vs. 32.4%; P=0.069). 981

982 Prevention of pouchitis: The results of a small study (40 participants) suggest that the 983 bacteriotherapy may be more effective than placebo for prevention of pouchitis (172). 984 Ninety per cent (18/20) of verum patients had no episode of acute pouchitis during the 985 twelve-month study compared to 60% (12/20) of placebo patients (RR 1.50). A GRADE 986 analysis indicated that the quality of evidence supporting this outcome was low due to 987 very sparse data (30 events). In contrast, Lactobacillus rhamnosus strain GG was not ef-988 fective in preventing relapses (173). ECCO guidelines suggest the use of "VSL#3" both 989 for maintenance of antibiotic-induced remission and for prevention of pouchitis in 990 adults (174) and in pediatric UC (175).

991 "VSL#3" refers only to the probiotic product used in the cited literature and equivalent992 products independent from the present product labeling.

993

#### 994 Recommendation 34A

- 995 Neither EN nor PN is recommended as primary therapy for maintaining remission
  996 in IBD.
- 997 Grade of recommendation GPP strong consensus (100 % agreement)
- 998

#### 999 Recommendation 34B

- 1000 **ONS or EN can be recommended in patients with CD in remission, if undernutri** 1001 **tion cannot be treated sufficiently by dietary counselling.**
- 1002 **Grade of recommendation GPP strong consensus (100 % agreement)**
- 1003

#### 1004 **Commentary for A/B**

Nutritional support has not been assessed as a maintenance therapy in UC, neither has
PN in CD. A recent systematic review of twelve RCTs and non-randomized cohort studies
(176) (1169 patients, including 95 children), most of good quality, showed that maintenance EN was as or more effective than the comparator (standard diet, 5-ASA or azathi-

25

1009 oprine) in preventing CD relapses over periods of six months to four years. The study 1010 with the lowest risk of bias compared supplemental (50%) EN with a regular diet in 51 adult CD patients (177). Patients in each arm of the study were on similar medications 1011 1012 (5-ASA or azathioprine). The study showed that in the EN group, nine of 26 patients 1013 (34%) had a relapse during a mean follow-up of 11.9 months, as compared with 16 of 25 1014 patients (64%) in the non-EN group (HR = 0.40; 95%CI 0.16 – 0.98; P < 0.01). The study 1015 of maintenance EN as an adjuvant to infliximab therapy has yielded conflicting results, 1016 with one negative (144) and two positive (178, 179) studies published so far.

1017 Elemental formulas have been the most studied. A systematic review was unable to 1018 show any significant difference in remission rate between elemental and polymeric for-1019 mulae (180). However, it found a lower adherence rate for elemental EN compared to an unrestricted diet. The European organizations for IBD and for pediatric gastroenterolo-1020 1021 gy and nutrition, ECCO and ESPGHAN, have advised on the possible use of partial maintenance EN in patients with very mild disease or a low risk of relapse, preferring 1022 polymeric feeds, with elemental feeds being advised only in the case of allergy to cow's 1023 1024 milk proteins (181).

1025

## 1026 **<u>Recommendation 35</u>**

Standard diet or ONS should be followed in patients with IBD in remission, giving
 attention to nutrition screening and generic nutritional support where needed.

- 1029 Grade of recommendation: GPP strong consensus (95 % agreement)
- 1030

## 1031 **Commentary**

Few dietary supplementations have been tested in maintenance of remission in IBD patients with clinical endpoints. An open label, parallel-group, multicenter, randomized clinical trial demonstrated in 105 UC patients in remission that plantago ovata seeds (10 g twice daily) were as efficient as mesalamine (500 mg thrice daily) in maintaining remission to one year (151). A Cochrane systematic review has analyzed six studies (1039 patients) of n-3 fatty acid supplementation (149): there was a marginal significant benefit of n-3 therapy on maintenance of remission.

1039

## 1040 **Recommendation 36:**

#### 1041 When more than 20 cm of distal ileum, whether or not in combination with the 1042 ileo-cecal valve, is resected, vitamin B12 shall be administered to patients with CD.

- 1043 **Grade of recommendation A strong consensus (100 % agreement)**
- 1044

## 1045 **Commentary**

1046 A recent systematic review has assessed the literature for prevalence, risk factors, eval-

1047 uation and management of vitamin B12 deficiency in IBD (182). Unresected UC does not

1048 predispose to low B12 levels or B12 deficiency. The prevalence of B12 deficiency in CD

- 1049 ranges from 5.6 to 38%. Resection of more than 30 cm of distal ileum, whether or not in
- 1050 combination with the ileo-cecal valve, will put the patient at risk for B12 deficiency. Re-

- section of less than 20 cm does not normally cause deficiency (183). Ileal CD is not inevi-
- tably associated with vitamin B12 deficiency (184, 185), but it is difficult to rule out its responsibility when more than 30-60 cm are involved (182). CD patients with ileal in-
- volvement and/or resection and/or clinical deficiency features should be screened year-
- 1055 ly for vitamin B12 deficiency (182).

1056 Patients with clinical deficiency should receive 1000 µg of vitamin B12 by intramuscular injection every other day for a week and then every month for life (186). Patients with 1057 1058 more than 20 cm of ileum resected should receive 1000 µg of vitamin B12 prophylacti-1059 cally also every month and indefinitely (186). Oral therapy may be as effective but is poorly explored in CD. A retrospective open-label non-randomized study of 36 CD pa-1060 tients has showed the oral route (1200 µg per day for 33, 2400 µg per day for three) to 1061 be effective in treating vitamin B12 deficiency (187). For now, parenteral supplementa-1062 1063 tion remains the reference, but oral supplementation may become standard in the com-1064 ing years.

1065

#### 1066 **Recommendation 37:**

# Selected IBD patients, e. g. those treated with sulphasalazine and methotrexate, should be supplemented with vitamin B9 / folic acid.

1069 **Grade of recommendation B – strong consensus (100 % agreement)** 

#### 1070 Commentary

1071 There are several causes for folate deficiency in IBD: low intake, malabsorption, excess 1072 folate utilization due to mucosal inflammation and medications. A combination of these

- 1072 for the deficiency of this vitamin. Drugs are most responsible
- 1074 for folate deficiency by inhibition of dihydrofolate reductase, an enzyme that catalyzes
- 1075 reduction of dihydrofolic acid to tetrahydrofolic acid (methotrexate) (188) or folate 1076 malabsorption (sulphasalazine) (189). Azathioprine and 6-mercaptopurine also induce
- 1077 macrocytosis but through myelosuppressive activity.
- 1078 A systematic review and meta-analysis of 10 studies reporting on 4517 patients found 1079 an overall protective effect for folic acid supplementation on the development of colo-1080 rectal cancer (pooled HR = 0.58; 95%CI 0.37 - 0.80) (190). An Italian study compared one month of supplementation with 15 mg of either folic or folinic acid in 30 IBD pa-1081 tients treated with sulphasalazine (191). Both were able to restore the body stores of 1082 1083 folate, but folinic acid was more efficient. The ECCO-ESPGHAN guidelines on the medical 1084 management of pediatric CD advise oral administration of folate in patients on methotrexate, 5 mg once weekly 24-72 hours after the methotrexate, or 1 mg daily for five 1085 1086 days per week (181). This panel recommends the same practice in adults.
- 1087

## 1088 **Recommendation 38A**

1089 In IBD patients who are pregnant, iron status and folate levels should be moni-1090 tored regularly and in the case of deficiencies, iron and/or vitamin B9/folic acid 1091 should be additionally supplemented.

- 1092 Grade of recommendation: GPP strong consensus (95 % agreement)
- 1093

#### 1094 **Recommendation 38B**

1095 **In IBD patients who are breastfeeding, nutritional status should be monitored** 1096 **regularly and in case of deficiencies, they should be supplemented** 

- 1097 Grade of recommendation: GPP strong consensus (100 % agreement)
- 1098

#### 1099 **Commentary for A/B**

1100 The consequences of anemia and those of neural tube defects (192), along with the fre-1101 quent deficiencies in IBD patients warrant regular screening for iron and folate deficien-1102 cies, respectively, during pregnancy, along with nutritional follow-up.

1103 There is little information available that is specific to the situation of the woman with 1104 IBD who is considering breastfeeding. However, there is no evidence of harm from the 1105 use of any nutritional intervention that is thought otherwise appropriate as part of the

- 1105 use of any nutritional intervention that is thought otherwise appropriate as part of the 1106 management of the new mother.
- 1107

#### 1108 **Recommendation 39**

1109 In all IBD patients, endurance training should be encouraged. In IBD patients with

1110 decreased muscle mass and/or muscle performance, appropriate physical activity

- 1111 should be recommended.
- 1112 Grade of recommendation: GPP strong consensus (95 % agreement)
- 1113

#### 1114 **Commentary**

The systematic review of 19 body composition studies reporting on 926 IBD patients
revealed a low fat-free mass in 28% of CD patients and in 13% of UC patients (193). Low
muscle mass, strength and performance have been reported in adult IBD cohorts
(194,195), similar findings have also been made in children (196). Sarcopenia was re-

- 1119 ported in 12% of IBD patients of mean age 31 years, associated with osteopenia (194).
- 1120 In a German study, 30 patients, aged 41±14 years, with mild to moderate IBD were ran-1121 domized to either supervised moderate-intensity running thrice a week for ten weeks or 1122 to a control group with no exercise. Health-related quality of life, reported as IBDQ total 1123 score, improved by 19% in the intervention group and 8% in the control group, with 1124 significant differences for the IBDQ social sub-scale that was significantly improved in
- 1125 the intervention group compared with controls (p = 0.023) (197).
- 1126 The reference treatment for sarcopenia, along with maintaining an adequate protein 1127 intake, is resistance training. This is what is advised in age-related sarcopenia (198). 1128 However, this hasn't been assessed in IBD patients. Still, the panel recommends pre-1129 scribing resistance training (weight-bearing exercises) in IBD patients with sarcopenia 1130 or features of sarcopenia (reduced muscle mass, strength and/or performance).
- 1131

#### 1132 **Recommendation 40**

## 1133 **Obese IBD patients should be advised to reduce weight only in phases of stable**

1134 remission and then according to current obesity guidelines.

#### 1135 **Grade of recommendation: GPP – strong consensus (100 % agreement)**

1136

#### 1137 Commentary

Overweight and obesity are nowadays the most frequent nutritional disorder in IBD pa-1138 tients. Their prevalence varies between countries, affecting 32.7% of 581 US adult IBD 1139 1140 patients (30.3% in CD patients and 35.2 in UC patients) (199) and 17% of 100 Irish adult CD patients (200). An US study of 1494 IBD patients (31.5% obese) found an association 1141 between obesity and its usual comorbidities, a poor quality of life and high C-reactive 1142 1143 protein levels (201). However, obesity was not associated with increased health care utilization or IBD-related surgery. No intervention study has addressed the treatment of 1144 1145 obesity in IBD patients. However, the high prevalence of both micronutrient deficiencies 1146 and sarcopenia, here indicating sarcopenic obesity, indicates that the patient on a re-1147 strictive diet is at risk of further deficiencies and muscle mass loss, especially in catabol-1148 ic states such as those associated with IBD flares. Therefore, the panel recommends 1149 against low-calorie diets in patients with active disease and recommends endurance 1150 training as the first step in any effort to lose weight.

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#### 1153 **References**

11541.Forbes A, Escher J, Hébuterne X, Kłęk S, Krznaric Z, Schneider S, et al. ESPEN guideline: Clinical1155nutrition in inflammatory bowel disease. Clin Nutr. 2017;36:321-47.

11562.Bischoff SC, Singer P, Koller M, Barazzoni R, Cederholm T, van Gossum A. Standard operating pro-<br/>cedures for ESPEN guidelines and consensus papers. Clin Nutr. 2015;34:1043-51.

11583.Hou JK, Abraham B, El-Serag H. Dietary Intake and Risk of Developing Inflammatory Bowel Dis-<br/>ease: A Systematic Review of the Literature. Am J Gastroenterol 2011;106:563–73.

4. Ananthakrishnan AN, Khalili H, Konijeti GG, Higuchi LM, de Silva P, Korzenik JR, et al. A prospective study of long-term intake of dietary fiber and risk of Crohn's disease and ulcerative colitis. Gastroenterology 2013;145:970-7.

11635.Li F, Liu X, Wang W, Zhang D. Consumption of vegetables and fruit and the risk of inflammatory1164bowel disease: a meta-analysis. Eur J Gastroenterol Hepatol. 2015;27:623-30.

Ananthakrishnan AN, Khalili H, Konijeti GG, Higuchi LM, de Silva P, Fuchs CS, et al. Long-term
intake of dietary fat and risk of ulcerative colitis and Crohn's disease. Gut 2014;63:776-84.

Tjonneland A, Overvad K, Bergmann MM, Nagel G, Linseisen J, Hallmans G, et al. Linoleic acid, a
dietary n-6 polyunsaturated fatty acid, and the aetiology of ulcerative colitis: a nested case-control study
within a European prospective cohort study. Gut 2009;58:1606-11.

11708.Klement E, Cohen RV, Boxman J, Joseph A, Reif S. Breastfeeding and risk of inflammatory bowel1171disease: a systematic review with meta-analysis. Am J Clin Nutr 2004; 80:1342-52.

Barclay AR, Russell RK, Wilson ML, Gilmour WH, Satsangi J, Wilson DC. Systematic review: the
role of breastfeeding in the development of pediatric inflammatory bowel disease. J Pediatr 2009;155:
421-6.

- 117510.Gearry RB, Richardson AK, Frampton CM, Dodgshun AJ, Barclay ML. Population-based cases con-<br/>trol study of inflammatory bowel disease risk factors. J Gastroenterol Hepatol 2010;25:325-33.
- 1177 11. Hansen TS, Jess T, Vind I, Elkjaer M, Nielsen MF, Gamborg M, et al. Environmental factors in inflammatory bowel disease: a case-control study based on a Danish inception cohort. J Crohns Colitis.
  2011;5:577-84.
- 118012.Guo AY, Stevens BW, Wilson RG, Russell CN, Cohen MA, Sturgeon HC, et al. Early life environment1181and natural history of inflammatory bowel diseases. BMC Gastroenterol. 2014;14:216.
- 1182 13. Ng SC, Tang W, Leong RW, Chen M, Ko Y, Studd C, et al. Asia-Pacific Crohn's and Colitis Epidemiology Study ACCESS Group. Environmental risk factors in inflammatory bowel disease: a population-based
  case-control study in Asia-Pacific. Gut. 2015;64:1063-71.

1185 14. ESPGHAN Committee on Nutrition, Agostoni C, Braegger C, Decsi T, Kolacek S, Koletzko B,
1186 Michaelsen KF, et al. Breast-feeding: A commentary by the ESPGHAN Committee on Nutrition. J Pediatr
1187 Gastroenterol Nutr. 2009;49:112-25.

- 1188 15. Nguyen GC, Munsell M, Harris ML. Nationwide prevalence and prognostic significance of clinically
  diagnosable protein-calorie malnutrition in hospitalized inflammatory bowel disease patients. Inflamm
  Bowel Dis 2008;14:1105-11.
- Sandhu A, Mosli M, Yan B, Wu T, Gregor J, Chande N, et al. Self-Screening for Malnutrition Risk in
  Outpatient Inflammatory Bowel Disease Patients Using the Malnutrition Universal Screening Tool (MUST).
  JPEN. J Parenter Enteral Nutr 2016;40:507-10.

1194 17. Gajendran M, Umapathy C, Loganathan P, Hashash JG, Koutroubakis IE, Binion DG. Analysis of
hospital-based emergency department visits for inflammatory bowel disease in the USA. Dig Dis Sci
2016;61:389-99.

1197 18. Ananthakrishnan AN, McGinley EL. Infection-related hospitalizations are associated with in-1198 creased mortality in patients with inflammatory bowel diseases. J Crohns Colitis 2013;7:107-12.

- 1199 19. Wallaert JB, De Martino RR, Marsicovetere PS, Goodney PP, Finlayson SR, Murray JJ, et al. Venous
  thromboembolism after surgery for inflammatory bowel disease: are there modifiable risk factors? Data
  from ACS NSQIP. Dis Colon Rectum 2012;55:1138-44.
- 1202 20. Ananthakrishnan AN, McGinley EL, Binion DG, Saeian K. A novel risk score to stratify severity of
   1203 Crohn's disease hospitalizations. Am J Gastroenterol 2010;105:1799-807.
- 1204 21. Vasseur F, Gower-Rousseau C, Vernier-Massouille G, Dupas JL, Merle V, Merlin B, et al. Nutritional
  1205 Status and Growth in Pediatric Crohn's Disease: A Population-Based Study. Am J Gastroenterol.
  1206 2010;105:1893-900.
- Hill RJ, Davies PS. You look all right to me: compromised nutritional status in paediatric patientswith ulcerative colitis. J Pediatr Gastroenterol Nutr 2013;56:385-9.
- Wiskin AE, Owens DR, Cornelius VR, Wootton SA, Beattie RM. Paediatric nutrition risk scores in
   clinical practice: children with inflammatory bowel disease. J Hum Nutr Diet 2012;25:319-22.
- 1211 24. Heuschkel R, Salvestrini C, Beattie RM, Hildebrand H, Walters T, Griffiths A. Guidelines for the
  1212 management of growth failure in childhood inflammatory bowel disease. Inflamm Bowel Dis
  1213 2008;14:839-49.
- 1214 25. Inoue M, Sasaki M, Takaoka A, Kurihara M, Iwakawa H, Bamba S, et al. Changes in energy metabolism after induction therapy in patients with severe or moderate ulcerative colitis. J Clin Biochem Nutr 2015;56:215-9.
- 1217 26. Sasaki M, Johtatsu T, Kurihara M, Iwakawa H, Tanaka T, Bamba S, et al. Energy expenditure in
  1218 Japanese patients with severe or moderate ulcerative colitis. J Clin Biochem Nutr 2010;47:32-6.
- 1219
  1220 Klein S, Meyers S, O'Sullivan P, Barton D, Leleiko N, Janowitz HD. The metabolic impact of active ulcerative colitis. Energy expenditure and nitrogen balance. J Clin Gastroenterol 1988;10:34-40.
- 122128.Stokes MA, Hill GL. Total energy expenditure in patients with Crohn's disease: measurement by1222the combined body scan technique. JPEN J Parenter Enteral Nutr 1993;17:3-7.
- 122329.Capristo E, Addolorato G, Mingrone G, Greco AV, Gasbarrini G. Effect of disease localization on the1224anthropometric and metabolic features of Crohn's disease. Am J Gastroenterol. 1998;93:2411-9.
- 30. Zoli G, Katelaris PH, Garrow J, Gasbarrini G, Farthing MJ. Increased energy expenditure in growing
   adolescents with Crohn's disease. Dig Dis Sci 1996;41:1754-9.
- 122731.Steiner SJ, Noe JD, Denne SC. Corticosteroids increase protein breakdown and loss in newly diag-1228nosed pediatric Crohn disease. Pediatric research 2011;70:484-8.
- 32. O'Keefe SJ, Ogden J, Rund J, Potter P. Steroids and bowel rest versus elemental diet in the treatment of patients with Crohn's disease: the effects on protein metabolism and immune function. JPEN.
  Journal of parenteral and enteral nutrition 1989;13:455-60.
- 33. Molodecky NA, Soon IS, Rabi DM, Ghali WA, Ferris M, Chernoff G, et al. Increasing incidence and
  prevalence of the inflammatory bowel diseases with time, based on systematic review. Gastroenterology
  2012;142:46-54.
- 1235 34. Hannon TS, Dimeglio LA, Pfefferkorn MD, Denne SC. Acute effects of enteral nutrition on protein
  1236 turnover in adolescents with Crohn disease. Ped Res 2007;61:356-60.
- 1237 35. Royall D, Jeejeebhoy KN, Baker JP, Allard JP, Habal FM, Cunnane SC, et al. Comparison of amino
  1238 acid v peptide based enteral diets in active Crohn's disease: clinical and nutritional outcome. Gut
  1239 1994;35:783-7.
- 1240 36. Griffiths RD, Hinds CJ, Little RA. Manipulating the metabolic response to injury. Br Med Bull 1241 1999;55:181-95.
- 1242 37. Royall D, Greenberg GR, Allard JP, Baker JP, Jeejeebhoy KN. Total enteral nutrition support improves body composition of patients with active Crohn's disease. JPEN J Parenter Enteral Nutr 1995;19:95-9.

- 38. Gerasimidis K, Edwards C, Stefanowicz F, Galloway P, McGrogan P, Duncan A, et al. Micronutrient
  status in children with IBD: true deficiencies or epiphenomenon of the systemic inflammatory response. J
  Pediatr Gastroenterol Nutr 2013;56:e50-1.
- 124839.Filippi J, Al-Jaouni R, Wiroth JB, Hébuterne X, Schneider SM. Nutritional deficiencies in patients1249with Crohn's disease in remission. Inflamm Bowel Dis. 2006;12:185-91.
- 40. Geerling BJ, Badart-Smook A, Stockbrügger RW, Brummer RJ. Comprehensive nutritional status in
   patients with long-standing Crohn disease currently in remission. Am J Clin Nutr.1998;67:919-26.
- 125241.Vagianos K, Bector S, McConnell J, Bernstein CN. Nutrition assessment of patients with inflamma-<br/>tory bowel disease. JPEN. Jour Jarenter Enteral NUtr. 2007;31:311-9.
- 1254 42. Santucci NR, Alkhouri RH, Baker RD, Baker SS. Vitamin and zinc status pretreatment and post1255 treatment in patients with inflammatory bowel disease. J Pediatr Gastroenterol Nutr 2014;59:455-7.
- 125643.Greenley RN, Stephens KA, Nguyen EU, Kunz JH, Janas L, Goday P, et al. Vitamin and mineral sup-1257plement adherence in pediatric inflammatory bowel disease. J Pediatr Psychol 2013;38:883-92.
- 1258 44. Dignass AU, Gasche C, Bettenworth D, Birgegård G, Danese S, Gisbert JP, et al. European Crohn's
  1259 and Colitis Organisation [ECCO]. European consensus on the diagnosis and management of iron deficiency
  1260 and anaemia in inflammatory bowel diseases. J Crohns Colitis. 2015;9:211-22.
- 126145.Wells CW, Lewis S, Barton JR, Corbett S. Effects of changes in haemoglobin level on quality of life1262and cognitive function in inflammatory bowel disease patients. Inflamm Bowel Dis 2006;12:123–30.
- 46. Evstatiev R, Marteau P, Iqbal T, Khalif IL, Stein J, Bokemeyer B, et al. FERGIcor, a randomized controlled trial on ferric carboxymaltose for iron deficiency anemia in inflammatory bowel disease. Gastroenterology 2011;141:846–53, e1–2.
- 126647.Benjamin JL, Hedin CR, Koutsoumpas A, Ng SC, McCarthy NE, Hart AL, et al. Randomised, double-1267blind, placebo-controlled trial of fructo-oligosaccharides in active Crohn's disease. Gut. 2011;60:923-9.
- 48. Pironi L, Arends J, Baxter J, Bozzetti F, Peláez RB, Cuerda C, et al. Home Artificial Nutrition &
  Chronic Intestinal Failure; Acute Intestinal Failure Special Interest Groups of ESPEN. ESPEN endorsed
  recommendations. Definition and classification of intestinal failure in adults. Clin Nutr. 2015;34:171-80.
- Hart JW, Bremner AR, Wootton SA, Beattie RM. Measured versus predicted energy expenditure in
  children with inactive Crohn's disease. Clinical Nutrition 2005;24:1047-55.
- 127350.Baker ML, Williams RN, Nightingale JM. Causes and management of a high-output stoma. Colorec-1274tal Dis. 2011;13:191-7.
- 1275 51. Grischkan D, Steiger E, Fazio V. Maintenance of home hyperalimentation in patients with high-1276 output jejunostomies. Arch Surg. 1979;114:838-41.
- 1277 52. Pironi L, Guidetti C, Incasa E, Poggioli G, Paganelli F, Merli C, et al. Oral rehydration solution containing rice maltodextrins in patients with total colectomy and high intestinal output. Int J Clin Pharmacol
  1279 Res. 2000;20:55-60.
- 1280 53. Nightingale JM, Lennard-Jones JE, Walker ER, Farthing MJ. Oral salt supplements to compensate
  1281 for jejunostomy losses: comparison of sodium chloride capsules, glucose electrolyte solution, and glucose
  1282 polymer electrolyte solution. Gut. 1992;33:759-61.
- 128354.Hu D, Ren J, Wang G, Li G, Liu S, Yan D, et al. Exclusive enteral nutritional therapy can relieve in-1284flammatory bowel stricture in Crohn's disease. J Clin Gastroenterol. 2014;48:790-5.
- 1285 55. Abraham BP, Prasad P, Malaty HM Vitamin D deficiency and corticosteroid use are risk factors for
  1286 low bone mineral density in inflammatory bowel disease patients. Dig Dis Sci 2014;59:1878-84.
- 1287 56. Bakker SF, Dik VK, Witte BI, Lips P, Roos JC, Van Bodegraven AA. Increase in bone mineral density
  in strictly treated Crohn's disease patients with concomitant calcium and vitamin D supplementation. J
  1289 Crohns Colitis. 2013;7:377-84.

1290 57. Lopes LH, Sdepanian VL, Szejnfeld VL, de Morais MB, Fagundes-Neto U. Risk factors for low bone
1291 mineral density in children and adolescents with inflammatory bowel disease. Dig Dis Sci. 2008;53:27461292 53.

1293 58. van Bodegraven AA, Bravenboer N, Witte BI, Dijkstra G, van der Woude CJ, Stokkers PC, et al.
1294 Dutch Initiative on Crohn and Colitis (ICC). Treatment of bone loss in osteopenic patients with Crohn's
1295 disease: a double-blind, randomised trial of oral risedronate 35 mg once weekly or placebo, concomitant
1296 with calcium and vitamin D supplementation. Gut. 2014;63:1424-30.

1297 59. Bernstein CN, Seeger LL, Anton PA, Artinian L, Geffrey S, Goodman W, et al. A randomized, placebo-controlled trial of calcium supplementation for decreased bone density in corticosteroid-using patients
with inflammatory bowel disease: a pilot study. Aliment Pharmacol Ther. 1996;10:777-86.

130060.Hebuterne X, Filippi J, Al-Jaouni R, Schneider S. Nutritional consequences and nutrition therapy in1301Crohn's disease. Gastroenterol Clin Biol 2009;33 Suppl 3:S235-44.

1302 61. Jacobsen O, Højgaard L, Hylander Møller E, Wielandt TO, Thale M, et al. Effect of enterocoated
1303 cholestyramine on bowel habit after ileal resection: a double blind crossover study. Br Med J (Clin Res Ed).
1304 1985;290:1315-8.

1305
62. Little KH, Schiller LR, Bilhartz LE, Fordtran JS. Treatment of severe steatorrhea with ox bile in an
ileectomy patient with residual colon. Dig Dis Sci. 1992;37:929-33.

1307 63. Westergaard H. Bile Acid malabsorption. Curr Treat Options Gastroenterol. 2007;10:28-33.

Hylander E, Jarnum S, Jensen HJ, Thale M. Enteric hyperoxaluria: dependence on small intestinal
resection, colectomy, and steatorrhoea in chronic inflammatory bowel disease. Scand J Gastroenterol.
1978;13:577-88.

131165.Andersson H, Filipsson S, Hultén L. Urinary oxalate excretion related to ileocolic surgery in pa-1312tients with Crohn's disease. Scand J Gastroenterol. 1978;13:465-9.

1313 66. Hueppelshaeuser R, von Unruh GE, Habbig S, Beck BB, Buderus S, Hesse A, et al. Enteric hyperox1314 aluria, recurrent urolithiasis, and systemic oxalosis in patients with Crohn's disease. Pediatr Nephrol.
1315 2012;27:1103-9.

131667.Charlebois A, Rosenfeld G, Bressler B. The Impact of Dietary Interventions on the Symptoms of1317Inflammatory Bowel Disease: A Systematic Review. Crit Rev Food Sci Nutr. 2016;56:1370-8

1318 68. Sigall-Boneh R, Pfeffer-Gik T, Segal I, Zangen T, Boaz M, Levine A. Partial enteral nutrition with a
1319 Crohn's disease exclusion diet is effective for induction of remission in children and young adults with
1320 Crohn's disease. Inflamm Bowel Dis. 2014; 20:1353-60.

1321 69. Rajendran N, Kumar D. Food-specific IgG4-guided exclusion diets improve symptoms in Crohn's
1322 disease: a pilot study. Colorectal Dis. 2011;13:1009-13.

132370.Riordan AM, Hunter JO, Cowan RE, Crampton JR, Davidson AR, Dickinson RJ, et al. Treatment of1324active Crohn's disease by exclusion diet: East Anglian multicentre controlled trial. Lancet. 1993;1325342:1131-4.

1326 71. Jones VA. Comparison of total parenteral nutrition and elemental diet in induction of remission of
1327 Crohn's disease. Long-term maintenance of remission by personalized food exclusion diets. Dig Dis Sci.
1328 1987;32:100S-107S.

1329
 72. Slonim AE, Grovit M, Bulone L. Effect of exclusion diet with nutraceutical therapy in juvenile
 1330 Crohn's disease. J Am Coll Nutr. 2009; 28:277-85.

1331 73. Oliva S, Di Nardo G, Ferrari F, Mallardo S, Rossi P, Patrizi G, et al. Randomised clinical trial: the
effectiveness of Lactobacillus reuteri ATCC 55730 rectal enema in children with active distal ulcerative
colitis. Aliment Pharmacol Ther. 2012;35:327-34.

1334 74. Miele E, Pascarella F, Giannetti E, Quaglietta L, Baldassano RN, Staiano A. Effect of a probiotic
preparation (VSL#3) on induction and maintenance of remission in children with ulcerative colitis. Am J
Gastroenterol 2009;104:437-43.

- 1337 75. Carter MJ, Lobo AJ, Travis SP. Guidelines for the management of inflammatory bowel disease in adults. Gut 2004;53 Suppl 5:V1-16.
- 1339 76. Valentini L, Schaper L, Buning C, Hengstermann S, Koernicke T, Tillinger W, et al. Malnutrition and
  impaired muscle strength in patients with Crohn's disease and ulcerative colitis in remission. Nutrition
  2008;24:694-702.
- 1342 77. Sakamoto N, Kono S, Wakai K, Fukuda Y, Satomi M, Shimoyama T, et al. Dietary risk factors for
  1343 inflammatory bowel disease: a multicenter case-control study in Japan. Inflamm Bowel Dis 2005;11:15463.
- 1345 78. Van Limbergen J, Haskett J, Griffiths AM, Critch J, Huynh H, Ahmed N, et al. Toward enteral nutri1346 tion for the treatment of pediatric Crohn disease in Canada: A workshop to identify barriers and enablers.
  1347 Can J Gastroenterol Hepatol 2015;29:351-6.
- 1348 79. Nguyen GC, Laveist TA, Brant SR. The utilization of parenteral nutrition during the in-patient
  1349 management of inflammatory bowel disease in the United States: a national survey. Aliment Pharmacol
  1350 Ther 2007;26:1499-507.
- 1351 80. Nguyen DL, Parekh N, Bechtold ML, Jamal MM. National trends and in-hospital outcomes of adult
  1352 patients with inflammatory bowel disease receiving parenteral nutrition support. JPEN J Parenter Enteral
  1353 Nutr. 2016;40:412-6.
- 1354 81. Dziechciarz P, Horvath A, Shamir R, Szajewska H. Meta-analysis: enteral nutrition in active
  1355 Crohn's disease in children. Aliment Pharmacol Ther 2007;26:795-806.
- 1356 82. Grover Z, Lewindon P. Two-Year Outcomes After Exclusive Enteral Nutrition Induction Are Supe1357 rior to Corticosteroids in Pediatric Crohn's Disease Treated Early with Thiopurines. Dig Dis
  1358 Sci 2015;60:3069-74.
- 1359 83. Grogan JL, Casson DH, Terry A, Burdge GC, El-Matary W, Dalzell AM. Enteral feeding therapy for
  1360 newly diagnosed pediatric Crohn's disease: a double-blind randomized controlled trial with two years
  1361 follow-up. Inflamm Bowel Dis. 2012;18:246-253.
- 136284.Li G, Ren J, Wang G, Hu D, Gu G, Liu S, et al. Preoperative exclusive enteral nutrition reduces the<br/>postoperative septic complications of fistulizing Crohn's disease. Eur J Clin Nutr. 2014;68:441-6.
- 136485.Smith MA, Smith T, Trebble T. Nutritional management of adults with inflammatory bowel dis-1365ease: practical lessons from the available evidence. Frontline Gastroenterology 2012;3:172-79.
- 136686.Lochs H, Dejong C, Hammarqvist F, Hebuterne X, Leon-Sanz M, Shulz T, et al. ESPEN Guidelines on1367Enteral Nutrition: Gastroenterology. Clin Nutr 2006; 25, 260-274.
- Fuchssteiner H, Nigl K, Mayer A, Kristensen B, Platzer R, Brunner B, et al. Nutrition and IBD: consensus of the Austrian working group of IBD (inflammatory bowel diseases) of the OGGH. Z Gastroenterol 2014;52:376–386.
- 1371 88. August D, Teitelbaum D, Albina J, Bothe A, Guenter P, Heitkemper M, et al. ASPEN Guidelines for
  1372 the Use of Parenteral and Enteral Nutrition in Adult and Pediatric Patients; JPEN, 2002;26:1SA-138SA.
- 137389.Matsui T, Sakurai T, Yao T. Nutritional therapy for Crohn's disease in Japan.J Gastroenter-1374ol. 2005;40 Suppl 16:25-31.
- 1375 90. Akobeng AK, Thomas AG. Enteral nutrition for maintenance of remission in Crohn's disease.1376 Cochrane Database Syst Rev 2007:CD005984.
- 1377 91. Nakahigashi M, Yamamoto T, Sacco R, Hanai H, Kobayashi F. Enteral nutrition for maintaining
  remission in patients with quiescent Crohn's disease: current status and future perspectives. Int J Colorectal Dis. 2016;31:1-7.
- Yamamoto T, Shiraki M, Nakahigashi M, Umegae S, Matsumoto K. Enteral nutrition to suppress
  postoperative Crohn's disease recurrence: a five-year prospective cohort study. Int J Colorectal
  Dis. 2013;28:335-40.
- 1383 93. Giannotta M, Tapete G, Emmi G, Silvestri E, Milla M. Thrombosis in inflammatory bowel diseases:
  1384 what's the link? Thromb J 2015; 13:14.

- 138594.Zezos P, Kouklakis G, Saibil F. Inflammatory Bowel Disease and thromboembolism. World J Gas-1386troenterol 2014;20:13863-78
- 1387 95. Bhakta A, Tafen M, Ahmed M, Ata A, Abraham C, Bruce D, et al. Risk of catheter-associated deep
  1388 venous thrombosis in inflammatory bowel disease. Dis Colon Rectum. 2014;57:1379-83.

138996.Ha C, Magowan S, Accortt NA, Chen J, Stone CD. Risk of arterial thrombotic events in inflammatory1390bowel disease. Am J Gastroenterol. 2009;104:1445-51.

- 1391 97. Papay P, Miehsler W, Tilg H, Petritsch W, Reinisch W, Mayer A, et al. Clinical presentation of venous thromboembolism in inflammatory bowel disease. J Crohns Colitis. 2013;7:723-9.
- 1393 98. Yan D, Ren J, Wang G, Liu S, Li J. Predictors of response to enteral nutrition in abdominal entero1394 cutaneous fistula patients with Crohn's disease. Eur J Clin Nutr. 2014;68:959-63.
- 1395 99. Visschers RG, Olde Damink SW, Winkens B, Soeters P, van Gemert WG. Treatment strategies in
  1396 135 consecutive patients with enterocutaneous fistulas. World J Surg. 2008;32:445-453.
- 1397 100. Llop JM, Cobo S, Padulles A, Farran L, Jodar R, Badia MB. Nutritional support and risk factors of appearance of enterocutaneous fistulas. Nutr Hosp 2012;27:213-8.
- 1399 101. Dignass A, Van Assche G, Lindsay JO, Lémann M, Söderholm J, Colombel JF, et al. The second Euro pean evidence-based Consensus on the diagnosis and management of Crohn's disease: Current management. J Crohn Colitis 2010;4:28–62.
- 1402 102. Forbes A, Goldesgeyme E, Paulon E. Nutrition in inflammatory bowel disease. J Parent Ent Nutr1403 2011;35:571-80.
- 1404103.Mowat C, Cole A, Windsor A, Ahmad T, Arnott I, Driscoll R, et al. Guidelines for the management of1405inflammatory bowel disease in adults. Gut. 2011;60:571-607.
- 1406104.Uchino M, Ikeuchi H, Matsuoka H, Matsumoto T, Takesue Y, Tomita N. Clinical features and man-<br/>agement of duodenal fistula in patients with Crohn's disease. Hepatogastroenterology. 2012;59:171-4.
- 1408105.Triantafillidis JK, Papalois AE. The role of total parenteral nutrition in inflammatory bowel dis-<br/>ease: current aspects. Scand J Gastroenterol. 2014;49:3-14.
- 1410106.Ravindran P, Ansari N, Young CJ, Solomon MJ. Definitive surgical closure of enterocutaneous fistu-1411la: outcome and factors predictive of increased postoperative morbidity. Colorectal Dis. 2014;16:209-18.
- 1412 107. Akobeng AK, Thomas AG. Refeeding syndrome following exclusive enteral nutritional treatment
   1413 in Crohn disease. J Pediatr Gastroenterol Nutr. 2010 51:364-6.
- 1414 108. Hernando A, Bretón I, Marín-Jimenez I, Menchén L. Refeeding syndrome in a patient with Crohn's
  1415 disease. J Clin Gastroenterol. 2008;42:430-1.
- 1416 109. Krznaric Z, Vranesic Bender D, Ljubas Keleric D, Brinar M. Wernicke's encephalopathy during
   1417 parenteral nutrition in a Crohn's disease patient. Nutrition. 2011 27:503-4
- 1418 110. Dignass A, Lindsay JO, Sturm A, Windsor A, Colombel JF, Allez M, et al. Second European evidencebased consensus on the diagnosis and management of ulcerative colitis Part 2: Current management. J
  1420 Crohn Colitis 2012;6:991–1030
- 1421 111. Salinas H, Dursun A, Konstantinidis I, Nguyen D, Shellito P, Hodin R, et al. Does preoperative total
  parenteral nutrition in patients with ulcerative colitis produce better outcomes? Int J Colorectal Dis.
  2012;27:1479-83.
- 1424112.Schwartz E. Perioperative parenteral nutrition in adults with inflammatory bowel disease: a re-1425view of the literature. Nutr Clin Pract. 2016;31:159-70.
- 1426 113. Weimann A, Braga M, Carli F, Higashiguchi T, Hübner M, Klek S, et al. ESPEN guideline: Clinical
  1427 nutrition in surgery. Clin Nutr 2017;36:623-650
- 1428114.Kuppinger D, Hartl WH, Bertok M, Hoffmann JM, Cederbaum J, Küchenhoff H, et al. Nutritional1429screening for risk prediction in patients scheduled for abdominal operations. Br J Surg 2012;99:728-737.

- 1430 115. Beattie AH, Prach AT, Baxter JP, Pennington CR. A randomised controlled trial evaluating the use
  of enteral nutritional supplements postoperatively in malnourished surgical patients. Gut 2000;46:8131432 818
- 1433 116. MacFie J, Woodcock NP, Palmer MD, Walker A, Townsend S, Mitchell CJ. Oral dietary supplements
  1434 in pre- and postoperative surgical patients: a prospective and randomized clinical trial. Nutrition
  1435 2000;16:723-728.
- 1436
  117. Espaulella J, Guyer H, Diaz-Escriu F, Mellado-Navas JA, Castells M, Pladevall M. Nutritional supplementation of elderly hip fracture patients. A randomized, double-blind, placebo-controlled trial. Age
  Ageing 2000; 29:425-431.
- 1439 118. Smedley F, Bowling T, James M, Stokes E, Goodger C, O'Connor O, et al. Randomized clinical trial of
  the effects of preoperative and postoperative oral nutritional supplements on clinical course and cost of
  care. Br J Surg 2004;91:983-990.
- 1442119.Burden S, Todd C, Hill J, Lal S. Pre-operative nutrition support in patients undergoing gastrointes-1443tinal surgery. Cochrane Database Syst Rev 2012; 11:CD008879.
- 1444 120. Braga M, Gianotti L, Gentilini O, Liotta S, Di Carlo V. Feeding the gut early after digestive surgery:
  1445 results of a nine-year experience. Clin Nutr 2002;21:59-65.
- 1446121.Daly JM, Bonau R, Stofberg P, Bloch A, Jeevanandam M, Morse M. Immediate postoperative jeju-<br/>nostomy feeding. Clinical and metabolic results in a prospective trial. Am J Surg 1987;153:198-206.
- 1448122. Delany HM, Carnevale N, Garvey JW, Moss GM. Postoperative nutritional support using needlecatheter feeding jejunostomy. Ann Surg 1977;186:165-170.
- 1450 123. Gabor S, Renner H, Matzi V, Ratzenhofer B, Lindenmann J, Sankin O, et al. Early enteral feeding
  1451 compared with parenteral nutrition after oesophageal or oesophagogastric resection and reconstruction.
  1452 Br J Nutr 2005;93:509-513
- 1453 124. Gupta V. Benefits versus risks: a prospective audit. Feeding jejunostomy during esophagectomy.
  1454 World J Surg 2009;33:1432-1438
- 1455 125. Kemen M, Senkal M, Homann HH, Mumme A, Dauphin AK, Baier J, et al. Early postoperative enter1456 al nutrition with arginine-omega-3 fatty acids and ribonucleic acid-supplemented diet versus placebo in
  1457 cancer patients: an immunologic evaluation of Impact. Crit Care Med 1995; 23:652-659.
- 1458
  126. Klein S, Kinney J, Jeejeebhoy K, Alpers D, Hellerstein M, Murray M, et al. Nutrition support in clinical practice: review of published data and recommendations for future research directions. Summary of a
  conference sponsored by the National Institutes of Health, American Society for Parenteral and Enteral
  Nutrition, and American Society for Clinical Nutrition. Am J Clin Nutr 1997; 66:683-706.
- 1462127.Veterans Affairs Total Parenteral Nutrition Cooperative Study Group. Perioperative total paren-1463teral nutrition in surgical patients. N Engl J Med 1991;325:525-532.
- 1464 128. Bozzetti F, Gavazzi C, Miceli R, Rossi N, Mariani L, Cozzaglio L, et al. Perioperative total parenteral
  1465 nutrition in malnourished, gastrointestinal cancer patients: a randomized, clinical trial. JPEN J Parenter
  1466 Enteral Nutr 2000;24:7-14.
- 1467 129. Shukla HS, Rao RR, Banu N, Gupta RM, Yadav RC. Enteral hyperalimentation in malnourishedsurgical patients. Indian J Med Res 1984;80:339-346.
- 1469 130. Von Meyenfeldt MF, Meijerink WJ, Rouflart MM, Builmaassen MT, Soeters PB. Perioperative nutri 1470 tional support: a randomised clinical trial. Clin Nutr 1992;11:180-186.
- 1471 131. Heyland DK, Montalvo M, MacDonald S, Keefe L, Su XY, Drover JW. Total parenteral nutrition in
  1472 the surgical patient: a meta-analysis. Can J Surg 2001;44:102-111.
- 1473 132. Andersen HK, Lewis SJ, Thomas S. Early enteral nutrition within 24h of colorectal surgery versus
  1474 later commencement of feeding for postoperative complications. Cochrane Database Syst Rev
  1475 2006 :CD004080.

1476 133. Lewis SJ, Andersen HK, Thomas S. Early enteral nutrition within 24 h of intestinal surgery versus
1477 later commencement of feeding: a systematic review and meta-analysis. J Gastrointest Surg 2009; 13:5691478 575

1479 134. Mazaki T, Ebisawa K. Enteral versus parenteral nutrition after gastrointestinal surgery: a system1480 atic review and meta-analysis of randomized controlled trials in the English literature. J Gastrointest Surg
1481 2008; 12:739-755

1482 135. Osland E, Yunus RM, Khan S, Memon MA. Early versus traditional postoperative feeding in pa1483 tients undergoing resectional gastrointestinal surgery: a meta-analysis. JPEN J Parenter Enteral Nutr
1484 2011;35:473-487.

1485136.Cohen AB, Lee D, Long MD, Kappelman MD, Martin CF, Sandler RS, et al. Dietary patterns and self-1486reported associations of diet with symptoms of inflammatory bowel disease. Dig Dis Sci 2013;58:1322-8.

1487137.Zvirbliene A, Kiudelis G, Zalinkevicius R, Kupcinskas L. [Dietary characteristics of patients with1488inflammatory bowel diseases]. Medicina (Kaunas) 2006;42:895-9.

1489 138. Banos Madrid R, Salama Benerroch H, Moran Sanchez S, Gallardo Sanchez F, Albadalejo Merono A,
et al. Lactose malabsorption in patients with inflammatory bowel disease without activity: would it be
necessary to exclude lactose products in the diet of all patients? Anales de Medicina Interna 2004;21:21214.

1493 139. Triggs CM, Munday K, Hu R, Fraser AG, Gearry RB, Barclay ML, et al. Dietary factors in chronic
inflammation: food tolerances and intolerances of a New Zealand Caucasian Crohn's disease population.
Mutat Res 2010;690:123-38.

1496 140. Jones VA, Dickinson RJ, Workman E, Wilson AJ, Freeman AH, Hunter JO. Crohn's disease: mainte-1497 nance of remission by diet. Lancet. 1985;2:177-80.

1498141.Turner D, Zlotkin SH, Shah PS, Griffiths AM. Omega 3 fatty acids (fish oil) for maintenance of re-<br/>mission in Crohn's disease. Cochrane Database Syst Rev 2009:CD006320.

1500 142. Cashman KD, Shanahan F. Is nutrition an aetiological factor for inflammatory bowel disease? Eur J
 1501 Gastroenterol Hepatol 2003;15:607-13.

1502143.Tanaka T, Takahama K, Kimura T, Mizuno T, Nagasaka M, Iwata K, et al. Effect of concurrent ele-1503mental diet on infliximab treatment for Crohn's disease. J Gastroenterol Hepatol 2006;21:1143-9.

1504144.Yamamoto T, Nakahigashi M, Umegae S, Matsumoto K. Prospective clinical trial: enteral nutrition1505during maintenance infliximab in Crohn's disease. J Gastroenterol 2010;45:24-9.

1506145.Maconi G, Ardizzone S, Cucino C, Bezzio C, Russo AG, Bianchi Porro G. Pre-illness changes in die-<br/>tary habits and diet as a risk factor for inflammatory bowel disease: a case-control study. World J Gastro-<br/>enterol 2010;16:4297-304.

1509 146. Esaki M, Matsumoto T, Hizawa K, Nakamura S, Jo Y, Mibu R, et al. Preventive effect of nutritional
therapy against postoperative recurrence of Crohn disease, with reference to findings determined by intra-operative enteroscopy. Scand J Gastroenterol 2005; 40:1431-7.

1512 147. Richman E, Rhodes JM. Review article: evidence-based dietary advice for patients with inflamma-1513 tory bowel disease. Aliment Pharmacol Ther. 2013;38:1156-71.

1514148.Cabré E, Mañosa M, Gassull MA. Omega-3 fatty acids and inflammatory bowel diseases - a system-1515atic review. Br J Nutr. 2012;107 Suppl 2:S240-52.

1516 149. Lev-Tzion R, Griffiths AM, Leder O, Turner D. Omega 3 fatty acids (fish oil) for maintenance of remission in Crohn's disease. Cochrane Database Syst Rev. 2014;2:CD006320.

1518150.Hallert C, Kaldma M, Petersson BG. Ispaghula husk may relieve gastrointestinal symptoms in ul-<br/>cerative colitis in remission. Scand J Gastroenterol. 1991;26:747-50.

1520 151. Fernández-Bañares F, Hinojosa J, Sánchez-Lombraña JL, Navarro E, Martínez-Salmerón JF, García1521 Pugés A, et al. Randomized clinical trial of Plantago ovata seeds (dietary fiber) as compared with mesala1522 mine in maintaining remission in ulcerative colitis. Spanish Group for the Study of Crohn's Disease and
1523 Ulcerative Colitis (GETECCU). Am J Gastroenterol. 1999;94:427-33.

1524 152. Hanai H, Kanauchi O, Mitsuyama K, Andoh A, Takeuchi K, Takayuki I, et al. Germinated barley 1525 foodstuff prolongs remission in patients with ulcerative colitis. Int J Mol Med. 2004;13:643-7.

1526 153. Brotherton CS, Taylor AG, Bourguignon C, Anderson JG. A high-fiber diet may improve bowel
1527 function and health-related quality of life in patients with Crohn disease. Gastroenterol Nurs.
1528 2014;37:206-16.

1529 154. Chiba M, Tsuji T, Nakane K, Komatsu M. High amount of dietary fiber not harmful but favorable1530 for Crohn disease. Perm J. 2015;19:58-61.

1531 155. Fujiya M, Ueno N, Kohgo Y. Probiotic treatments for induction and maintenance of remission in
 inflammatory bowel diseases: a meta-analysis of randomized controlled trials. Clin J Gastroenterol
 2014;7:1-13.

1534 156. Kruis W, Fric P, Pokrotnieks J, Lukas M, Fixa B, Kascak M, et al. Maintaining remission of ulcerative colitis with the probiotic Escherichia coli Nissle 1917 is as effective as with standard mesalazine. Gut
2004;53:1617-23.

1537 157. Floch MH, Walker WA, Sanders ME, Nieuwdorp M, Kim AS, Brenner DA, et al. Recommendations
1538 for Probiotic Use--2015 Update: Proceedings and Consensus Opinion. J Clin Gastroenterol 2015;49 Suppl
1539 1:S69-73.

1540 158. Ishikawa H, Matsumoto S, Ohashi Y, Imaoka A, Setoyama H, Umesaki Y, et al. Beneficial effects of
probiotic bifidobacterium and galacto-oligosaccharide in patients with ulcerative colitis: a randomized
controlled study. Digestion 2011;84:128-33.

1543
159. Yoshimatsu Y, Yamada A, Furukawa R, Sono K, Osamura A, Nakamura K, et al. Effectiveness of
probiotic therapy for the prevention of relapse in patients with inactive ulcerative colitis. World J Gastroenterol 2015;21:5985-94.

1546 160. Meini S, Laureano R, Fani L, Tascini C, Galano A, Antonelli A, et al. Breakthrough Lactobacillus
rhamnosus GG bacteremia associated with probiotic use in an adult patient with severe active ulcerative
colitis: case report and review of the literature. Infection 2015;43:777-81.

1549 161. Vahabnezhad E, Mochon AB, Wozniak LJ, Ziring DA. Lactobacillus bacteremia associated with probiotic use in a pediatric patient with ulcerative colitis. J Clin Gastroenterol 2013;47:437-9.

1551 162. Prantera C, Scribano ML, Falasco G, Andreoli A, Luzi C. Ineffectiveness of probiotics in preventing
recurrence after curative resection for Crohn's disease: a randomised controlled trial with Lactobacillus
GG. Gut 2002;51:405-9.

- 1554 163. Schultz M, Sartor RB. Probiotics and inflammatory bowel diseases. Am J Gastroenterol 1555 2000;95:S19-21.
- 1556 164. Guslandi M, Giollo P, Testoni PA. A pilot trial of Saccharomyces boulardii in ulcerative colitis. Eur J
   1557 Gastroenterol Hepatol 2003;15:697-8.
- 1558 165. Rolfe VE, Fortun PJ, Hawkey CJ, Bath-Hextall F. Probiotics for maintenance of remission in Crohn's
   1559 disease. Cochrane Database Syst Rev 2006:CD004826.

1560 166. Campieri M, Rizzello F, Venturi A, Poggioli G, Ugolini F, Helwig U. Combination of antibiotic and
 probiotic treatment is efficacious in prophylaxis of post operative recurrence of Crohn's disease: a ran domized controlled study vs mesalazine. Gastroenterology 2000;118:A4179.

1563 167. Garcia Vilela E, De Lourdes De Abreu Ferrari M, Oswaldo Da Gama Torres H, Guerra Pinto A, et al.
1564 Influence of Saccharomyces boulardii on the intestinal permeability of patients with Crohn's disease in
1565 remission. Scand J Gastroenterol 2008;43:842-8.

1566 168. Singh S, Stroud AM, Holubar SD, Sandborn WJ, Pardi DS. Treatment and prevention of pouchitis
1567 after ileal pouch-anal anastomosis for chronic ulcerative colitis. Cochrane Database Syst Rev.
1568 2015;11:CD001176.

1569 169. Mimura T, Rizzello F, Helwig U, Poggioli G, Schreiber S, Talbot IC, et al. Once daily high dose probiotic therapy (VSL#3) for maintaining remission in recurrent or refractory pouchitis. Gut 2004;53:108–
1571 14.

1572 170. Gionchetti P, Rizzello F, Venturi A, Brigidi P, Matteuzzi D, Bazzocchi G, et al. Oral bacteriotherapy
1573 as maintenance treatment in patients with chronic pouchitis: a double-blind, placebo-controlled trial.
1574 Gastroenterology 2000;119:305-9.

1575 171. Tursi A, Brandimarte G, Papa A, Giglio A, Elisei W, Giorgetti GM, et al. Treatment of relapsing mildto-moderate ulcerative colitis with the probiotic VSL#3 as adjunctive to a standard pharmaceutical treatment: a double-blind, randomized, placebo-controlled study. Am J Gastroenterol 2010;105:2218-27.

1578 172. Gionchetti P, Rizzello F, Helwig U, Venturi A, Lammers KM, Brigidi P, et al. Prophylaxis of pouchi1579 tis onset with probiotic therapy: a double-blind, placebo-controlled trial. Gastroenterology.
1580 2003;124:1202-9.

1581 173. Kuisma J, Mentula S, Kahri A, Kahri A, Saxelin M, Farkkila M. Effect of Lactobacillus rhamnosus GG1582 on ileal pouch inflammation and microbial flora. Aliment Pharmacol Ther 2003;17:509-515.

1583 174. Biancone L, Michetti P, Travis S, Escher JC, Moser G, Forbes A, et al. European evidence-based
1584 Consensus in the management of ulcerative colitis: special situations. J Crohns Colitis 2008;2:63-92.

1585 175. Turner D, Levine A, Escher JC, Griffiths AM, Russell RK, Dignass A, et al. European Crohn's and
1586 Colitis Organization; European Society for Paediatric Gastroenterology, Hepatology, and Nutrition. Man1587 agement of pediatric ulcerative colitis: joint ECCO and ESPGHAN evidence-based consensus guidelines. J
1588 Pediatr Gastroenterol Nutr. 2012;55:340-61.

- 1589176.El-Matary W, Otley A, Critch J, Abou-Setta AM. Enteral Feeding Therapy for Maintaining Remis-1590sion in Crohn's Disease: A Systematic Review. JPEN J Parenter Enteral Nutr. 2015;41:550-561.
- 1591 177. Takagi S, Utsunomiya K, Kuriyama S, Yokoyama H, Takahashi S, Iwabuchi M, et al. Effectiveness of
  an 'half elemental diet' as maintenance therapy for Crohn's disease: A random-ized-controlled trial. Aliment Pharmacol Ther 2006; 24: 1333-40
- 1594 178. Hirai F, Ishihara H, Yada S, Esaki M, Ohwan T, Nozaki R, et al. Effectiveness of concomitant enteral
  nutrition therapy and infliximab for maintenance treatment of Crohn's disease in adults. Dig Dis Sci.
  2013;58:1329-34.
- 1597 179. Sazuka S, Katsuno T, Nakagawa T, Saito M, Saito K, Matsumura T, et al. Concomitant use of enteral
  nutrition therapy is associated with sustained response to infliximab in patients with Crohn's disease. Eur
  J Clin Nutr. 2012 Nov;66:1219-23.
- 1600 180. Tsertsvadze A, Gurung T, Court R, Clarke A, Sutcliffe P. Clinical effectiveness and costeffectiveness of elemental nutrition for the maintenance of remission in Crohn's disease: a systematic
  review and meta-analysis. Health Technol Assess. 2015;19:1-138.
- 1603 181. Ruemmele FM, Veres G, Kolho KL, Griffiths A, Levine A, Escher JC, et al. Consensus guidelines of
   1604 ECCO/ESPGHAN on the medical management of pediatric Crohn's disease. J Crohns Colitis. 2014;8:1179 1605 207.
- 1606 182. Battat R, Kopylov U, Szilagyi A, Saxena A, Rosenblatt DS, Warner M, et al. Vitamin B12 deficiency
  1607 in inflammatory bowel disease: prevalence, risk factors, evaluation, and management. Inflamm Bowel Dis.
  2014;20:1120-8.
- 1609 183. Duerksen DR, Fallows G, Bernstein CN. Vitamin B12 malabsorption in patients with limited ileal
   1610 resection. Nutrition. 2006;22:1210-3.
- 1611 184. Headstrom PD, Rulyak SJ, Lee SD. Prevalence of and risk factors for vitamin B(12) deficiency in patients with Crohn's disease. Inflamm Bowel Dis. 2008;14:217-23.
- 1613 185. Yakut M, Ustün Y, Kabaçam G, Soykan I. Serum vitamin B12 and folate status in patients with in1614 flammatory bowel diseases. Eur J Intern Med. 2010;21:320-3.
- 1615 186. Stabler SP. Clinical practice. Vitamin B12 deficiency. N Engl J Med. 2013;368:149–160.
- 1616
  187. Plener I, Ferguson C, Kashkooli S, Saibil F. Oral B12 replacement in Crohn's disease is B12 by
  1617 injection obsolete? Aliment Pharmacol Ther. 2014;40:1365-6.

- 1618 188. Hornung N, Ellingsen T, Stengaard-Pedersen K, Poulsen JH. Folate, homocysteine, and cobalamin
  1619 status in patients with rheumatoid arthritis treated with methotrexate, and the effect of low dose folic acid
  1620 supplement. J Rheumatol 2004;31:2374–81.
- 1621 189. Halsted CH, Gandhi G, Tamura R. Sulphasalazine inhibits the absorption of folates in ulcerative colitis. N Engl J Med 1981;305:1513–7.
- 1623 190. Burr NE, Hull MA, Subramanian V. Folic Acid Supplementation May Reduce Colorectal Cancer Risk
  1624 in Patients With Inflammatory Bowel Disease: A Systematic Review and Meta-Analysis. J Clin Gastroenter1625 ol. 2017;51:247-253.
- 1626 191. Pironi L, Cornia GL, Ursitti MA, Dallasta MA, Miniero R, Fasano F, et al. Evaluation of oral admin1627 istration of folic and folinic acid to prevent folate deficiency in patients with inflammatory bowel disease
  1628 treated with salicylazosulfapyridine. Int J Clin Pharmacol Res. 1988;8:143-8.
- 1629 192. Honein MA, Paulozzi LJ, Mathews TJ, Erickson JD, Wong LY. Impact of folic acid fortification of the
  1630 US food supply on the occurrence of neural tube defects. JAMA. 2001;285:2981–6.
- 1631 193. Bryant RV, Trott MJ, Bartholomeusz FD, Andrews JM. Systematic review: body composition in adults with inflammatory bowel disease. Aliment Pharmacol Ther. 2013;38:213-25.
- 1633 194. Bryant RV, Ooi S, Schultz CG, Goess C, Grafton R, Hughes J, et al. Low muscle mass and sarcopenia:
  1634 common and predictive of osteopenia in inflammatory bowel disease. Aliment Pharmacol Ther.
  1635 2015;41:895-906.
- 1636
  195. Wiroth JB, Filippi J, Schneider SM, Al-Jaouni R, Horvais N, Gavarry O, et al. Muscle performance in patients with Crohn's disease in clinical remission. Inflamm Bowel Dis. 2005;11:296-303.
- 1638 196. Werkstetter KJ, Ullrich J, Schatz SB, Prell C, Koletzko B, Koletzko S. Lean body mass, physical activity and quality of life in paediatric patients with inflammatory bowel disease and in healthy controls. J
  1640 Crohns Colitis. 2012;6:665-73.
- 1641 197. Klare P, Nigg J, Nold J, Haller B, Krug AB, Mair S, et al. The impact of a ten-week physical exercise
  program on health-related quality of life in patients with inflammatory bowel disease: a prospective randomized controlled trial. Digestion. 2015;91:239-47.
- 1644 198. Cruz-Jentoft AJ, Landi F, Schneider SM, Zúñiga C, Arai H, Boirie Y et al. Prevalence of and interven1645 tions for sarcopenia in ageing adults: a systematic review. Report of the International Sarcopenia Initiative
  1646 (EWGSOP and IWGS). Age Ageing. 2014;43:748-59.
- 1647 199. Flores A, Burstein E, Cipher DJ, Feagins LA. Obesity in Inflammatory Bowel Disease: A Marker of
  1648 Less Severe Disease. Dig Dis Sci. 2015;60:2436-45.
- 1649
  200. Nic Suibhne T, Raftery TC, McMahon O, Walsh C, O'Morain C, O'Sullivan M. High prevalence of
  overweight and obesity in adults with Crohn's disease: associations with disease and lifestyle factors. J
  1651
  Crohns Colitis. 2013;7:e241-8.
- 201. Seminerio JL, Koutroubakis IE, Ramos-Rivers C, Hashash JG, Dudekula A, Regueiro M, et al. Impact
  of Obesity on the Management and Clinical Course of Patients with Inflammatory Bowel Disease. Inflamm
  Bowel Dis. 2015;21:2857-63.
- 1655









**General advices** 

R8: There is no "IBD diet" that can be generally recommended to promote remission in IBD patients with active disease.

R11: In IBD patients (adults and children) with active disease and those who are steroid-treated, serum calcium and 25(OH) vitamin D should be monitored and supplemented if required to help prevent low bone mineral density. Osteopenia and osteoporosis should be managed according to current osteoporosis guidelines.

R13: Exclusion diets cannot be recommended to achieve remission in active CD, even if the patient suffers from individual intolerances.

**Probiotics** 

R14A: Probiotic therapy using Lactobacillus reuteri or "VSL#3", but not necessarily other probiotics, can be considered for use in patients with mild to moderate UC for the induction of remission.

R14B: Probiotics should not be used for treatment of active CD.



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#### Active disease - Surgical aspects of nutrition - General **Elective surgery?** R23A: In most elective surgery cases, pre-operative fasting R23B: In emergency surgery patients medical nutrition (EN, from midnight should not be performed - instead, an PN) should be initiated if the patient is malnourished at the Pre-OP enhanced recovery (ERAS) protocol can be used. time of surgery or if oral diet cannot be recommenced within 7 days after surgery. . Nutritional deficiency? R24B: Patients who do not meet R24A: Patients who do not meet their R24C: If malnutrition is diagnosed, then energy and/or protein needs from normal IBD surgery should be delayed for 7-14 their energy and/or protein needs food should be encouraged to take oral from normal food plus ONS days whenever possible, and that time Peri-OP should be used for intensive medical nutritional supplements (ONS) during the should receive EN during the perioperative period. perioperative period. nutrition. R27B: In the early phase after R27A: Normal food intake or EN can be proctocolectomy or colectomy, water and commenced early after surgery in most Colectomy Post-OP electrolytes shall be administered to IBD patients in the postoperative phase. assure haemodynamic stability.

Figure 8



#### Remission - General

#### Recommended

R28: All IBD patients in remission should undergo counselling by a dietician as part of the multidisciplinary approach to improve nutritional therapy and to avoid malnutrition and nutrition-related disorders.

R29: No specific diet needs to be followed during remission phases of IBD.

R35: Standard diet or ONS should be followed in patients with IBD in remission, giving attention to nutrition screening and generic nutritional support where needed.

R34B: ONS or EN can be recommended in patients with CD in remission, if undernutrition cannot be treated sufficiently by dietary counselling.

R39: In all IBD patients, endurance training should be encouraged. In IBD patients with decreased muscle mass and/or muscle performance, appropriate physical activity should be recommended.

R40: Obese IBD patients should be advised to reduce weight only in phases of stable remission and then according to current obesity guidelines.

#### Not recommended

R30: Supplementation with omega-3 fatty acids should not be advised to support maintenance of remission in patients with IBD.

R31: Non-specific high fiber diets should not normally be recommended for maintenance of remission in IBD

R34A: Neither EN nor PN is recommended as primary therapy for maintaining remission in IBD.

