

Enteral feeding reduces metabolic activity of the intestinal microbiome in Crohn's disease: Observational study

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30 **Abstract**

31 Background

32

33 Enteral feeding will induce remission in as many as 80-90% of compliant patients with active Crohn's
34 Disease (CD) but its method of action remains uncertain. This study was designed to examine its effects
35 on the colonic microbiome.

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37 Method

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39 Healthy volunteers and patients with CD followed a regimen confined to enteral feeds alone for one or
40 two weeks respectively. Chemicals excreted on breath or in faeces were characterised at the start and at
41 the end of the feeding period by gas chromatography mass spectrometry (GC/MS).

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43 Results

44 One week of feeding in healthy volunteers caused significant changes in stool colour and deterioration in
45 breath odour, together with increased excretion of phenol and indoles on the breath. Feeding for two
46 weeks in patients with CD produced significant improvements in symptoms and a decrease in the
47 concentration of C-reactive protein. The faecal concentrations of microbial products including short chain
48 fatty acids (SCFAs), and potentially toxic substances including 1-propanol, 1-butanol and the methyl and
49 ethyl esters of SCFAs showed significant falls.

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51 Conclusion

52 A significant change occurs in the production of microbial metabolites after enteral feeding in both
53 healthy volunteers and patients with CD. Many of those detected in CD are toxic and may feasibly lead

54 to the immunological attack on the gut microbiota, which is characteristic of IBD. The reduction in the
55 production of such metabolites after enteral feeding may be the reason for its effectiveness in CD.
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80 **Introduction**

81 Despite the increasing frequency of Crohn's disease, its treatment remains unsatisfactory. Many of the
82 therapeutic agents used have unpleasant or even dangerous side effects and some are very expensive. The
83 continuing perception of CD as a relapsing and remitting disorder emphasises the difficulty in
84 maintaining long term control. A complete cure remains elusive.

85
86 Reports of a positive response to dietary manipulation in CD have emerged from several sources.¹⁻⁸ 2-4
87 weeks of total enteral feeding has been reported to reduce remission in 85-90% of compliant patients
88 suffering active CD.¹⁻⁶ Lack of understanding of the method of action of enteral feeds in CD has
89 however, discouraged their use.²⁻⁸

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91 Enteral feeds are nutritionally complete liquid mixtures of pre-digested foods presenting nitrogen as
92 amino acids, oligopeptides or a single protein, carbohydrates as simple sugars, typically malto-dextrins,
93 and fat as a single oil, (eg. Rapeseed oil), together with minerals and vitamins. They are nutritionally
94 complete.

95
96 Suggestions as to the method of action of enteral feeding are many, but it is now known that bowel rest⁷
97 and the reduction of potential food allergens²⁻⁴ are incorrect. Enteral feeding is unlikely to have
98 therapeutic benefit by producing immunosuppression as it is ineffective in the treatment of ulcerative
99 colitis.³ Reduction in inflammation can be detected before any improvement in nutritional state begins,⁸
100 and the suggestion that dietary particles might be important was not supported by a controlled trial.⁹

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102 The increasing evidence that inflammation in CD is provoked by an immune response targeted against the
103 intestinal microbiome implies that manipulation of the metabolic activity of the microbiota might have a
104 role in the treatment of this disease^{10,12} We and others have recently demonstrated an association between
105 Crohn's disease and the profile volatile organic compounds obtained from breath and faecal headspace

samples^{16,34}. These measurements are a useful indication of changes in the gut microbiome, being simple, rapid and non-invasive. We have used this approach here in a study of the effects of enteral feeding. It has been suggested that food intolerance, as distinct from food allergy, might reflect an interaction between unabsorbed food residues and the intestinal microbiome.¹³ As the nutrients contained in enteral feeds are absorbed high in the small intestine, they supply little in the way of energy substrates to micro-organisms in the lower bowel. This might lead to changes in microbial metabolism which in turn could lead to a reduction in inflammation. The present studies were designed to investigate this possibility.

Methods

Study 1 Healthy volunteers

Volunteers were recruited from students of either sex aged 18-65 years, at Cranfield University who were in good health and eating a normal diet. A total of 12 subjects was recruited aged 23-32, of which 8 were female. Subjects suffering conditions possibly requiring specific diets e.g. irritable bowel syndrome (IBS), or coeliac disease were excluded. Other exclusions were pregnancy or lactation, a course of antibiotics in the previous six weeks, bacterial products such as pro- or pre- biotics and any chronic medication other than oral contraceptives.

Subjects were randomly allocated to take either E028 extra (Nutricia UK Liverpool), or Modulen-IBD (Nestle Ltd, Croydon UK), for 7 days with all other foodstuffs excluded except water *ad libitum*.. Nutritional requirements were calculated for each individual using Schofield's equation.¹⁴

After 7 days subjects returned to normal diets for 21 days before commencing the alternative enteral feed for a further 7 days. The two feeds were administered 4 weeks apart in order that they were taken at the same stage of the menstrual cycles of female volunteers. During enteral feeding, subjects were asked to record how much feed they consumed and to complete symptom score sheets recording on a daily basis stool frequency, consistency and colour and any changes in breath odour. Weights were recorded and

131 breath samples taken before the study and after each week of enteral feeding. This trial was an open,
132 randomised controlled study performed at Cranfield University and approved by the ethics committee of
133 Cranfield University and the NHS Cambridge local research ethics committee.

134 Volunteers were provided with a sheet depicting a range of faecal colours ranging from dark brown to
135 bright green (copies supplied to the editor) and asked to assess the stool colour, consistency and
136 frequency. They were asked to record daily changes in breath odour which was assessed subjectively on
137 a scale from 1 (odourless) to 4 (extremely unpleasant).

138 Bio-VOC samplers were used according to manufacturer's recommendations to obtain a one-litre end-
139 tidal breath sample after breakfast on the first day of each feeding period. Samples were injected onto
140 Thermal desorption tubes containing 1:1 Tenax TA and Carbotrap adsorbents (Markes International,
141 Llantrisant, UK).

142 **Study 2 Patients with Crohn's disease**

143 Patients aged 18-65 years were recruited in the department of Gastroenterology, Addenbrooke's Hospital,
144 Cambridge. A total of 17 patients each provided a faecal sample before treatment with enteral feed
145 E028extra and again when they went into remission. At recruitment, all had symptoms of active disease.
146 The diagnosis of CD was made by standard diagnostic criteria and the severity of symptoms was assessed
147 using the Harvey and Bradshaw Index¹⁵. The concentration of C-reactive protein (CRP) in serum
148 samples obtained at each visit was determined by the Biochemistry Department of Addenbrooke's
149 Hospital to provide an objective measure of disease activity.

150
151 Any patients who had received antibiotics in the previous 6 weeks were excluded. Some were taking
152 medication including 5-aminosalicylic acid compounds and/or azathioprine which had been insufficient
153 to control their symptoms, but none had received previous dietary treatment. They were asked to continue
154 such medication during the period of feeding with elemental diet. Non-fasting morning samples of faeces
155 were obtained before starting two weeks treatment with E028 extra (Nutricia Liverpool UK) with

amounts again being calculated by Schofield's equation. A further faecal sample was obtained at the end of this period. Samples were delivered to the hospital on the same day as passed with a maximum delay before freezing of 4 hours. They were stored at -40°C until transferred to the laboratory for analysis.

Ethical permission for this study was granted by the Leeds West LREC (Ref: 07/Q1205/39).

Laboratory analysis

An internal standard solution comprising 50 ng deuterated (D8) toluene (Supelco Cat no 48,593) in methanol was added to each tube according to the manufacturer's instructions (Markes International Ltd, Llantrisant, UK). Head space samples were analysed by automated thermal desorption gas chromatography/mass spectrometry. A Perkin Elmer system was used for analysis combining a TurboMass MS 4.1 Autosystem XL GC and Automatic Thermal Desorption system (ATD 400 PerkinElmer, Wellesley MA). The gas carrier was CP-grade helium (BOC gases Guildford UK) passed through a combined trap for removal of hydrocarbons, oxygen and water vapour. A wall-coated Zebron ZB624 chromatographic column was used with dimensions 60 x .04 x 0.25mm (internal diameter), the liquid phase comprising a 0.25 µm layer of 6% cyanopropylphenyl and 94% methylpolysiloxane.

Thermal desorption tubes were initially purged for 2 minutes to remove air and water vapour and then desorbed for 5 minutes at 300°C. The automatic thermal desorption valve temperature was set at 180°C and TD tubes were desorbed onto the secondary cold trap, which was initially maintained at 30°C. Once desorption was complete, the secondary trap was heated to 320°C using the fastest available heating rate and then maintained for 5 minutes. The effluent was transferred to the gas chromatograph through a transfer line heated to 210°C. The gas chromatograph oven was maintained at 50°C for 4 minutes after injection and then raised at a rate of 10°C/min to 220°C and then held for 9 minutes. Eluted products were transferred to the mass spectrometer via a line heated to 240°C. Electron ionisation (70eV) was used. Full scan mode was selected with mass-to-charge ratios from 33 to 350 m/z with a scan time of 0.3 second and 0.1 second interscan delay to produce a total ion count (TIC) chromatogram.

Study 2

Samples were transferred to the laboratory packed in dry ice inside insulated containers and on arrival were stored at -80°C until analysis.

Aliquots (5ml) of the defrosted samples were placed in gas sampling bags which were then sealed and filled with hydrocarbon-free air and incubated for 10 minutes at body temperature. A portable air pump was then used to draw 500ml of headspace through TD tubes packed with 50% Carbotrap and 50% Tenax. Full details have been published elsewhere ¹⁶

Data and statistical analysis

Study 1

Compound identification was achieved using Automated Mass Spectral Deconvolution and Identification (AMDIS version 2.62) software and the National Institute of Standards and Technology mass spectral library. Quantification was achieved by comparing the area of each compound peak with the peak area associated with the known amount of d8 toluene.

Concentration data proved to be heavily right-skewed, therefore a non-parametric approach was adopted. A McNemar test was used to determine whether the probability of a compound to be present before or after the diet was significant. When present a Wilcoxon Rank Test was used to see if the compound was present in different quantities. Raw TIC data (i.e. a matrix of time vs. ion abundance) were also subjected to Principal Components Analysis (PCA)²⁷ using Matlab (version 6/5 Mathworks Inc USA incorporating functions from the PLS Toolbox version 2.0 Eigenvector Research Inc USA).

Study 2

Compound identification and quantification were carried out as for study 1. In any given faecal headspace sample, automated mass spectral deconvolution and identification (AMDIS) would identify between 100-300 different compounds and it was therefore found necessary to select a subset of those we observed to render statistical analysis tractable. Three approaches were followed to provide a list of what we have termed 'candidate compounds'. The list comprised first compounds that appeared to be most abundant

from inspection of the results obtained using AMDIS; second compounds that appeared to discriminate between patient groups by visual inspection of a subset of pre treatment sample chromatograms and third compound selected on the basis of a search of the relevant literature. An initial generic list was made including short-chain fatty acids (SCFAs) and their derivatives, phenolic compounds and indoles and sulfides. This list was then refined according to publications dealing more explicitly with VOC profiles in disease. A final list of compounds was obtained in this way.

Results

Study 1

Of the 12 volunteers recruited, two females withdrew before the feeding commenced. During the first feeding period 2 withdrew after 2 days feeding, one (female having E028) because of persistent hunger and the other (male having Modulen-IBD) because of insomnia attributed to an empty stomach. Eight subjects completed the first phase. A further subject (male Modulen-IBD) withdrew after 4 days in the second phase because of malaise and headaches.

Stool consistency and frequency showed no change. There was a consistent change in stool colour from browns towards green on E028 extra ($r=0.639$, $p<0.05$ Spearman test), and a similar but less marked effect was seen after Modulen-IBD ($r=0.598$, $p<0.05$). Faecal colour had returned to normal by the start of the second feeding period.

All subjects showed deterioration in odour on E028 extra and 5 out of 6 on Modulen-IBD. One volunteer did not record his breath changes on a daily basis. A Spearman test showed a significant difference between the odour of the breaths of the volunteers before they started and the last day of the diet (E028 extra $r=0.575$ $p<0.05$, Modulen-IBD $r=0.574$ $p<0.05$). Subjects breath odour had returned to normal at the

start of the second feeding period. Numerical results are presented as mean with upper and lower quartiles. The frequency distributions for all compounds were found to be highly skewed with a proportion of nondetects; therefore, a nonparametric statistical approach was adopted.

Over 140 compounds were seen in the breath analysis including aldehydes, ketones, saturated and non-saturated hydrocarbons, organic acids, alkenes, alcohols and furans. The compounds also varied between volunteers. As at least one third of compounds were known to be environmental contaminants, e.g. benzene, toluene, xylene, we concentrated on two marker compounds known to be bacterial metabolites, phenol and indole.

The mean alveolar gradient for indole on a normal diet was $0.034 \pm \text{SD } 0.029$. There is little change following Modulen-IBD $0.041 \pm \text{SD } 0.028$ (NS). After E028 it rose to $0.149 \pm \text{SD } 0.099$ (NS). The differences between the values after diet did not differ significantly from those before, but the alveolar gradient after E028 was significantly higher than that after Modulen-IBD ($P < 0.03$).

The mean level of alveolar gradient for phenol on the breath on a normal diet was $0.024 \pm \text{SD } 0.017$. After Modulen-IBD it rose to $0.055 \pm \text{SD } 0.025$ (NS). After E028 the levels were $0.229 \pm \text{SD } 0.152$ ($p < 0.05$). The increase after E028 was significantly greater to that after Modulen-IBD $P = 0.035$. After 3 weeks of normal eating, breath chemicals had in every case returned to levels indistinguishable from those present at the start of the first period of enteral feeding.

Results

Study 2:

At the start of treatment all 17 patients had active disease as confirmed by a Harvey and Bradshaw index of >6 and raised concentration of C-reactive protein (CRP) in the blood. 9 patients were receiving no medication, 4 were taking 5ASA compounds, 2 were taking 5ASA with Azathioprine, 1 taking

258 Azathioprine alone and 1 taking Azathioprine and Prednisolone. Patients were asked to continue the same
259 medication throughout the study and this was not changed in any way, remission being achieved in all
260 cases by the addition of enteral feed. The mean Harvey & Bradshaw (H&B) before treatment was $6.88 \pm$
261 $SD\ 2.93$ falling to $4 \pm SD\ 5.50$ after treatment, ($p < 0.05$). The initial mean CRP was $36.0 \pm SD\ 41.3\text{mg/L}$
262 falling to $8.11 \pm SD\ 3.59$ after treatment ($p < 0.05$).

263

264 The results of GC/MS faecal analysis are summarised in Table 1. Many compounds of known bacterial
265 origin were present in the initial sample. These included propanoic and butanoic acids, para-cresol,
266 indole, dimethyl disulphide and phenol. The concentrations of the SCFAs fell dramatically after enteral
267 feeding. No difference was discerned in the fall of concentrations of bacterial metabolites in those
268 subjects receiving enteral feeds alone, and those who continued their previous medication. Thus the
269 results of all the patients were analysed together.

270

271 There were also however, a number of potentially toxic compounds present. These included the alcohols,
272 1-propanol and 1-butanol as well as the methyl and ethyl esters of propanoic acid and butanoic acid.
273 After treatment, the amounts of these compounds also fell significantly. The SCFA-esters disappeared
274 virtually completely and there was a significant fall in the concentration of 1-propanol and 1-butanol.
275 However, other chemicals including those derived by bacterial breakdown of amino acids, phenol and
276 indole did not change significantly (table 1).

277

278 **Discussion**

279

280 The present study demonstrates changes in chemicals of microbial origin in both healthy controls and in
281 patients with CD after administration of enteral feeds. Our first study confirms reports of stool colour
282 change during treatment with the development of breath odour. It is probable that this was the result of
283 the cessation of the normal microbial breakdown of biliverdin (green) to stercobilin (brown).

284

285 We also attempted to assess bacterial activity by determination on the breath of known bacterial
286 metabolites that might be absorbed into the blood stream from the colon. Many chemicals are present in
287 breath and urine and we detected 140. Their origins of many are poorly understood. We therefore
288 concentrated on changes in the excretion of two chemicals whose synthesis by the microbiota is well
289 understood, namely phenol and indole.^{14,15}

290

291 Phenol and indole are produced by the microbial conversion of tyrosine and tryptophan respectively .
292 Much less is produced when carbohydrate fermentation is continuing in the colon.. Conversely, when
293 carbohydrate was withdrawn from the diet, phenol production from endogenous protein sources such as
294 intestinal secretions and exfoliated cells was increased ^{17,18}

295

296 In the present study, phenol and indole identified on the breath showed a significant increase in
297 concentration after feeding with Modulen-IBD and an even greater increase after E028extra, which
298 rapidly returned to base line on resumption of a normal diet. This is consistent with a switch in colonic
299 fermentation to a protein-based pattern, as an effect of ingesting carbohydrate in the form of
300 maltodextrins - simple sugars that are absorbed high in the small intestine - rather than complex
301 carbohydrates that may pass down to be fermented by the colonic flora. Indole is malodorous and may
302 contribute to the unpleasant breath odour reported by our volunteers.

303

304 The term ‘enterometabolic disease’ has been suggested for non-infective conditions arising from
305 abnormal fermentation by the colonic microbiota ¹³. Patients with IBS have a similar abnormal gut flora
306 to that seen in CD.^{11,20,23} and have a markedly increased excretion of a bacterial product, hydrogen. This
307 was dramatically reduced, with highly significant reduction in symptoms, when patients were switched
308 from a standard diet to an exclusion diet, suggesting that the diet reduced microbial activity²⁴. Support
309 for this concept was provided by the demonstration of reduced hydrogen excretion in patients with IBS ,

again with significant improvement in symptoms, when microbial activity was reduced by administration of antibiotics or by enteral feeding.²⁵

It seems possible that CD like IBS may be an 'entero-metabolic disorder',¹³ and that enteral feeding is effective because it reduces the metabolic activity of an abnormal colonic flora?

There is strong evidence that the host microflora provokes an immunological response in CD. Duchmann and his colleagues showed that monocytes from the peripheral blood and the lamina propria were activated when incubated with preparations of faecal bacteria from other subjects, but not by such preparations derived from the faeces of the host. Monocyte activation occurred only when host faeces was incubated with cells obtained from the lamina propria from sites of active CD. No activation was seen in monocytes obtained from areas where no active CD was present, suggesting that monocytes in areas of active CD were specifically targeted against the host microflora.¹⁰

This finding has been supported by later studies that demonstrated that the great majority of microorganisms found in the faeces of patients with IBD were coated with immunoglobulin, including IgA, IgG and IgM, whereas in normal subjects or those with IBS, less than 20% were so affected¹².

Furthermore, a significant reduction in the number of microorganisms coated with immunoglobulin was seen after 2 weeks treatment with corticosteroids in UC, and a similar response occurred in CD after a two week course of elemental feeding. This suggested that the immune response to the flora had been significantly reduced, an interpretation supported by the finding that patients with CD and UC in long term remission had similar numbers of coated bacteria to those seen in healthy controls.¹²

No specific pathogen has as yet been confirmed as being the cause of CD, but it has been demonstrated that the faecal flora is abnormal with an overgrowth of facultative anaerobes and reduction in the numbers of important beneficial species such as *Lactobacilli* and *Bifidobacter*.^{11,20} Although previous studies of the effects of enteral feeding on the composition of the bacterial flora in CD, had been inconclusive,^{21,22}

a recent study of the entire gut mucosal microbiome in a child with CD before and after nutritional therapy showed that the flora, initially markedly abnormal, returned after therapy, to a pattern very similar to that found in a healthy control ²⁶. Likewise, it has also been shown that enteral nutrition in CD may reduce the levels of certain bacteria within the *Firmicutes*. These bacteria are important producers of SCFAs and this report is in keeping with our discovery of reduced SCFA production. ²⁷

Unfortunately, it was not possible in the present study to perform complex studies of changes in the gut microbiome, but changes in bacterial metabolites serve as valuable markers of its metabolic activity. SCFAs have an important function in the colon especially butanoic acid which is a major source of nutrition for colonocytes.¹⁹ They are produced by the microbial fermentation of undigested complex carbohydrates entering the caecum and the fall in faecal SCFA concentration found after enteral feeding in our patients with CD was consistent with reduction in colonic fermentation.

Such a reduction in fermentation might be beneficial if it resulted in less production of toxic metabolites. There were highly significant falls in the concentrations of number of chemicals including 1-propanol, p-cresol, phenol, 1-butanol, dimethyl disulphide and fatty acid ethyl esters (Table 1). These are known to be toxic chemicals which we have shown not to be present in the stools of healthy volunteers. ¹⁶ It seems possible that the production of such chemicals might be a factor initiating an immune attack on the microflora. This could lead to coating of microflora with immunoglobulin – a suggestion which has been supported by the significant reduction in bacterial coating seen after 2 weeks feeding with enteral feeds.^{12,16}

Similar toxic chemicals also appear in UC, but in contrast to CD, **do not fall** after enteral feeding, but only after successful treatment by immunosuppression with prednisolone¹⁶. Although evidence on the role of diet in UC remains weak, this suggests that the microflora in UC differs from that in CD in that it derives its nutritional requirements, not from food residues, but from other substances present in the large

intestine – possibly mucus or intestinal secretions. It is therefore feasible, that the production of toxic chemicals resulting from abnormal bacterial metabolism, may be an important factor in the initiation of an immune attack on the microflora in inflammatory bowel disease.

Reference List

1. O'Morain C, Segal AW, Levi AJ, Elemental diet as primary treatment of acute Crohn's disease: A controlled trial BMJ 1984 **288**: 1859-62
2. Middleton SJ, Rucker JT, Kirby GA, Riordan AM, Hunter JO Long chain triglycerides reduce the efficacy of enteral feeds in patients with active Crohn's disease. Clinical Nutrition 1995 **14** 229-236
3. King TS, Woolner JT, Hunter JO Dietary treatment of Crohn's disease Review article – The Dietary management of Crohn's disease Aliment. Pharm. Toxicol **11** 17-31 1997
4. Walker-Smith J (2001) The role of enteral feeding in Crohn's disease of childhood Minerva Pediatr. **52** 277-9
5. Gupta K, Noble A, Kachelries KE, Albenberg L, Kelsen JR, Grossman AB, Baldassano RN A novel enteral nutrition protocol for the treatment of pediatric Crohn's disease. Inflammatory Bowel Diseases 2013 **19** 1374-8
6. Brown AC, Roy M, Does evidence exist to include dietary therapy in the treatment of Crohn's disease? Expert Review of Gastroenterology and Hepatology 2010 **4** 191-215

- 387 7. Greenberg GR, Fleming CR, Jeejeebhoy KN, Rosenberg IH, Sales D, Tremaine Controlled
388 trial of bowel rest and nutritional support in the management of Crohn's disease. *WJ. Gut.* 1988
389 Oct; **(10)**:1309-15
390
- 391 8. Teahon K, Pearson M, Smith T, Bjarnason I, Alterations in nutritional status and disease activity
392 during treatment of Crohn's disease with elemental diet *Scan J Gastroenterol.* 1995 **30** 54-60
393
- 394 9. Lomer MC, Grainger SL, Ede R et al Lack of efficacy of a reduced microparticle diet in a
395 multicentered trial of patients with active Crohn's disease. *Euro J Gastroenterol Hepatol* 2005
396 **17** 377-84
397
- 398 10. Duchmann R, Kaiser I, Hermann E, Mayet W, Meyer zum Buschenfelde KE and KH Tolerance
399 exists towards intestinal flora but is broken in active Inflammatory Bowel Disease *Clin. Exp.*
400 *Immunol.* 1995 **102** 448-455
401
- 402 11. Albenberg LG, Lewis JD, Wu GD, Food and the Gut microbiota in inflammatory bowel disease;
403 a critical connection. *Current Opinion in Gastroenterology* 2012 **28** 314-20
404
- 405 12. Van der Waaij LA, Kroese FG, Visser A, Nelis GF, Westenveld BD, Jansen PL, Hunter JO
406 Immunoglobulin of faecal bacterial in Inflammatory Bowel Disease *Eur. J Gastroenterol*
407 *Hepatol.* 2004 **16** 669-74
408
- 409 13. Hunter JO Food Allergy – or entero metabolic disorder *Lancet* 1991 **338** 495-6
410
- 411 14. Schofield WN Predicting Basal metabolic rate, new standards and review of previous work.
412 *Hum Nutr Clin Nutr* 1985 **39C**: 5-41

- 413
- 414 15. Harvey RF, Bradshaw JM, A simple index of Crohn's disease activity Lancet 1980 **i** 514
- 415
- 416 16. Walton C, Fowler DP, Turner C, Jia W, Whitehead RN, Griffiths L, Dawson C, Waring RH,
- 417 Ramsden DB, Cole JA, Cauchi M, Bessant C, Hunter JO. Analysis of volatile organic
- 418 compounds of bacterial origin in chronic gastrointestinal diseases. *Inflamm Bowel Dis.* 2013 **19**
- 419 2069-78
- 420
- 421 17. Cummings JH, Hill MJ, Bone ES, Branch WJ, Jenkins DJA The effect of meat protein and
- 422 dietary fiber on colonic function and metabolism II Bacterial metabolites in faeces and urine.
- 423 *Am J Clin Nutr* 1979 **32** 2094-2101
- 424
- 425 18. Macfarlane GT, Cummings JH, Allison C Protein degradation by human intestinal bacteria J
- 426 *Gen Microbiol* 1986 **132** 1647-56
- 427
- 428 19. Roediger WE, Oxidative and synthetic functions of n-Butyrate in colonocytes *Dis Colon*
- 429 *Rectum* 1992 **35** 511-12
- 430
- 431 20. Sartor RB, Therapeutic manipulation of the enteric microflora in inflammatory bowel diseases;
- 432 antibiotics, probiotics and prebiotics. *Gastroenterology* 2004 **126** 1620-33
- 433
- 434 21. Kaakoush NO, Day AS, Leach ST, Lemberg DA, Nielsen S Mitchell HM. Effect of exclusive
- 435 enteral nutrition on children with newly diagnosed Crohn's disease. *Clin Trans Gastroenterol.*
- 436 2015; 6, e71; doi: 10.1038/ctg.2014.21
- 437 22. den Besten G, van Eunen K, Groen AK, Venema K, Reijngoud D-J, Bakker B. The role of
- 438 short-chain fatty acids in the interplay between diet, gut microbiota, and host energy
- 439 metabolism. 2013 *J. Lipid Res.* 54: 2325-2340.
- 440 23. Garner CE, Smith S, Costello BD, White P, Spencer R, Probert CSJ, Ratcliffe NM. Volatile
- 441 organic compounds from feces and their potential for diagnosis of gastrointestinal disease. 2007

442 Faseb J. 21(8): 1675-1688.

443 24. Rehman A, Lepage P, Nolte A, Helmig S, Schreiber S, Ott SJ. Transcriptional activity of the
444 dominant gut mucosal microbiota in chronic inflammatory bowel disease patients. 2010 J. Med.
445 Microbiol. 59(9): 1114-1122.

446 25. Bodelier AGL, Smolinska A, Baranska A, Dallinga JW, Mujagic Z, Vanhees K, van der Heuvel
447 T, Masclee AAM, Jonkers D, Pierik MJ, van Schooten FJ. Volatile Organic Compounds in
448 Exhaled Air as Novel Marker for Disease Activity in Crohn's Disease: A Metabolomic
449 Approach 2015 Inflamm. Bowel Dis. 21(8): 1776-1785.

450 26. D'Argenio V, Precone V, Casaburi G, Miele E, Martinelli M, Staiano A, et al An altered gut
451 microbiome profile in a child affected by Crohn's disease normalized after nutritional therapy.
452 AmJ Gastroenterol 2013 **108** 851-852

453 27. Shiga H, Jajiura T, Shinozaki J, Suzuki M, Takagi S, Kinouchi Y, Takahashi S, Shimosegawa T,
454 Changes of faecal microbiota in Crohn's disease treated with an elemental diet and total
455 parenteral nutrition. Digestive Liver disease 2012 **44** 736-42.

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459
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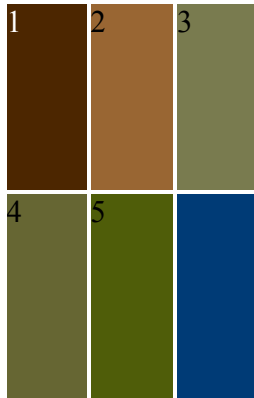
Table 1 Changes in faecal chemicals before and after elemental feeding in patients with CD.

Compound	VOC concentration (ng/l)		
	Median (lower quartile, upper quartile)	Pre-treatment	Post-treatment p-value
acetone	57 (38, 128)	80 (50, 104)	0.435
propanoic acid	169 (0, 328)	12 (0, 84)	0.031*
butanoic acid	1110 (316, 1506)	24 (0, 104)	0.001*
1-propanol	229 (41, 892)	36 (0, 233)	0.025*
propanoic acid, ethyl ester	19 (0, 117)	0 (0, 15)	0.008*
butanoic acid, methyl ester	19 (7, 121)	0 (0, 1)	0.013*
butanoic acid, ethyl ester	46 (4, 255)	0 (0, 15)	0.008*
p-cresol	518 (118, 1160)	480 (144, 1051)	0.687
indole	118 (54, 146)	20 (0, 128)	0.125
dimethyl disulphide	83 (34, 683)	39 (0, 140)	0.113
1-butanol	99 (57, 256)	58 (0, 199)	0.030*
butanoic acid, 3-methyl	147 (48, 504)	0 (0, 45)	0.015*
phenol	64 (16, 102)	24 (10, 177)	0.332

Concentrations of volatile organic compounds (VOCs) in fecal headspace from healthy controls and volunteers diagnosed with Crohn's disease (CD) before and after treatment. Median and upper and lower quartile values are shown. Differences were examined by Wilcoxon matched pairs test, $p < 0.05$ (two-sided) being considered significant and indicated by an asterisk.

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472 Copy of the stool colour chart provided to volunteers in study 1. The colours are numbered from brown
473 (1) to increasing greenness (5) the blue is just for the slide background. This is not for publication but to
474 be made available for any readers who may enquire about it.

475

476 **SUPPORTING INFORMATION**

477 Declaration of funding interests:

478 The work for this study was funded by the Wellcome Trust Grant No. 080238/Z/06/Z. Professor Hunter
479 has acted as a consultant and received research grants from Nutricia (UK) Liverpool and from Nestle.

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481 **STROBE STATEMENT**

482 All items on the strobe checklist have been checked and confirmed to be included in this paper.



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Enteral feeding reduces metabolic activity of the intestinal microbiome in Crohn's disease: an observational study

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