



OPEN ACCESS

# The gut microbiota and host health: a new clinical frontier

Julian R Marchesi,<sup>1,2</sup> David H Adams,<sup>3</sup> Francesca Fava,<sup>4</sup> Gerben D A Hermes,<sup>5,6</sup> Gideon M Hirschfield,<sup>3</sup> Georgina Hold,<sup>7</sup> Mohammed Nabil Quraishi,<sup>3</sup> James Kinross,<sup>8</sup> Hauke Smidt,<sup>5</sup> Kieran M Tuohy,<sup>4</sup> Linda V Thomas,<sup>9</sup> Erwin G Zoetendal,<sup>5,6</sup> Ailsa Hart<sup>10</sup>

For numbered affiliations see end of article.

## Correspondence to

Dr Ailsa Hart, IBD Unit, St Mark's Hospital and Imperial College London, Watford Road, London HA13UJ, UK; ailsa.hart@nhs.net

Received 20 May 2015  
Revised 14 July 2015  
Accepted 16 July 2015  
Published Online First  
2 September 2015

## ABSTRACT

Over the last 10–15 years, our understanding of the composition and functions of the human gut microbiota has increased exponentially. To a large extent, this has been due to new 'omic' technologies that have facilitated large-scale analysis of the genetic and metabolic profile of this microbial community, revealing it to be comparable in influence to a new organ in the body and offering the possibility of a new route for therapeutic intervention. Moreover, it might be more accurate to think of it like an immune system: a collection of cells that work in unison with the host and that can promote health but sometimes initiate disease. This review gives an update on the current knowledge in the area of gut disorders, in particular metabolic syndrome and obesity-related disease, liver disease, IBD and colorectal cancer. The potential of manipulating the gut microbiota in these disorders is assessed, with an examination of the latest and most relevant evidence relating to antibiotics, probiotics, prebiotics, polyphenols and faecal microbiota transplantation.

integrate an individual's microbiota into some form of personalised healthcare and, by better understanding its role, treat an individual's diseases more efficiently and in a more targeted fashion. With a more complete understanding of the disease process, we will be able to more accurately stratify different disease states and determine whether or not the gut microbiota is a potential therapeutic target which we can modulate in order to treat specific diseases.

This review gives a much needed update on current understanding of the gut microbiota in GI diseases and metabolic disorders, and gives an insight into how this might impact on clinical practice. The evidence for the preventive and therapeutic benefit of different ways of modulating the gut microbiota, such as probiotics, prebiotics, antibiotics and faecal microbiota transplantation (FMT) (see [box 2](#)), is reviewed.

## CURRENT UNDERSTANDING OF THE GUT MICROBIOTA

In the last decade, several large-scale projects, for example, the human microbiome project, have investigated the microbiota of a variety of bodily niches, including the skin as well as the oral, vaginal and nasal cavities.<sup>2</sup> While some of these are relatively easy to access, the GI tract remains a challenging environment to sample, and to describe. Currently the majority of research is focused on the gut microbiota, since this is where the greatest density and numbers of bacteria are found, with most data being derived from faecal samples and, to a lesser extent, mucosal biopsies. While it is relatively easy to obtain fresh faecal samples, the information obtained from them does not represent the complete picture within the gut. From a number of limited studies, we know that the small intestine contains a very different abundance and composition of bacteria, with much more dynamic variation compared with the colon.<sup>3</sup> The colonic microbiota is largely driven by the efficient degradation of complex indigestible carbohydrates but that of the small intestine is shaped by its capacity for the fast import and conversion of relatively small carbohydrates, and rapid adaptation to overall nutrient availability. While faeces are not an ideal proxy for the GI tract, they do give a snapshot of the diversity within the large intestine. Furthermore, the majority of the data comes from North American and European studies with very few studies in Asia, Africa or South America. Hence we have a somewhat biased view of the gut microbiota.

## INTRODUCTION

Imagine the scenario: a scientist at a conference claims to have found a new organ in the human body. It is comparable to the immune system in as much as it is made up of a collection of cells, it contains a 100 times more genes than the host, is host-specific, contains heritable components, can be modified by diet, surgery or antibiotics, and in its absence nearly all aspects of host physiology are affected. While this may seem far-fetched, it is the current situation in which we find ourselves. We now realise that the human microbiota is an overlooked system that makes a significant contribution to human biology and development. Moreover, there is good evidence that humans co-evolved a requirement for their microbiota.<sup>1</sup>

In the past decade, partly because of high resolution observational studies using next-generation sequencing technologies and metabolite profiling (see [box 1](#)), the gut microbiota has become associated with promotion of health and the initiation or maintenance of different GI and non-GI diseases. As we enter the postmetagenomic era, we need to move away from simple observations to determine what are merely correlations and what are causal links—and focus efforts and resources on the latter. This postmetagenomic era is starting to provide new therapeutic targets based on a better understanding of how the microbiota interacts with the host's physiology. Ultimately, we aim to



Open Access  
Scan to access more  
free content



CrossMark

To cite: Marchesi JR, Adams DH, Fava F, *et al.* Gut 2016;**65**:330–339.

**Box 1 A short primer of microbiology (see also Lepage et al)<sup>4</sup>****Dysbiosis**

A disturbance or imbalance in a biological system, for example, changes in the types and numbers of bacteria in the gut which may lead to developing different diseases, such as IBD.

**Inflammasomes**

Recently discovered multiprotein complexes that are involved in a wide range of inflammatory processes including programmed cell death (pyroptosis), in response to the recognition of microbial and danger signals.

**Lipopolysaccharide (LPS)**

A major component of the outer membrane of Gram-negative bacteria; an endotoxin. Now implicated as a driver of inflammation and associated with onset of certain diseases.

**Lipoteichoic acid**

A major component of the outer membrane of Gram-positive bacteria; an endotoxin. Now implicated as a driver of inflammation and associated with onset of certain diseases.

**Metabonome/metabolome**

A profile of the chemicals in a tissue or sample, for example the urine metabolome. This profile represents a snapshot in time of what chemicals are present in the sample.

**Metagenomics**

A method which allows us to create catalogues of what the bacteria can do based on the genes that they have.

**Microbiome**

A collection of different microbes and their functions or genes found in an environmental habitat. Different parts of the body have different microbiomes, for example, the skin microbiome is different to the gut microbiome, but they are all part of the human microbiome.

**Microbiota**

The types of organisms that are present in an environmental habitat, whether they are bacteria, viruses or eukaryotes.

**'Omic' methods**

A term which describes a set of methods, such as genomics, metabonomics, metagenomics, etc, which we use to explore the interactions between the bacteria in the gut and the host.

**Pathobiont**

A commensal organism that can cause disease when specific genetic or environmental conditions are altered in the host.

**Phenome**

A collection of measurable features that define an individual.

This rapid increase in interest in the microbiome has also been driven by the application of multi-'omic' technologies; we refer the reader to Lepage et al<sup>4</sup> for more detailed explanation of these (see also [box 1](#)).

**What do we know about the gut microbiota?**

Bearing in mind the limitations above, the GI tract is often seen as a two phylum system (the Firmicutes and Bacteroidetes) although it should be noted that members of at least 10 different phyla can also have important functional contributions (see [box 3](#)). We are also very bacteria-centric when we look at the gut microbiota; only a handful of papers have looked at the viral component (or virome) and micro-eukaryotes (protozoa and fungi). When the gut microbiota of relatively large cohorts of individuals (eg, more than 100) is analysed, it can be seen

**Box 2 Potential therapies aimed at modulation of the gut microbiota**

**Probiotics:** Live microorganisms that, when administered in adequate amounts, confer a health benefit on the host.<sup>107 108</sup> Examples include strains of the genera *Bifidobacterium* and *Lactobacillus*. Probiotics can have multiple interactions with the host,<sup>109</sup> including competitive inhibition of other microbes, effects on mucosal barrier function and interaction with antigen presenting dendritic cells.<sup>72</sup>

**Prebiotics:** A selectively fermented ingredient that results in specific changes in the composition and/or activity of the GI microbiota, thus conferring benefit(s) upon host health.<sup>110</sup>

Prebiotics are usually non-digestible carbohydrates, oligosaccharides or short polysaccharides, with inulin, oligofructose, galactofructose, galacto-oligosaccharides and xylo-oligosaccharides being some of the most intensively studied.

**Faecal microbiota transplantation:** The introduction of gut bacteria from a healthy donor into a patient, through transfer of an infusion of a faecal sample via nasogastric tube, nasoduodenal tube, rectal enema or the biopsy channel of a colonoscope.<sup>111</sup>

that the ratio of the Firmicutes:Bacteroidetes is not the same in all individuals. Currently we do not know the significance of being at either end of this continuum, especially as a large shift in the relative abundance of a group of organisms translates to a modest change in bacterial numbers. Yet there is evidence that depletion of a single species, for example, *Faecalibacterium prausnitzii*, belonging to the Firmicutes phylum, has been associated with IBD.<sup>5</sup> But in the scientific literature, we see counter-arguments for any involvement of this species in IBD.<sup>6</sup> This disparity highlights the current status of understanding. We

**Box 3 A primer in taxonomics**

In order to classify bacteria we have adopted the Linnaean system, which comprises hierarchies into which an organism is placed.

For example humans are classified at the species level as *Homo sapiens*, which are members of the genus *Homo*, family *Hominidae*, order Primates, class Mammalia, phylum Chordata and finally kingdom Eukaryota. As one moves up through the different taxonomic levels, from species to kingdom, greater numbers of organisms become associated with each other. In life there are three kingdoms, the Bacteria, Archaea and Eukaryota, with the majority of bacterial-like (or prokaryotes) being classified within the Bacteria and Archaea. For example the gut commensal and sometime pathogenic species *Escherichia coli* is found in the kingdom Bacteria; phylum Proteobacteria; class Gammaproteobacteria; order *Enterobacteriales*; family *Enterobacteriaceae* and finally genus *Escherichia*. Thus when we refer to phyla or a phylum, we are usually describing very large collections of related organisms. In the large intestine of healthy adults the two most dominant phyla are the Firmicutes (comprised mainly of Gram-positive clostridia) and Bacteroidetes (comprised mainly of Gram-negative bacteria such as the species *Bacteroides fragilis*).

know that the gut microbiota is essential to the proper function and development of the host but we are unsure which are key-stone species and whether the microbiota's function is more important than any individual member of the community. But this is too simplistic a view. In several cases, strain differences within a species can be the difference between being a pathogen/pathobiont and being a probiotic: for example, *Escherichia coli* is associated with IBD and colorectal cancer (CRC)<sup>7 8</sup> yet an *E. coli* strain is used as a probiotic.

In fact, five phyla represent the majority of bacteria that comprise the gut microbiota. There are approximately 160 species in the large intestine of any individual<sup>9</sup> and very few of these are shared between unrelated individuals. In contrast, the functions contributed by these species appear to be found in everybody's GI tract, an observation that leads us to conclude that function is more important than the identity of the species providing it. Yet differences in the gut microbiota may matter because these may result in differences in the effectiveness of a function. For example, while the ability to synthesise short chain fatty acids (SCFAs) is found in all humans,<sup>10</sup> their amounts can vary.

## DIETARY MODULATION OF THE GUT MICROBIOTA

### Metabolic activities of the gut microbiota

Carbohydrate fermentation is a core activity of the human gut microbiota, driving the energy and carbon economy of the colon. Dominant and prevalent species of gut bacteria, including SCFA-producers, appear to play a critical role in initial degradation of complex plant-derived polysaccharides,<sup>11</sup> collaborating with species specialised in oligosaccharide fermentation (eg, bifidobacteria), to liberate SCFAs and gases which are also used as carbon and energy sources by other more specialised bacteria (eg, reductive acetogens, sulfate-reducing bacteria and methanogens).<sup>12</sup> Efficient conversion of complex indigestible dietary carbohydrates into SCFA serves microbial cross-feeding communities and the host, with 10% of our daily energy requirements coming from colonic fermentation. Butyrate and propionate can regulate intestinal physiology and immune function, while acetate acts as a substrate for lipogenesis and gluconeogenesis.<sup>13</sup> Recently, key roles for these metabolites have been identified in regulating immune function in the periphery, directing appropriate immune response, oral tolerance and resolution of inflammation, and also for regulating the inflammatory output of adipose tissue, a major inflammatory organ in obesity.<sup>14</sup> In the colon, the majority of this carbohydrate fermentation occurs in the proximal colon, at least for people following a Western style diet. As carbohydrate becomes depleted as digesta moves distally, the gut microbiota switches to other substrates, notably protein or amino acids. Fermentation of amino acids, besides liberating beneficial SCFAs, produces a range of potentially harmful compounds. Some of these may play a role in gut diseases such as colon cancer or IBD. Studies in animal models and in vitro show that compounds like ammonia, phenols, *p*-cresol, certain amines and hydrogen sulfide, play important roles in the initiation or progression of a leaky gut, inflammation, DNA damage and cancer progression.<sup>15</sup> On the contrary, dietary fibre or intake of plant-based foods appears to inhibit this, highlighting the importance of maintaining gut microbiome carbohydrate fermentation.<sup>16</sup> Recognition of carbohydrate fermentation as a core activity of the gut microbiota provides the scientific basis for rational design of functional foods aimed at improving gut health and also for impacting on microbiota activities linked to systemic host physiology through newly recognised interkingdom axes of communication such as the gut:liver axis, the gut:brain axis and the gut:brain:skin axis.<sup>17</sup>

### Three 'P's' for gut health: probiotics, prebiotics and polyphenols

A number of dietary strategies are available for modulating either the composition or metabolic/immunological activity of the human gut microbiota: probiotics, prebiotics and polyphenols are among the most well established.<sup>18</sup>

There are many examples of positive results with different probiotic strains against a range of disease states in animal models, however the human data are equivocal. This may partly be due to poor study design and poor choice of strain. However, there is also a persistent lack of understanding as to the very nature of probiotics, which cannot be considered a 'class' of bioactives, amenable to traditional efficacy assessments such as the meta-analysis (unless restricted to one strain), since they are all unique living organisms and their health-promoting traits are strain-specific. Rarely have probiotic strains been selected with specific mechanisms of effect in mind; this has led to conflicting observations and damaged the reputation of this area of science. A few exceptions do exist, most notably the work of Jones *et al* who selected a bile salt-hydrolysing *Lactobacillus reuteri* strain, to study its ability to reduce cholesterol levels in hypercholesterolaemic individuals. In two well powered, randomised, placebo-controlled and double-blinded studies, they demonstrated that ingestion of this strain significantly lowered total and low density lipoprotein (LDL)-cholesterol. Moreover, they suggested an underlying novel mechanism linked to reduced fat absorption from the intestine<sup>19</sup> via the nuclear receptor farnesoid X receptor (FXR).<sup>20</sup>

Prebiotics represent a specific type of dietary fibre that when fermented, mediate measurable changes within the gut microbiota composition, usually an increase in the relative abundance of bacteria thought of as beneficial, such as bifidobacteria or certain butyrate producers. As with probiotics, despite convincing and reproducible results from animal studies showing efficacy in prevention or treatment of many diseases (eg, IBD, IBS, colon cancer, obesity, type 2 diabetes (T2D) and cardiovascular disease), the data in humans remain ambiguous. Fewer well powered or well designed clinical studies have been conducted with prebiotics compared with probiotics, and there may be an issue with prebiotic dose. Human studies rarely, if ever, employ prebiotics. A prebiotic is shown to be efficacious in animal studies: typically 10% w/w of the diet, which in humans equates to about 50 g per day.<sup>18</sup> However, as we learn more about the ecology of the gut microbiota, it is becoming clear that the prebiotic concept has tapped into the underlying fabric of the gut microbiota as a primarily saccharolytic and fermentative microbes community evolved to work in partnership with its host's digestive system to derive energy and carbon from complex plant polysaccharides which would otherwise be lost in faeces.

Polyphenols are a diverse class of plant secondary metabolites, often associated with the colour, taste and defence mechanisms of fruit and vegetables. They have long been studied as the most likely class of compounds present in whole plant foods capable of affecting physiological processes that protect against chronic diet-associated diseases. The gut microbiota plays a critical role in transforming dietary polyphenols into absorbable biologically active species, acting on the estimated 95% of dietary polyphenols which reach the colon.<sup>21</sup> Recent studies show that dietary intervention with polyphenol extracts, most notably dealcoholised red wine polyphenol extract and cocoa-derived flavanols, modulate the human gut microbiota towards a more 'health-promoting profile' by increasing the relative abundance of bifidobacteria and lactobacilli. These data again raise the possibility

that certain functional foods tap into the underlying ecological processes regulating gut microbiome community structure and function, contributing to the health of the gut microbiota and its host.<sup>22</sup>

### OBESITY-RELATED DISEASES AND THE GUT MICROBIOTA

Starting around 2004, the hallmark studies of Gordon *et al* demonstrated a potential relationship between the gut microbiome and development of an obese phenotype. An increase in relative abundance of Firmicutes and a proportional decrease in Bacteroidetes were associated with the microbiota of obese mice,<sup>23</sup> which was confirmed in a human dietary intervention study demonstrating that weight loss of obese individuals (body mass index, BMI > 30) was accompanied by an increase in the relative abundance of Bacteroidetes.<sup>24</sup> Nevertheless, based on most human studies, the obesity-associated decrease in the ratio of Bacteroidetes to Firmicutes (B:F) remains controversial.<sup>24–25</sup> This is likely due to heterogeneity among human subjects with respect to genotype and lifestyle. Recent studies have identified diet, especially fat, as a strong modulator of the microbiota, particularly in inbred and age-standardised laboratory animals. The sources of variation in the microbiota are mainly limited to the experimental diets used, and there is growing evidence that the high fat intake rather than obesity per se had a direct effect on the microbiota and linked clinical parameters.<sup>26</sup> However, in humans the microbiome is exposed to fundamentally different 'environmental' factors in obese and lean individuals that go beyond BMI alone, including diet<sup>26</sup> and host hormonal factors.<sup>27</sup> In addition, the aetiology of obesity and its metabolic complications, including low grade inflammation, hyperlipidaemia, hypertension, glucose intolerance and diabetes, reflect the complex interactions of these multiple genetic, behavioural and environmental factors.<sup>28</sup> Lastly, the accuracy of BMI as an indicator for obesity is limited; 25% of obese people could in fact be regarded as metabolically 'healthy' (ie, with normal lipid and glucose metabolism).<sup>29</sup> Therefore, linking GI tract microbial composition directly and exclusively to obesity in humans will remain challenging due to the various confounding factors within the heterogeneous population.

This complexity has led to a shift from treating obesity as a single phenotype, to attempts at correlating microbial signatures to distinct or multiple features associated with (the development of) metabolic syndromes such as T2D. Recently, two (meta) genome-wide association studies were performed, with 345 Chinese individuals<sup>30</sup> and 145 European women.<sup>31</sup> In both studies, de novo generated metagenomic species-level gene clusters were employed as discriminant markers which, via mathematical modelling, could better differentiate between patients and controls with higher specificity than a similar analysis based on either human genome variation or other known risk factors such as BMI and waist circumference. At the functional level, membrane transporters and genes related to oxidative stress were enriched in the microbiota of patients,<sup>31</sup> while butyrate biosynthesis was decreased.<sup>30</sup> Although both studies observed high similarities in microbial gene-encoded functions, the most discriminant metagenomic species-level gene clusters differed between the cohorts (*Akkermansia* did not contribute to the classification in the European cohort whereas *Lactobacillus* showed no contribution in the Chinese study population), indicating that diagnostic biomarkers could be specific to the population studied.

In another metagenomic study, a bimodal distribution of microbial gene richness in obese individuals was observed, stratifying individuals as High Gene Count or Low Gene Count

(HGC and LGC).<sup>32</sup> HGC individuals were characterised by higher prevalence of presumed anti-inflammatory species such as *E. prausnitzii*, and an increased production potential of organic acids (including butyrate). In contrast, LGC individuals showed higher relative abundance of potentially proinflammatory *Bacteroides* spp and genes involved in oxidative stress response. Remarkably, only biochemical obesity-associated variables, such as insulin resistance, significantly correlated with gene count while weight and BMI did not, underscoring the inadequacy of BMI as an indicator for 'Obesity and its Associated Metabolic Disorders' (OAMD).<sup>33</sup> An accompanying paper demonstrated that a diet-induced weight-loss intervention significantly increased gene richness in the LGC individuals which was associated with improved metabolic status.<sup>34</sup> Although gene richness was not fully restored, these findings support the reported link between long-term dietary habits and the structure of the gut microbiota.<sup>31</sup> It also suggests permanent adjustment of the microbiota may be achieved through diet.

Most studies involving the microbiome have been solely correlative but recently a causal relationship was established between host glucose homeostasis and gut microbial composition. FMT from lean donors to individuals with metabolic syndrome significantly increased their insulin sensitivity.<sup>33</sup> The transplant produced an increase in faecal butyrate concentrations, microbial diversity and the relative abundance of bacteria related to the butyrate-producing *Roseburia intestinalis*.

Together, these studies produce a body of evidence that the microbiome plays a role in host energy homeostasis and the establishment and development of OAMD, although the exact mechanisms remain obscure. Previous contradictory findings might be attributed to miscellaneous approaches,<sup>35</sup> and also heterogeneity in genotype, lifestyle and diet of humans combined with the complex aetiology of OAMD. Nonetheless, a clearer picture is emerging. The gut of individuals with OAMD is believed to harbour an inflammation-associated microbiome, with a lower potential for butyrate production and reduced bacterial diversity and/or gene richness. Although the main cause of OAMD is excess caloric intake compared with expenditure, differences in gut microbial ecology might be an important mediator and a new therapeutic target or a biomarker to predict metabolic dysfunction/obesity in later life.

### LIVER DISEASE AND THE GUT MICROBIOTA

The liver receives 70% of its blood supply from the intestine via the portal vein, thus it is continually exposed to gut-derived factors including bacterial components, endotoxins (lipopolysaccharide, flagellin and lipoteichoic acid) and peptidoglycans. Multiple hepatic cells, including Kupffer cells, sinusoidal cells, biliary epithelial cells and hepatocytes, express innate immune receptors known as pathogen-recognition-receptors that respond to the constant influx of these microbial-derived products from the gut.<sup>36</sup> It is now recognised that the gut microbiota and chronic liver diseases are closely linked. Characterising the nature of gut dysbiosis, the integrity of the gut barrier and mechanisms of hepatic immune response to gut-derived factors is potentially relevant to development of new therapies to treat chronic liver diseases.<sup>37</sup> Furthermore the field of bile acid signalling has thrown open the concept of the gut:liver axis as being active and highly regulated.<sup>38</sup>

### Non-alcoholic fatty liver disease

The pathophysiology of NAFLD is multifactorial with strong genetic and environmental contributions. Recent evidence demonstrates that gut microbiota dysbiosis can result in the

development of obesity-related non-alcoholic fatty liver disease (NAFLD), and patients with NAFLD have small intestinal bacterial overgrowth and increased intestinal permeability.<sup>39</sup> In the 1980s, development of non-alcoholic steatohepatitis (NASH) and small intestinal bacterial overgrowth was observed in humans after intestinal bypass and, interestingly, regression of hepatic steatosis after metronidazole treatment, suggesting a possible role for the gut bacteria in NAFLD.<sup>40</sup> Disruption of the murine inflammasomes (see box 1) is associated with an increase in Bacteroidetes and reduction in Firmicutes and results in severe hepatic steatosis and inflammation.<sup>41</sup> Faecal microbiota analysis of patients with NAFLD and NASH has produced variable results due to significant variation of patient demographics, severity of liver disease and methodology. A lower proportion of *Ruminococcaceae* was noted in patients with NASH compared with healthy subjects<sup>42</sup> and a study which characterised gut microbiota of children with NASH, obesity and healthy controls showed that patients with NASH had a higher proportion of *Escherichia* compared with other groups.<sup>43</sup> Patients with NAFLD also have increased gut permeability suggesting that translocation of bacteria or microbe derived products into the portal circulation contributes to the pathogenesis.<sup>39</sup>

### Alcoholic liver disease

Since not all alcoholics develop liver injury, it appears that chronic alcohol abuse is necessary but not sufficient to cause liver dysfunction. Numerous animal model and human observational studies indicate that gut bacterial products like endotoxin may mediate inflammation and function as cofactors for the development of alcohol-related liver injury.<sup>44</sup> Serum endotoxin levels are elevated in humans and rats with alcoholic liver disease, and monocytes from alcoholics are primed to produce cytokines after endotoxin exposure. Alcohol causes intestinal bacterial overgrowth in humans and bacterial numbers were significantly higher in jejunal aspirates from patients with chronic alcohol abuse compared with controls, with similar findings in patients with alcohol-induced cirrhosis.<sup>45</sup> The degree of overgrowth correlates with the severity of cirrhosis. Tsukamoto-French model mice fed intragastrically with alcohol for 3 weeks showed increased relative abundance of Bacteroidetes and *Akkermansia* spp and a reduction in *Lactobacillus*, *Leuconostoc*, *Lactococcus* and *Pediococcus* while control mice showed a relative predominance of Firmicutes.<sup>46</sup> Patients with alcoholic liver disease also show increased gut permeability, allowing translocation of bacteria and bacterial products to the liver.<sup>47</sup>

### Autoimmune liver diseases

These consist of primary sclerosing cholangitis (PSC), primary biliary cirrhosis (PBC) and autoimmune hepatitis and represent at least 5% of all chronic liver diseases. They are presumed autoimmune conditions but the expectation is that the gut microbiota is relevant to pathogenesis, particularly because (A) PSC is associated with IBD and aberrant lymphocyte tracking, and (B) significant gut:liver axes exist through bile acid signalling. Patients with PSC develop a distinct form of IBD thus understanding the relationship between PSC and IBD is essential in uncovering the pathogenesis of PSC, which remains largely undetermined. However, it is likely that in genetically susceptible individuals, intestinal bacteria could trigger an abnormal or inadequate immune response that eventually leads to liver damage and fibrosis. Recently it was shown that patients with PSC have distinct gut microbiota. Analysis of colon biopsy microbiota revealed that patients with PSC-IBD and IBD

showed reduced abundance of *Prevotella* and *Roseburia* (a butyrate-producer) compared with controls.<sup>48–49</sup> Patients with PSC-IBD had a near-absence of *Bacteroides* compared with patients with IBD and control patients, and significant increases in *Escherichia*, *Lachnospiraceae* and *Megasphaera*. Randomised controlled trials (RCTs) investigating antibiotic therapy in PSC have shown these to be superior in improving biochemical surrogate markers and histological parameters of disease activity compared with ursodeoxycholic acid alone.<sup>50</sup> In a recent prospective paediatric case series, oral vancomycin was shown to normalise or significantly improve liver function tests.<sup>51</sup> There is evidence that mucosal integrity is compromised in patients with PSC, supporting the traditional leaky gut hypothesis of microbe-derived products translocating to the liver and biliary system to trigger an inflammatory reaction.<sup>52</sup> It was also demonstrated that tight junctions of hepatocytes were impaired in patients with PSC and infusion of non-pathogenic *E. coli* into portal circulation caused portal fibrosis in animal models.<sup>53</sup> These findings collectively suggest that bacterial antigens translocate across a leaky and possibly inflamed gut wall into the portal and biliary system to induce an abnormal immune response and contribute to PSC pathogenesis.

PBC is a chronic cholestatic liver disease with an uncertain aetiology. It is generally believed to be an autoimmune disease triggered by environmental factors in individuals with genetic susceptibility. As yet, there have been no studies directly characterising the gut microbiota in patients but molecular mimicry has been suggested as a proposed mechanism for the development of autoimmunity in PBC, with serum antibodies of patients cross-reacting with conserved bacterial pyruvate dehydrogenase complex component E2 (PDC-E2) homologues of *E. coli*, *Novosphingobium aromaticivorans*, *Mycobacterium* and *Lactobacillus* species. Hence it has been speculated that these bacteria (of possible GI origin) may initiate molecular mimicry and development of PBC in genetically susceptible hosts.<sup>54</sup>

### Modulation of the microbiota as a therapy in liver disease

Probiotics have shown promise in ameliorating liver injury by reducing bacterial translocation and hepatic inflammation.<sup>55</sup> A recent meta-analysis concluded that probiotics can reduce liver aminotransferases, total cholesterol, tumour necrosis factor  $\alpha$  and improve insulin resistance in patients with NAFLD.<sup>56</sup> A recent study in patients with cirrhosis with ascites showed that the probiotic VSL#3 significantly reduced portal hypertension.<sup>57</sup> A further study evaluated the role of FMT in modulating liver disease by transferring the NAFLD phenotype from mice with liver steatosis to germ-free mice.<sup>58</sup> There remains a need for detailed descriptive and interventional studies focused on bacterial diversity and mechanisms linking gut dysbiosis with inflammatory, metabolic and autoimmune/biliary liver injury.

### IBD AND THE GUT MICROBIOTA

Early studies implicating bacteria in IBD pathogenesis focused on identifying a potential culprit that could initiate the inflammatory cascade typical of IBD. Many organisms have been proposed: *Mycobacterium avium* subsp *paratuberculosis* and a number of Proteobacteria including enterohepatic *Helicobacter*, non-*jejuni/coli* *Campylobacter* and adherent and invasive *E. coli*. The focus has recently shifted with the realisation that the gut microbiota as a whole is altered in IBD. The concept of an altered gut microbiota or dysbiosis is possibly the most significant development in IBD research in the past decade. A definitive change of the normal gut microbiota with a

breakdown of host-microbial mutualism is probably the defining event in IBD development.<sup>59</sup>

Changes in the gut microbiota have been repeatedly reported in patients with IBD, with certain changes clearly linked to either Crohn's disease (CD) or UC: the most consistent change is a reduction in Firmicutes.<sup>60</sup> This has been balanced by reports of increased levels of Bacteroidetes phylum members,<sup>61</sup> although a reduction in Bacteroidetes has also been reported.<sup>62</sup> There is a suggestion that there may be spatial reorganisation of the *Bacteroides* species in patients with IBD, with *Bacteroides fragilis* being responsible for a greater proportion of the bacterial mass in patients with IBD compared with controls.<sup>63</sup>

Reduction in the Firmicutes species *E. prausnitzii* has been well documented in patients with CD, particularly those with ileal CD, although an increase in *E. prausnitzii* has been shown in a paediatric cohort, suggesting a more dynamic role for the species that merits further study.<sup>64</sup> Other studies have also demonstrated a decrease in Firmicutes diversity, with fewer constituent species detected in patients with IBD compared with controls.<sup>65</sup> Changes in the two dominant phyla, Firmicutes and Bacteroidetes, are coupled with an increase in abundance of members of the Proteobacteria phylum, which have been increasingly found to have a key role in IBD.<sup>66</sup> Studies have shown a shift towards an increase in species belonging to this phylum, suggesting an aggressor role in the initiation of chronic inflammation in patients with IBD.<sup>67</sup> More specifically, increased numbers of *E. coli*, including pathogenic variants, have been documented in ileal CD.<sup>68</sup> The IBD metagenome contains 25% fewer genes than the healthy gut with metaproteomic studies showing a correlative decrease in proteins and functional pathways.<sup>69</sup> Specifically, ileal CD has been shown to be associated with alterations in bacterial carbohydrate metabolism and bacterial-host interactions, as well as human host-secreted enzymes.<sup>69</sup> A detailed investigation of functional dysbiosis during IBD built on this by including inferred microbial gene content from 231 subjects and an additional 11 metagenomes.<sup>70</sup> This study identified enrichment in microbial pathways for oxidative stress tolerance, immune evasion and host metabolite uptake, with corresponding depletions in SCFA biosynthesis and typical gut carbohydrate metabolism and amino acid biosynthetic processes. Intriguingly, similar microbial metabolic shifts have been observed in other inflammatory conditions such as T2D,<sup>30</sup> suggesting a common core gut microbial response to chronic inflammation and immune activation. In addition, recent work suggests a role for viruses in IBD, with a significant expansion of *Caudovirales* bacteriophage in patients.<sup>71</sup>

### Modulation of the microbiota as a therapy in IBD

Several clinical trials have examined the approach of modulating the microbiota in patients with IBD, many of which predate the 'omics' era. Such trials provide a 'proof of concept' for the importance of the role of the gut microbiota in IBD, but marrying up individual approaches with the complex multifactorial nature of IBD remains a challenge, particularly in addressing the different phenotypes and genotypes of disease and the different 'phases' of the disease process: for example, prophylaxis, maintenance of remission, treatment of relapses.

### Ulcerative colitis

In terms of probiotic research, one of the largest clinical trials in IBD was the use of *E. coli* Nissle 1917 in the setting of remission maintenance in UC. Patients (n=327) were assigned to a double-blind, double-dummy trial to receive either the probiotic or mesalazine.<sup>72</sup> Both treatments were deemed equivalent with

regards to relapse. *E. coli* Nissle is now considered an effective alternative to 5-aminosalicylate for remission maintenance in UC.<sup>73</sup> There are two published clinical trials of the multistrain probiotic VSL#3 in the setting of mild to moderate flares of UC.<sup>74</sup> Both demonstrate that high doses improve disease activity scores but whether such improvements in scores are clinically meaningful for patients, particularly compared with other treatment options, remains to be clarified. An alternative approach is transplantation of the whole gut microbiota from a healthy donor: FMT. In IBD, a recent systematic review and meta-analysis has shown that of nine cohort studies, eight case studies and one randomised controlled trial, overall 45% (54/119) achieved clinical remission. When only cohort studies were analysed 36% achieved clinical remission.<sup>75</sup> Since that meta-analysis, two randomised controlled trials in UC show discrepant results. One trial, in which two faecal transplants were given via the upper GI route, showed no difference in clinical or endoscopic remission between the faecal transplant group and the control group (given autologous stool).<sup>76</sup> A second trial, in which patients with UC were randomised to weekly faecal enemas from healthy donors or placebo enemas for 6 weeks, demonstrated remission in a greater percentage of patients given FMT compared with the control group (given water enema).<sup>77</sup> There are unanswered questions regarding mode of delivery, frequency of delivery and optimal donor/host characteristics.

### Crohn's disease

Antibiotics demonstrate efficacy in particular groups of patients with CD but some antibiotics may be detrimental, showing a complex interplay between host and microbiota. Patients who have had a resection for CD have a decreased rate of endoscopic and clinical recurrence when metronidazole or ornidazole are used as prophylactic therapies.<sup>78</sup> Several studies have assessed the specific role of antimycobacterial therapies in CD treatment but overall results are disappointing. There is no clinically relevant evidence base for the use of probiotics in CD and in terms of prebiotics, although an open label trial of fructo-oligosaccharide in CD showed promise,<sup>79</sup> and a subsequent randomised placebo-controlled trial of fructo-oligosaccharide did not support any clinical benefit.<sup>80</sup>

### Pouchitis

Restorative proctocolectomy with ileal-pouch anal anastomosis is the operation of choice for patients with UC requiring surgery. Pouchitis has an incidence of up to 50% of patients although it is a significant clinical problem for only about 10%. Antibiotics are used as primary therapy; if single antibiotics fail, dual antibiotics used for longer periods of time or antibiotics tailored to the microbiota in an individual patient can be used. VSL#3 reduced the risk of disease onset and maintained an antibiotic-induced disease remission in pouchitis.<sup>81</sup> A meta-analysis has shown that VSL#3 significantly reduced the clinical relapse rates for maintaining remission in patients with pouchitis.<sup>82</sup>

### CRC AND THE GUT MICROBIOTA

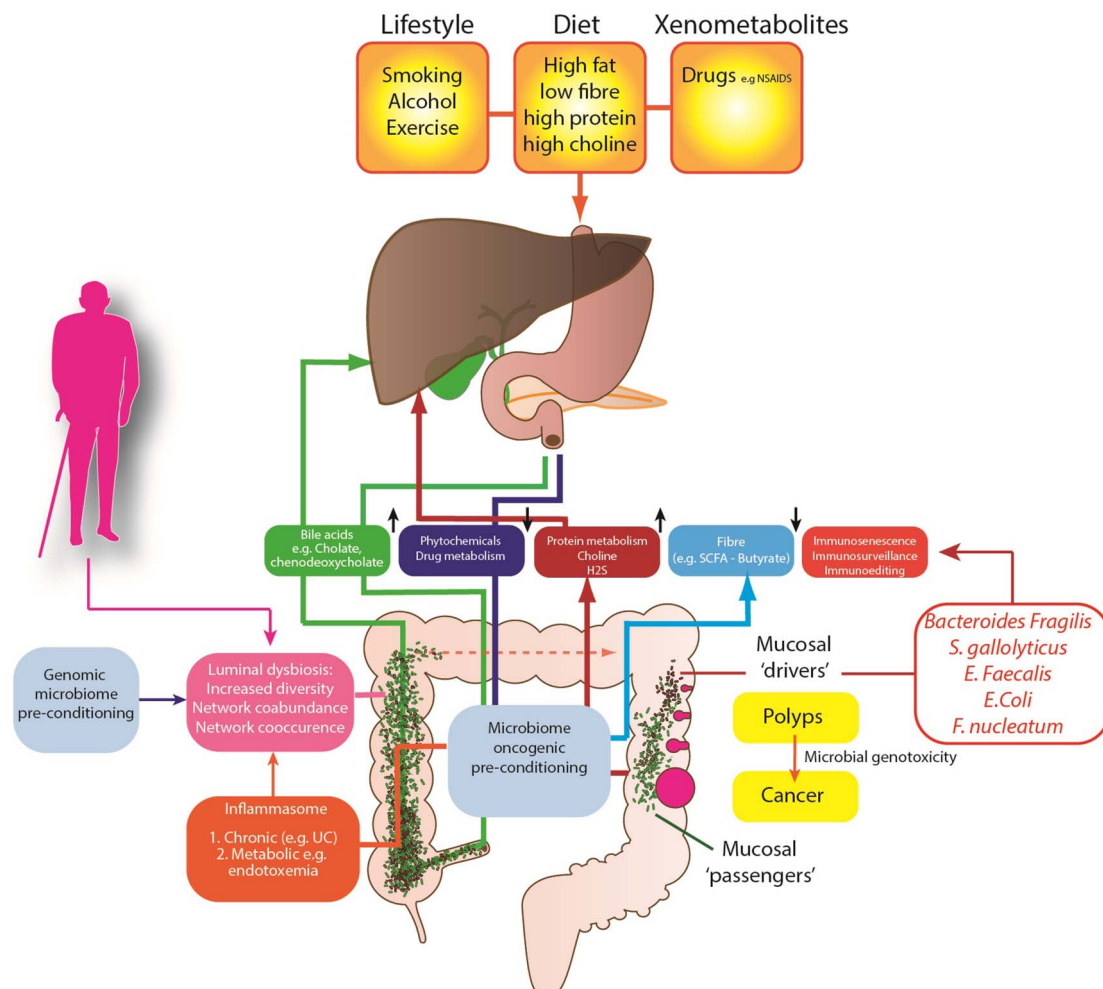
Many microbiome studies have focused on colitis-associated cancers<sup>83</sup> or rodent preclinical models.<sup>84</sup> Despite this, there is increasing evidence that the colonic microbiota plays an important role in the cause of sporadic CRC.<sup>85</sup> Reduced temporal stability and increased diversity has been shown for the faecal microbiota of subjects with established CRC and polyposis,<sup>86</sup> and now metagenomic and metatranscriptomic studies have identified an individualised oncogenic microbiome and specific

bacterial species that selectively colonise the on-tumour and off-tumour sites.<sup>87</sup>

Several competing theories of the microbial regulation of CRC have emerged (figure 1) to explain these observations. The keystone-pathogen hypothesis<sup>88</sup> and the  $\alpha$ -bug hypothesis both state that certain, low abundance microbiota members (such as enterotoxigenic *B. fragilis*) possess unique virulence traits, which are pro-oncogenic and remodel the microbiome and in turn promote mucosal immune responses and colonic epithelial cell changes.<sup>89</sup> Tjalsma *et al*<sup>90</sup> have also proposed the ‘driver-passenger’ model for CRC: a first hit by indigenous intestinal bacteria (‘bacterial drivers’), which drive the DNA damage that contributes to CRC initiation. Second, tumorigenesis induces intestinal niche alterations that favour the proliferation of opportunistic bacteria (‘bacterial passengers’). For example, CRCs have an increased enrichment of opportunistic pathogens and polymicrobial Gram-negative anaerobic bacteria<sup>91</sup> but it is not yet clear whether these opportunistic pathogens merely benefit from the CRC microenvironment or influence disease progression. However, colonic polyps demonstrate higher bacterial diversity and richness when compared with control patients, with higher abundance of mucosal Proteobacteria and lower abundance of Bacteroidetes.<sup>92</sup> This may in part be explained by the mucosal defensive strategies designed to manage the commensal microbiota. For example,  $\alpha$ -defensin expression is significantly increased in adenomas resulting in an increased antibacterial activity compared with normal mucosa.<sup>93</sup>

At present, human studies have involved small patient numbers, with evidence of sampling heterogeneity, limited tumour phenotyping and oncological data. Despite this, a small number of specific pathobionts have now been linked with adenomas and CRC including *Streptococcus gallolyticus*,<sup>94</sup> *Enterococcus faecalis*<sup>95</sup> and *B. fragilis*.<sup>84</sup> *E. coli* is also overexpressed on CRC mucosa; it expresses genes that confer properties relevant to oncological transformation including M cell translocation, angiogenesis and genotoxicity.<sup>96</sup> Enrichment of *Fusobacterium nucleatum* has also been identified in adenoma versus adjacent normal tissue and is more abundant in stools from CRC and adenoma cases than in healthy controls. *F. nucleatum*'s fadA, a unique adhesin, allows it to adhere to and invade human epithelial cells, eliciting an inflammatory response<sup>97</sup> and stimulating cell proliferation.<sup>98</sup> Novel mechanisms from previously unassociated bacteria are also being described to explain how bacterial proteins target proliferating stem-progenitor cells. For example, AvrA, a pathogenic product of *Salmonella*, has been shown to activate  $\beta$ -catenin signals and enhance colonic tumorigenesis.<sup>99</sup>

Work has also focused on the metabolic function of the gut microbiome and dietary microbiome interactions in the aetiology of CRC. It is likely that the metabolism of fibre is critical to this. Metagenomic analyses have consistently identified a reduction of butyrate-producers in patients with CRC,<sup>100</sup> a finding replicated in animals.<sup>101</sup> The microbiome also plays an important role in the metabolism of sulfate, through



**Figure 1** Proposed mechanisms of the gut microbiome in colon cancer aetiology.

assimilatory sulfate-reduction to produce cysteine and methionine, and dissimilatory sulfate-reduction to produce hydrogen sulfide (H<sub>2</sub>S). H<sub>2</sub>S is likely to contribute to CRC development, as colonic detoxification of H<sub>2</sub>S is also reduced in patients with CRC; it also induces colonic mucosal hyperproliferation.<sup>102</sup> There is also evidence that differences in host genotype, which affect the carbohydrate landscape of the distal gut, interact with diet to alter the composition and function of resident microbes in a diet-dependent manner.<sup>103</sup> Therefore it is possible that patients genetically predisposed to CRC have a modified metabolically active microbiome, which is determined by their genes and by their family environment and dietary habits. There is other evidence from global studies of cancer risk, that the microbiome is important in cancer risk.

African Americans possess a colon dominated by *Bacteroides*, while in Africans *Prevotella* are more abundant.<sup>104</sup> African Americans, who are at high risk of CRC, may have evolved a CRC-microbiota moulded by dietary habits and environmental exposures. Critically, mucosal Ki67 expression (a biomarker for cancer risk) may decrease or increase within 2 weeks of either a high fibre (>50 g/day) dietary intervention in African Americans or a high fat, high protein low fibre Westernised diet in African subjects. This short-term intervention leads to reciprocal changes in luminal microbiome co-occurrence network structures that overwhelm interindividual differences in microbial gene expression. Specifically, an animal-based diet increases the abundance of bile-tolerant microorganisms (*Alistipes*, *Bilophila* and *Bacteroides*) and decreases the levels of Firmicutes that metabolise dietary plant polysaccharides (*Roseburia*, *Eubacterium rectale* and *Ruminococcus bromii*).<sup>105 106</sup>

## CONCLUSIONS

In the past decade, interest in the human microbiome has increased considerably. A significant driver has been the realisation that the commensal microorganisms that comprise the human microbiota are not simply passengers in the host, but

may actually drive certain host functions as well. In sterile rodents, we see the dramatic impact that removing the microbiota has on nearly all aspects of the host's ability to function normally. This review highlights some key disease areas in which the microbiota and its microbiome are thought to have not just an association, but also a key modulatory role (table 1). By better understanding the mechanisms and contribution the microbiota make to these diseases, we hope to develop novel therapeutics and strategies to modulate the microbiota to treat or prevent disease. Additionally, in some instances it may be possible to use the microbiome to detect gut-related diseases before conventional diagnostics can. In the future we hope to use this information to stratify patients more accurately and for more efficient treatment. A body of evidence also points to the gut microbiota being an environmental factor in drug metabolism, for example, inactivation of the cardiac drug digoxin by *Eggerthella lenta* in the gut. Thus, if we are to realise the vision of a personalised healthcare revolution, we must explore how the microbiome fits with this notion.

### Author affiliations

<sup>1</sup>School of Biosciences, Museum Avenue, Cardiff University, Cardiff, UK

<sup>2</sup>Centre for Digestive and Gut Health, Imperial College London, London, UK

<sup>3</sup>NIHR Biomedical Research Unit, Centre for Liver Research, University of Birmingham, Birmingham, UK

<sup>4</sup>Nutrition and Nutrigenomics Group, Department of Food Quality and Nutrition, Research and Innovation Centre, Trento, Italy

<sup>5</sup>Laboratory of Microbiology, Wageningen University, Wageningen, The Netherlands

<sup>6</sup>Top Institute Food and Nutrition (TIFN), Wageningen, The Netherlands

<sup>7</sup>Division of Applied Medicine, School of Medicine and Dentistry, University of Aberdeen, Institute of Medical Sciences, Aberdeen, UK

<sup>8</sup>Section of Computational and Systems Medicine, Faculty of Medicine, Imperial College London, London, UK

<sup>9</sup>Yakult UK Limited, Middlesex, UK

<sup>10</sup>IBD Unit, St Mark's Hospital and Imperial College London, London, UK

**Acknowledgements** This review was commissioned by the Gut Microbiota for Health expert panel of the British Society of Gastroenterology.

**Contributors** AH, JRM, DHA, GDAH, LVT, FF, GMH and GH, MNQ, HS, KMT, EGZ and JK contributed to the conception/design of the work, drafting the work and revising it critically for important intellectual content and final approval of the version published.

**Competing interests** AH has lectured for Yakult.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Open Access** This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

## REFERENCES

- Muegge BD, Kuczynski J, Knights D, *et al*. Diet drives convergence in gut microbiome functions across mammalian phylogeny and within humans. *Science* 2011;332:970–4.
- Segata N, Haake SK, Mannon P, *et al*. Composition of the adult digestive tract bacterial microbiome based on seven mouth surfaces, tonsils, throat and stool samples. *Genome Biol* 2012;13:R42.
- Zoetendal EG, Raes J, van den Bogert B, *et al*. The human small intestinal microbiota is driven by rapid uptake and conversion of simple carbohydrates. *ISME J* 2012;6:1415–26.
- Lepage P, Leclerc MC, Joossens M, *et al*. A metagenomic insight into our gut's microbiome. *Gut* 2013;62:146–58.
- Sokol H, Seksik P, Furet JP, *et al*. Low counts of *Faecalibacterium prausnitzii* in colitis microbiota. *Inflamm Bowel Dis* 2009;15:1183–9.
- Gerasimidis K, Bertz M, Hanske L, *et al*. Decline in presumptively protective gut bacterial species and metabolites are paradoxically associated with disease improvement in pediatric Crohn's disease during enteral nutrition. *Inflamm Bowel Dis* 2014;20:861–71.
- Bonnet M, Buc E, Sauvanet P, *et al*. Colonization of the human gut by *E. coli* and colorectal cancer risk. *Clin Cancer Res* 2014;20:859–67.

**Table 1** Key insights into the influence of the gut microbiota on GI and liver diseases

Focus	Insights
The gut microbiota	<ul style="list-style-type: none"> <li>▶ The gut microbiota is host-specific and variable</li> <li>▶ Loss of diversity has a negative impact on health; conservation of key microbial functions is even more important</li> </ul>
Dietary modulation	<ul style="list-style-type: none"> <li>▶ Indigestible carbohydrates are the 'food' of the gut microbiota</li> <li>▶ Probiotics, prebiotics and polyphenols can promote gut health via the microbiota</li> </ul>
Obesity	<ul style="list-style-type: none"> <li>▶ The gut microbiome is an environmental factor in obesity</li> <li>▶ Unknown functions in the microbiome can be transferred and recapitulate or treat obesity and its associated metabolic disorders</li> </ul>
Liver disease	<ul style="list-style-type: none"> <li>▶ Bacterial dysbiosis can drive hepatitis</li> <li>▶ Bacterial products can cause inflammation in the liver</li> </ul>
IBD	<ul style="list-style-type: none"> <li>▶ Alterations of the microbiota lie at the core of IBD pathogenesis; these may be driven by host genetics and/or environmental factors</li> <li>▶ Targeting the microbiota remains an attractive option to treating the disease cause</li> </ul>
Colorectal cancer	<ul style="list-style-type: none"> <li>▶ Pathobionts such as <i>Fusobacterium nucleatum</i> are overexpressed in adenomas and cancers of the colon.</li> <li>▶ 'High-risk' (high fat, high protein) diets modulate this risk through gut microbiome co-metabolic processes.</li> <li>▶ The gut microbiota may drive the first DNA damage either via specific proteins or metabolites</li> <li>▶ Diet can play a large role in shaping the composition of the microbiota and it thus affects risk of developing the disease</li> </ul>



- 8 Thorkildsen LT, Nwosu FC, Avershina E, *et al.* Dominant fecal microbiota in newly diagnosed untreated inflammatory bowel disease patients. *Gastroenterol Res Pract* 2013;2013:636785.
- 9 Rajilic-Stojanovic M, de Vos WM. The first 1000 cultured species of the human gastrointestinal microbiota. *FEMS Microbiol Rev* 2014;38:996–1047.
- 10 Louis P, Young P, Holtrop G, *et al.* Diversity of human colonic butyrate-producing bacteria revealed by analysis of the butyryl-CoA:acetate CoA-transferase gene. *Environ Microbiol* 2010;12:304–14.
- 11 Flint HJ, Scott KP, Louis P, *et al.* The role of the gut microbiota in nutrition and health. *Nature Rev Gastroenterol Hepatol* 2012;9:577–89.
- 12 Ze X, Le Mougou F, Duncan SH, *et al.* Some are more equal than others: the role of “keystone” species in the degradation of recalcitrant substrates. *Gut Microbes* 2013;4:236–40.
- 13 Macfarlane GT, Macfarlane S. Fermentation in the human large intestine: its physiologic consequences and the potential contribution of prebiotics. *J Clin Gastroenterol* 2011;45(Suppl):S120–7.
- 14 Arpaia N, Campbell C, Fan X, *et al.* Metabolites produced by commensal bacteria promote peripheral regulatory T-cell generation. *Nature* 2013;504:451–5.
- 15 Windey K, De Preter V, Verbeke K. Relevance of protein fermentation to gut health. *Mol Nutr Food Res* 2012;56:184–96.
- 16 Tang WH, Wang Z, Levison BS, *et al.* Intestinal microbial metabolism of phosphatidylcholine and cardiovascular risk. *New Engl J Med* 2013;368:1575–84.
- 17 Clarke G, Stilling RM, Kennedy PJ, *et al.* Mini-review: gut microbiota: the neglected endocrine organ. *Molec Endocrinol* 2014;28:1221–38.
- 18 Tuohy K, Del Rio D, eds. *Diet-microbe interactions in the gut. Effects on human health and disease.* Elsevier Science Publishing Co. Inc., 2014.
- 19 Jones ML, Martoni CJ, Prakash S. Cholesterol lowering and inhibition of sterol absorption by *Lactobacillus reuteri* NCIMB 30242: a randomized controlled trial. *Eur J Clin Nutr* 2012;66:1234–41.
- 20 Li F, Pang X, Krausz KW, *et al.* Stable isotope- and mass spectrometry-based metabolomics as tools in drug metabolism: a study expanding temporal pharmacology. *J Proteome Res* 2013;12:1369–76.
- 21 Clifford MN. Diet-derived phenols in plasma and tissues and their implications for health. *Planta Medica* 2004;70:1103–14.
- 22 Queipo-Ortuno MI, Boto-Ordóñez M, Murri M, *et al.* Influence of red wine polyphenols and ethanol on the gut microbiota ecology and biochemical biomarkers. *Am J Clin Nutr* 2012;95:1323–34.
- 23 Ley RE, Backhed F, Turnbaugh P, *et al.* Obesity alters gut microbial ecology. *Proc Natl Acad Sci USA* 2005;102:11070–5.
- 24 Ley RE, Turnbaugh PJ, Klein S, *et al.* Microbial ecology: human gut microbes associated with obesity. *Nature* 2006;444:1022–3.
- 25 Duncan SH, Lohley GE, Holtrop G, *et al.* Human colonic microbiota associated with diet, obesity and weight loss. *Int J Obes* 2008;32:1720–4.
- 26 Claesson MJ, Jeffery IB, Conde S, *et al.* Gut microbiota composition correlates with diet and health in the elderly. *Nature* 2012;488:178–84.
- 27 Romero-Corral A, Somers VK, Sierra-Johnson J, *et al.* Accuracy of body mass index in diagnosing obesity in the adult general population. *Int J Obesity* 2008;32:959–66.
- 28 Zupancic ML, Cantarel BL, Liu Z, *et al.* Analysis of the gut microbiota in the old order Amish and its relation to the metabolic syndrome. *PLoS ONE* 2012;7:e43052.
- 29 Blüher M. The distinction of metabolically ‘healthy’ from ‘unhealthy’ obese individuals. *Curr Opin Lipidol* 2010;21:38–43.
- 30 Qin J, Li Y, Cai Z, *et al.* A metagenome-wide association study of gut microbiota in type 2 diabetes. *Nature* 2012;490:55–60.
- 31 Karlsson FH, Tremaroli V, Nookaew I, *et al.* Gut metagenome in European women with normal, impaired and diabetic glucose control. *Nature* 2013;498:99–103.
- 32 Le Chatelier E, Nielsen T, Qin J, *et al.* Richness of human gut microbiome correlates with metabolic markers. *Nature* 2013;500:541–6.
- 33 Vriee A, Van Nood E, Holleman F, *et al.* Transfer of intestinal microbiota from lean donors increases insulin sensitivity in individuals with metabolic syndrome. *Gastroenterol* 2012;143:913–6.e7.
- 34 Cottillard A, Kennedy SP, Kong LC, *et al.* Dietary intervention impact on gut microbial gene richness. *Nature* 2013;500:585–8.
- 35 Hermes GD, Zoetendal EG, Smidt H. Molecular ecological tools to decipher the role of our microbial mass in obesity. *Benef Microbes* 2015;6:61–81.
- 36 Adams DH, Eksteen B, Curbishley SM. Immunology of the gut and liver: a love/hate relationship. *Gut* 2008;57:838–48.
- 37 Son G, Kremer M, Hines IN. Contribution of gut bacteria to liver pathobiology. *Gastroenterol Res Pract* 2010;2010:1–13.
- 38 de Aguiar Vallim TQ, Tarling EJ, Edwards PA. Pleiotropic roles of bile acids in metabolism. *Cell Metab* 2013;17:657–69.
- 39 Miele L, Valenza V, La Torre G, *et al.* Increased intestinal permeability and tight junction alterations in nonalcoholic fatty liver disease. *Hepatology* 2009;49:1877–87.
- 40 Drenick EJ, Fislis J, Johnson D. Hepatic steatosis after intestinal bypass—prevention and reversal by metronidazole, irrespective of protein-calorie malnutrition. *Gastroenterol* 1982;82:535–48.
- 41 Henao-Mejia J, Elinav E, Jin C, *et al.* Inflammation-mediated dysbiosis regulates progression of NAFLD and obesity. *Nature* 2012;482:179–85.
- 42 Raman M, Ahmed I, Gillevet PM, *et al.* Fecal microbiome and volatile organic compound metabolome in obese humans with nonalcoholic fatty liver disease. *Clin Gastroenterol Hepatol* 2013;11:868–75.e1–3.
- 43 Zhu L, Baker SS, Gill C, *et al.* Characterization of gut microbiomes in nonalcoholic steatohepatitis (NASH) patients: a connection between endogenous alcohol and NASH. *Hepatology* 2013;57:601–9.
- 44 Nanji AA, Khettry U, Sadrzadeh SM, *et al.* Severity of liver injury in experimental alcoholic liver disease. Correlation with plasma endotoxin, prostaglandin E2, leukotriene B4, and thromboxane B2. *Am J Pathol* 1993;142:367–73.
- 45 Pande C, Kumar A, Sarin SK. Small-intestinal bacterial overgrowth in cirrhosis is related to the severity of liver disease. *Aliment Pharmacol Ther* 2009;29:1273–81.
- 46 Yan AW, Fouts DE, Brandl J, *et al.* Enteric dysbiosis associated with a mouse model of alcoholic liver disease. *Hepatology* 2011;53:96–105.
- 47 Yan AW, Schnabl B. Bacterial translocation and changes in the intestinal microbiome associated with alcoholic liver disease. *World J Hepatol* 2012;4:110–18.
- 48 Quraishi SM, Adams D, Pallen M, *et al.* Probing the microbiota in PSC: the gut adherent microbiota of PSC-IBD is distinct to that of IBD and controls. *Hepatology* 2014;60:267A.
- 49 Rossen NG, Fuentes S, Boonstra K, *et al.* The mucosa-associated microbiota of PSC patients is characterized by low diversity and low abundance of uncultured Clostridiales II. *J Crohn's Colitis* 2015;9:342–8.
- 50 Tabibian JH, Talwalkar JA, Lindor KD. Role of the microbiota and antibiotics in primary sclerosing cholangitis. *BioMed Res Int* 2013;2013:389537.
- 51 Davies YK, Cox KM, Abdullah BA, *et al.* Long-term treatment of primary sclerosing cholangitis in children with oral vancomycin: an immunomodulating antibiotic. *J Pediatr Gastroenterol Nutr* 2008;47:61–7.
- 52 Björnsson E, Cederborg A, Akvist A, *et al.* Intestinal permeability and bacterial growth of the small bowel in patients with primary sclerosing cholangitis. *Scand J Gastroenterol* 2005;40:1090–4.
- 53 Kono K, Ohnishi K, Omata M, *et al.* Experimental portal fibrosis produced by intraportal injection of killed nonpathogenic *Escherichia coli* in rabbits. *Gastroenterol* 1988;94:787–96.
- 54 Mattner J, Savage PB, Leung P, *et al.* Liver autoimmunity triggered by microbial activation of natural killer T cells. *Cell Host Microbe* 2008;3:304–15.
- 55 Lata J, Jurankova J, Kopacova M, *et al.* Probiotics in hepatology. *World J Gastroenterol* 2011;17:2890–6.
- 56 Ma YY, Li L, Yu CH, *et al.* Effects of probiotics on nonalcoholic fatty liver disease: a meta-analysis. *World J Gastroenterol* 2013;19:6911–18.
- 57 Rincon D, Vaquero J, Hernando A, *et al.* Oral probiotic VSL#3 attenuates the circulatory disturbances of patients with cirrhosis and ascites. *Liver Int* 2014;34:1504–12.
- 58 Le Roy T, Llopis M, Lepage P, *et al.* Intestinal microbiota determines development of non-alcoholic fatty liver disease in mice. *Gut* 2013;62:1787–94.
- 59 Hold GL, Smith M, Grange C, *et al.* Role of the gut microbiota in inflammatory bowel disease pathogenesis: what have we learnt in the past 10 years? *World J Gastroenterol* 2014;20:1192–210.
- 60 Peterson DA, Frank DN, Pace NR, *et al.* Metagenomic approaches for defining the pathogenesis of inflammatory bowel diseases. *Cell Host Microbe* 2008;3:417–27.
- 61 Walker AW, Sanderson JD, Churcher C, *et al.* High-throughput clone library analysis of the mucosa-associated microbiota reveals dysbiosis and differences between inflamed and non-inflamed regions of the intestine in inflammatory bowel disease. *BMC Microbiol* 2011;11:7.
- 62 Frank DN, St Amand AL, Feldman RA, *et al.* Molecular-phylogenetic characterization of microbial community imbalances in human inflammatory bowel diseases. *Proc Natl Acad Sci USA* 2007;104:13780–5.
- 63 Swidsinski A, Weber J, Loening-Baucke V, *et al.* Spatial organization and composition of the mucosal flora in patients with inflammatory bowel disease. *J Clin Microbiol* 2005;43:3380–9.
- 64 Hansen R, Russell RK, Reiff C, *et al.* Microbiota of *de-novo* pediatric IBD: increased *Faecalibacterium prausnitzii* and reduced bacterial diversity in Crohn's but not in ulcerative colitis. *Am J Gastroenterol* 2012;107:1913–22.
- 65 Manichanh C, Rigottier-Gois L, Bonnaud E, *et al.* Reduced diversity of faecal microbiota in Crohn's disease revealed by a metagenomic approach. *Gut* 2006;55:205–11.
- 66 Mukhopadhyay I, Thomson JM, Hansen R, *et al.* Detection of *Campylobacter concisus* and other *Campylobacter* species in colonic biopsies from adults with ulcerative colitis. *PLoS ONE* 2011;6:e21490.
- 67 Lupp C, Robertson ML, Wickham ME, *et al.* Host-mediated inflammation disrupts the intestinal microbiota and promotes the overgrowth of Enterobacteriaceae. *Cell Host Microbe* 2007;2:204.
- 68 Barnich N, Denizot J, Darfeuille-Michaud A. E. coli-mediated gut inflammation in genetically predisposed Crohn's disease patients. *Pathol Biol (Paris)* 2013;61:e65–9.

- 69 Erickson AR, Cantarel BL, Lamendella R, *et al.* Integrated metagenomics/metaproteomics reveals human host-microbiota signatures of Crohn's disease. *PLoS ONE* 2012;7:e49138.
- 70 Morgan XC, Tickle TL, Sokol H, *et al.* Dysfunction of the intestinal microbiome in inflammatory bowel disease and treatment. *Genome Biol* 2012;13:R79.
- 71 Norman JM, Handley SA, Baldrige MT, *et al.* Disease-specific alterations in the enteric virome in inflammatory bowel disease. *Cell* 2015;160:447–60.
- 72 Kruis W, Fris P, Pokrotnieks J, *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.
- 73 Dignass A, Lindsay JO, Sturm A, *et al.* Second European evidence-based consensus on the diagnosis and management of ulcerative colitis part 2: current management. *J Crohn's Colitis* 2012;6:991–1030.
- 74 Tursi A, Brandimarte G, Papa A, *et al.* Treatment of relapsing mild-to-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.
- 75 Colman RJ, Rubin DT. Fecal microbiota transplantation as therapy for inflammatory bowel disease: a systematic review and meta-analysis. *J Crohn's Colitis* 2014;8:1569–81.
- 76 Rossen NG, Fuentes S, van der Spek MJ, *et al.* Findings from a randomized controlled trial of fecal transplantation for patients with ulcerative colitis. *Gastroenterology* 2015;149:110–18.e4.
- 77 Moayyedi P, Surette MG, Kim PT, *et al.* Fecal microbiota transplantation induces remission in patients with active ulcerative colitis in a randomized, controlled trial. *Gastroenterology* 2015;149:102–9.e6.
- 78 Rutgeerts P, Hiele M, Geboes K, *et al.* Controlled trial of metronidazole treatment for prevention of Crohn's recurrence after ileal resection. *Gastroenterol* 1995;108:1617–21.
- 79 Lindsay JO, Whelan K, Stagg AJ, *et al.* Clinical, microbiological, and immunological effects of fructo-oligosaccharide in patients with Crohn's disease. *Gut* 2006;55:348–55.
- 80 Benjamin JL, Hedin CR, Koutsoumpas A, *et al.* Randomised, double-blind, placebo-controlled trial of fructo-oligosaccharides in active Crohn's disease. *Gut* 2011;60:923–9.
- 81 Gionchetti P, Rizzello F, Helwig U, *et al.* Prophylaxis of pouchitis onset with probiotic therapy: a double-blind, placebo-controlled trial. *Gastroenterol* 2003;124:1202–9.
- 82 Shen J, Zuo ZX, Mao AP. Effect of probiotics on inducing remission and maintaining therapy in ulcerative colitis, Crohn's disease, and pouchitis: meta-analysis of randomized controlled trials. *Inflamm Bowel Dis* 2014;20:21–35.
- 83 Arthur JC, Perez-Chanona E, Muhlbauer M, *et al.* Intestinal inflammation targets cancer-inducing activity of the microbiota. *Science* 2012;338:120–3.
- 84 Wu S, Rhee KJ, Albesiano E, *et al.* A human colonic commensal promotes colon tumorigenesis via activation of T helper type 17T cell responses. *Nat Med* 2009;15:1016–22.
- 85 Ou J, Carbonero F, Zoetendal EG, *et al.* Diet, microbiota, and microbial metabolites in colon cancer risk in rural Africans and African Americans. *Am J Clin Nutr* 2013;98:111–20.
- 86 Scanlan PD, Shanahan F, Clune Y, *et al.* Culture-independent analysis of the gut microbiota in colorectal cancer and polyposis. *Environ Microbiol* 2008;10:789–98.
- 87 Marchesi JR, Dutilh BE, Hall N, *et al.* Towards the human colorectal cancer microbiome. *PLoS ONE* 2011;6:e20447.
- 88 Hajishengallis G, Darveau RP, Curtis MA. The keystone-pathogen hypothesis. *Nature Rev Microbiol* 2012;10:717–25.
- 89 Sears CL, Pardoll DM. Perspective: alpha-bugs, their microbial partners, and the link to colon cancer. *J Infect Dis* 2011;203:306–11.
- 90 Tjalsma H, Boleij A, Marchesi JR, *et al.* A bacterial driver-passenger model for colorectal cancer: beyond the usual suspects. *Nature Rev Microbiol* 2012;10:575–82.
- 91 Warren RL, Freeman DJ, Pleasance S, *et al.* Co-occurrence of anaerobic bacteria in colorectal carcinomas. *Microbiome* 2013;1:16.
- 92 Shen XJ, Rawls JF, Randall T, *et al.* Molecular characterization of mucosal adherent bacteria and associations with colorectal adenomas. *Gut Microbes* 2010;1:138–47.
- 93 Pagnini C, Corleto VD, Mangoni ML, *et al.* Alteration of local microflora and alpha-defensins hyper-production in colonic adenoma mucosa. *J Clin Gastroenterol* 2011;45:602–10.
- 94 Boleij A, Tjalsma H. The itinerary of *Streptococcus gallolyticus* infection in patients with colonic malignant disease. *Lancet Infect Dis* 2013;13:719–24.
- 95 Wang X, Yang Y, Huycke MM. Commensal bacteria drive endogenous transformation and tumour stem cell marker expression through a bystander effect. *Gut* 2015;64:459–68.
- 96 Prorok-Hamon M, Friswell MK, Alswied A, *et al.* Colonic mucosa-associated diffusely adherent afaC+ *Escherichia coli* expressing IpfA and pks are increased in inflammatory bowel disease and colon cancer. *Gut* 2014;63:761–70.
- 97 Han YW, Shi W, Huang GT, *et al.* Interactions between periodontal bacteria and human oral epithelial cells: *Fusobacterium nucleatum* adheres to and invades epithelial cells. *Infect Immun* 2000;68:3140–6.
- 98 Rubinstein MR, Wang X, Liu W, *et al.* *Fusobacterium nucleatum* promotes colorectal carcinogenesis by modulating E-cadherin/beta-catenin signaling via its FadA adhesin. *Cell Host Microbe* 2013;14:195–206.
- 99 Lu R, Wu S, Zhang YG, *et al.* Enteric bacterial protein AvrA promotes colonic tumorigenesis and activates colonic beta-catenin signaling pathway. *Oncogenesis* 2014;3:e105.
- 100 Wu N, Yang X, Zhang R, *et al.* Dysbiosis signature of fecal microbiota in colorectal cancer patients. *Microb Ecol* 2013;66:462–70.
- 101 Zhu Q, Jin Z, Wu W, *et al.* Analysis of the intestinal lumen microbiota in an animal model of colorectal cancer. *PLoS ONE* 2014;9:e90849.
- 102 Carbonero F, Benefiel AC, Alizadeh-Ghamsari AH, *et al.* Microbial pathways in colonic sulfur metabolism and links with health and disease. *Front Physiol* 2012;3:448.
- 103 Kashyap PC, Marcobal A, Ursell LK, *et al.* Genetically dictated change in host mucus carbohydrate landscape exerts a diet-dependent effect on the gut microbiota. *Proc Natl Acad Sci USA* 2013;110:17059–64.
- 104 Wu GD, Chen J, Hoffmann C, *et al.* Linking long-term dietary patterns with gut microbial enterotypes. *Science* 2011;334:105–8.
- 105 David LA, Maurice CF, Carmody RN, *et al.* Diet rapidly and reproducibly alters the human gut microbiome. *Nature* 2014;505:559–63.
- 106 O'Keefe SJ, Li JV, Lahti L, *et al.* Fat, fibre and cancer risk in African Americans and rural Africans. *Nature Commun* 2015;6:6342.
- 107 World Health Organization, Food and Agricultural Organization of the United Nations. *Health and nutritional properties of probiotics in food including powder milk with live lactic acid bacteria*. FAO Nutrition Paper, 2001;85.
- 108 Hill C, Guarner F, Reid G, *et al.* Expert consensus document. The International Scientific Association for Probiotics and Prebiotics consensus statement on the scope and appropriate use of the term probiotic. *Nat Rev Gastroenterol Hepatol* 2014;11:506–14.
- 109 Ng SC, Hart AL, Kamm MA, *et al.* Mechanisms of action of probiotics: recent advances. *Inflamm Bowel Dis* 2009;15:300–10.
- 110 Gibson GR, Roberfroid MB. Dietary modulation of the human colonic microbiota: introducing the concept of prebiotics. *J Nutr* 1995;125:1401–12.
- 111 NICE (National Institute for Health and Care Excellence). Faecal microbiota transplant for recurrent *Clostridium difficile* infection. *NICE Interventional Procedure Guidance* 2014:485.