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ORIGINAL ARTICLE

International Cancer Microbiome Consortium consensus statement on the role of the human microbiome in carcinogenesis

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ABSTRACT

Objective In this consensus statement, an international panel of experts deliver their opinions on key questions regarding the contribution of the human microbiome to carcinogenesis.

Design International experts in oncology and/or microbiome research were approached by personal communication to form a panel. A structured, iterative, methodology based around a 1-day roundtable discussion was employed to derive expert consensus on key questions in microbiome-oncology research.

Results Some 18 experts convened for the roundtable discussion and five key questions were identified regarding: (1) the relevance of dysbiosis/an altered gut microbiome to carcinogenesis; (2) potential mechanisms of microbiota-induced carcinogenesis; (3) conceptual frameworks describing how the human microbiome may drive carcinogenesis; (4) causation versus association; and (5) future directions for research in the field. The panel considered that, despite mechanistic and supporting evidence from animal and human studies, there is currently no direct evidence that the human commensal microbiome is a key determinant in the aetiopathogenesis of cancer. The panel cited the lack of large longitudinal, cohort studies as a principal deciding factor and agreed that this should be a future research priority. However, while acknowledging gaps in the evidence, expert opinion was that the microbiome, alongside environmental factors and an epigenetically/genetically vulnerable host, represents one apex of a tripartite, multidirectional interactome that drives carcinogenesis.

Conclusion Data from longitudinal cohort studies are needed to confirm the role of the human microbiome as a key driver in the aetiopathogenesis of cancer.

INTRODUCTION

The last decade has seen a surge in the number of studies of the gut microbiome and cancer, facilitated by next-generation sequencing platforms, 'omics' technologies and advanced bioinformatics approaches. In contrast to the human genome which is inherited, largely static and which consists of approximately 20000 genes, the human microbiome is acquired, changes dynamically throughout life and contains approximately 10 to 20 million non-redundant genes.¹

Significance of this study

What is already known on this subject?

- Putative pathogens such as human papilloma virus and *Helicobacter pylori* have been demonstrated to be aetiological agents in a variety of cancers.
- Humans possess anatomic niche-specific microbiomes which are involved in key aspects of host physiology and pathology.
- Data from in vitro, animal and human studies suggest that the human microbiome may be involved in the aetiopathogenesis of cancer.

What are the new findings?

- This is the first published expert consensus on the role of the microbiome in cancer.
- Experts considered that there is currently no direct evidence that the human commensal microbiome is a key determinant in the aetiopathogenesis of cancer.
- However, expert opinion was that the microbiome is one apex of a tripartite, multidirectional interactome alongside environmental factors and an epigenetically/genetically vulnerable host that combine to cause cancer.

How might it impact on clinical practice in the foreseeable future?

- Clinicians should appreciate the current status of microbiome research with regards to the aetiology of cancer as it seems likely that manipulation of the microbiome may offer a therapeutic strategy for the prevention or treatment of cancer.

The human microbiota was once viewed as comprising either pathogens, causing infectious disease, or passive bystanders. In appreciating the beneficial functions of commensal microorganisms, we have departed from this dichotomy to embrace a novel, subtler paradigm in which host microorganisms contribute to both physiology and pathology. Commensal microorganisms, by definition, do not harm their host. Similarly, pathogens are defined by their ability to cause disease.



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Key Terms	
Microbiota	All the organisms in a specified niche
Microbiome	The combined genetic material from all organisms in a specified niche
Metagenome	All the genetic material present in a biological sample
16S rRNA Taxonomic Profiling	Specific amplification and sequencing of a hypervariable section of the gene encoding the 16S ribosomal RNA subunit. Sequence conservation in this region varies according to taxonomy and can be used as a "molecular fingerprint" to infer relative abundance of organisms in a sample.
Metabolome	The complete set of metabolites found in a sample
Symbiont	An organism living closely with another. Symbiotic relationships can be: <ul style="list-style-type: none"> Mutualistic: both organisms benefit Commensalistic: one organism benefits and the other is not harmed or helped Parasitic: on organism benefits by harming another Amensalistic: one organism harms another without benefitting
Pathogen	A microorganism that can cause disease

Figure 1 Key terms in the microbiome literature.

However, we now appreciate that there is a large and poorly defined middle ground occupied by 'pathogens' coexisting with a disease free-host (eg, *Helicobacter pylori*) and the so-called 'commensals' that may contribute to disease under certain conditions. This article is principally concerned with the latter cohort, which may be considered *dynamic symbionts*—a group of organisms whose symbiotic nature may vary along a spectrum from mutualism and commensalism to parasitism and amensalism. For simplicity, we will refer to this group of dynamic symbionts as the *human microbiome* (figure 1).

Genetic and environmental factors do not fully explain individual cancer risk—why do some (largely genetically identical) individuals who have similar lifestyles and ages get cancer while others do not? The random distribution of DNA replicative errors is part of the explanation,² but the significant interindividual variation in microbiome composition and function, coupled with known and emerging potentially carcinogenic microbial-host interactions, suggest that the human microbiome may also be an important cofactor driving cancer risk. Evidence supporting this hypothesis has begun to accumulate and, furthermore, the human microbiome appears to have importance in relation to cancer prognosis³ and treatment.⁴

The International Cancer Microbiome Consortium (ICMC) was established in 2017 in recognition of the emerging significance of the human microbiome in oncology. The ICMC comprises academic and clinical experts in oncology and the microbiome and its aim is to promote education and international research collaboration. In its inaugural meeting, ICMC hosted a 1-day roundtable discussion to draw together current research and establish an expert consensus regarding the role of the human microbiome in carcinogenesis.⁵ Key topics (figure 2) were identified and debated before generating the written document, which was subsequently revised by experts to produce the final consensus.

METHODS

The aim of this consensus statement was to perform a qualitative assessment of the current evidence relevant to the hypothesis that the human microbiome plays a causative role in the aetiopathogenesis of cancer. Discussion and conclusions were weighted towards bacteria (as opposed to viruses or fungi) because bacteria accounted for most of the available data. Discussion was not restricted to the gut and all cancer types were considered, though again, the gut microbiome and colorectal cancer (CRC) feature prominently because of biases within the literature. Given that our aim

was to convey current expert opinion on a novel area of science, as opposed to a specific clinical question, a structured, iterative, 'pseudo-Delphi' methodology was employed rather than a formal Delphi methodology. This structured approach is summarised in figure 2. The core participants (AJS, JLA, CAM, JMK and JRM) identified contributors based on their expertise in the microbiome, surgical and medical oncology (upper and lower GI cancer, urological cancers and oncogynaecology) and bioinformatics and invited them to contribute to the consensus by personal communication. Some 18 experts from Canada, China, Europe and the USA agreed to form the expert panel. In stage 1, panel members agreed, by email, on six key topics for discussion centred on the theme of the microbiome and carcinogenesis. In stage 2, panel members were assigned to each of these topics according to their expertise and led the debate and discussion on each topic during a 1-day face-to-face roundtable discussion which was recorded and minuted. In stage 3, the core participants reviewed the discussion and distilled it into five key questions to be addressed in the consensus statement. These key questions reflect the roundtable discussion rather than the *a priori* identified topics directly and were chosen because of their perceived interest to the academic and clinical community and their depth of discussion during the face-to-face meeting:

1. How does the concept of 'dysbiosis' relate to carcinogenesis?
2. What are the broad molecular mechanisms by which the human microbiome may be involved in the aetiopathogenesis of cancer?
3. What are the conceptual frameworks that best describe the promotion of carcinogenesis by the human microbiome?
4. Is the relationship between the human microbiome and the aetiopathogenesis of cancer causative or associative?
5. What are the key future directions for research in this field?

Extensive quotes from the discussion are available (see supplement) while selected quotes are included in figures alongside the discussion pertaining to each key question (figures 3–7). In stage 4, the core participants drew on the roundtable debate and suggested literature to draft statements and supporting discussion in response to each key question. Statements were distributed electronically to the expert panel who rated the strength of evidentiary support and their personal level of agreement with each statement according to Likert scales (figure 2, stage 5). Some 16 (out of the 18 present at the panel discussion) experts responded and statements were assigned the modal response (number given in brackets, see supplement for full responses). Panel members also had the opportunity to revise the content of the consensus statement. During stage 6, the expert panel reviewed and revised the manuscript to arrive at consensus over multiple iterations.

RESULTS AND DISCUSSION

Question one: How does the concept of dysbiosis relate to carcinogenesis (table 1)?

Table 1 How does the concept of dysbiosis relate to carcinogenesis?

Statement	Evidentiary support	Expert agreement
With respect to carcinogenesis, 'dysbiosis' should be considered a persistent departure of the host microbiome from the health-associated, homeostatic state, towards a cancer promoting and/or sustaining phenotype.	Weak evidence from human studies (9)	Agree (10)
At the present time, there is no accepted quantitative definition of a 'normal' microbiome.	Strong evidence from human studies (7)	Agree (8)

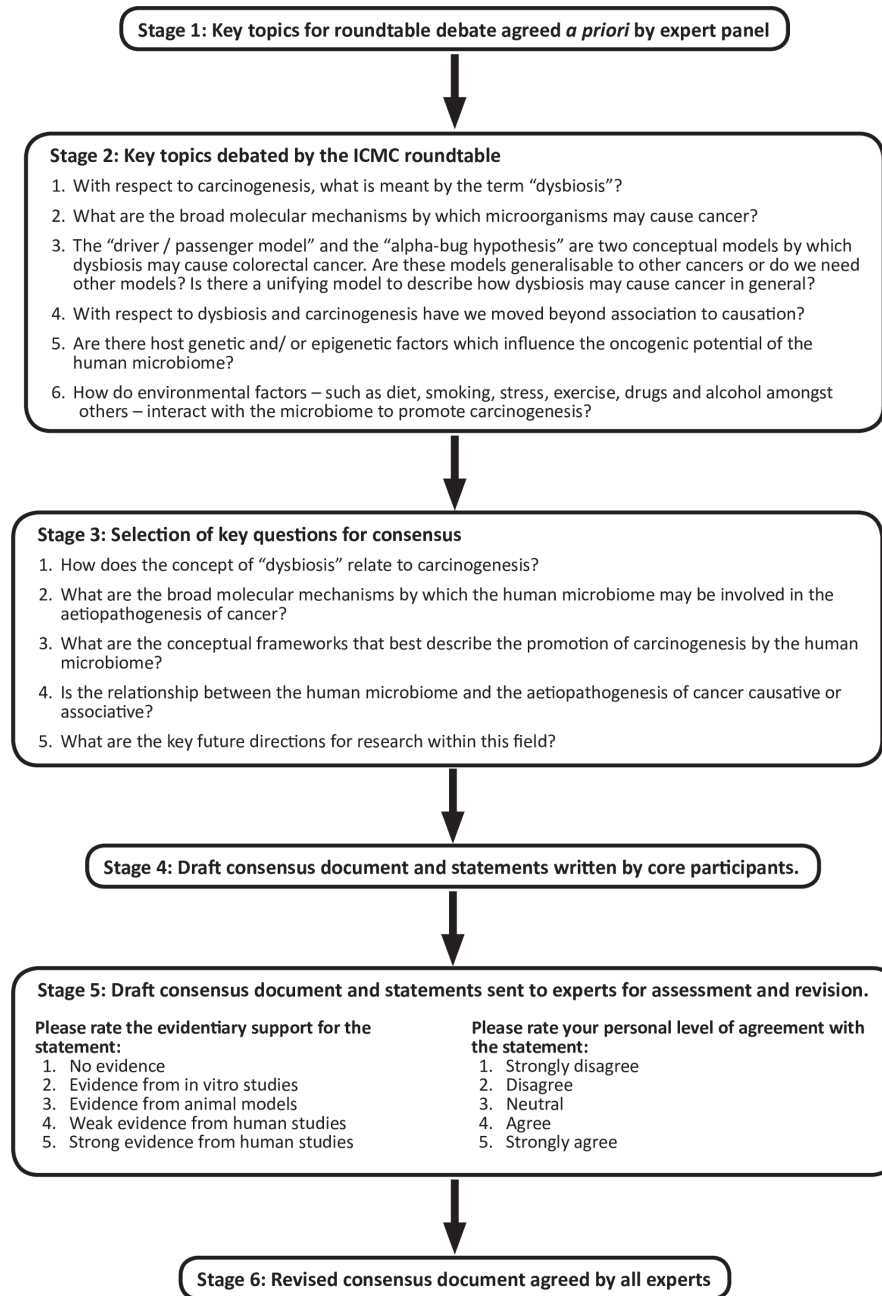


Figure 2 A multistage, structured, iterative methodology was employed. ICMC, International Cancer Microbiome Consortium.

When considering the influence of the microbiome on disease states,⁶ the term dysbiosis is frequently invoked within the microbiome literature and is mostly based on ‘relative abundance’

inferred from 16S rRNA gene sequencing data. However, the term is often loosely applied, poorly defined and has drawn criticism from the scientific community.⁷ We therefore sought to address the concept of dysbiosis specifically within the context of carcinogenesis. Dysbiosis has been defined as an *abnormality*—in composition and/or function—of the host symbiotic microbial ecosystem that exceeds its restitutive capacity and has negative effects on the host.⁸ However, dysbiosis is likely host specific and disease specific—a microbiome may be dysbiotic in one individual, but not in another and/or may promote one pathology, but not another. Therefore, dysbiosis is not an absolute and cannot be defined outside of the context of the host and the disease in question. The broad hypothesis that the expert consensus explored is that *abnormalities in the host microbiome composition and/or function—or dysbiosis—are associated with the aetiopathogenesis of cancer*. But how can we elucidate these

How does the concept of “dysbiosis” relate to carcinogenesis?

Professor Hans Verstraelen
“You can’t talk about dysbiosis... if you don’t know what a healthy microbiome is.”

Professor Julian Marchesi
“At the moment we can’t define a healthy microbiome. We can define a continuum, on which people sit but unless we can follow those individuals over time and capture their [associated] metadata we can’t say that microbiome for you as an individual is a risk for a, b and c.”

Dr Claire Merrifield
“The early microbiome is so key in developing the immune system... which is tolerant to your microbiome. So your microbiome is always going to be unique. I don’t know that we can [define] one healthy microbiome.”

Dr Alasdair Scott
“Can we define “normal” not in terms of... ecology... but in functional terms? A healthy microbiome would be commensal, immune tolerant, diverse, resistant to change, tumour suppressive. There might be lots of ecologies which would give the [same] result. Any departure from that you could define as dysbiosis.”

Figure 3 Select quotes on dysbiosis and the microbiome.

abnormalities when even consensus regarding what comprises a ‘normal’ microbiome remains elusive? For example, although there can be said to be a ‘core microbiome’ at the phylum level (consisting primarily of *Bacteroidetes* and *Firmicutes*), there is significant difference between *apparently* healthy individuals at lower taxonomic levels.⁹ We stress ‘apparently’ because the long lag time associated with the development of most cancers makes it difficult to be sure that an individual is as ‘healthy’ as they seem without long-term follow-up. Interindividual variation in microbiome composition far exceeds variation associated with a given disease state. Such compositional variation may arise from the degree of functional redundancy displayed by the microbiome. For example, numerous microorganisms can produce short chain fatty acids (SCFA, eg, butyrate)—harbouring any of these organisms may be considered normal. Conversely, some functions are more conserved and may even be strain specific. 16S rRNA taxonomic profiling cannot resolve this detail and instead researchers are considering whether there is a core metagenome or metabolome.¹⁰

We suggest that such a functional approach is of more utility in discussing normality and dysbiosis. With respect to the aetiopathogenesis of cancer, we propose that dysbiosis should be considered a persistent departure of the host microbiome from the health-associated homeostatic state (consisting of mutualists and commensals), towards a cancer promoting and/or sustaining phenotype (parasitism or amensalism). This dysbiosis is specific to the individual and thus can only be defined by prospective longitudinal analysis. For example, based on the current literature, the health-associated gut microbiome can be said to have several core features. It is diverse, resilient to short-term environmental pressures and has sufficient plasticity to adapt to the benefit of the host in the face of longer term pressures.¹¹ The health-associated microbiome should synergise with the host to drive beneficial immune responses¹² and metabolic mutualism.¹³ Finally, the microbiome should have a tumour-suppressant effect on the host.^{14 15} Departure from these core features can be considered dysbiotic and may have the potential to incite or sustain cancer. It should be noted that the health-associated microbiomes of other niches will have different core features.

Question two: What are the broad molecular mechanisms by which the human microbiome may be involved in the aetiopathogenesis of cancer (table 2)?

Table 2 What are the broad molecular mechanisms by which the human microbiome may be involved in the aetiopathogenesis of cancer?

Statement: the mechanisms by which the microbiome may initiate and/or drive carcinogenesis can be classified into:	Evidentiary support	Expert agreement
Genomic integration	Strong evidence from human studies (9)	Agree (8)
Genotoxicity	Evidence from animal studies (6)	Agree (9)
Inflammation	Strong evidence from human studies (7)	Agree (8)
Immunity	Evidence from animal studies (6)/weak evidence from human studies (6)	Agree (9)
Metabolism	Strong evidence from human studies (7)	Agree (13)

At the molecular level, the mechanisms by which microorganisms can contribute to carcinogenesis are multiple and varied and have been extensively discussed elsewhere.¹⁶ The mechanisms

may broadly be categorised into genomic integration, genotoxicity, inflammation, immunity and metabolism. Many of these mechanisms are likely not evolved to cause pathologies in the host but rather do so as collateral damage, as the organism optimises its chances of survival. Ultimately, all mechanisms converge on a final common pathway of prolonged host cell survival, enhanced replicative capacity and dedifferentiation.

Genomic integration

Integration of microbial DNA into the host genome is a virulence mechanism for multiple viruses that are causally associated with the development of cancer. Cervical cancer caused by human papilloma virus (HPV-16 and HPV-18) is a classic example. Insertion of two HPV genes, E6 and E7, into the host genome in cervical cells confers a survival advantage as their protein derivatives bind to and inactivate tumour-suppressor gene products (p53 and pRB).¹⁷ Note that this mechanism relies on transcription of viral proteins which exerts a carcinogenic effect on the host. This mechanism contrasts with *insertional mutagenesis* which describes the deregulation of host gene expression caused by insertion of the exogenous genetic material. If the host genes in question are tumour suppressor genes or oncogenes, cellular transformation to an oncogenic phenotype can ensue. Insertional mutagenesis is commonly associated with retroviruses and has been proposed as contributing to the oncogenicity of human T-cell lymphotropic virus.¹⁸

Genotoxicity

Genotoxicity describes structural DNA damage, for example, strand breaks, adducts, deletions and rearrangements. If host DNA damage does not lead to cell death, it may affect tumour suppressor genes or oncogenes with carcinogenic effects. Cytolethal distending toxin (CDT) and colibactin are two well-characterised genotoxins. CDT is produced by *Escherichia coli* and *Campylobacter jejuni* (among others) and induces double-strand DNA breaks via its DNase activity.¹⁹ CDT-deficient strains have attenuated carcinogenic potential in murine CRC models.²⁰ Members of the *Enterobacteriaceae* family can also produce colibactin which induces DNA strand breaks and has been associated with human CRC.²¹ Aside from specific toxins, bacterial metabolites may also exert genotoxic effects. Reactive oxygen species (produced by *Porphyromonas* sp) and hydrogen sulphide (produced by *Bilophila* and *Fusobacterium*) are two examples that have been associated with colorectal neoplasia.^{22 23}

Inflammation

Inflammation is a central feature of carcinogenesis regardless of the aetiological agent and is the principal oncogenic mechanism underlying numerous well-defined, causal, microbial associations with cancer.²⁴ Microbial virulence factors induce chronic inflammation of host tissue, stimulating cellular proliferation

What are the broad molecular mechanisms by which the human microbiome may be involved in the aetiopathogenesis of cancer?

Dr Stephen O’Keefe

“There really is very good evidence that you can prevent most cancers by changing your diet. Now, whether that’s induced by changes in the microbiota and metabolites... well, that’s conjecture.”

Professor Julian Marchesi

“I think you can... find a potential hypothesis where microbiomes in the gut could influence any cancer [outside of the gastrointestinal tract].”

“We can talk about the class one carcinogens, like HPV, *Helicobacter pylori*... a pathogen causing a known cancer. There are molecular mechanisms behind each of those. They have effector molecules... virulence factors. They interfere with some aspect of cellular biochemistry and immunology... Then you have... the metabolite-driven and the carcinogen-driven [cancers]. So, *Desulfovibrio piger* – would you call it a pathogen? No, but it produces a potent genotoxin. *E.coli* producing colibactin – is that a pathogen?”

Professor Rex Gaskins

“In our own work... microbes are closely associated with the mucosa but they’re not adherent. They are making a mutagen. Adhesion is not a requirement.”

Figure 4 Select quotes on mechanisms of microbiome-induced carcinogenesis. HPV, human papilloma virus.

which can ultimately become deregulated and, combined with failure of apoptosis, result in a malignant phenotype.²⁵ These effects may be mediated by microbial interaction with specific host cell intracellular signalling pathways. For example, *Fusobacterium nucleatum* (*F. nucleatum*; associated with CRC) can induce activation of the nuclear factor- κ B (NF- κ B) pathway²⁶ while both *F. nucleatum* and *Bacteroides fragilis* possess virulence factors which negatively regulate E-cadherin, activating WNT/ β -catenin signalling and driving cell proliferation.^{27–28} Enterotoxigenic *B. fragilis* (ETBF) secretes *B. fragilis* toxin and stimulates a TH-17/IL-17-dependent colitis and promotes tumourigenesis.²⁹ In addition to direct, niche-organ-specific effects, there is also evidence that microbe-associated molecular patterns can induce proinflammatory effects in remote organs via their interactions with host pattern recognition receptors, such as toll-like (TLR) and nucleotide-binding oligomerization domain-like (NOD) receptors. Interaction between bacterial lipopolysaccharide and TLR4 results in the downstream activation of cell survival pathways and has been cited as a mechanism by which the intestinal microbiome may contribute to carcinogenesis outside the GI tract.³⁰

Immunity

The interactions between the human microbiota and the immune system are well documented and have significant implications for a wide range of pathologies from atopy and autoimmunity to cancer.^{31–32} The host immune system plays a key role in preventing carcinogenesis by inducing cell death in abnormal host cells with neoplastic potential. The microbiome may interfere with this process at multiple levels. HIV has tropism towards CD4⁺ T lymphocytes, impairing the host's ability to detect potentially neoplastic cells and increasing the rate of carcinogenesis. *F. nucleatum* expresses the Fap2 cell surface protein which interacts with T and Natural Killer cells to suppress antitumour cytotoxicity.³³ Although the microbiota may suppress host immunity, in the healthy state, it is hypothesised that microbiota-immune crosstalk facilitates the maintenance of an immune 'tone' promoting basal anticancer immunosurveillance. Various mechanisms are proposed including broadening the T cell receptor repertoire and enhancing the intensity of immune responses.³⁴

Metabolism

In addition to the immune system, metabolism is the second key level at which the host and the microbiota interact.¹³ The human microbiome is replete with genes which influence the metabolism of dietary vitamins and nutrients, xenobiotics and host-derived compounds such as bile acids. Bacterial metabolism appears to be a crucial cofactor that underpins the observed associations between diet and various cancers.³⁵ Pertinent to CRC, gut bacterial fermentation of dietary fibre to SCFA, such as butyrate, is believed to play an important role in suppressing oncogenesis via its anti-inflammatory and antiproliferative effects.¹⁵ In contrast, bacterial metabolism of bile acids and proteins can result in the formation of carcinogenic aromatic amines and sulphides.³⁵ The microbiome also plays a substantial xenometabolic role which includes the formation of ultimate carcinogenic end products, acetaldehyde production from alcohol for example.³⁶ However, the effects of microbial metabolites are likely to be context dependent on host factors. For example, microbial butyrate production has been suggested to induce CRC in mice deficient in the *MSH2* gene (which codes a protein playing a key role in DNA mismatch repair) by driving hyperproliferation of colonocytes.³⁷

Question three: What are the conceptual frameworks that best describe the promotion of carcinogenesis by the human microbiome (table 3)?

Table 3 What are the conceptual frameworks that best describe the promotion of carcinogenesis by the human microbiome?

Statement	Evidentiary support	Expert agreement
With respect to the pathogenesis of CRC, 'the driver-passenger' model ⁴⁰ accounts for key observations from mechanistic studies and investigations of the on-tumour versus off-tumour microbiome.	Weak evidence from human studies (9)	Agree (8)
The role of the microbiome in the aetiopathogenesis of cancer can be conceptualised as one apex of a tripartite, multidirectional interactome alongside the environment and an epigenetically/genetically vulnerable host.	Weak evidence from human studies (7)	Strongly agree (9)

Traditional carcinogens are generally conceptualised as initiating agents, causing DNA damage and inciting carcinogenesis. This conceptual framework can also be applied to well-established microbial carcinogens which induce DNA damage either directly via genomic integration (eg, HPV) or indirectly via immunosuppression (eg, HIV) and/or chronic inflammation (eg, HCV). However, our understanding of how the dynamic symbionts of the human microbiome may contribute to carcinogenesis is less well established and are largely drawn from the investigation of CRC. Based on in vitro and murine in vivo data, Sears and Pardoll proposed the 'alpha-bug hypothesis' in which key members of the host microbiome possess specific virulence traits that are both directly oncogenic and that also remodel the microbial community towards an oncogenic phenotype.³⁸ They cite *ETBF* as a candidate 'alpha-bug', causing colonic epithelial cell damage directly via *B. fragilis* toxin and indirectly by modifying mucosal immune function and intracellular signalling events.^{27–29} In support of this hypothesis, investigators have demonstrated that high abundance of ETBF in colonic tissues is associated with early-stage carcinogenic lesions.³⁹ However, numerous studies of the on-tumour microbiome show large variation and fail to find consistent overabundance of putative alpha bugs in carcinoma tissue. In recognition of this observation, Tjalsma *et al* proposed a refinement to the alpha-bug hypothesis dubbed the 'driver-passenger model'.⁴⁰ They agreed that certain bacterial *drivers* (akin to alpha bugs) are responsible for the initial epithelial DNA damage that incites oncogenesis. However, they propose that the oncogenic process creates a unique tumour microenvironment that favours the proliferation of opportunistic *passenger*

What are the conceptual frameworks that best describe the promotion of carcinogenesis by the human microbiome?

Professor Daniel Rosenberg

"The idea that 20 years of exposure to one of these metabolic products [e.g. secondary bile acids] is causing colon cancer isn't easy for me to accept. [You said] bile acids are potent carcinogens – we're all exposed to secondary bile acids for many many decades of our lives and not many of us get colon cancer."

Dr Stephen O'Keefe

"There's a threshold effect."

Professor Rex Gaskins

"In the context of a genotoxin, you're constantly repairing and it only takes one hit. So... over time... I would suggest that's why the colon cancer onset is at a late age. You constantly repair and then, at some point, with a susceptible genetic background... Different alleles are going to affect susceptibility."

Dr John Alverdy

"It's a... bidirectional... conversation. And it's iterative... We should use the term "a vulnerable host" because we do know that lifestyle affects cancer. We ought to make it "interactome" – lifestyle, vulnerable host and pathogen/microbiome. This interaction can lead to the acquisition of a carcinogenic [microbiome], which, if there is a failure of [host] resistance... can [lead to the development of] a cancer."

Figure 5 Select quotes on conceptual frameworks describing microbiome-induced carcinogenesis.

bacteria that gradually outcompete the driver species. Thus, the colorectal tumour microbiome undergoes a temporal evolution correlating with tumour stage. This hypothesis is supported by observational data regarding bacterial communities present on colorectal tumours of varying stages.^{41 42} The ‘driver–passenger model’ does not exclude bacterial passengers from playing an active role in cancer progression. *Fusobacterium* sp have been consistently found to colonise CRC tissue in numerous studies and represent an archetypal passenger bacterium.^{43 44} Colorectal tumours are rich in amino acids, an essential substrate for *Fusobacterium* sp,⁴⁵ and express ligands for *Fusobacterium* sp cell-surface receptors of which may explain their overabundance.⁴⁶ *Fusobacterium* sp may also promote tumour progression via interactions with intracellular signalling pathways and immune cells.^{26 28}

Although these models account for key observations from mechanistic studies and investigations of the on-tumour microbiome, they perhaps fall short of addressing the aetiopathogenesis of cancer at the macrolevel. For example, how do environmental exposures, such as diet, contribute to cancer risk? What is the impact of host genetic factors? Are these independent risk factors for cancer development or do they act in consort with the microbiome? In consideration of these questions, we propose a broad framework for modelling the role of the microbiome in the development of cancer based on the concept of the *interactome*. In this model, we consider carcinogenesis to be the ultimate outcome from harmful, tripartite, multidirectional interactions between the microbiome, the environment and the epigenetically/genetically vulnerable host. The relative contributions of these three apices will vary among cancer subtypes. In the case of CRC, for example, it seems likely that the microbiota is necessary, but not sufficient for carcinogenesis; a hypothesis supported by chemical (environment)⁴⁷ or genetic (vulnerable host)⁴⁸ rodent models of CRC that display significantly attenuated or no tumour development in germ-free settings. Bacterial *driver functions* (such as toxin, hydrogen sulphide or (reduced) butyrate production) are the key transducers of environmental cues (such as diet) that can incite carcinogenesis in a vulnerable host (eg, specific TLR genetic polymorphisms). Progression of a neoplasm may depend on continued exposure to environmental stimuli, maladaptive/adaptive changes in microbiome function and/or host response.

Question four: Is the relationship between the human microbiome and the aetiopathogenesis of cancer causative or associative (table 4)?

Table 4 Is the relationship between the human microbiome and the aetiopathogenesis of cancer causative or associative?

Statement	Evidentiary support	Expert agreement
At the single-organism level, the role of microorganisms as aetiological agents in carcinogenesis is well established.	Strong evidence from human studies (10)	Strongly agree (8)
There are plausible mechanisms by which the human microbiome may cause cancer.	Evidence from animal studies (8)	Strongly agree (10)
There is a causal relationship between the human microbiome and the aetiopathogenesis of some cancers.	Weak evidence from human studies (8)	Agree (10)

Despite a re-evaluation that considered modern sequence-based bacterial identification,⁴⁹ Koch’s postulates cannot be applied to prove the causation of cancer by the host microbiome because it is not an ‘infectious agent’. The development of cancer

Is the relationship between the human microbiome and the aetiopathogenesis of cancer causative or associative?

Dr James Kinross

“I don’t think there is debate about [the role of *H. pylori* in gastric cancer.]”

“[Regarding the microbiome,] There is causation. [However], we need more [longitudinal] clinical data before a clinical audience [will] buy into this on a large scale.”

“What is lacking from the evidence that has been presented is a really large-scale ambitious clinical study that... recruits patients... and follows them from childhood all the way through longitudinally. We need something [like this] in cancer if we are ever going to get to the bottom of [causation].”

“I understand why we have these animal models and I understand their value and their function but I just worry... how translatable they really are to the complexity of a human.”

Professor Christian Jobin

“With a cautious “yes” I think we could probably make some claim of causation as far as colorectal cancer [is concerned].”

Figure 6 Select quotes on the potential causal relationship between the human microbiome and carcinogenesis.

is determined by the cumulative effects of a wide variety of risk factors and protective factors on an individual. Despite this complexity, at the single organism level, the role of microorganisms as aetiological agents (or risk factors) in carcinogenesis is well established. Numerous viruses, including human herpesvirus 8 select virotypes of HPV, hepatitis B and C, Epstein-Barr virus and HIV are classified as class 1 carcinogens by the International Agency for Research on Cancer (Lyon, France). *H. pylori*, schistosomes and liver flukes are also known to be carcinogenic. Such causal relationships are founded on Bradford Hill’s criteria which can be broadly categorised as: direct evidence from studies that a relationship between an exposure and an outcome is causal; mechanistic evidence that connects the exposure and outcome and parallel evidence that supports the causal relationship suggested in a study.⁵⁰ For example, direct evidence for the causal relationship between *H. pylori* and gastric cancer (controversial for some time) comes from Uemura *et al*’s landmark prospective cohort study demonstrating that gastric cancer developed in approximately 3% of individuals infected with *H. pylori* compared with none of the uninfected individuals.⁵¹ This finding has been supported by numerous studies demonstrating mechanisms by which *H. pylori* contributes to gastric carcinogenesis including chronic inflammation and direct genotoxic effects mediated by a variety of defined virulence factors and effectors (eg, CagA, cagPAI and VacA).⁵² Parallel evidence comes from randomised trials demonstrating a reduction in the incidence of gastric cancer following *H. pylori* eradication.⁵³ It is notable that not all individuals infected with *H. pylori* develop gastric cancer. Infection alone is not sufficient to cause disease; there are other factors which govern an individual’s susceptibility.

In contrast, a causal relationship between the human symbiotic microbiome and the development of cancer is not firmly established. The nature of the microbiome presents several challenges for proving causation. The microbiome—including the virome, mycobiome and other organisms—is orders of magnitude more complex than a discrete risk factor such as smoking or colonisation by a single organism. The microbiome is dynamic, changing with age, environmental exposures (eg, diet, antibiotics and some surgeries) and the development of cancer itself. While a causal relationship has been posited for numerous cancer types, the role of the gut microbiome in colorectal carcinogenesis has received the most attention and offers the strongest evidence. We have already summarised the mechanistic evidence supporting causation. Parallel evidence comes from murine studies in which germ-free mice chemically or genetically predisposed to intestinal neoplasia, develop fewer tumours compared with conventionally housed counterparts, while carcinogenesis can be potentiated by exposing mice to specific bacteria or stool

from patients with CRC.^{26 54} Human studies have demonstrated that the CRC microbiome differs from that of nearby normal tissue with reduced diversity and altered community structure with lower relative abundance of potentially protective taxa (eg, *butyrate-producing Roseburia*) and increased abundance of taxa with potentially procarcinogenic phenotypes (eg, *Fusobacterium*, *Bacteroides*, *Campylobacter*, *Escherichia* and *Porphyromonas*).^{43 44 55–57} However, it is currently unclear whether this altered community structure is a risk factor for CRC development or rather evolves secondarily to the unique tumour microenvironment. Studies have shown a linear relationship between microbiome composition and tumour stage, moving through the adenoma-carcinoma sequence, and also with other histological tumour features such as lymphovascular and perineural invasion.^{41 42} These findings are at least consistent with causation but could also be argued to reflect changes in the tumour microenvironment. However, uninvolved, normal colonic mucosa from patients with CRC also demonstrates taxonomic differences (eg, abundance of sulfidogenic *Bilophila wadsworthia*) compared with healthy mucosa from controls suggesting a microenvironment-independent underlying field change.⁵⁸ Nonetheless, direct evidence supporting causation in a human population is lacking as current studies have been cross sectional with single-time-point sampling—there are no prospective, longitudinal data from large cohort studies. The evidence for other cancer types and microbiome niches is generally weaker than that for the gut microbiome and CRC and is therefore also unproven.

Question five: What are the key directions for future research to develop our understanding of the role of the microbiome in carcinogenesis (table 5)?

Table 5 What are the key directions for future research to develop our understanding of the role of the microbiome in carcinogenesis?

Statement: key areas for future development with respect to the investigation of the microbiome and carcinogenesis are:	Expert agreement
Large, international cohort studies	Strongly agree (10)
Prospective longitudinal sampling	Strongly agree (14)
More focus on interventional, rather than purely observational studies	Strongly agree (6)/agree (6)
Integration of microbiome analysis with other oncological research projects	Strongly agree (12)
Standardisation and transparency in reporting microbiome research	Strongly agree (10)

With the aims of making a more definitive statement regarding causality and generating true translational impact, the consortium suggested several key directions for future work on the microbiome and cancer.

Large, longitudinal international cohort studies

Proving causation in cancer is difficult because of the lag between exposure and phenotype and the complex interactive nature of potential causative factors. However, these difficulties can be overcome by conducting large, international, longitudinal cohort studies as exemplified by the European Prospective Investigation into Cancer and Nutrition (EPIC)⁵⁹ and Framingham⁶⁰ studies. The execution of such studies within the microbiome field is not without challenge⁶¹ but is necessary to move from testing association to causation.

Increased focus on interventional studies

Observational studies have provided valuable knowledge regarding microbiome composition and function in cancer but

What are the key directions for future research to develop our understanding of the role of the microbiome in carcinogenesis?

Professor Julian Marchesi

"It is an old problem, this concept of standardisation in biology... sometimes you run the risk of just measuring the standardisation process itself and not real biology."

Dr David Hughes

"Existing epidemiological studies] have not really looked specifically at the microbiome – this is something we can tap into."

Dr Stephen O'Keefe

"The problem with [microbiome research] is that too many people are doing observational studies."

Professor David Cunningham

"What we need is longitudinal information so we can reach a view on what [is] a "normal" or "optimal" microbiome."

Figure 7 Select quotes regarding future directions in the field.

if we are to translate this to patient benefit, more emphasis must be placed on interventional research. We need to develop our understanding of how manipulation of the microbiome can impact cancer. For example, early studies have suggested that antibiotic use may increase cancer risk.⁶²

Integration with other oncology research

One way in which to deliver the above three goals is to consider the integration of microbiome research into oncology studies and trials. It is relatively common for clinical trials to have a basic/translational aspect that frequently delves into mechanistic aspects of disease. Given the potential involvement of the host microbiome in the aetiopathogenesis of cancer, cancer diagnosis as well as the efficacy and toxicity of chemotherapy, immunotherapy and radiotherapy, microbiome analysis would be a pertinent angle of inquiry in many oncology trials. Primary or secondary cancer prevention studies with interventions such as dietary change are prime examples.

Standardisation and transparency in reporting microbiome research

Microbiome science has captured the imaginations of scientists, clinicians and the public alike but often blinds consumers with complex 'omics' methodologies while simultaneously failing to deliver on its translational potential. This has precipitated criticism of the entire field⁶³. Reporting guidelines have become *de rigueur* in many aspects of basic and clinic research and the panel would welcome the application of similar frameworks to microbiome research. Transparent reporting of experimental and analytical methods alongside detailed metadata is essential to facilitate reproducibility.

CONCLUSION

In this consensus statement, we have reflected on current expert opinion regarding the crucial question of whether the human microbiome plays a causative role in the aetiopathogenesis of cancer, with a focus on CRC. While cancer causation by specific microorganisms (such as *H. pylori*) is beyond doubt, proving causation for the human microbiome in the broader sense is more challenging. There are plausible mechanisms and ample parallel or supporting evidence from *in vitro*, murine and cross-sectional human studies. However, direct evidence from large longitudinal cohort studies is lacking and should be a key focus in the future. Until then, the expert panel considers that the case in support of a causative role for the human microbiome in the aetiopathogenesis of cancer remains unproven, but a majority nevertheless agrees with the hypothesis. The panel considers the microbiome to be one aspect of an interactome with an epigenetically/genetically vulnerable host and the environment. However, the host genotype is largely fixed and evidence suggests that the microbiome may be a key player in modulating/mediating the response to environmental pressures. In this context, the microbiome

presents a very exciting opportunity to better understand cancer and how it might be prevented.

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REFERENCES

- Lloyd-Price J, Mahurkar A, Rahnavard G, et al. Strains, functions and dynamics in the expanded Human Microbiome Project. *Nature* 2017;550:61–6.
- Tomasetti C, Li L, Vogelstein B, et al. Somatic mutations, cancer etiology, and cancer prevention. *Science* 2017;355:1330–4.
- Mima K, Nishihara R, Qian ZR, et al. *Fusobacterium nucleatum* in colorectal carcinoma tissue and patient prognosis. *Gut* 2016;65:1973–80.
- Zitvogel L, Ma Y, Raouf D, et al. The microbiome in cancer immunotherapy: Diagnostic tools and therapeutic strategies. *Science* 2018;359:1366–70.
- Scott AJ, Merrifield CA, Alexander JL, et al. Highlights from the Inaugural International Cancer Microbiome Consortium Meeting (ICMC), 5–6 September 2017, London, UK. *Ecancermedicalscience*;11:791.
- Carding S, Verbeke K, Vipond DT, et al. Dysbiosis of the gut microbiota in disease. *Microb Ecol Health Dis* 2015;26:26191.
- Hooks KB, O'Malley MA. Dysbiosis and Its Discontents. *MBio* 2017;8.
- Levy M, Kolodziejczyk AA, Thaiss CA, et al. Dysbiosis and the immune system. *Nat Rev Immunol* 2017;17:219–32.
- Human Microbiome Project C. Structure, function and diversity of the healthy human microbiome. *Nature* 2012;486:207–14.
- Zierer J, Jackson MA, Kastenmüller G, et al. The fecal metabolome as a functional readout of the gut microbiome. *Nat Genet* 2018;50:790–5.
- Lozupone CA, Stombaugh JJ, Gordon JJ, et al. Diversity, stability and resilience of the human gut microbiota. *Nature* 2012;489:220–30.
- Belkaid Y, Hand TW. Role of the microbiota in immunity and inflammation. *Cell* 2014;157:121–41.
- Nicholson JK, Holmes E, Kinross J, et al. Host-gut microbiota metabolic interactions. *Science* 2012;336:1262–7.
- Bhatt AP, Redinbo MR, Bultman SJ. The role of the microbiome in cancer development and therapy. *CA Cancer J Clin* 2017;67:326–44.
- Rios-Covián D, Ruas-Madiedo P, Margolles A, et al. Intestinal Short Chain Fatty Acids and their Link with Diet and Human Health. *Front Microbiol* 2016;7:185.
- Schwabe RF, Jobin C. The microbiome and cancer. *Nat Rev Cancer* 2013;13:800–12.
- Münger K, Baldwin A, Edwards KM, et al. Mechanisms of human papillomavirus-induced oncogenesis. *J Virol* 2004;78:11451–60.
- Cook L, Melamed A, Yaguchi H, et al. The impact of HTLV-1 on the cellular genome. *Curr Opin Virol* 2017;26:125–31.
- Nesic D, Hsu Y, Stebbins CE. Assembly and function of a bacterial genotoxin. *Nature* 2004;429:429–33.
- He Z, Gharaibeh RZ, Newsome RC, et al. *Campylobacter jejuni* promotes colorectal tumorigenesis through the action of cytolethal distending toxin. *Gut* 2019;68:289–300.
- Buc E, Dubois D, Sauvanet P, et al. High prevalence of mucosa-associated *E. coli* producing cyclomodulin and genotoxin in colon cancer. *PLoS One* 2013;8:e56964.
- Attene-Ramos MS, Wagner ED, Plewa MJ, et al. Evidence that hydrogen sulfide is a genotoxic agent. *Mol Cancer Res* 2006;4:9–14.
- Huycke MM, Gaskins HR. Commensal bacteria, redox stress, and colorectal cancer: mechanisms and models. *Exp Biol Med* 2004;229:586–97.
- Francescone R, Hou V. Microbiome, inflammation, and cancer. *Cancer J* 2014;20:181–9.
- Pagano JS, Blaser M, Buendia MA, et al. Infectious agents and cancer: criteria for a causal relation. *Semin Cancer Biol* 2004;14:453–71.
- Kostic AD, Chun E, Robertson L, et al. *Fusobacterium nucleatum* potentiates intestinal tumorigenesis and modulates the tumor-immune microenvironment. *Cell Host Microbe* 2013;14:207–15.
- Wu S, Rhee KJ, Zhang M, et al. *Bacteroides fragilis* toxin stimulates intestinal epithelial cell shedding and gamma-secretase-dependent E-cadherin cleavage. *J Cell Sci* 2007;120:1944–52.
- Rubinstein MR, Wang X, Liu W, et al. *Fusobacterium nucleatum* promotes colorectal carcinogenesis by modulating E-cadherin/β-catenin signaling via its FadA adhesin. *Cell Host Microbe* 2013;14:195–206.
- Wu S, Rhee KJ, Albesiano E, et al. A human colonic commensal promotes colon tumorigenesis via activation of T helper type 17 T cell responses. *Nat Med* 2009;15:1016–22.
- Dapito DH, Mencin A, Gwak GY, et al. Promotion of hepatocellular carcinoma by the intestinal microbiota and TLR4. *Cancer Cell* 2012;21:504–16.
- Hooper LV, Littman DR, Macpherson AJ. Interactions between the microbiota and the immune system. *Science* 2012;336:1268–73.
- Routy B, Le Chatelier E, Derosa L, et al. Gut microbiome influences efficacy of PD-1-based immunotherapy against epithelial tumors. *Science* 2018;359:91–7.
- Gur C, Ibrahim Y, Isaacson B, et al. Binding of the Fap2 protein of *Fusobacterium nucleatum* to human inhibitory receptor TIGIT protects tumors from immune cell attack. *Immunity* 2015;42:344–55.
- Zitvogel L, Ayyoub M, Routy B, et al. Microbiome and Anticancer Immunosurveillance. *Cell* 2016;165:276–87.
- O'Keefe SJ, Li JV, Lahti L, et al. Fat, fibre and cancer risk in African Americans and rural Africans. *Nat Commun* 2015;6:6342.
- Seitz HK, Stickel F. Molecular mechanisms of alcohol-mediated carcinogenesis. *Nat Rev Cancer* 2007;7:599–612.
- Belcheva A, Irrazabal T, Robertson SJ, et al. Gut microbial metabolism drives transformation of MSH2-deficient colon epithelial cells. *Cell* 2014;158:288–99.
- Sears CL, Pardoll DM. Perspective: alpha-bugs, their microbial partners, and the link to colon cancer. *J Infect Dis* 2011;203:306–11.
- Purcell RV, Pearson J, Aitchison A, et al. Colonization with enterotoxigenic *Bacteroides fragilis* is associated with early-stage colorectal neoplasia. *PLoS One* 2017;12:e0171602.
- Tjalsma H, Boleij A, Marchesi JR, et al. A bacterial driver-passenger model for colorectal cancer: beyond the usual suspects. *Nat Rev Microbiol* 2012;10:575–82.
- Kinross J, Mirnezami R, Alexander J, et al. A prospective analysis of mucosal microbiome-metabonomic interactions in colorectal cancer using a combined MAS 1HNMR and metataxonomic strategy. *Sci Rep* 2017;7:8979.
- Flanagan L, Schmid J, Ebert M, et al. *Fusobacterium nucleatum* associates with stages of colorectal neoplasia development, colorectal cancer and disease outcome. *Eur J Clin Microbiol Infect Dis* 2014;33:1381–90.

- 43 Marchesi JR, Dutilh BE, Hall N, *et al.* Towards the human colorectal cancer microbiome. *PLoS One* 2011;6:e20447.
- 44 Castellarin M, Warren RL, Freeman JD, *et al.* *Fusobacterium nucleatum* infection is prevalent in human colorectal carcinoma. *Genome Res* 2012;22:299–306.
- 45 Dai ZL, Wu G, Zhu WY. Amino acid metabolism in intestinal bacteria: links between gut ecology and host health. *Front Biosci* 2011;16:1768–86.
- 46 Abed J, Emgård JE, Zamir G, *et al.* Fap2 Mediates *Fusobacterium nucleatum* Colorectal Adenocarcinoma Enrichment by Binding to Tumor-Expressed Gal-GalNAc. *Cell Host Microbe* 2016;20:215–25.
- 47 Uronis JM, Mühlbauer M, Herfarth HH, *et al.* Modulation of the intestinal microbiota alters colitis-associated colorectal cancer susceptibility. *PLoS One* 2009;4:e6026.
- 48 Kado S, Uchida K, Funabashi H, *et al.* Intestinal microflora are necessary for development of spontaneous adenocarcinoma of the large intestine in T-cell receptor beta chain and p53 double-knockout mice. *Cancer Res* 2001;61:2395–8.
- 49 Fredricks DN, Relman DA. Sequence-based identification of microbial pathogens: a reconsideration of Koch's postulates. *Clin Microbiol Rev* 1996;9:18–33.
- 50 Howick J, Glasziou P, Aronson JK. The evolution of evidence hierarchies: what can Bradford Hill's 'guidelines for causation' contribute? *J R Soc Med* 2009;102:186–94.
- 51 Uemura N, Okamoto S, Yamamoto S, *et al.* *Helicobacter pylori* infection and the development of gastric cancer. *N Engl J Med* 2001;345:784–9.
- 52 Ishaq S, Nunn L. *Helicobacter pylori* and gastric cancer: a state of the art review. *Gastroenterol Hepatol Bed Bench* 2015;8:S6–S14.
- 53 Fuccio L, Zagari RM, Eusebi LH, *et al.* Meta-analysis: can *Helicobacter pylori* eradication treatment reduce the risk for gastric cancer? *Ann Intern Med* 2009;151:121–8.
- 54 Wong SH, Zhao L, Zhang X, *et al.* Gavage of Fecal Samples From Patients With Colorectal Cancer Promotes Intestinal Carcinogenesis in Germ-Free and Conventional Mice. *Gastroenterology* 2017;153:1621–33.
- 55 Yu J, Feng Q, Wong SH, *et al.* Metagenomic analysis of faecal microbiome as a tool towards targeted non-invasive biomarkers for colorectal cancer. *Gut* 2017;66:70–8.
- 56 Kostic AD, Gevers D, Pedamallu CS, *et al.* Genomic analysis identifies association of *Fusobacterium* with colorectal carcinoma. *Genome Res* 2012;22:292–8.
- 57 Flemer B, Lynch DB, Brown JM, *et al.* Tumour-associated and non-tumour-associated microbiota in colorectal cancer. *Gut* 2017;66:633–43.
- 58 Yazici C, Wolf PG, Kim H, *et al.* Race-dependent association of sulfidogenic bacteria with colorectal cancer. *Gut* 2017;66:1983–94.
- 59 Riboli E, Kaaks R. The EPIC Project: rationale and study design. European Prospective Investigation into Cancer and Nutrition. *Int J Epidemiol* 1997;26(Suppl 1):6S–14.
- 60 Mahmood SS, Levy D, Vasan RS, *et al.* The Framingham Heart Study and the epidemiology of cardiovascular disease: a historical perspective. *Lancet* 2014;383:999–1008.
- 61 Vandeputte D, Tito RY, Vanleeuwen R, *et al.* Practical considerations for large-scale gut microbiome studies. *FEMS Microbiol Rev* 2017;41:S154–S67.
- 62 Cao Y, Wu K, Mehta R, *et al.* Long-term use of antibiotics and risk of colorectal adenoma. *Gut* 2018;67:672–8.
- 63 Hanage WP. Microbiology: Microbiome science needs a healthy dose of scepticism. *Nature* 2014;512:247–8.