Abstract
Evidence supports associations between human gut microbiome variation and multiple health outcomes and diseases. Despite compelling results from in vivo and in vitro models, few findings have been translated into an understanding of modifiable causal relationships. Furthermore, epidemiological studies have been unconvincing in their ability to offer causal evidence due to their observational nature, where confounding by lifestyle and behavioural factors, reverse causation and bias are important limitations. Whilst randomized controlled trials have made steps towards understanding the causal role played by the gut microbiome in disease, they are expensive and time-consuming. This evidence that has not been translated between model systems impedes opportunities for harnessing the gut microbiome for improving population health. Therefore, there is a need for alternative approaches to interrogate causality in the context of gut microbiome research.

The integration of human genetics within population health sciences have proved successful in facilitating improved causal inference (e.g., with Mendelian randomization [MR] studies) and characterising inherited disease susceptibility. MR is an established method that employs human genetic variation as natural “proxies” for clinically relevant (and ideally modifiable) traits to improve causality in observational associations between those traits and health outcomes. Here, we focus and discuss the utility of MR within the context of human gut microbiome research, review studies that have used this method and consider the strengths, limitations and challenges facing this research. Specifically, we highlight the requirements for careful examination and interpretation of derived causal estimates and host (i.e., human) genetic effects themselves, triangulation across multiple study designs and inter-disciplinary collaborations. Meeting these requirements will help support or challenge causality of the role played by the gut microbiome on human health to develop new, targeted therapies to alleviate disease symptoms to ultimately improve lives and promote good health.
Keywords
microbiome, human genetics, Mendelian randomization, causality

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Author roles: Wade KH: Conceptualization, Data Curation, Investigation, Methodology, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; Hall LJ: Conceptualization, Writing – Original Draft Preparation, Writing – Review & Editing

Competing interests: No competing interests were disclosed.

Grant information: KHW is supported by the Elizabeth Blackwell Institute for Health Research, University of Bristol and the Wellcome Trust Institutional Strategic Support Fund [204813]. LJH is supported by a Wellcome Trust Investigator Award [100974], the Biotechnology and Biological Sciences Research Council (BBSRC) Institute Strategic Programme Gut Microbes and Health [BB/R012490/1] and its constituent project(s) [BBS/E/F/000PR10353 and BBS/E/F/000PR10356].

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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How to cite this article: Wade KH and Hall LJ. Improving causality in microbiome research: can human genetic epidemiology help? [version 2; peer review: 1 approved] Wellcome Open Research 2020, 4:199 https://doi.org/10.12688/wellcomeopenres.15628.2

First published: 10 Dec 2019, 4:199 https://doi.org/10.12688/wellcomeopenres.15628.1
List of abbreviations
AUC = area under the curve; BMI = body mass index;
CAD = coronary artery disease; CKD = chronic kidney disease;
FGFP = Flemish Gut Flora Project; GECCO = Genetics and
Epidemiology of Colorectal Cancer Consortium; GWAS = genome-
wide association study; IBD = inflammatory bowel disease;
LD = linkage disequilibrium; MI = myocardial infarction;
MR = Mendelian randomization; OR = odds ratio; RCT = random-
ized controlled trial; RR = risk ratio; SCFA = short chain fatty
acid; SD = standard deviation; SNP = single nucleotide polymor-
phism; TMAO = trimethylamine-N-oxide; T2D = type 2 diabetes

Introduction
Evidence from microbiome-wide studies has highlighted rela-
tionships between the gut microbiome and many complex traits
and diseases – from dietary composition, obesity, rheuma-
toid arthritis and type 2 diabetes (T2D) to Alzheimer’s disease,
Parkinson’s disease and depression14,15. For example, the lower
diversity and relative abundances of bacteria within the Bacter-
oideetes vs. Firmicutes phyla in obese vs. lean individuals has
been observed in studies of mice and humans, with cross-
sectional, longitudinal and experimental designs18,19. Several stud-
ies have also demonstrated that the relative bacterial abundances
in the Bacteroidetes order increases and Firmicutes order decreases
with low-calorie diets (e.g., through fat or carbohydrate
restriction) or surgery-induced weight loss in obese individu-
als, whereby the gut microbiota composition becomes similar
to that of their lean counterparts19–21. Ostensibly, these stud-
ies suggest that the manipulation of the human gut microbiome
(e.g., not only through diet and surgery but also via the intake
of pre- or pro-biotics, antibiotic usage or faecal microbiome
transplants) may have potential as an approach to develop new,
targeted therapies and treatments to reduce disease in the
population.

However, the design of human studies has been largely obser-
vational (with additional potential for experimental biases in
sample collection, storage and analysis) and, owing to this,
there are the many inconsistencies within the literature, casting
doubt on the reliability of existing findings. Moreover, causality
in these relationships is often difficult to ascertain, with a
concerning lack of robust evidence able to discern correlation
from causation (despite being called for 14–16). It is particularly
alarming that, despite this lack of evidence, and with much scepticism14,
there is a growing market for commercial initiatives
targeting the microbiome as a consumer-driven intervention (e.g.,
ubioMe, VioMe, BIOHM and Atlas BioMed), where
companies ask for, obtain and sequence faecal samples from
consumers and prescribe “personalised” nutritional information
based on the, often, only sample. Furthermore, there has
been an increase in clinical recommendation of pro-biotics –
“live microorganisms that, when administered in adequate
amounts, confer a health benefit on the host”18 – for treating
various diseases or following antibiotic prescription19–21. How-
ever, with the development of such efforts, it is important to
recognise that the gut microbiome is a dynamic and complex
ecosystem; therefore, careful investigation of the off-target
effects of any treatment or intervention intended to alter one
or a small number of specific bacteria is required.

Some of the current literature comprising in vivo and in vitro
experiments has provided promising results, which have been
supported by small-scale observational studies within humans.
However, many studies have failed to be translated between
model organisms, and studies within humans have been
unconvincing in their ability to provide evidence for causality
in these relationships (even those with compelling results from
in vivo and in vitro models). Despite the few examples that have
proved successful in their consistency between model organisms
and their clinical application in humans (e.g., faecal microbio-
me transplantation in cases of recurrent Clostridiodies difficile
infection, which has a global success rate of over 80%22),
evidence that has not been translated between model organism-
isms impedes any opportunity for harnessing the gut microbiome
for reducing the burden of disease in the population and has
induced scepticism in its causal relevance in human health23,24.

Reasons for these discrepancies between and within model
organisms include the challenges in the increasing volume of
high-dimensional multi-omic data produced and, specifically,
how these are integrated using complex bioinformatics and
incorporated into traditional study designs, alongside the sensitive
experimental models that aim to replicate disease traits in
humans. Whilst murine models have played a key role in the
emerging gut microbiota research field (given the inability to
research all questions within humans), there are important inter-
study variations due to experimental design (e.g., sample collec-
tion and processing), environmental conditions (i.e., differential
microbiome composition between rodent housing facilities),
genetic differences and chosen analyses25. Despite some comp-
pelling examples exhibiting consistency and providing mecha-
nistic understanding of these relationships within humans and
animals (e.g., in malnutrition and obesity26–27) and between
in silico and in vitro models28, it is still debatable as to whether
animal models of the human gut microbiome (and methods
used such as germ-free or gnotobiotic mice) are translatable
to humans, particularly with all scientific questions29,30.

In addition to the limitations of in vitro and in vivo
experimental design – currently fuelling the “bottom-up”
approach for assessing causality between model systems – there
are limitations of the observational human epidemiological study
designs. Single-sample observational epidemiological studies
(i.e., population samples in cross-section) or case-control stud-
ies suffer confounding by lifestyle and behavioural factors, biases
(e.g., error in measures of the gut microbiome and non-random
or unrepresentative selection of participants) within and between
studies and may not be generalizable. Indeed, whilst both murine
and human studies have provided support for a relationship
between the gut microbiome and diseases such as inflamma-
tory bowel diseases (IBDs) and T2D14,16, these study designs
usually include the assessment of differential gut microbiome
compositions within clinical patients (i.e., those who already have
the disease of interest) compared to controls. Meta-analyses of studies like these can provide some insight into consistency and robustness of potential findings, which may lead to greater precision of observed associations (e.g., with metagenomic signatures within colorectal cancer cases\(^3\)), but the direction of causality in these relationships is ambiguous. Specifically, is it that differences in the gut microbiome protect/exacerbate a disease or is it the disease state itself that is leading to variation in the gut microbiome composition? Furthermore, if evidence of a likely causal effect of a component of the gut microbiome on the risk of a disease is provided, this does not imply that the same is true for the progression of that disease (and vice versa). Specifically, treating individuals who suffer from, say, IBD with a pre-/pro-biotic that promotes/contains bacteria found to be associated with the risk of developing IBD may not lead to better prognosis of IBD after diagnosis.

Given that the gut microbiome is a dynamic system, assessment of the prospective changes in this system alongside the development of disease and variation in health outcomes is required. Longitudinal cohort studies have developed some understanding in the transitional association of the gut microbiome and specific health outcomes over a relatively short period of time (e.g., with childhood obesity, type 1 diabetes and adult weight gain\(^3\)). However, these are still limited by traditional observational epidemiological complications (i.e., confounding and bias) if not designed or conducted well and, usually, by statistical power given usually small sample sizes.

Randomized controlled trials (RCTs) in this area have made some steps towards understanding the causal role played by the gut microbiome in disease. However, many mainly focus on alleviating symptoms within patients who have an established condition\(^3\). RCTs have focused on the influence of diet, pre- and pro-biotics, or antibiotics on the gut microbiome and related traits and have presented an array of conclusions ranging from a preventative to detrimental role of these interventions. However, most of these RCTs consist of fewer than 50 participants, who tend to be selected based on disease status and who are often on different medications, which are difficult to control\(^2\). Only a handful of registered trials to date have completed with tangible results and more have been terminated, suspended or withdrawn\(^2\). Given the current literature, trials focusing on disease prevention in healthy individuals or those understanding how variations within the gut microbiome can promote good health are imperative. Whilst larger efforts are ongoing (e.g., as of July 2019, there were approximately 650 RCTs of the gut microbiome still recruiting), such trials are likely not feasible to answer every scientific question and are importantly expensive, time-consuming and sometimes unethical, particularly within a healthy human population. The application of alternative causal inference methods in this context are needed to improve causal inference and help elucidate the role played by the gut microbiome in human health and disease.

MR

MR is an approach that uses human genetic variation (usually single nucleotide polymorphisms [SNPs], identified in genome-wide association studies [GWAS]) to act as a “proxy” measure for exposures of interest (e.g., here, the gut microbiome)\(^4\). Provided a number of key assumptions are met (Figure 1), these genetic variants can be argued to have properties that

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**Figure 1. Framework, assumptions and example of Mendelian randomization (MR) in the context of gut microbiome research.** (A) MR relies on the following three core assumptions: (1) the genetic variant(s) being used as an instrument (Z) is associated with the exposure (X); (2) the genetic variant(s) are independent of measured and unmeasured confounders (U) of the association between the exposure (X) and outcome (Y); and (3) there is no independent pathway between the genetic variant(s) and outcome (Y) other than through the exposure (X) – known as horizontal pleiotropy or the exclusion restriction criteria. (B) Example of MR applied to understanding the causal role played by Bifidobacterium and obesity using the rs4988235 SNP (i.e., the lactase persistence genetic variant within the MCM6 locus) as an instrument (see text for discussion).
approximate those of “instruments” and thus can be used to estimate the causal effect of a trait on disease or health outcome. Theoretically analogous to arms of an RCT, genetic variants used in MR are largely independent of confounding factors, due to the random nature of their allocation within a population in the absence of any population stratification. These genetic variants are also not modified by the later development of disease or health outcome and, with very accurate genotyping being commonplace, measurement error is largely reduced. Therefore, at a population level, the portion of variance in the modifiable trait explained by human genetic variants (unlike the direct measurement of the trait itself) can be used to model situations that are free of the aforementioned limitations that would otherwise weaken causal inference in observational studies.

MR expands upon traditional genetic analyses (such as candidate gene studies, GWAS or aggregating genome-wide variation to inform characterisation of disease susceptibility) by harnessing randomly allocated genetic variation identified as being associated with variation in a particular trait to specifically interrogate the causal implications of that trait on a disease or health outcome, in a manner comparable to RCTs. Such analyses can be done at a fraction of the time and cost required for a large-scale RCT, by exploiting data from pre-existing and large-scale genetic studies, and can be used to ask many of the scientific questions that may not be feasible or ethical with RCTs. Particularly demonstrative of the advantages of the MR paradigm is the example of selenium and prostate cancer (Box 1). Given the inconsistencies between studies aimed at estimating the causal role of the gut microbiome in human health and disease, MR provides the opportunity to assess causality in observed associations not only between the gut microbiome and health outcomes but also the impact of various traits on the gut microbiome itself, without the need for costly RCTs or lab-based study designs in the first instance. Results derived from the application of MR in this context also provides a potential mechanism to direct the prioritisation of characteristics of the gut microbiome as interventional targets (e.g., via dietary regulation or using pre- and pro-biotics), to inform clinical and public health guidelines and to improve population health in an efficient and cost-effective manner. For a full description and definitions of terminology, methods and assumptions specific to MR, please see the online MR Dictionary.

**Current applications of MR**

Our understanding of the host (i.e., human) genetic contribution to the gut microbiome has primarily arisen from candidate gene studies and genome-wide screens in model organisms. In recent years, and with the advent of higher-throughput technologies that are able to capture measurements of the gut microbiome at scale, several GWASs in humans have been conducted to further uncover host genetic variation that shapes the gut microbiome. Together, these initial GWASs have identified associations between more than 100 human genetic variants associated with constituents of the gut microbiome (e.g., microbial diversity, taxon abundance and community structure). However, there has been limited overlap of identified host genetic variants implicated as being associated with the gut microbiome across studies. With the existence of host genetic variants associated with the gut microbiome, we and others have applied MR to appraise causality in the relationships between the gut microbiome and human health. However, it is of upmost importance for careful examination and interpretation of MR-derived causal estimates, host genetic effects and the assessment of other benefits that the integration of human genetics to this field may provide in appraising causality. The current applications of MR in microbiome research are described below, with pertinent limitations shared between these applications discussed.

**Microbiota genera and ischemic heart disease, T2D and risk factors**

In 2018, Yang *et al.* used MR to assess the causal effect of 27 component genera of the gut microbiome on ischemic heart disease, T2D, adiposity, lipid levels and insulin resistance using human genetic variation that had previously been associated with these particular bacterial taxa measured with 16S rRNA sequencing. In this case, host SNPs (i.e., human genetic variation) associated with these 27 genera were obtained from previous studies, with highly correlated SNPs removed based on linkage disequilibrium (LD; based on an $r^2 \geq 0.8$). These were also cross-referenced with Ensembl and the GWAS catalog to remove potentially pleiotropic SNPs (i.e., those that may have an effect on the outcome other than through the exposure of interest) and to reduce possibility of invalidating the third MR assumption (Figure 1).
By using MR to analyse the causal impact of these 27 genera on cardiometabolic disease and related traits, authors found evidence that a greater relative abundance of bacteria in the *Bifidobacterium* genus was associated with a 1.5% lower odds of ischemic heart disease (odds ratio [OR]: 0.99; 95% CI: 0.97–1.00), a 0.01 standard deviation (SD) lower body mass index (BMI; 95% CI: 0.01–0.02) and a 0.03 SD lower level of Lipoprotein cholesterol (95% CI: 0.02–0.03).

This application of MR provides compelling evidence of the causal impact of the gut microbiome on cardiometabolic traits. However, these results were not robust when testing whether the effects of these genetic variants were independent of *Bifidobacterium* (i.e., “horizontal pleiotropy” — Figure 1), suggesting that this bacterial genus may not have been the sole driver of these seemingly beneficial metabolic effects. Additionally, many of the associations between the human genetic variants and the gut microbiome used in the MR analyses were not replicated either within or between the studies from which they were obtained51–53,55,57, questioning the validity of using them in MR analyses, as they may not be reliably associated with the exposure (i.e., invalidating the first MR assumption, Figure 1).

### Short-chain fatty acids and metabolic diseases

In 2019, authors of another study based on a collection of 952 normoglycemic individuals from the LifeLines-DEEP cohort with genetic, metagenomic sequences and faecal short-chain fatty acid (SCFA) levels performed bi-directional MR to assess the association of 245 metagenomic features describing functionality of the gut microbiome (2 of which were linked to SCFA production, 57 unique taxa and 186 pathways) with 17 metabolic and anthropometric traits58–60. Genetic variants were chosen by conducting a GWAS within 952 individuals from the LifeLines-DEEP cohort. A fairly lenient p-value threshold of $1 \times 10^{-5}$ (identified from an independent cohort of 445 individuals) was used to define host genetic variants independently associated with functional features of the gut microbiome and explained most variance in the same feature.

Authors found that the microbial functional pathway characterised from metagenomic sequencing involved in 4-aminobutanoate (GABA) degradation (PWY-5022), of which the SCFAs butyrate and acetate are products, was associated with improved insulin response after an oral glucose-tolerance test (characterised by the ratio of the areas under the curve (AUC) for measured insulin and glucose levels, $\text{AUC}_\text{insulin} / \text{AUC}_\text{glucose}$). Specifically, by using MR to assess the impact of functionality of the gut microbiome, authors found that each SD increase in the abundance of the PWY-5022 pathway was associated with a 0.16 mU/mmol increase in the $\text{AUC}_\text{insulin} / \text{AUC}_\text{glucose}$ (95% CI: 0.08–0.24), which was robust to MR methods that test validity with regards to horizontal pleotropy. In taxonomic analyses, the bacteria most correlated with the PWY-5022 functional pathway were *Eubacterium rectale* and *Roseburia intestinalis* (both of the Clostridiales order), species known to produce butyrate.61 The proposed mechanism explaining these results suggested that host genetic variation influences the gut microbiome composition to modulate GABA degradation, thus, increasing the ability of the pancreas to secrete insulin in response to a glucose challenge.

Whilst the metagenomic features authors used provided more insight into the functionality of the gut microbiome (over and above measuring relative abundances with 16S RNA sequencing), authors were unable to test the relationship between these features and circulating levels of the SCFAs (e.g., butyrate and propionate), as these were not measured in the study sample. Similar to Yang *et al.*, the associations between the genetic variants used in MR analyses and the functional features of the gut microbiome were not replicated in other studies. Additionally, much of the GWAS summary-level data that authors used was adjusted for other covariates (mainly BMI), which may induce false correlations between the exposure and outcome via a certain type of selection bias (i.e., collider bias) and, in the most extreme cases, this can reverse the direction of the causal effect estimate (which was observed in this study)64,65.

### Microbiota-derived metabolites and cardiometabolic health

A further study conducted by Jia *et al.* used MR to examine the association between the trimethylamine-N-oxide (TMAO) metabolite, produced by processes specific to gut bacteria when metabolising choline from high-fat foods such as eggs and beef, and its predecessors with both continuous measures of cardiometabolic health and diseases66. This study was motivated by the observational epidemiological literature, suggesting that choline, TMAO (a derivative of choline) and carnitine are associated with an × risk of heart disease and other cardiometabolic diseases, hypothesised through their atherosclerotic effects in blood vessels66,67. The authors undertook a bi-directional MR analysis to unpick the direction of association of circulating levels of these metabolites with traits relating to adiposity, glycaemic profile, lipids and kidney function alongside diseases including T2D, coronary artery disease (CAD), myocardial infarction (MI), stroke, atrial fibrillation and chronic kidney disease (CKD)68. Genetic variants used in MR analyses as instruments for each of four metabolites (choline, TMAO, carnitine and betaine) were obtained from a GWAS of 217 blood-based metabolites in 2,076 individuals of European descent from the Framingham Heart Study (Offspring Cohort)69 and chosen based on a lenient threshold of “suggestive" genome-wide significance ($P<5 \times 10^{-5}$). Given the number of tests being performed, authors set an *a priori* Bonferroni-corrected threshold ($P<0.0005$) to detect evidence for association.

By using MR to assess the causal role of gut microbiome-derived metabolites and cardiometabolic health, authors found some evidence to suggest that higher circulating levels of choline increased the risk of T2D (OR: 1.84 per 10 units; 95% CI: 1.00, 3.42) and higher circulating levels of betaine reduced the risk of T2D (OR per 10 units: 0.68; 95% CI: 0.48, 0.95). There was little evidence to suggest that any metabolite had a causal role on the continuous measures of cardiometabolic health. In the reverse direction, there was evidence suggesting that a higher liability to T2D may play a causal role in increasing levels of TMAO (0.13 units; 95% CI: 0.06, 0.20). Most of these findings were consistent across multiple MR methods, which test robustness with regards to horizontal pleotropy (i.e., the third MR assumption, Figure 1).

Results presented by Jia *et al.* provided further insight into the functional relevance of the gut microbiome on cardiometabolic disease, performing power calculations and utilising a...
selection of appropriate sensitivity analyses that appraise validity of derived causal estimates. These results suggested that, by altering levels of these microbiome-dependent metabolites, there may be an opportunity to modify the potential causal impact of the gut microbiome and its metabolic functionality to reduce the risk of T2D. However, it is important to clarify firstly that authors tested the relationship of circulating levels of metabolites with cardiometabolic traits and not characteristics of the gut microbiome itself. Indeed, whilst constituent gut microbiota do produce these metabolites naturally and are associated with their circulating levels, such metabolites can be introduced into the bloodstream via other means (e.g., supplementation and diet). Thus, the direct causal impact of gut microbiome in this instance is uncertain and may, in fact, be irrelevant. Similar to the other applications of MR described above, the genetic variants used as instruments for the microbiota-derived metabolites were not replicated in the original GWAS from which they were obtained and were selected based on a fairly lenient threshold (similar to Sanna et al.89), questioning the reliability of their use.

Genome-wide association of gut microbiome variation and causal inference analyses

As is evident from the current applications of MR in the context of microbiome research, one of the main limitations is the limited overlap and replication of identified host genetic variants associated with the gut microbiome across studies. Most recently, a GWAS of the gut microbiome characterised with 16S rRNA sequencing was conducted by Hughes et al. in over 3800 individuals from three independent studies, including a discovery sample comprising the Flemish Gut Flora Project and two German replication samples. Within this analysis, 13 SNPs reached conventional levels of genome-wide significance ($P<2.5\times10^{-8}$), with some showing low heterogeneity between studies, and two of these reached a strict study-level threshold ($P<1.57\times10^{-10}$). This novel and persistent collection of results provided some insight into the validity of using host genetic variation and genome-wide information on both the exposure and outcome, these methods will become more feasible with the growing availability of large-scale GWASs of the gut microbiome and other traits.

These results provided evidence for causality between five microbial traits and seven outcomes, including evidence for a causal role of bacteria within the Butyricicoccus genus on IBDs and bacteria within the Firmicutes phylum on waist circumference. The strongest association indicated that presence (vs. absence) of bacteria within the Dialister genus decreased the risk of Alzheimer’s disease (risk ratio [RR] with a doubling of the genetic liability to presence vs. absence of Dialister: 0.81; 95% CI: 0.73, 0.90). In the reverse direction, there were also evidence for causal relationships between four phenotypes and three microbial traits, including evidence for a causal role of a higher liability to Parkinson’s disease, T2D and Crohn’s disease on bacteria within the Firmicutes phylum. The strongest result suggested that individuals with Alzheimer’s disease were more likely to carry bacteria in the Dialister genus within their gut (RR for a doubling of the genetic liability to Alzheimer’s disease: 1.81; 95% CI: 1.13-2.89).

Focusing on the components of this latter result, the bi-directional analyses of the causal relationship between bacteria within the Dialister genus and Alzheimer’s disease may seemingly be somewhat contradictory (i.e., presence of Dialister reducing the risk of Alzheimer’s disease but Alzheimer’s disease presence increasing likelihood of Dialister). However, there are two important points to note. Firstly, there may well be a true protective role of the Dialister bacteria on the onset of Alzheimer’s disease (hence the inverse effect), which is supported by a recent study proposing this very notion. Secondly, the single genetic variant used as an instrument in the MR analysis assessing the impact of Dialister bacteria on Alzheimer’s disease is within SORL1, a gene characterised as being associated with Alzheimer’s disease itself. Therefore, the question arises as to whether the SNP is indeed a valid instrument for bacteria within the Dialister genus (as it may be a pleiotropic SNP, independently associated with Alzheimer’s disease) or whether the mechanism by which the SNP influences Alzheimer’s disease is through its impact on bacteria within the Dialister genus. At the present time, is difficult to discern without further biological, functional and mechanistic knowledge. Therefore, there is a requirement for careful examination and interpretation of the host (i.e., human) genetic effects on these microbial traits before using them in such applied analyses.

One proposed mechanism for examining these complexities and unpicking the link between the human gut microbiome and various health outcomes is utilizing the plethora of human genetic epidemiological methods and sensitivity analyses that specifically explore the validity of host genetic variation in MR analyses. For example, as proposed by Richardson et al.31, methods such as colocalization, bivariate genetic fine mapping and bi-directional MR may provide some distinction between reverse causality and either direct or LD-induced horizontal pleiotropy (Figure 2). As these methods (particularly colocalization and genetic fine mapping) require individual-level and genome-wide information on both the exposure and outcome, these methods will become more feasible with the growing availability of large-scale GWASs of the gut microbiome and other traits.

Indeed, the application of such methods in a recent study provided some insight into the validity of using host genetic variants associated with the human gut microbiome in examining the relationship between gut microbiome variation and colorectal cancer (unpublished). For this, the summary statistics from the 13 SNPs associated with microbial traits derived from 16S rRNA sequencing reported by Hughes et al. were combined with those from the Genetics and Epidemiology of Colorectal Cancer Consortium (GECCO) in a two-sample MR analysis. Results provided evidence that the presence (vs. absence) of uncharacterised bacterial genera within the Bacteroidales order increased the risk of colorectal cancer by approximately 8%.
(95% CI: 2-15%), with no strong evidence that the SNP used as an instrument was associated with other traits or the outcome itself, reducing the likelihood of horizontal pleiotropy.

Whilst these are first steps in the right direction of understanding the utility of host genetic variation in microbiome research for improved causality, there is much room for improvement. Importantly, when appreciating the complexity of these relationships, the integration of human genetics, genetic epidemiological techniques and causal inference methodologies to the field of microbiome research holds great potential.

**Limitations of the MR approach**

With the advent of the application of MR within the context of gut microbiome research, and the growing data available for such analyses, there are important limitations and complexities to these applied epidemiological analyses that need to be acknowledged and addressed. The general limitations of MR have been summarised before\(^ {41,43}\), but those most pertinent to microbiome research are currently the lack of robust and reliable genetic variants associated with the gut microbiome and its functionality and the complexity of mechanisms by which host genetic variants impact these microbial traits. This latter complication includes the possibility that identified genetic variants associated with components of the gut microbiome are also associated with the outcome of interest in an MR study through independent mechanisms (Figure 2). In addition, many of the current studies utilizing MR have used lenient \( p \)-value thresholds to define the set of included genetic variants, leading to concerns in their instrumental variable quality.

As the pool of increasingly larger-scale GWASs and meta-analyses grow (e.g., most imminently with the MiBioGen initiative\(^ {72}\)), the number of genetic variants associated with the various characterisations of the microbiome (i.e., bacteria-specific metabolites, bacterial taxa or functional features) will also likely grow. A greater number of genetic variants robustly associated with features of the gut microbiome will further enable the application of the continuously developing plethora of MR methods that require multiple instruments to investigate the effect of confounding, mediation, pleiotropy and invalidation of MR assumptions. However, it seems clear that the environmental contribution to the gut microbiome will be much greater than the host genetic contribution\(^ {8}\). At the very least, this calls for greater sample sizes in individual cohort studies and consortia for adequate statistical power within MR analyses testing the causal role of these environmental cues on gut microbiome variation (or, indeed, the role of the gut microbiome on disease). It is important to note that the mechanisms by which these genetic variants are associated with the gut microbiome need to be carefully considered and will rely on comprehensive functional and biological experiments and knowledge from both animal and human models that will only be possible with interdisciplinary collaboration. Furthermore, the application of MR will only be relevant to the components of the gut microbiome that are detectibly heritable and, where this is not the case, alternative approaches to interrogate causality in relationships between the gut microbiome and health traits will be required.

Understanding the complex mechanisms that link genetic variation with the gut microbiome will be particularly important.
when interpreting results obtained from MR analyses (Figure 2). As an example of this, in the GWAS conducted by Hughes et al.\textsuperscript{81}, bacteria in the *Bifidobacterium* genus were associated with the well-characterized rs4988235 lactase persistence variant at the *MCM6* locus, which is common in European populations and the only persistent signal among existing microbiome-wide GWASs. Each additional copy of the lactase persistent allele decreases the relative bacterial abundance of *Bifidobacterium*, where individuals predisposed to be lactose tolerant are likely to have a reduced average *Bifidobacterium* bacterial abundance within their gut. This observation is supported biologically as species and strains of *Bifidobacterium* can metabolise lactose (preferentially over other simple sugars), where individuals who are lactose intolerant have higher levels of *Bifidobacterium* to aid digestion, and where supplementation of *Bifidobacterium* has been shown to reduce lactose intolerance in humans\textsuperscript{73,74}. In this study, by using the rs4988235 variant as an instrument in MR analyses to interrogate the causal role of bacteria in the *Bifidobacterium* genus in the aetiology of the collection of metabolic, inflammatory and neurological traits, there was suggestive evidence of causal effects with adiposity-related traits. Specifically, a greater relative bacterial abundance of *Bifidobacterium* had a potentially causal role in lowering waist circumference, BMI and waist–hip ratio (where the association with BMI was consistent with that described by Yang et al.\textsuperscript{75}).

However, it is currently difficult to determine whether this relationship with adiposity-related traits is the direct product of variations in the relative abundance of *Bifidobacterium* (i.e., via immune modulation or the production of SCFAs, for example\textsuperscript{75,76}) or the direct impact of rs4988235 variation on adiposity via other exposures such as dietary composition (e.g., milk intake), independently of *Bifidobacterium* (i.e., horizontal pleiotropy – Figure 2D). Therefore, whilst there may appear to be a causal effect of *Bifidobacterium* on adiposity measures, this observation may be an artefact resulting in the independent impact of the rs4988235 on both *Bifidobacterium* and adiposity, which is difficult to discern without further functional knowledge of the host genetic variants being used as instruments for the gut microbiome in MR analyses. This ambiguity is particularly pertinent at a time where there are few genetic variants reliably associated with characteristics of the gut microbiome.

In addition to these MR-specific limitations, it is worth revisiting the general limitations with microbiome research that have likely driven the limited overlap of genetic variants consistently associated with gut microbiota between microbiome-wide GWASs. These include (but are certainly not limited to) differences in protocols/standards for sample collection and storage, DNA extraction method (including chosen hypervariable region for PCR and sequencing methods\textsuperscript{77}), PCR primers, and amplicon vs. shotgun sequencing\textsuperscript{83,79}. Whilst 16S rRNA amplicon sequencing is useful in providing insight into the types of bacteria present within samples, one particular issue in current studies is the limited resolution afforded by this technology. However, as studies using 16S sequencing combine for undertaking harmonized GWASs (e.g., with the MiBioGen initiative), the power afforded by larger sample sizes will be invaluable for understanding the host genetic contribution to the gut microbiome. This, in combination with future studies using data from complementary technologies (e.g., shotgun metagenomics achieving strain-level resolution, metatranscriptomics, proteomics and metabolomics) able to provide more refined measures of components and functionality of the gut microbiome, will enable a more comprehensive understanding of the role played by these microorganisms in health and disease and the mechanisms by which these occur. In addition, whilst most studies have focused on the bacterial component of the gut microbiome, it is important to interrogate the causal role that other integral microorganisms (i.e., fungi, viruses and archaea) play in the development and progression of host disease and health outcomes over the lifecourse.

**Conclusions**

MR is an established approach that uses human genetic variation to estimate causal associations in observational epidemiological relationships and can be used to provide further insight into the causal relevance of the gut microbiome in human health and disease. The applications discussed here currently flag the potential of MR analyses following the growing collection of genetic association data for the human gut microbiome, but there are also important issues likely to arise with a naïve integration of complex GWAS results to understanding causes of health outcomes without in-depth knowledge of the host genetic variants themselves and the performance and pitfalls of MR methodology\textsuperscript{41,90}. Whilst MR was motivated to confer certain advantages over traditional epidemiological study designs, like any study design within epidemiology, it is not exclusively adequate to conclusively demonstrate causality. Therefore, there is a continued need for triangulation across multiple traditional epidemiological approaches and interdisciplinary collaboration to support or challenge causality of the role played by the gut microbiome on human health and to understand the mechanisms by which these relationships occur. Such partnerships are necessary to maximise translation into the development of new, targeted therapies to alleviate disease symptoms to ultimately improve lives, and promote good health whilst preventing ill health\textsuperscript{81}.

**Data availability**

No data are associated with this article.

**Author contributions**

KHW conceptualised and wrote the initial draft of the manuscript. LJH and KHW revised and approved the final version.

**Acknowledgements**

We would also like to thank Tom Battram and Nic Timpson for their input on initial drafts, from whom we have obtained permission to acknowledge in this manuscript.
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Open Peer Review

Current Peer Review Status: ✔

Version 2

Reviewer Report 21 April 2020

https://doi.org/10.21956/wellcomeopenres.17347.r38333

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I thank the authors for incorporating my suggestions.
I only want to get back again at point 4, because I think the explanation given on the study doesn't fully reflect the original design. I suggest the following:

“Genetic variants were chosen by conducting a GWAS within 952 individuals from the LifeLines-DEEP cohort. A fairly lenient p-value threshold of 1x10-5 was used to define host genetic variants independently associated with functional features of the gut microbiome. Genetic variants selected at this threshold explained most variance of the same feature in an independent cohort of 445 individuals, compared to variants selected with different thresholds”

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Genetic mapping, Statistical Genetics, Mendelian randomisation methods, population based microbiome studies.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 21 Apr 2020

Kaitlin Wade, University of Bristol, Bristol, UK

We have incorporated the changes, as requested by the Reviewer.

Competing Interests: N/A
Serena Sanna

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This review is clearly written and comprehensive. It explains the urgent need in identifying causal relationships between microbiome and diseases before moving to real clinical application and to commercial kits, and then introduces the Mendelian Randomization approach that has been recently used in this context. The authors present findings and limitations of these studies, and also more in general the pros and cons of Mendelian Randomization in application to microbiome.

I have few minor suggestions for further improvement:

- **Introduction.** When discussing the potential role of microbiome modulation as therapy, I think it is important to notice that despite causality can be proved (either by MR or mechanistic studies in mice/humans), we have to 1) remember that gut microbiome is an ecosystem and thus we need to also investigate carefully the side effects of any type intervention intended to module one specific bacteria and 2) remember that evidence for causal relationship with a disease doesn’t necessarily mean that the scenario is reversible, in other words, that modulating that specific bacteria/function will not necessarily cure the disease (while remains a valuable information in terms of prevention).

- **MR paragraph.** Please remark that MR can be run using limited time and reduced costs because it takes advantage of pre-existing large scale genetic studies (otherwise the cost and time are not that limited, even if still less than many RCT).

- **MR paragraph – limitations.** Please add a note that MR is a great source but only applicable to those bacteria that are modulated by genetic of the host – only a small fraction of bacteria are, so alternative approaches to detect causality are still much needed.

- **Page 6,** please amend the details of the study by Sanna et al. (2019). In this study, the number of individuals used to carry out the GWAS was 952. An additional independent cohort of 445 individuals was used to derive the optimal threshold for selecting the instruments; specifically this was chosen as the threshold that selected the set of independent variants that, when used in a polygenic risk score in the independent cohort, explained the most of the variance of the same microbiome feature. [This point explains my answer “partly” to the question “Are all factual statements correct and adequately supported by citations?”]
Page 7 second paragraph. Please add the note that some of the complimentary approaches to inform MR (for example the bivariate finemapping utilized by Richardson et al.) – require individual level data of both the exposure and the outcome, and thus are currently impractical to apply to diseases, as there are not such large scale studies of microbiome on case-control studies.

Page 2, last sentence of the second paragraph. I suggest to add a sentence remarking that to solve this ambiguity, large studies are extremely needed as they will likely lead to multiple associated SNPs, which will help MR to better investigate effect of confounders.

Is the topic of the review discussed comprehensively in the context of the current literature? Yes

Are all factual statements correct and adequately supported by citations? Partly

Is the review written in accessible language? Yes

Are the conclusions drawn appropriate in the context of the current research literature? Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Genetic mapping, Statistical Genetics, Mendelian randomisation methods, population based microbiome studies.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Kaitlin Wade, University of Bristol, Bristol, UK

We wanted to thank the Reviewer for their kind words on the writing and understanding of the importance of this area. We hope that this report provides understandable insight into the Mendelian randomization approach and how this may help further integrate epidemiology and microbiology and interrogate causal relationships between the microbiome and health outcomes. We also thank the Reviewer on their suggestions for improvement, which we have addressed below. For each comment made the Reviewer (in bold) we have responded and added the italicized excerpt.

1. Introduction. When discussing the potential role of microbiome modulation as therapy, I think it is important to notice that despite causality can be proved (either by MR or mechanistic studies in mice/humans), we have to 1) remember that gut microbiome is an ecosystem and thus we need to also investigate carefully the side effects of any type intervention intended to module one specific bacteria and 2) remember that evidence for causal relationship with a disease doesn’t necessarily mean that the scenario is reversible, in other words, that modulating that specific bacteria/function will not necessarily cure the disease (while remains a valuable information in terms of prevention).
The Reviewer makes an excellent point regarding the complexity of the gut microbiome and the implication of this complexity within possible therapies and interventions. We have added a sentence to the introduction to incorporate this point, as follows:

“However, with the development of such efforts, it is important to recognise that the gut microbiome is a dynamic and complex ecosystem; therefore, careful investigation of the off-target effects of any treatment or intervention intended to alter one or a small number of specific bacteria is required.”

Additionally, the Reviewer’s point regarding implications of findings to disease prognosis is valuable and, indeed, an important consideration when interpreting findings from all epidemiological studies focusing on disease risk in comparison to disease prognosis. We totally agree that, even if evidence points to a causal impact of the gut microbiome on the risk of a disease, this certainly does not imply that the same is true for the progression of that disease (i.e., treating individuals who suffer from a disease with, say, a pre-/pro-biotic that promotes/contains bacteria associated with the risk of that disease may not lead to a better prognosis – or, as the Reviewer states, cure that disease). This is an important distinction to make and is only resolved by specifically assessing the association between the gut microbiome and disease progression, prognosis or survival (rather than disease risk). As it stands, the application of Mendelian randomization to the study of disease progression is in its infancy, with methods being developed to address the additional caveats of studying populations selected on case-only status, as outlined by Paternoster et al. (e.g., that proposed by Dudbridge et al. and the Slope-Hunter method proposed by Mahmoud et al.). To address this comment, we have stated the following to emphasize this point.

“Furthermore, if evidence of a likely causal effect of a component of the gut microbiome on the risk of a disease is provided, this does not imply that the same is true for the progression of that disease (and vice versa). Specifically, treating individuals who suffer from, say, IBD with a pre-/pro-biotic that promotes/contains bacteria found to be associated with the risk of developing IBD may not lead to better prognosis of IBD after diagnosis.”

2. MR paragraph. Please remark that MR can be run using limited time and reduced costs because it takes advantage of pre-existing large scale genetic studies (otherwise the cost and time are not that limited, even if still less than many RCT).

We have edited a sentence in the MR paragraph to clarify this point:

“Such analyses can be done at a fraction of the time and cost required for a large-scale RCT, by exploiting data from pre-existing and large-scale genetic studies, and can be used to ask many of the scientific questions that may not be feasible or ethical with RCTs[45, 46].”

3. MR paragraph – limitations. Please add a note that MR is a great source but only applicable to those bacteria that are modulated by genetic of the host – only a small fraction of bacteria are, so alternative approaches to detect causality are still much needed.

We have added the following sentence to the appropriate section:

“Furthermore, the application of MR will only be relevant to the components of the gut microbiome
that are detectibly heritable and, where this is not the case, alternative approaches to interrogate causality in relationships between the gut microbiome and health traits will be required.”

4. Page 6, please amend the details of the study by Sanna et al. (2019). In this study, the number of individuals used to carry out the GWAS was 952. An additional independent cohort of 445 individuals was used to derive the optimal threshold for selecting the instruments; specifically this was chosen as the threshold that selected the set of independent variants that, when used in a polygenic risk score in the independent cohort, explained the most of the variance of the same microbiome feature. [This point explains my answer “partly” to the question “Are all factual statements correct and adequately supported by citations?”]

We thank the Reviewer for clarifying the sample sizes on this discussed paper. We have edited the referenced sentence as follows in light of this information:

“Genetic variants were chosen by conducting a GWAS within 952 individuals from the LifeLines-DEEP cohort. A fairly lenient p-value threshold of 1x10^{-5} (identified from an independent cohort of 445 individuals) was used to define host genetic variants independently associated with functional features of the gut microbiome and explained most variance in the same feature.”

5. Page 7 second paragraph. Please add the note that some of the complimentary approaches to inform MR (for example the bivariate finemapping utilized by Richardson et al.) – require individual level data of both the exposure and the outcome, and thus are currently impractical to apply to diseases, as there are not such large scale studies of microbiome on case-control studies.

We have added the following sentence to reflect this.

“As these methods (particularly colocalization and genetic fine mapping) require individual-level and genome-wide information on both the exposure and outcome, these methods will become more feasible with the growing availability of large-scale GWASs of the gut microbiome and other traits.”

6. Page 2, last sentence of the second paragraph. I suggest to add a sentence remarking that to solve this ambiguity, large studies are extremely needed as they will likely lead to multiple associated SNPs, which will help MR to better investigate effect of confounders.

The Reviewer makes an important point and we have added the following sentence to a paragraph in the limitations section to address this:

“A greater number of genetic variants robustly associated with features of the gut microbiome will further enable the application of the continuously developing plethora of MR methods that require multiple instruments to investigate the effect of confounding, mediation, pleiotropy and invalidation of MR assumptions.”

**Competing Interests:** N/A