A unified catalog of 204,938 reference genomes from the human gut microbiome

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Comprehensive, high-quality reference genomes are required for functional characterization and taxonomic assignment of the human gut microbiota. We present the Unified Human Gastrointestinal Genome (UHGG) collection, comprising 204,938 non-redundant genomes from 4,644 gut prokaryotes. These genomes encode >170 million protein sequences, which we collated in the Unified Human Gastrointestinal Protein (UHGP) catalog. The UHGP more than doubles the number of gut proteins in comparison to those present in the Integrated Gene Catalog. More than 70% of the UHGG species lack cultured representatives, and 40% of the UHGP lack functional annotations. Intraspecies genomic variation analyses revealed a large reservoir of accessory genes and single-nucleotide variants, many of which are specific to individual human populations. The UHGG and UHGP collections will enable studies linking genotypes to phenotypes in the human gut microbiome.
Results

More than 200,000 human gut microbial genomes in the UHGG catalog. We first gathered all prokaryotic isolate genomes and MAGs from the human gut microbiome (publicly available as of March 2019). We compiled the isolate genomes from the Human Gastrointestinal Bacteria Culture Collection (HBC)19 and the Culturable Genome Reference (CGR)21, as well as cultured human gut genomes available in the NCBI22, PATRIC23 and IMG24 repositories, which include genomes from several other large studies1,2,23. In addition, we included all of the gut MAGs generated in Pasolli et al.29 (CIBIO), Almeida et al.28 (EBI) and Nayfach et al.16 (HGM). To standardize the genome quality across all sets, we used thresholds of >50% genome completeness and <5% contamination, combined with an estimated quality score (completeness×5×contamination) > 50. The final numbers of genomes matching these criteria were 734 (HBC), 1,519 (CGR), 651 (NCBI), 7,744 (PATRIC/IMG), 137,474 (CIBIO), 87,386 (EBI) and 51,489 (HGM), resulting in a total of 286,997 genome sequences (Fig. 1a and Supplementary Table 1). These represented 204,938 nonredundant genomes on the basis of a Mash26 distance threshold of 0.001 (99.9% nucleotide identity) and only considering one genome per species per sample to account for the fact that the three large MAG studies analyzed many samples in common. Genomes were recovered in samples from a total of 31 countries across six continents (Africa, Asia, Europe, North America, South America and Oceania), but the majority originated from samples collected in China, Denmark, Spain and the United States (Fig. 1b).

To determine how many species were included in this gut reference collection, we clustered all 286,997 genomes using a multistep distance-based approach (Methods) with an average nucleotide identity (ANI) threshold of 95% over at least a 30% alignment fraction (AF)25. The clustering procedure resulted in a total of 4,644 inferred prokaryotic species (4,616 bacterial and 28 archaeal; Supplementary Table 2). We found the species clustering results to be highly consistent with those previously obtained16,18,20 (Supplementary Table 3). The best quality genome from each species cluster was selected as its representative on the basis of genome completeness, minimal contamination and assembly N50 (with isolate genomes always given preference over MAGs), and the final set was used to generate the UHGG catalog (Fig. 1c). Of the 4,644 species-level genomes, 3,207 were >90% complete (interquartile range, IQR = 87.2–98.8%) and <5% contaminated (IQR = 0.0–1.34%), with 573 of these having the rRNA operon satisfied the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species. 38 from MAGs) satisfy the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species. 38 from MAGs) satisfy the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species. 38 from MAGs) satisfy the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species. 38 from MAGs) satisfy the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species. 38 from MAGs) satisfy the ‘high quality’ criteria set for MAGs by the Genomic Standards Consortium28. The rRNA operon for these species is specific and representative of the species.
Toolkit30 (GTDB-Tk; Extended Data Fig. 2), a standardized taxonomic framework based on a concatenated protein phylogeny representing >140,000 public prokaryote genomes, fully resolved to the species level (see Methods for details on the taxonomy nomenclature used). However, over 60% of the gut genomes could not be assigned to an existing species, confirming that the majority of the UHGG species lack representation in current reference databases.

To obtain further insights into the quality of the UHGG genomes, we inferred the level of strain heterogeneity within each MAG with CMseq32. The median strain heterogeneity (proportion of polymorphic positions) of the UHGG MAGs was 0.06% (IQR = 0.01–0.25%; Extended Data Fig. 1c and Supplementary Table 1), which is below the 0.5% threshold defined previously32 to distinguish medium- from high-quality MAGs. We believe that this additional metric on strain heterogeneity is a useful complement to the standard completeness and contamination estimates, providing further evidence of the overall high quality of the genomes included here.

Comparison of species recovered in individual studies. We investigated how many of the 4,644 gut species were found in the different study collections to determine their level of overlap and reproducibility, as well as the ratio between cultured and uncultured species (Fig. 2a). Each of the large MAG studies used a different assembly and binning approach: the CIBIO study used metaSPAdes31 and MetaBAT 2 (ref. 32) for assembling and binning sequencing runs previously merged by sample; the HGM study used MEGAHIT33 to assemble runs merged by sample and applied a combination of MaxBin 2 (ref. 34), MetaBAT 2 (ref. 32), CONCOCT35 and DAS Tool36 for binning and refinement; and the EBI study used metaSPAdes31 and MetaBAT 2 (ref. 32) for assembling and binning individual runs without merging by sample. Despite these methodological differences, the largest intersection found was between these collections of MAGs, with the same 1,081 species detected independently in the CIBIO, EBI and HGM datasets, but not in any of the cultured genome studies. By restricting the analysis to genomes recovered from 1,554 samples common to all three MAG studies, we found that 93–97% of the species from each set were detected in at least one other MAG collection and 79–86% were detected across all three (Extended Data Fig. 3a). A similar level of species overlap was observed when comparing studies on a per-sample basis (Extended Data Fig. 3b). Furthermore, conspecific genomes recovered from the same samples across different studies had a median ANI and AF of 99.9% and 92.1%, respectively (94.5% AF with ≥90% complete genomes and 86.6% AF with medium-quality genomes; Extended Data Fig. 3c). These results suggest that the large-scale studies of human gut MAGs37,38,39 generally recovered highly similar genomes. However, the smaller AF values detected among genomes that were <90% complete suggest that caution is needed when using medium-quality genomes in downstream analyses.

Rarefaction analysis indicated that the number of uncultured species detected has not reached a saturation point, meaning that additional species remain to be discovered (Fig. 2b). However, these most likely represent rarer members of the human gut microbiome, as the number of species is closer to saturating when only considering those with at least two conspecific genomes.

We also investigated the intersection between the three large culture-based datasets: the HBC, CGR and NCBI (which contains gut genomes from the HMP). Unlike the MAGs, the majority of cultured species were unique within a single collection (486/698; 70%), with only 70 (10%) common to all three collections (Extended Data Fig. 3d). This may be due to varied geographic sampling between the collections (Asia, Europe and North America) or might highlight the stochastic nature of culture-based studies.

Most gut microbial species lack isolate genomes. We found that 3,750 (81%) of the species in the UHGG catalog did not have a representative in any of the human gut culture databases. To extend the search to isolate genomes from other environments or lacking information on the isolation source, we compared the UHGG catalog to all NCBI RefSeq isolate genomes. We identified an additional set of 438 species closely matching cultured genomes (88 from human body sites, 29 from other animals, 3 from plants and the remainder (318) from unknown sources), leaving 3,312 (71%) UHGG species as uncultured (Supplementary Table 2).

By calculating the number of genomes contained within each cultured and uncultured human gut species, we found that species containing isolate genomes represented the largest clusters, while those exclusively encompassing MAGs tended to be the rarest, as discussed previously32,33,34. For example, only 2 of the 25 largest bacterial clusters were exclusively represented by MAGs (Fig. 2c), with 1,212 uncultured species represented by a single genome (80% of which originated from samples only analyzed in one of the MAG studies; Extended Data Fig. 4). The bacterial species most represented in our collection were Agathobacter rectalis (recently reclassified from Eubacterium rectale2), Escherichia coli D and Bacteroides uniformis (Fig. 2c, Extended Data Fig. 5a and Supplementary Table 2), whereas the most frequently recovered archaeal species was Methanobrevibacter smithii, with 608 genomes found across all six continents (Extended Data Fig. 6). We inferred the level of geographic diversity of each species by calculating the Shannon diversity index on the proportion of samples in which each species was found per continent. The largest species clusters displayed similarly high levels of geographic distribution, indicating that the most highly represented species were not restricted to individual locations (Fig. 2c and Extended Data Fig. 5b).

We determined how representative the UHGG catalog is of the human gut microbial diversity by mapping 1,005 independent metagenomic datasets against the 4,644 UHGG species (Fig. 2d and Supplementary Table 4). Using Kraken 2 (ref. 38), we obtained a median classification rate of 85.9% (IQR = 83.5–88.1%). Notably, this corresponded to a median improvement of 155% over the standard RefSeq database. The increase in classification rate was most pronounced in non-Western samples from Cameroon, Ethiopia, Ghana and Tanzania, highlighting the potential of the UHGG catalog to improve the study of microbiome diversity from these understudied populations.

The phylogenetic distribution of the 4,616 bacterial (Fig. 3a) and 28 archaean (Extended Data Fig. 6) species revealed that uncultured species exclusively represented 66% and 31% of the phylogenetic diversity of Bacteria and Archaea, respectively, with several phyla lacking cultured representatives (Fig. 3b). The four largest monophyletic groups lacking cultured genomes were the 4C28d-15 order (167 species, recently proposed as the novel order Comantemales ord. nov.40; Fig. 3c), order RF39 (139 species), family CAG-272 (88 species) and order Gastranaerophilales (67 species). While none have been successfully cultured, several have been described in the literature, including for RF39 (ref. 41) and Gastranaerophilales (previously classified as a lineage in the Melainabacteria42), which are characterized by highly reduced genomes with numerous auxotrophies. This analysis suggests that, despite recent culture-based studies31,32,33,34, much of the diversity in the gut microbiome remains uncultured, including several large and prevalent clades.

Expanding the set of proteins in the human gut microbiome. Metagenomic approaches have the ability to leverage gene content information not only for more precise taxonomic analysis but also to predict the functional capacity of individual species of interest in comparison to marker-gene-based methods (for example, relying solely on the 16S rRNA gene or a limited number of diagnostic genes). We built the UHGP catalog with a total of 625,255,473 full-length protein sequences predicted from the 286,997 analyzed genomes herein. These were clustered at 50% (UHGP-50), 90%
(UHGP-90), 95% (UHGP-90) and 100% (UHGP-100) amino acid identity, generating between 5 to 171 million protein clusters (Fig. 1c and Extended Data Fig. 7a). While the number of UHGP-95 and UHGP-90 clusters showed a steady increase as a function of the number of nonredundant genomes analyzed. Curves are depicted both for all the UHGG species and after excluding singleton species (represented by only one genome). c, Number of nonredundant genomes detected per species (left) alongside the degree of geographic diversity (calculated with the Shannon diversity index; right). Only the 25 most represented species clusters are depicted. d, Left, proportion of metagenomic reads from 1,005 independent datasets classified with Kraken 2 against the UHGG species representatives. Right, the degree of classification improvement provided over the standard Kraken 2 RefSeq database. The following correspond to the number of datasets analyzed per country: Cameroon, n = 54; Ethiopia, n = 25; Germany, n = 56; Ghana, n = 40; India, n = 105; Italy, n = 50; Luxembourg, n = 26; Russia, n = 4; Tanzania, n = 61; United Kingdom, n = 210; United States, n = 374. Box lengths represent the IQR of the data, and whiskers extend to the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively.

To determine how comprehensive the UHGG is when compared to existing human gut gene catalogs, we combined the UHGP-90 (n = 13,910,025 protein clusters) with the IGC, a collection of 9.9 million genes from 1,267 gut metagenome assemblies, which we grouped into 7,063,981 protein clusters at 90% protein identity (referred to as IGC-90). Nearly all samples used to generate the IGC were also included in the UHGP catalog (except for 59 transcriptome datasets), but the latter was generated from a larger and more geographically diverse metagenomic dataset (including samples from Africa, South America and Oceania). Combining the UHGP-90 and IGC-90 resulted in a set of 15.2 million protein clusters, with an overlap of 5.8 million sequences (Fig. 4b). This revealed that 81% of the IGC is represented in the UHGP catalog, with the missing 19% likely representing fragments of prokaryotic genomes that are <50% complete and viral or eukaryotic sequences, plasmids or other sequences not binned into MAGs. In fact, only 0.2% (n = 34,070 clusters) of the UHGP-90 was predicted to be of viral origin (on the basis of eggNOG annotations), as compared to the 5% estimate obtained in a previous human gut gene catalog included in the IGC.

Most notably, though, the UHGP provided an increase of 115% in coverage of the gut microbiome protein space over the IGC (from 7,063,981 to a total of 15,217,595 protein clusters).
We also compared the read mapping rate using the same 1,005 metagenomic samples tested against the UHGG catalog (Supplementary Table 4). Even though the classification rate was substantially higher when using the UHGG catalog than with RefSeq, the increase with the UHGG-90 over IGC-90 was more modest (median of 5%; Extended Data Fig. 7b). These results suggest that, although the UHGP collectively encompasses a much larger number of protein clusters, most of the newly added proteins are at lower abundance/prevalence within individual samples. However, as the UHGP was generated from individual genomes and not from their original unbinned metagenome assemblies, our catalog also has the advantage of providing a direct link between each gene cluster and its genome of origin. To this end, we have also generated high-quality subsets of the UHGP-95, UHGP-90 and UHGP-50 consisting of protein clusters where at least two proteins from different genomes belonging to the same species were retrieved (UHGP-95-HQ, \( n = 10,798,224 \); UHGP-90-HQ, \( n = 8,082,122 \); UHGP-50-HQ, \( n = 3,088,278 \)). This clustering criterion was used to control for the presence of contaminating sequences within each MAG and for the possibility that multiple copies of the same protein-coding sequence may be present in one genome. The UHGP ultimately allows the combination of individual genes with their genomic context for an integrated study of the gut microbiome.

**Functional capacity of the human gut microbiota.** We used the eggNOG\(^4\), InterPro\(^4\), COG\(^4\) and KEGG\(^4\) annotation schemes to capture the full breadth of functions within the UHGP. However, we found that 41.5% of UHGP-100 was poorly characterized, as 27.3% lacked a match to any database and a further 14.2% only had a match to a COG with no known function (Fig. 4c). On the basis of the distribution of COG functions, the most highly represented categories were related to amino acid transport and metabolism, cell wall/membrane/envelope biogenesis and transcription.

We further leveraged the set of 171 million proteins derived from the human gut genomes to explore the functional diversity within each of the UHGG species. Protein sequences from all conspecific genomes were clustered at 90% amino acid identity to generate a pan-genome for each species. Analysis of the functional capacity of the UHGG species pan-genomes identified a total of 363 KEGG modules encoded by at least one species (Extended Data Fig. 8a and
We analyzed the pan-genome size for each species in relation to the number of conspecific genomes to look for differences in intraspecific gene richness. We observed distinct patterns across different gut phyila, with species from various Firmicutes clades showing the highest rates of gene gain (Fig. 5a). There was wide variation in the proportion of core genes between species even among clades with more than 1,000 genomes (Fig. 5b), with a median core genome proportion (percentage of core genes among all genes in the representative genome) estimated at 66% (IQR = 59.6–73.9%).

To distinguish the functions encoded in the core and accessory genes, we analyzed their associated annotations. Core genes were well covered, with a median of 96%, 94%, 92% and 69% of the genes assigned with an eggNOG, InterPro, COG and KEGG annotation, respectively (Fig. 5c). In contrast, the accessory genes had a significantly higher proportion of unknown functions (P < 0.001) with key metabolic functions involved in nucleotide, amino acid and lipid metabolism, as well as other housekeeping functions (for example, related to translation and ribosomal structure; Fig. 5d). In contrast, accessory genes had a much greater proportion of COGs without a known function and of genes involved in replication and recombination, which are typically found in mobile genetic elements (MGEs; Fig. 5d). A significant number of accessory genes were related to defense mechanisms, which encompass not only general mechanisms of antimicrobial resistance (AMR) such as ABC transporter efflux pumps but also systems targeted toward invading MGEs (for example, CRISPR–Cas and restriction modification systems against bacteriophages). These results highlight the potential of this resource to provide better understanding of the dynamics of chromosomally encoded

Patterns of intraspecies genomic diversity. With the protein annotations and pan-genomes inferred for each of the UHGG species, we explored their intraspecies core and accessory gene repertoire. Only near-complete genomes (≥90% completeness) and species with at least ten independent conspecific genomes were analyzed. The overall pattern of gene frequency within each of the 781 species considered here showed a distinctive bimodal distribution (Extended Data Fig. 9), with most genes classified as either core or rare (that is, present in ≥90% or <10% of conspecific genomes, respectively). We analyzed the pan-genome size for each species in relation to the number of conspecific genomes to look for differences in intraspecies gene richness. We observed distinct patterns across different gut phyila, with species from various Firmicutes clades showing the highest rates of gene gain (Fig. 5a). There was wide variation in the proportion of core genes between species even among clades with more than 1,000 genomes (Fig. 5b), with a median core genome proportion (percentage of core genes among all genes in the representative genome) estimated at 66% (IQR = 59.6–73.9%).

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AMR within the gut and allow deciphering of the extent to which the microbiome may be a source of both known and novel resistance mechanisms.

We next investigated intraspecies single-nucleotide variants (SNVs) within the UHGG species. We generated a catalog consisting of 249,435,699 SNVs from 2,489 species with three or more conspecific genomes (Fig. 6a). For context, a previously published catalog contained 10.3 million single-nucleotide polymorphisms from 101 gut microbiome species\textsuperscript{65}. Of note, more than 85% of these SNVs were exclusively detected in MAGs, whereas only 2.2% were exclusive to isolate genomes (Fig. 6b). We found the overall pairwise SNV density between MAGs to be higher than that observed between isolate genomes (Fig. 6c). This was irrespective of the level of strain heterogeneity of the MAGs, as there was no correlation between SNV density and the degree of strain heterogeneity estimated with CMseq (Extended Data Fig. 10). Next, we assigned the detected SNVs to the

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**Fig. 5 | Pan-genome diversity patterns within the gut microbiome.** a. Normalized pan-genome size as a function of the number of conspecific genomes. Regression curves were generated for each phylum, with the corresponding coefficients of determination indicated next to each curve and the shaded regions representing the 95% confidence level intervals. The following correspond to the number of species considered for each phylum: Actinobacteriota, \( n = 66 \); Bacteroidota, \( n = 122 \); Firmicutes, \( n = 90 \); Firmicutes A, \( n = 325 \); Firmicutes C, \( n = 44 \); Proteobacteria, \( n = 65 \); Verrucomicrobiota, \( n = 13 \). b. Fraction of the core genome for each species according to the number of conspecific genomes (left) and as a histogram (right), colored by phylum. The horizontal dashed line represents the median value across all species. c. Proportion of core and accessory genes (\( n = 781 \) species) classified with various annotation schemes, alongside the percentage of genes lacking any functional annotation. Box lengths represent the IQR of the data, and whiskers extend to the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively. A two-tailed Wilcoxon rank-sum test was performed to compare the classification between the core and accessory genes (**\( P < 0.001 \)). d. Comparison of the functional categories assigned to the core and accessory genome (**\( P < 0.001 \)).
continent of origin of each genome and observed that 36% of the SNVs were continent specific. Notably, genomes with a European origin contributed to the most exclusive SNVs (Fig. 6d). However, genomes from Africa contributed over three times more variation on average than European or North American genomes. Pairwise SNV analysis also supported a higher cross-continent SNV density, especially between genomes from Africa and Europe (Fig. 6e). SNV analysis also supported a higher cross-continent SNV density, especially between genomes from Africa and Europe (Fig. 6e).

Resource implementation. Both the UHGG and UHGP catalogs are available as part of a new genome layer within the MGnify website, where summary statistics of each species cluster and their functional annotations can be interactively explored and downloaded (see ‘Data availability’ for more details). We have generated a Bitsliced Genomic Signature Index (BIGSI) of the UHGG catalog, which allows users to interactively query sequence fragments <5 kb in length to search for similar sequences in this collection.

We plan to periodically update the resource (approximately every 6–12 months) as new genomes are generated and made publicly available. MAGs will be retrieved from the European Nucleotide Archive (ENA), where a new MAG analysis class was recently implemented. Genomes (MAGs or isolates) will be incorporated in the resource either as new species or by replacing uncultured reference genomes with better quality versions. We will adopt a versioning system whereby previous iterations of the catalog will still be accessible after subsequent updates to ensure reproducibility.

Discussion

We have generated a unified sequence catalog representing over 200,000 genomes and 171 million protein sequences of the human gut microbiome. Of the 4,644 species contained in the UHGG
catalog, 71% lack a cultured representative, meaning that the majority of microbial diversity in the catalog remains to be experimentally characterized. During preparation of our manuscript, a new collection of almost 4,000 cultured genomes from 106 gut species was released10, which will be incorporated in future versions of the resource. As 96% of these genomes were reported to have a species representative in the culture collections included here, we do not anticipate that this dataset will provide a substantial increase in the number of species discovered. Nevertheless, our analyses suggest that additional uncultured species from the human gut microbiome are yet to be discovered, highlighting the importance and need for culture-based studies. Furthermore, given the sampling bias toward populations from China, Europe and the United States, we expect that many under-represented regions still contain substantial uncultured diversity.

By comparing recently published large datasets of uncultured genomes11,12,13, we were able to assess the reproducibility of the results from each study. We show that, despite the different assembly, binning and refinement procedures used in the three studies, almost all of the same species and similar strains were recovered independently when using a consistent sample set. Although these results increase confidence in the use of MAGs, new methods for metagenome assembly, binning and quality control continue to be developed to overcome existing limitations, meaning that improved versions of the MAGs included here will likely be generated in the future.

With the establishment of this massive sequence catalog, it is evident that a large portion of the species and functional diversity within the human gut microbiome remains uncharacterized. Moreover, knowledge of the intraspecies diversity of many species is still limited owing to the presence of a small number of conspecific genomes. Having this combined resource can help guide future studies and prioritize targets for further experimental validation. Using the UHGG or UHGP catalogs, the community can now screen for the prevalence and abundance of species or genes in a large panel of intestinal samples and in specific clinical contexts. By pinpointing particular taxonomic groups with biomedical relevance, more targeted approaches could be developed to improve understanding of their role in the human gut. The functional predictions generated for the species pan-genomes could also be leveraged to develop new culturing strategies for isolation of candidate species. Target-enrichment methods such as single-cell14 and/or bait-capture hybridization15 approaches could also be applied. Given the large uncultured diversity still remaining in the human gut microbiome, having a high-quality catalog of all currently known species substantially enhances the resolution and accuracy of metagenome-based studies. Therefore, the presented genome and protein catalogs represent a key step toward a hypothesis-driven, mechanistic understanding of the human gut microbiome.

**Online content**

Any methods, additional references, Nature Research report- ing summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41587-020-0603-3.

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Methods

**Genome collection.** We compiled all the prokaryotic genomes publicly available as of March 2019 that were sampled from the human gut. To retrieve isolate genomes, we surveyed the IMG, NCBI, and PATRIC databases for genome sequences annotated as having been isolated from the human gastrointestinal tract. We complemented this set with bacterial genomes belonging to two recent culture collections: the HBC and CGR. To avoid including duplicate entries due to redundancy between reference databases, we combined genomes obtained from the PATRIC and IMG repositories and added only those without an identical genome in the set extracted from NCBI, HBC, and CGR. This was determined by comparing isolate genomes between different databases using Mash v2.1 (ref. 19; mash dist function) and only selecting one genome among those estimated to be identical (Mash distance of 0). MAGs (that is, uncultured genomes) were obtained from Pasolli et al. (CIBIO), Almeida et al. (EBI) and Nayfach et al. (HGM). For the CIBIO set, only genomes retrieved from samples collected from the intestinal tract were used. MAGs for the CIBIO set were first retrieved from the five large human gut studies (refs. 16–18). These were further enriched with data obtained using the ENA API (https://www.ebi.ac.uk/ena/portal/api) and the NCBI E-utilities (http://eutils.ncbi.nlm.nih.gov/). Metadata on the isolate genomes from IMG and PATRIC were retrieved using the GOLD system and the PATRIC FTP website (ftp://ftp.patriscb.org/patric2/current_release/RELEASE_NOTES_genome_metadata), respectively. We only extracted metadata on the geographic origin of each genome, as other factors such as disease status and demographic information were missing from most of the samples.

**Assessing genome quality.** Assembly statistics were calculated with the `stats.sh` script from BBMap v38.75 (https://sourceforge.net/projects/bbmap/). Genome quality (completeness and contamination) was estimated with CheckM v1.0.11 (ref. 20) using the ‘lineage_wf’ workflow to select only genomes that passed the following criteria: > 50% genome completeness, < 5% contamination and an estimated quality score (completeness × 5 - contamination) > 50. We also searched for the presence of rRNAs in each genome with the `rnam中小型 function of INFERNAL v1.1.2 (ref. 21) with options ‘-z Z1000 -hmmonly -cut_ga -noali -blout’ against the Rfam55 covariance models for the 5S, 16S and 23S rRNAs. tRNAs of the standard 20 amino acids were identified with tRNAscan-SE v2.0 (ref. 22) with options ‘-A -Q’ for archaeal species and ‘-B’ for species belonging to bacterial lineages.

To investigate the level of strain heterogeneity represented within each MAG, we used the CMseq tool (https://github.com/SegataLab/cmseq) as previously described24. Briefly, metagenomic reads from the sample used to generate the MAG were aligned to the respective MAG using bowtie v2.2.3 (ref. 25), with the resulting alignments file indexed and sorted with samtools v1.5 (ref. 25). The level of strain heterogeneity was estimated with the ‘polymorph.py’ script from the CMseq package by calculating the number of nonsynonymous substitutions detected out of all positions mapped with a depth of coverage of at least 10 reads and base quality of at least 30 (a minimum of 100 positions were needed to estimate strain heterogeneity).

**Species clustering.** We clustered the total set of 286,997 genomes at each estimated species level (ANI ≥ 95%; ref. 26) using dRep v2.2.4 (ref. 27) with the following options: ‘-p 0.9 -a 0.95 -nc 0.30 -cm larger’ and ‘-b 0.9 -Q’ for archaeal species and ‘-B’ for species belonging to bacterial lineages. To reduce the final computational load, we used the CMseq tool from BBMap v38.75 (https://sourceforge.net/projects/bbmap/) to calculate the score. The score was calculated as follows:

\[
\text{Score} = \text{CMP} - 5 \times \text{CNT} + 0.5 \times \log(N50) \]

where CMP represents the completeness level, CNT is the estimated contamination and N50 is the assembly contiguity characterized by the minimum contig size in which half of the total genome sequence is contained. The genome with the highest score was chosen as the species representative, with cultured genomes prioritized over uncultured genomes (that is, if a MAG had a higher score than an isolate genome, the latter would still be chosen as the representative). To further investigate the within-species population diversity, we calculated pairwise distances for all conspecific genomes using Mash v2.1 (ref. 19; default sketch size). From these results, we generated individual distance trees for each species using the ‘complete’ hierarchical clustering method implemented in the Fastcluster R package28. We calculated the number of clusters recovered using a distance cutoff of 0.03 (97% ANI) and 0.01 (99% ANI).

**Evaluating reproducibility of the methods.** The species clusters inferred here were compared with those previously generated in human gut MAG studies16,18,20 from a common set of genomes. Similarity between species clusters was estimated using the adjusted Rand index (ARI) computed in the Scikit-learn Python package29. This metric considers both the number of clusters and cluster membership to compute a similarity score ranging from 0 to 1.

Conspecific genomes recovered in the same metagenomic samples but in different studies were compared with FastANI v1.1 (ref. 30) with default parameters to obtain both the maximum AF and ANI for each pairwise comparison.

**Inferring cultured status.** To determine cultured status, the UHGG species representatives were searched against NCBI ReSeQ release 93 after excluding uncultured genomes (that is, metagenome-assembled or single-cell amplified genomes). Genome alignments were performed in two stages: (1) Mash v2.1 (ref. 21) was used as an initial screen (using the function ‘mash dist’) to identify the most similar ReSeQ genome to each of the UHGG species and (2) ‘dnadiff’ from MUMmer v4.0.1bata2 (ref. 31) was subsequently used to compute whole-genome ANI for the genome pairs. A species was considered to have been cultured if it contained a cultured gut genome from the UHGG catalog or it matched an isolate ReSeQ genome with at least 95% ANI over at least 30% of the genome length.

Genome alignments and the NCBI FTP site were used to obtain the accession numbers of the most similar RefSeq genomes for each MAG. ANI for the genome pairs. A species was considered to have been cultured if it contained a cultured gut genome from the UHGG catalog or it matched an isolate ReSeQ genome with at least 95% ANI over at least 30% of the genome length.

**Calculating the number of conspecific genomes.** For an accurate assessment of the number of nonredundant genomes belonging to each species, we de-replicated all conspecific genomes at a 99.9% ANI threshold using dRep with options ‘-pa 0.999 –SkipSecondary’. Furthermore, the frequency of each species was only counted once per sample to avoid cases where the same genome was recovered multiple times because of overlapping samples between the three MAG studies.

**Estimating geographic diversity.** A geographic diversity index was estimated to assess how widely distributed each species was. We calculated the Shannon diversity index on the proportion of samples in which each species was found per continent. This metric combines both richness and evenness, such that the level of estimated diversity is highest in species found across all continents at a similar proportion.

**Metagenomic read mapping.** A set of 1,005 metagenomic datasets from 14 studies (Supplementary Table 4) were retrieved from ENA and used to perform read mapping against the genome (UHGG) and protein (UHGP) catalogs. Only studies that were not used to generate the UHGG or UHGP catalogs were included. Reads were quality filtered and trimmed using TrimGalore v0.6.0 (https://github.com/FelixKrueger/TrimGalore), and human contamination was removed by aligning the reads with BWA MEM v0.7.16a1181 (ref. 28; default options) against human genome GRC38. Filtered reads were then mapped using Kraken v2.0.8-beta (with default settings) against a custom database of the UHGG catalog available from the MGnify FTP site (ftp://ftp.ebi.ac.uk/pub/databases/metagenomics/magnify_genomes/), and the standard ReSeQ database (release 96). Bracken databases of the UHGG catalog for read lengths of 50, 100, 150 and 250 bp were also generated and have been made available from the MGnify FTP site. Classification improvement was calculated on a per-sample basis as (proportion of reads classified with UHGG)−proportion of reads classified with ReSeQ)/proportion of reads classified with ReSeQ × 100. DIAMOND v0.9.21.122 (ref. 28) was used to translate and map the reads against the IGC-90 and UHGP-90 protein catalogs using the ‘blastx’ function with options ‘-id 90 -evalue 1e-6 -k 1 --max-ends 1’.

**Phylogenetic analyses.** Taxonomic annotation of each species representative was performed with GTDB-TK v0.3.1 (refs. 29,30; database release 04-RS89) using the ‘classify_wf’ function and default parameters. To use consistent species boundaries between the genome clustering and taxonomic classification procedures, genomes were assigned at the species level if the ANI to the closest GTDB-TK species representative genome was ≥ 95% and the AF was ≥ 30%. In this taxonomy scheme, genera and species names with an alphabetic suffix indicate taxa that are polyphyletic or need to be subdivided on the basis of taxonomic rank normalization according to the current GTDB reference tree. The lineage containing the type strain retains the suffix (valid) name, and all other lineages are given alphabetic suffixes, indicating that the common name of the genus is a placeholder name that need to be replaced in due course. Taxon names above the rank of genus appended with an alphabetic suffix indicate groups that are not monophyletic in the GTDB reference tree but for which there exists alternative evidence that they are monophyletic groups. We also generated NCBI taxonomy annotations for each species-level genome on the basis of its placement in the GTDB tree, using the `gtdb_to_ncbi_majority_vote.py` script available in the GTDB-TK repository (https://github.com/Ecogenomics/GTDB-TK).

Maximum-likelihood trees were generated de novo using the protein sequence alignments produced by GTDB-TK: we used IQ-TREE v1.6.11 (ref. 32) to build a phylogenetic tree of the 4,616 bacterial and 28 archaeal species. The best fit model was automatically selected by `ModelFinder` on the basis of the Bayesian information criterion (BIC) score. The LG+F+I model was chosen for building the bacterial tree, while the LG+F+R4 model was used for the archaeal phylogeny. Trees were visualized and annotated with Interactive Tree Of Life (iTOl) v4.4.2 (ref. 33). Phylogenetic diversity (PD) was estimated by the sum of branch lengths, with the amount that was exclusive to uncultured species calculated as
Data availability

 Genome assemblies of the UHGG catalog have been deposited in the European Nucleotide Archive under study accession ERP116715. The UHGG, UHPG and SNV catalogs are available from the MGnify FTP site (http://ftp.ebi.ac.uk/pub/databases/metagenomics/mgnify_genomes/) alongside functional annotations, pan-genome results and custom Kraken 2/Bracken databases of the UHGG catalog. These data, together with the BIGSI search index of the UHGG catalog, can also be accessed interactively via the MGnify website at https://www.ebi.ac.uk/metagenomics/genomes. Mash distance trees have been generated for each individual species cluster and are available at both the MGnify website and the associated FTP site.

Code availability

 The workflow used to generate the genome and protein catalogs, alongside the pan-genome and functional annotations, is described in a Common Workflow Language (CWL) pipeline at https://github.com/EBI-Metagenomics-genomes-pipeline. Scripts used to generate the SNV catalogs are available at https://github.com/ezhu/snv_analysis_almeida2019.

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SNV analyses

 A total of 2,489 species with at least three conspecific genomes were used to generate a catalog of SNVs. For each species, we mapped all conspecific genomes to the species genome with options '-q r' to exclude chance- and repeat-induced alignments. Thereafter, we identified SNVs using the 'show-snps' program. Single-base insertions and deletions were not counted as SNVs. Each SNV locus was included in the catalog only when the alternate allele was detected in at least two conspecific genomes. The final SNV catalog was generated by unifying the SNV coordinates on the basis of the position in the respective representative genome. The SNV entries in the catalog were characterized as genome type or continent specific on the basis of whether the alternate allele could be found solely in genomes from a specific genome type or continent. The number of continent-specific SNVs was normalized by the number of genomes from the corresponding continent to estimate the contribution per genome type in the continent-specific SNV discoveries.

Similar programs and parameters were used for the pairwise genome alignment, but in this case only near-complete genomes (>90% completeness) and species with at least ten independent conspecific genomes were considered. Because of the high computational demand, pairwise alignments of species with at least ten independent conspecific genomes were performed between almost 113,000 genomes from 909 species. For each pairwise comparison, a total of 29,283,684 pairwise genome alignments were performed with options '-q r' to exclude chance- and repeat-induced alignments. Single-base insertions and deletions were not counted as SNVs. Each SNV locus was included in the catalog only when the alternate allele was detected in at least two conspecific genomes. The final SNV catalog was generated by unifying the SNV coordinates on the basis of the position in the respective representative genome. The SNV entries in the catalog were characterized as genome type or continent specific on the basis of whether the alternate allele could be found solely in genomes from a specific genome type or continent. The number of continent-specific SNVs was normalized by the number of genomes from the corresponding continent to estimate the contribution per genome type in the continent-specific SNV discoveries.

Reported Summary

 Further information on research design is available in the Nature Research Reporting Summary linked to this article.
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**Author contributions**

A.A., S.N., N.C.K. and R.D.F. conceived the study. A.A. performed the genome clustering, annotations and read mapping, compared study sets, carried out the pan-genome analyses, built the BIGSI index and drafted the manuscript. S.N. provided feedback and performed phylogenetic, rarefaction and clustering analyses, as well as the comparison with RefSeq. M. Boland and M. Beracochea built the resource implementation within the MGnify website. F.S. built the protein catalog and performed the comparison with the IGC. Z.J.S. generated the SNV catalog and performed related analyses according to genome type and geographic origin. K.S.P. provided feedback and funding and contributed to the SNV analyses. E.S. wrote the CWL pipeline implementation. D.H.P and P.H. provided feedback and assisted in the species taxonomic classification. N.S. provided feedback and funding and contributed to the generation of the protein catalog. N.C.K. and R.D.F. supervised the work and provided feedback and funding. All authors read, edited and approved the final manuscript.

**Competing interests**

F.S. is an employee of Enterome. P.H. is a cofounder and is director of Microba Life Sciences Ltd. D.H.P. is a consultant to Microba Life Sciences Ltd. R.D.F. is a consultant to Microbiotica Pty Ltd.

**Additional information**

Extended data is available for this paper at https://doi.org/10.1038/s41587-020-0603-3.

Supplementary information is available for this paper at https://doi.org/10.1038/s41587-020-0603-3.

Correspondence and requests for materials should be addressed to A.A. or R.D.F.

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Extended Data Fig. 1 | Genome quality of species representatives. 

**a**, Completeness and contamination scores for each of the 4,644 species representatives, colored by their quality classification category. Medium quality: >50% completeness; near complete: ≥90% completeness; high-quality: >90% completeness, presence of 5S, 16S and 23S rRNA genes, as well as at least 18 tRNAs. All genomes have a quality score (QS = completeness - 5 × contamination) above 50. 

**b**, Number of species according to different completeness and contamination criteria.

**c**, Distribution of the level of strain heterogeneity (proportion of non-synonymous substitutions) estimated for the species-level MAGs using CMseq. Dashed vertical line corresponds to the threshold defined in Pasolli, et al. to distinguish medium- from high-quality MAGs.
Extended Data Fig. 2 | Taxonomy composition of the bacterial and archaeal species. **a.** Taxonomic affiliation of the 4,616 bacterial species detected. Data is partitioned by taxonomic rank, with only the five most highly represented taxa per rank depicted in the legend. **b.** Taxonomic affiliation of the 28 archaeal species detected, partitioned by taxonomic rank.
Extended Data Fig. 3 | Species overlap across study sets. a, Number of species found across the three metagenome-assembled genome sets, ordered by their level of overlap. Only those genomes recovered from the 1,554 metagenomic samples used by all three studies were considered in this analysis. b, Distribution of the proportion of species recovered per sample (n = 1,554) in each study set out of all species recovered across all three studies in the same samples. Box lengths represent the IQR of the data, and the whiskers the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively. c, Estimated aligned fractions and average nucleotide identities (ANI) between conspecific genomes obtained in the same sample but in different MAG studies. Results for medium-quality genomes are illustrated in the top panel, whereas those for near complete (≥90% completeness) genomes are represented in the lower panel. Vertical dashed lines denote the median values. d, Number of species identified in three culture-based studies and their degree of overlap. The NCBI study set consists mainly of genomes from the Human Microbiome Project (HMP).
Extended Data Fig. 4 | Quality and sample origin of uncultured singleton species. 

**a**, Genome completeness and contamination estimates of the 1,212 uncultured species represented by a single genome. Box lengths represent the IQR of the data, and the whiskers the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively.  

**b**, Proportion of the 1,212 singleton species, by study set, that originated from samples analysed in one, two or three of the MAG studies (CIBIO, EBI and HGM).
Extended Data Fig. 5 | Species frequency and geographical diversity. **a**, Number of nonredundant genomes retrieved from the 50 most highly represented species in the UHGG catalog. Each species is colored by its assigned phylum according to the figure legend. **b**, Geographical diversity estimated using the Shannon index in relation to the number of nonredundant genomes from each species containing more than one genome (n = 2,786). Percentage values represent the estimated diversity normalized by the maximum theoretical value (considering an equal distribution of samples across the six major continents — Africa, Asia, Europe, North America, South America and Oceania). The Spearman’s rank correlation coefficient and P value (calculated with the Spearman’s test) are depicted in the graph. Predicted values represent the random geographical distribution of equivalent numbers of genomes observed for each species. Dashed horizontal line indicates the median observed value for species with more than one genome.
**Extended Data Fig. 6 | Diversity of the gut archaeal species detected.** Phylogenetic tree of the 28 archaeal species detected in the human gut. Tips are labelled with the corresponding species representative code and colored according to its cultured status. The taxonomic affiliation (family), geographical distribution, number of nonredundant genomes and total pan-genome size are represented next to the tree.
Extended Data Fig. 7 | UHGP cluster size and mapping rate. a, Cumulative distribution curve of the number and size of the gene clusters of the UHGP-95 (n = 20,239,340), UHGP-90 (n = 13,907,849) and UHGP-50 (n = 4,735,546). Dashed vertical lines indicate the cluster size below which 90% of the gene clusters can be found. b, Proportion of metagenomic reads from 1,005 independent datasets aligned with DIAMOND against the combined clusters of UHGP-90 and IGC-90 (left). The degree of classification improvement provided over the IGC-90 alone is represented in the right panel. The following represents the number of datasets analysed per country: Cameroon, n = 54; Ethiopia, n = 25; Germany, n = 56; Ghana, n = 40; India, n = 105; Italy, n = 50; Luxembourg, n = 26; Russia, n = 4; Tanzania, n = 61; United Kingdom, n = 210; United States, n = 374. Box lengths represent the interquartile range (IQR) of the data, and the whiskers the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively.
Extended Data Fig. 8 | Functional annotation of gut microbiome species. **a**, Functional profiles of the UHGG species pan-genomes (rows) according to 363 KEGG modules (columns). Numbers of genes matching each module were normalized to centered log ratios after imputing values with zero counts. Species are colored according to phylum. KEGG modules and species were hierarchically clustered using the Ward's criterion method. **b**, Proportion of each species pan-genome, partitioned by phylum, without any assignment to the eggNOG, InterPro, COG or KEGG databases (left). Proportion of the pan-genome with a match to the carbohydrate-active enzymes (CAZy) database (right). Sample size (number of species) of each phylum is indicated in parentheses \((n = 4,644\) total species). Box lengths represent the IQR of the data, and the whiskers the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively.
Extended Data Fig. 9 | Gene frequency distribution within the species-level clusters. a, Distribution of the number of genes found per fraction of conspecific genomes. Only near-complete genomes (≥90% completeness) were considered in the analysis. b, Number of core genes detected based on the threshold of genomes per species used to classify as core. Vertical dashed line represents the 90% threshold used in this study.
Extended Data Fig. 10 | SNV density and MAG strain heterogeneity. a. Correlation between the SNV density calculated among MAGs and their level of strain heterogeneity estimated with CMseq (n = 268,994 comparisons). A Pearson correlation test was performed to determine the correlation coefficient and P value. Colors denote density of data points (increasing from dark purple to yellow). b. Comparison of pairwise SNV density between isolates (n = 808,331 comparisons) and between MAGs with <0.01% (n = 2,923,610 comparisons) and <0.1% strain heterogeneity (n = 13,634,222 comparisons). A two-tailed Wilcoxon rank-sum test was performed to assess statistical significance and further adjusted for multiple comparisons using the Benjamini-Hochberg correction (**P <0.001). Box lengths represent the IQR of the data, and the whiskers the lowest and highest values within 1.5 times the IQR from the first and third quartiles, respectively.
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Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection

- European Nucleotide Archive (ENA); National Center for Biotechnology Information (NCBI); Integrated Microbial Genomes & Microbiomes (IMG/M); Pathosystems Resource Integration Center (PATRIC)

Data analysis

- CheckM v1.0.11; INFERNAL v1.1.2; iRNASCAN-SE v2.0; dRep v2.2.4; FastANI v1.1; Python v2.7 and v3.6; R v3.5; GTDB-Tk v0.3.1; Mash v2.1; MUMmer v4.0.0beta2; IQ-TREE v1.6.11; BiGSI v0.3.8; Prokka v1.13.3; Prodigal v2.6.3; Roary v3.12.0; MMseq2 v1.5.1; eggNOG-mapper v2; InterProScan v5.35-74.0; Kraken v2.0.8-beta; Bracken v2.5;Trim Galore v0.6.0; BWA v0.7.16a-r1181; bowtie v2.2.3; samtools v1.5; DIAMOND v0.9.21.122; BBMap v38.75; CMseq (https://github.com/SegataLab/cmseq); iTOL v4.4.2; gdb_to_ncbi_majority_vote.py (https://github.com/Ecogenomics/GTDBTk/).

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Data

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All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:
- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

Genome assemblies of the UHGG have been deposited in the European Nucleotide Archive under study accession ERP116715. The UHGG, UHGP and SNV catalogs are available from the MGnify FTP site (http://ftp.ebi.ac.uk/pub/databases/metagenomics/mgnify_genomes/) alongside functional annotations, pan genome results and custom Kraken 2/Bracken databases of the UHGG. These data together with the BiGSI search index of the UHGG can also be accessed interactively via the MGnify website: https://www.ebi.ac.uk/metagenomics/genomes. Mash distance trees have been generated for each individual species cluster and are available both in the MGnify website and the associated FTP site.
Field-specific reporting

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Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size
No sample size calculation was performed, as the aim was to include all publicly available human gut microbial genomes available (n = 286,997 genomes as of March 2019).

Data exclusions
Genomes were filtered to select those with an estimated completeness >50%, contamination <5% and quality score (completeness × 5 * contamination) >50. Quality criteria for exclusion was pre-established based on existing literature and current standards in the field.

Replication
By comparing recently published large datasets of uncultured genomes, we were able to assess the reproducibility of the results from each study. We showed that despite the different assembly, binning and refinement procedures employed in the three studies, almost all of the same species and similar strains were recovered independently when using a consistent sample set.

Randomization
Randomization was not relevant to this study, as we analysed publicly available data.

Blinding
Not relevant to this study, as we analysed publicly available data.

Reporting for specific materials, systems and methods

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Materials & experimental systems

| n/a | Involved in the study |
|-----|-----------------------|
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| ☒   | Eukaryotic cell lines |
| ☒   | Palaeontology         |
| ☒   | Animals and other organisms |
| ☒   | Human research participants |
| ☒   | Clinical data         |

Methods

| n/a | Involved in the study |
|-----|-----------------------|
| ☒   | ChIP-seq              |
| ☒   | Flow cytometry        |
| ☒   | MRI-based neuroimaging |