Unique Features of Ethnic Mongolian Gut Microbiome revealed by metagenomic analysis

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The human gut microbiota varies considerably among world populations due to a variety of factors including genetic background, diet, cultural habits and socioeconomic status. Here we characterized 110 healthy Mongolian adults gut microbiota by shotgun metagenomic sequencing and compared the intestinal microbiome among Mongolians, the Hans and European cohorts. The results showed that the taxonomic profile of intestinal microbiome among cohorts revealed the Actinobaceria and Bifidobacterium were the key microbes contributing to the differences among Mongolians, the Hans and Europeans at the phylum level and genus level, respectively. Metagenomic species analysis indicated that Faecalibacterium prausnitzii and Coprococcus come were enrich in Mongolian people which might contribute to gut health through anti-inflammatory properties and butyrate production, respectively. On the other hand, the enriched genus Collinsella, biomarker in symptomatic atherosclerosis patients, might be associated with the high morbidity of cardiovascular and cerebrovascular diseases in Mongolian adults. At the functional level, a unique microbial metabolic pathway profile was present in Mongolian's gut which mainly distributed in amino acid metabolism, carbohydrate metabolism, energy metabolism, lipid metabolism, glycan biosynthesis and metabolism. We can attribute the specific signatures of Mongolian gut microbiome to their unique genotype, dietary habits and living environment.

The gut microbiota (GM) is recognized as a human co-evolutionary partner that facilitates nutritional acquisition and immune modulation, and helps maintain host homeostasis in response to profound lifestyle changes1–3. Researchers have delineated the structural and functional configurations of gut metagenomes in different world populations by next generation sequencing4–7. The functional GM layout may represent host genotypic characteristics and reflect an adaptive ecosystem response to the host diet and cultural habits.

The microbiota of the Mongolian population is of particular interest to researchers, because Mongolia encompasses a uniquely wide range of environmental conditions and ethnogeographical cohorts. Moreover, the Mongol Empire was the world’s largest contiguous empire, exerted a major influence that greatly enhanced the cultural exchange between Mongolia and Europe that took place during the Middle Ages, when Mongolians intermingled with various populations in the Eurasian continent. Importantly, it has been reported that genetically, more than 20% people in the world are related to Mongolians8. Mongolians living in pastureland, such as Khentii Province, maintains a traditional nomadic lifestyle (Fig. 1A), whereas those in urban areas, such as Ulan Bator (the capital of Mongolia) and TUW Province (the suburbs of the capital), have adopted an urban lifestyle due to rapid modernization and economic development. Chinese Mongolians living in Hohhot and Xilingol have inter-married with ethic Han Chinese, which may generate some differences from their brethren in the north at the genomic level. Nevertheless, all ethnic Mongolians consume similar diet, including traditional cheese (Fig. 1B) and red meat (Fig. 1C) as their main food.

In our previous study, by employing 454 pyrosequencing technology, we described the profiles of gut microbiota of Mongolians from Mongolia and China. At the genus level, Prevotella of the Firmicutes phylum was the

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most abundant genus, and the amounts of Bacteroides, Faecalibacterium, Oscillibacter, Roseburia, Clostridium, Coprococcus, Ruminococcus, Alitipes, Parabacteroides, Catenibacterium, Subdoligranulum and Eubacterium all exceeded 1%, and we found seasonal effects on intestinal microbiota were more distinct in rural Mongolians. From previous research, we found the intestinal microbiota of Mongolians exhibited some unique features. However, little is known about the functional gene and metabolic pathway of the intestinal microbiota of Mongolians. It has been reported that prevalence of cardiovascular and cerebrovascular diseases in Mongolian adults was significantly higher than individuals of other ethnic groups and that these diseases became the leading cause of mortality in Mongolians. Some previous research attributes the aforementioned chronic metabolic diseases to the Mongolian dietary tradition of high consumption of red meat and liquor. With the development of human microbiome project and next generation sequencing, great research effort has been dedicated to elucidating the relationship between intestinal microbiota and various human metabolic diseases. The approach may also generate great insight by exploring the potential connection between cardiovascular and cerebrovascular diseases and the intestinal microbiota.

Here we characterized 110 healthy Mongolian adults gut microbiota by shotgun metagenomic sequencing and compared the intestinal microbiome among Mongolians, the Hans and European cohorts. (i) The taxonomic profile of intestinal microbiome among cohorts revealed the Actinobacteria and Bifidobacterium were the key microbes contributing to the differences among Mongolians, the Hans and Europeans at the phylum and genus level, respectively. (ii) We constructed a non-redundant microbial catalogue for Mongolian contained 1,491,813 genes, and merged present catalogue with other published healthy gut microbial catalogues of the Hans and European cohorts. The final merged non-redundant catalogue contained 5,127,164 genes, and 283,401 genes were common for all individuals. (iii) At the functional level, a unique microbial metabolic pathway profile was present in Mongolian’s gut which mainly distributed in amino acid metabolism, carbohydrate metabolism, energy metabolism, lipid metabolism, glycan biosynthesis and metabolism. We can attribute the specific signatures of Mongolian gut microbiome to their unique genotype, dietary habits and living environment.

### Results

**Taxonomic characterization.** In present research, we characterized the fecal microbiota of 110 Mongolian individuals and compared the structure of intestinal microbiota among the cohorts from Hohhot, Xinlingol, TUW, Ulan Bator and Khentii. The sampling sites included a substantial part of relative densely populated places of ethnic Mongolian settlements, including areas in China and Mongolia. As a result, five representative Mongolian settlements were chosen to collect samples, including Ulan Bator city (35 healthy adults), TUW province (16 healthy adults) and the Khentii pasturing area (12 healthy adults) of Mongolia, Hohhot city (22 healthy adults) and Xilingol pasturing area (25 healthy adults) of China. Principal coordinates analysis (PCoA) based on the Bray–Curtis distance of species abundance (Fig. 2A) revealed the structure of intestinal microbiota between the Han Chinese’s and the European’s were similar, but significantly different from the Mongolian’s. The bacteria alpha diversity based on Shannon index (Fig. 2C) indicated the abundance of intestinal microbiota in Mongolian’s exceeded 1%, and we found seasonal effects on intestinal microbiota were more distinct in rural Mongolians.

To further explore the gut microbial community features of Mongolian population, we compared the results with that of previously reported samples and carried out taxonomic assignment for the metagenomic data using Metaphlan2 programme. Results were presented in the phylogenetic tree in Fig. 3. For all of these Mongolian subgroups, Actinomyces, Rothia, Catenibacterium, Phascolarctobacterium, Sutterella and Lactobacillus genera were remarkably enriched. On the contrary, Holdemania, Peptostreptococcaceae, Potyviridae or Anaerostipes were rarely detected in all Mongolian samples (Fig. 3). Taxonomic profile of the gut microbiome among the populations (Fig. 4) also revealed that Actinobacteria and Bifidobacterium were the key components contributing to the differences among Mongolians, the Hans and Europeans at the phylum level and genus level, respectively.

**Metagenomic species (MGS) associated with Mongolians.** To explore features of the gut microbiota in Mongolians, we compared the genes abundance in 110 Mongolians and 268 Hans, whereby we identified 115,783 genes with significant abundance differences (Wilcoxon rank-sum, Benjamin–Hochberg q-value < 0.001): 78,345 were more abundant in the Hans, and 37,435 in the Mongolians. Likewise, the similarity

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**Figure 1. Country-wide sampling highlights distinct features of Mongolia gut microbiota.** (A) Traditional Mongolian yurt in pasturing area. (B,C) Mongolian’s staple foods of such as cheese and red meat are drying in the air.
A comparison between Mongolians and Europeans was also carried out, which identified 127,540 genes: 66,030 were enriched in the Europeans and 61,510 in the Mongolians.

The differentially abundant genes with high correlation were clustered into MGS\(^{16,17}\) based on their abundance in all samples. We found 50 MGS to be differentially abundant between the two groups, 26 and 24 enriched in Hans individuals and Mongolian subjects, respectively (Fig. 5 and Supplementary Table 3). Based on the taxonomic characterization of the MGS significantly enriched in Mongolians, we found three Faecalibacterium prausnitzii, two identified at strain level and one at species level, which has anti-inflammatory properties and found to be associated with ‘healthy’ microbiome in previous research\(^ {16,18}\). One other MGS enriched in Mongolians was identified at species levels as Coprococcus comes ATCC 27758, which might contribute to gut health through butyrate production. Interestingly, Collinsella aerofaciens, which is enriched in Mongolians, was reportedly enriched in patients with symptomatic atherosclerosis\(^ {12}\). For the comparison of Mongolians and Europeans, we found 29 and 16 MGS were enriched in European individuals and Mongolian subjects, respectively (Fig. S2). Remarkably, like the comparison between the Hans and Mongolians, Faecalibacterium prausnitzii, Bifidobacterium adolescentis and Collinsella aerofaciens were enriched in Mongolians.

Another intriguing question we focused on was the correlation between the Mongolians diet and their intestinal microbiota. By transforming food frequency questionnaire to the nutrition information, we calculated the Spearman’s rank correlation coefficient between the Mongolian-enriched MGS and the dietary nutrition factors (Fig. 6 and Fig. S3). From the figure, a significant positive (\(P < 0.05\)) correlation can be observed between the protein, K, Zn, Fe and VB1 and the species Collinsella aerofaciens. Additionally, the elements Se and Mg exhibited significant positive (\(P < 0.05\)) correlation with the species Bifidobacterium adolescentis.

### Gene catalogue of gut microbes

We sequenced the fecal metagenome of 110 Mongolian individuals using Illumina platform and obtained 725 gigabases (Gb) raw data, with an average of 6.54 Gb for each sample. After quality control, we got an average of 6.33 Gb data for each, totalling 696 Gb of high-quality data which was free of adaptor and human DNA contaminants. Based on these data, we constructed a Mongolian gene catalogue of gut microbes. We sequenced the fecal metagenome of 110 Mongolian individuals using Illumina platform and obtained 725 gigabases (Gb) raw data, with an average of 6.54 Gb for each sample. After quality control, we got an average of 6.33 Gb data for each, totalling 696 Gb of high-quality data which was free of adaptor and human DNA contaminants. Based on these data, we constructed a Mongolian gene catalogue of gut microbes.
catalogue distributed in five areas (Khentii pasturing area, TUW province, Ulan Bator, Hohhot City and Xilingol pasturing area) using the methodology developed by MetaHIT. Samples from Ulan Bator had the most abundant gene catalogue, containing 768,497 genes. Khentii set contained 400,178 genes, representing the fewest sample. Xilingol, Hohhot and TUW catalogue contained 512,067,702,521 and 433,515 genes, respectively. The merged non-redundant catalogue contained 1,491,813 genes. The Hohhot, Khentii, TUW, Ulan Bator and Xilingol catalogue sets contained 137,496, 99,760, 110,428, 277,758 and 242,637 unique genes, respectively.

Furthermore, we compared the Mongolian catalogue with four other published healthy gut microbial catalogues: ShenZhen (T2D), HangZhou (LC), American (HMP), European (MetaHIT)²,¹⁰,¹¹,¹⁵. To facilitate this comparison, we predicted genes of the four healthy groups above using the same criteria. The Mongolian gene catalogue contained 1,418,578 genes, the American catalogue 2,243,320 genes, the European catalogue 1,711,862 genes, the HangZhou catalogue 1,183,368 genes and the ShenZhen catalogue 2,067,885 genes. In total, 283,401 genes were common to all catalogues. The Mongolian, American, European, HangZhou, and ShenZhen sets contained 599,370, 1015,117, 591,857, 233,036 and 777,465 unique genes, respectively. The final merged non-redundant catalogue contained 5,127,164 genes (Fig. S1).

**Functional characterization.** To explore the functional feature of the gut microbiota in Mongolian, we annotated the gene catalogue by KEGG and eggNOG. At the functional level, it can be observed the functional structures of intestinal microbiota among Mongolian, the Hans and Europeans were significantly different (Fig. 7B), as a unique KEGG orthologue group (KO) profile was present in Mongolian’s gut. Similarly to taxonomic diversity, Mongolian’s gut and European’s gut show higher diversity than Han Chinese’s in functional level (Fig. 7C). At pathway level, Mongolians enriched KOs was mainly distributed in amino acid metabolism,
carbohydrate metabolism, energy metabolism, lipid metabolism, glycan biosynthesis and metabolism (Fig. 8A,C).

In the amino acid metabolism, mainly enriched pathway were alanine, aspartate and glutamate metabolism, cysteine and methionine metabolism, lysine biosynthesis and lysine degradation (Fig. 8B,D). Combined the abundance analysis of amino acid metabolic pathways for each species, *Faecalibacterium prausnitzii*, *Prevotella copri* which were enriched in Mongolians, seemed to be the primary contributors to the amino acid metabolism, especially the L-lysine biosynthesis and L-iso-leucine biosynthesis (Fig. 9). It potentially indicated some functional enrichment might be related to the enrichment of some specific species. The enriched amino acid metabolism and energy metabolism functions were associated with Mongolians’ high meat and fermentation uptake (Fig. S6). Moreover, we identified significantly different (reporter score $> 1.6$) abundant metabolic models between Mongolian and the Hans or Europeans by calculating the reporter scores. The majority of the models that were differentially abundant in the Mongolian subjects coincided with the observed changes in the metabolic pathway which mainly belong to the metabolic of carbohydrate, amino and nucleotide sugar, phosphotransferase and aminoacyl-tRNA and the biosynthesis of the methane and sulfide (Fig. S4).

Since we observed the carbohydrate metabolic between Mongolians and the Hans was significantly different. We further explored the potential for complex carbohydrate degradation in Mongolian and the Hans gut metagenomes, we screened for carbohydrate-active enzymes (CAZymes) in assembled contigs. As a result, Mongolian GM showed a higher diversity of CAZymes than that of the Hans GM ($P < 0.0001$, Wilcoxon rank-sum test; Fig. S5). This demonstrates an increased capacity for complex carbohydrate metabolism in the Mongolian GM.

**Discussion**

In present research, we observed several distinct features present in the gut microbiota of ethnic Mongolian from two countries, Mongolia and China. Taxonomically, the genus *Collinsella* was significantly enriched in Mongolians, and a significant positive correlation was observed between protein uptaken in diet with this genus (Fig. 6). It was reported that the *Collinsella* was considered as the biomarker in symptomatic atherosclerosis patients. In previous study, the intestinal microbiota of patients of symptomatic atherosclerotic and healthy controls were compared. In genus level, it is indicated the abundance of the *Collinsella* was increased in patients with symptomatic atherosclerosis, whereas *Roseburia* and *Eubacterium* were enriched in healthy controls. Additionally, the *Collinsella*, *Enterobacteriaceae* and *Streptococcus* were considered as the potential pro-inflammatory GM components enriched in the T2D patients, which resulted in the rapid decline of the SCFA in gut. Accordingly, we can infer the high protein uptake in Mongolians diet correlates with robust growth of the intestinal *Collinsella*, which may increase the risk of the morbidity of cardiovascular and cerebrovascular diseases in Mongolian adults.

On the other hand, the *Lactobacillus* and *Bifidobacterium* commonly considered as probiotics were largely detected in Mongolians gut. As we known, the *Lactobacillus* and *Bifidobacterium* are major benefit microbes
in human gut. They exhibit a range of health-promoting effects, including the regulation of intestinal micro-
bial homeostasis, the inhibition of pathogens in the intestinal mucosa, the modulation of local and systemic
immune responses, the production of vitamins, and the bioconversion of a number of dietary compounds
into bioactive molecules. Various species of Lactobacillus played a key role in yoghurt fermentation. And in
previous study, researchers had demonstrated that the Lactobacillus was able to promote the colonization of
Bifidobacterium. Thus, the widely existence of the genera Lactobacillus and Bifidobacterium may be related to
Mongolians dietary habits of fermented milk consumption. Additionally, we found that the genus Bifidobacteri-
exhibited significant positive correlation with the dietary selenium (Se). The interaction between the Se and the
genus Bifidobacterium had been reported in previous study, and the supplementary Se in MRS ager was able to
promote the growth of the Bifidobacterium.

The increased pathway abundance of carbohydrate, amino acid and nucleotide sugar, and phosphotransferase
and aminoacyl-tRNA of intestinal microbiota in Mongolians suggested the more vigorous microbial growth and
metabolic stage in Mongolian’s gut compared with that of Han Chinese. The aminoacyl-tRNA was able to com-
bine with its corresponding animal acid, and transferred the animal acid to the ribosome. In present research,
we found the biosynthesis of aminoacyl-tRNA was enhanced which indicated an overall enhanced level of micro-
bial gene expression in the gut of Mongolians. Moreover, the up-regulation of the metabolic pathway of phospho-
transferase provide the source of energy for microbe’s proliferation. On one hand, the robust metabolism of gut
microbes in Mongolians was able to promote the digestion and absorption of food in intestine. On the other hand,
attributed to the stronger metabolism, intestinal microbes would produce more metabolic products including
beneficial or harmful factors which had a greater impact on human health.

Accordingly, we found that the biosynthesis of the methane and sulfide in intestinal microbiota of Mongolians
were significantly higher than that of other populations. The two compounds were recognized as the typical
inflammatory factor in human gut. Previous researches had demonstrated the perniciousness of the methane and
sulfide in intestine of T2D and IBD patients. As such, it is tempting to speculate that the high morbidity of

Figure 5. Gut MGS in the Mongolian and the Hans individuals. (A) The Heatmap of 25 ‘tracer’ genes
abundance for each MGS in the Mongolians (110 individuals) and the Hans (268 individuals). Individuals are
represented along the horizontal axis, sorted by increasing abundance of the Hans enriched MGS. Abundance of
genes in rows is indicated by color gradient (white, not detected), and the enrichment significance is shown with
Benjamin–Hochberg q value. (B) Co-occurrence network of enriched MGS in Mongolian (n = 24) and the Hans
individuals (n = 26), respectively. Each node represents one MGS, and two nodes are linked if Spearman’s rank
 correlation > 0.6, using the edge width to represent the correlation strength. The node size is proportional to
the mean relative abundance of MGS in the respective population. Nodes were colored based the phylogenetic
order level.
cardiovascular and cerebrovascular diseases in Mongolian adults were pertinent to their relatively strong biosynthesis of the methane and sulfide in intestinal microbiota.

In Mongolian diet, meat (including mutton and beef) is the major source of protein, and the Mongolians consume a significantly greater amount of animal protein than Han Chinese. Intestinal microbiota played the key role in animal protein metabolism. Within this microflora, the genera *Clostridium* and *Bacteroides*, which were extensively detected in the Mongolians gut, have been mainly associated with the metabolism of animal proteins, a variety of amino acids and saturated fat acids. Accordingly, we noticed the pathways for degradation of protein and the biosynthesis of the amino acid were enriched in the intestinal microbiota of Mongolians.

**Materials and Methods**

**Sample information.** In present research, a total of 110 Mongolians were recruited for sampling. Among these participants, 63 Mongolians were from Mongolia and the other 47 were from Inner Mongolia of China. For participants from Mongolia, 35 volunteers lived a typical modern lifestyle in Ulan Bator, the capital of Mongolia, 16 volunteers lived in the TUW province which covers the suburbs of Ulan Bator, and 12 volunteers lived in the Khentii pasturing area, a typical Mongolian steppe. For the 47 volunteers from Inner Mongolian of China, 22 volunteers lived in Hohhot, the capital of Inner Mongolia, which lived an urban lifestyle. The others lived in Xilingol pasturing area, a typical Mongolian grassland like Khentii. After obtaining the written informed consent, we collected habitual long-term dietary information from all participants using a food frequency questionnaire and transformed them to the nutrition information (Supplementary Table 1). The study protocol was approved by the Ethical Committee of the Inner Mongolia Agriculture University (Hohhot, China). Sampling and all subsequent steps described in the Materials and Methods have been conducted in accordance with the approved guidelines.

**DNA sequencing.** DNA was extracted from faecal samples using the QIAampDNA Stool MiniKit as previously described. The metagenomic DNA libraries were constructed with 2 μg genome DNA according to the Illumina TruSeq DNA Sample Prep v2 Guide, with an average of 500 bp insert size. The quality of all libraries was evaluated using an Agilent bioanalyzer with a DNA LabChip 1000 kit. Sequencing was performed by Illumina Hiseq2500.

**Metagenomic data for comparison.** The public gut microbial metagenomic data used in this study include: (i) 83 healthy Hans fecal samples (Hangzhou), which were downloaded from EBI under accession number ERP005860; (ii) 185 healthy Hans fecal samples (Shenzhen), which were downloaded from NCBI under
accession number SRA04564614; (ii) 99 healthy European fecal samples from the MetaHIT project, which were downloaded from EBI under accession number ERA000116⁵.

**Quality control of reads.** Raw paired-end reads were processed quality control using the following criteria: (1) reads with adaptor were removed; (2) reads were trimmed from the 3' end using a quality threshold of 30; (3) reads containing more than 50% bases with low quality (Q30) were removed; (4) Reads short than 70 bp were removed; (5) reads that mapped to human genome were removed. For alignment, SOAPaligner 2.21 27 was used by parameter of '-m 100 -x 1000', which are "reporting all repeat hits, Minimal insert size of 100 bp, Maximal insert size of 1000". For detailed information on the meaning of each option above, we refer to http://soap.genomics.org.cn/soapaligner.html. The resulting high-quality reads were used for the further analysis.

**De novo assembly and gene catalogue construction.** High-quality reads were used to assembly with SOAPdenovo 28 (version 2.04) using parameters of '-M 3 -u -L 100 -d 1 -F -k 53', which are "mergeLevel of 3, un-masking contigs with high/low coverage before scaffolding, minContigLen of 100 and KmerFreqCutoff of 1, filling gaps in scaffold, k-mer length of 53". For detailed information on the meaning of each option above, we refer to http://soap.genomics.org.cn/soapdenovo.html. The resulting scaffolds were cut into contigs at ambiguous Ns and the contigs longer than 500 bp were saved. All these contigs were applied for gene prediction.

We use MetaGeneMark 31 (version 3.26) to identify ORFs from the contigs of each sample using a length threshold of 100 bp (Supplementary Table 2). Then the non-redundant gene catalogue was constructed by pairwise comparison of all the predicted ORFs with CD-HIT 32 (version 4.5.7) and the redundant genes were removed using a sequence identity cut-off of 0.95 and aligned length covered over 90% of the shorter sequence. The final non-redundant Mongolian gene catalogue contains 1,491,813 microbial genes, with an average length of 808 bp.

In light of some low-abundance microbes that were not detected in the limited sequencing data, we combined with the previously constructed gene catalogue, including MetaHIT gene catalogue², HMP gene catalogue⁴, T2D gene catalogue¹⁴ and LC gene catalogue¹³ to build a non-redundant gene catalogue for further analyses. The ORFs

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**Figure 7. Function of Mongolian gut microbiomes distinctive to the European and the Hans population.** (A) PCoA plot with Bray–Curtis distances generated from KEGG Orthologue (Enzyme) profiling. (B) PCoA plot with Bray–Curtis distances generated from eggNOG Orthologue profiling. (C) Gene diversity of Mongolians, Europeans, and the Hans.
in the contigs of healthy individuals from above studies were predicted using MetaGeneMark and merged with Mongolian gene catalogue into a final one with the same criteria of 90% coverage of shorter gene and 95% identity. Finally, we got a non-redundant human gut gene catalogue which contains 4,998,380 genes, with an average length of 763 bp.

**Taxonomic and gene profiling.** We use MetaPhlAn2\(^{33}\) to produce organism abundance profiling with default parameters, which relied on about 1 million unique clade-specific marker genes identified from about 17,000 reference genomes. 1,036 microorganisms were mapped to the marker genes, include 9 Archaea, 963 Bacteria, 18 Eukaryota and 46 Viruses.

Relative abundances of the genes were determined with the procedure introduced in Qin N \textit{et al.} Nature 2014\(^{13}\). When calculating the abundance of genes, the high quality reads from each sample were aligned against the gene catalogue by using SOAPalign 2.21\(^{15}\) with parameters of ‘-r 2 -m 100 -x 1000’ and only the both paired-end reads which could be mapped to a same gene were accepted. For alignment, SOAPaligner 2.21 was used by parameter of ‘-m 100 -x 1000’, which are “reporting all repeat hits, Minimal insert size of 100 bp, Maximal insert size of 1000”. For detailed information on the meaning of each option above, we refer to http://soap.genomics.org.cn/soapaligner.html.

**MGS analysis.** For the comparison of the faecal microbiome between Mongolians and the Hans, gene markers with differentially abundance were identified (Benjamin–Hochberg q-value < 0.001). The comparison between Mongolians and Europeans was carried out in the same way. To cluster genes into Metagenomic Species (MGS), we followed the method described by Le Chatelier\(^{36}\) and Nielsen\(^{15}\). We clustered the genes with Spearman correlation coefficient (rho) > 0.8 using single-linkage clustering and then fused the clusters with more than 25 genes which had a Spearman correlation coefficient (rho) > 0.8.

The taxonomically annotation of MGS were performed as previously describe\(^{11}\). MGS was assigned to a taxonomic level from strain to super kingdom level when >90% of its genes had a best hit to the same phylogenetic group using blast with >95% identity and >90% overlap of query.

To constructed the co-occurrence network of MGS, we computed the Spearman correlation coefficient between MGS using their abundances and clustered the MGS according to the Spearman's correlation. The co-occurrence network of MGS was then visualized by Cytoscape3.0.2.

**KEGG, EggNOG and CAZy analysis.** Putative amino acid sequences of the predicted genes were aligned against the proteins in eggNOG 3.0 database\(^{34}\) and KEGG database\(^{35}\) (release 2014-12-09) using BLAST. Each gene was assigned to one or more OG(s) or EC(s) by the highest annotated scoring hit(s) containing at least one
HSP scoring over 60 bits. To construct the profiling of OGs in eggNOG and ECs in KEGG database, we accumulated the relative abundance of genes from the same OG or EC using the methods introduced in Qin J. et al. Nature 2010. To identify the differentially enriched KO modules, we computed their reporter scores from the Z-scores of individual KOs. A module with a reporter score of Z > 1.6 was defined as differentially enriched module.

CAZymes (Carbohydrate-Active Enzymes) were predicted from amino acid sequences by against to family-specific HMM of CAZymes in dbCAN database using Hmmscan program in HMMER 3.0 package. The raw sequencing data for all samples have been deposited to Sequence Read Archive (http://www.ncbi.nlm.nih.gov/sra/) under accession SRP080787.

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Acknowledgements
We sincerely thank all the Mongolian volunteers for their participation. This work was supported by National Natural Science Foundation of China (NO. 31430066), National Natural Science Foundation of China (NO. 81301475) and Natural Science Foundation of Zhejiang Province (NO. LR15H030002).

Author Contributions
H.Z. led the project and H.Z., N.Q. and W.L. conceived and designed the project, J.Z., X.X., W.H. and Q.H. collected the samples. B.Z. performed DNA extraction experiments. B.Z. and C.J. performed library construction and sequencing. N.Q. designed the analysis. C.W., C.J. and Z.L. analyzed the data. N.Q. and S.C. did the functional annotation analyses. J.Z. and C.W. wrote the initial manuscript with significant contributions from S.C., N.Q. and C.J. and critical input from all other authors.

Additional Information
Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Liu, W. et al. Unique Features of Ethnic Mongolian Gut Microbiome revealed by metagenomic analysis. Sci. Rep. 6, 34826; doi: 10.1038/srep34826 (2016).

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Corrigendum: Unique Features of Ethnic Mongolian Gut Microbiome revealed by metagenomic analysis

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Scientific Reports 6:34826; doi: 10.1038/srep34826; published online 06 October 2016; updated on 04 January 2017

The original version of this Article contained errors in the spelling of the authors Bin Zhou, which was incorrectly given as Bing Zhou, and Xiaoxia Xi, which was incorrectly given as Xiaoxia XI.

This Article also contained typographical errors in the Results section under the subheading 'Metagenomic species (MGS) associated with Mongolians'.

“whereby we identified 115,786 genes with significant abundance differences”.

now reads

“whereby we identified 115,783 genes with significant abundance differences”.

In addition, under the subheading “Functional characterization”.

“At the functional level, it can be observed the functional structures of intestinal microbiota among Mongolian, the Hans and Europeans were significantly different (Fig. 7A), as a unique KEGG orthologue group (KO) profile was present in Mongolian's gut. Similarly to taxonomic diversity, Mongolian's gut and European's gut show higher diversity than Han Chinese's in functional level (Fig. 7B)”.

Now reads

“At the functional level, it can be observed the functional structures of intestinal microbiota among Mongolian, the Hans and Europeans were significantly different (Fig. 7B), as a unique KEGG orthologue group (KO) profile was present in Mongolian's gut. Similarly to taxonomic diversity, Mongolian's gut and European's gut show higher diversity than Han Chinese's in functional level (Fig. 7C)”.

Finally, Figure legend 2D.

“Shannon diversity index of Mongolians, Europeans, and the Hans at the species level”.

now reads:

“Shannon diversity index of the Mongolians located in Hohhot, Xinlingol, TUW, Ulan Bator and Khentii at the species level”.

These errors have now been corrected in the PDF and HTML versions of the Article.
