Echinococcus granulosus Infection Results in an Increase in Eisenbergiella and Parabacteroides Genera in the Gut of Mice

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Cystic echinococcosis (CE) is a chronic infectious disease caused by Echinococcus granulosus. To confirm whether the infection impacts on the gut microbiota, we established a mouse model of E. granulosus infection in this study whereby BALB/c mice were infected with micro-cysts of E. granulosus. After 4 months of infection, fecal samples were collected for high-throughput sequencing of the hypervariable regions of the 16S rRNA gene. Sequence analysis revealed a total of 13,353 operational taxonomic units (OTUs) with only 40.6% of the OTUs having genera reference information and 101 of the OTUs were significantly increased in infected mice. Bioinformatics analysis showed that the common core microbiota were not significantly changed at family level. However, two genera (Eisenbergiella and Parabacteroides) were enriched in the infected mice (PAMOVA < 0.05) at genus level. Functional analysis indicated that seven pathways were altered in the E. granulosus Infection Group compared with the Uninfected Group. Spearman correlation analysis showed strong correlations of IgG, IgG1 and IgG2a with nine major genera. E. granulosus cyst infection may change the gut microbiota which may be associated with metabolic pathways.

KEYWORDS: Echinococcus granulosus, cystic echinococcosis, mice, microbiome, immunoglobins

INTRODUCTION

Cystic echinococcosis (CE) is a cosmopolitan zoonosis caused by the cystic stage of the dog tapeworm Echinococcus granulosus (McManus et al., 2012). The disease causes serious health problems and economic losses, especially in Central Asia (including western China), northern Africa and South America (McManus et al., 2012). E. granulosus requires two hosts [an intermediate host including sheep, goats, cattle or wild herbivores and a definitive host such as dogs (or wolves and other carnivores)] to complete its life-cycle. Humans also become infected as an incidental host by ingesting eggs released from E. granulosus in carnivore feces. After hatching in the stomach and
The gut microbiota play an important role in human health (Chen et al., 2017) impacting on metabolism, immunity, development and the behavior of the host (Thaiss et al., 2016). In addition, microbiota components are impacted by medical conditions such as cancer (Jensen et al., 2015; O’Keefe, 2016; Sonnenburg and Backhed, 2016). Similar changes occur in experimental models as well (Gkouskou et al., 2014; Yu et al., 2018). Studies showed that helminth infection in the gut induces typical Th2 immune responses which may control the microbiota in the gut of mice (Ramanan et al., 2016; Guernier et al., 2017; Wegener Parfrey et al., 2017). However, it is not known whether *E. granulosus* infection impacts on the gut microbiota of humans or mice. Mice have been used for *E. granulosus* larval infection including primary (Zhang et al., 2001) and secondary infection (Gottstein, 2001; Mourglia-Ettlin et al., 2016). Mice models play an important role in studies of developmental biology and host specificity in echinococcosis (Nakaya et al., 2006). Recently, mouse models were successfully used for drug screening and development (Elissondo et al., 2007; Wang et al., 2017). To increase the success of secondary infection, we developed a method using micro-cysts cultured in vitro to infect mice (Zhang et al., 2005), and obtained more than 70% of cyst recovery from 50 PSC-generated cysts.

In this study, BALB/C mice were infected with micro-cysts of *E. granulosus* and their fecal samples were collected for sequencing the variable regions of 16S rRNA genes of gut commensal bacteria to determine their composition and diversity. We show that *E. granulosus* impacted on the gut microbiota of the mice with microbiota changes likely being associated with the altered host immune status in infected individuals.

**MATERIALS AND METHODS**

**Ethics Statement**

The protocols for using mice in the study were approved by the Ethics Committee of The First Affiliated Hospital of Xinjiang Medical University (FAH-XMU, Approval No. IACUC-201206250003). The “Guidelines for the Care of Laboratory Animals” by the Ministry of Science and Technology of the People’s Republic of China (2006) were rigidly followed in the use of these animals.
TABLE 1 | The number and average size of cysts in the *E. granulosus* infection group.

|        | Eg1 | Eg2 | Eg3 | Eg4 | Eg5 | Eg11 | Eg12 | Eg13 | Eg14 | Eg15 | Eg31 | Eg32 | Eg33 | Eg35 |
|--------|-----|-----|-----|-----|-----|------|------|------|------|------|------|------|------|------|
| Number of cysts | 12  | 15  | 17  | 37  | 21  | 16   | 17   | 14   | 15   | 11   | 12   | 8    | 23   | 13   |
| Average size of cysts (cm) | 0.58 | 0.54 | 0.58 | 0.72 | 0.74 | 0.55 | 0.67 | 0.42 | 0.49 | 0.71 | 0.55 | 0.75 | 1.01 | 0.77 |

*Eg, mice infected with *E. granulosus.*

**FIGURE 1** | IgG antibody isotypes and subtypes in sera of mice infected with *E. granulosus.* *p < 0.05, **p < 0.01, ***p < 0.005, ****p < 0.001.

The V3–V4 hypervariable region of the 16S rRNA gene was amplified by PCR and sequenced; the length of the V3–V4 hypervariable region was approximately 469 bp. Amplicon pools were prepared for sequencing with AMPure XT beads (Beckman Coulter Genomics, Danvers, MA, United States) and quantification with the Library Quantification Kit for Illumina (Kapa Biosciences, Woburn, MA, United States), respectively. The libraries were sequenced on 300PE MiSeq runs.

**Bioinformatics and Statistical Analysis**

Mothur (version 1.39.5) was used to assemble the paired FASTQ files (Schloss et al., 2011). The selected quality DNA sequences were confirmed using the following criteria: (1) no contaminant sequences, (2) containing no ambiguous bases, (3) the size length ≥350 bp, (4) containing no chimeric sequences, and (5) primers were trimmed. The average length of selected DNA sequences was 414 bp (350–446 bp). The selected DNA sequences were then grouped into operational taxonomic units (OTUs) by comparing with SILVA reference databases (V128) (Quast et al., 2013) at 97% similarity. The minimum reads number of samples (24,097) was used for data normalization. Community richness, evenness and diversity analysis (Shannon, Simpson, Shannoneven, Simpsonenven, ACE, Chao and Good’s coverage) were analyzed using the Mothur *T*-test (with 95% confidence
Spearman rank correlation algorithm. A coefficient of correlation was calculated to estimate the relationship between bacterial genera and antibody isotypes. Statistical analysis was performed using the R statistical package through the STAMP (Parks et al., 2019) software. Differences were determined using STAMP (Parks et al., 2019) software commands “classify.otu” and “make.biom”. PICRUSt (Langille et al., 2013) was also performed to detect abundance taxa at the phylum level (80% threshold) (Wang et al., 2007) based on the RDP (Cole et al., 2009) classifier. LEfSe (Segata et al., 2011) was also performed to detect abundance taxa using “metastats” command. Differences among the two groups were also assessed using Analysis of Molecular Variance (AMOVA) in Mothur. Microbiome functions were analyzed using PICRUSt based on the KEGG (AMOV A) in Mothur. Microbiome groups were also assessed using Analysis of Molecular Variance (AMOVA) in Mothur. Microbiome functions were analyzed using PICRUSt (Langille et al., 2013) based on the KEGG pathways by normalizing the 16S rRNA copy numbers. The input file (biom file) of PICRUSt was calculated using the Mothur software command “classify.otu” and “make.biom”, and then the input file was uploaded to the online PICRUSt for function analysis. Differences were determined using STAMP (Parks et al., 2014).

**Correlation of Antibody Isotypes and Bacteria**

Statistical analysis program-R Package was performed to calculate the coefficient relationship between bacterial genera present and immunoglobulin isotypes using the non-parametric Spearman rank correlation algorithm. A coefficient of >0.68 or <−0.68 was considered to represent strong correlation (Taylor, 1990).

**RESULTS**

**Infection and Blood Serum Isotypes**

In this study 14 mice were transplanted with 35 micro-cysts of *E. granulosus*. All the mice were successfully infected with an average number of 16 (SD ± 7.0) cysts and an average size of 6.4 mm (0.1–21 mm) in diameter (Table 1). Serological antibody tests showed that these infected mice had a predominantly IgG1 antibody response against HCF antigens, followed by IgG2b, IgG2a and IgG3 (Figure 1), indicating *E. granulosus* infection induced a predominant Th2 response.

**Bacterial Populations in the Stool Samples**

Stool samples from the 25 mice were collected for gastrointestinal microbiota analysis, including 14 samples from the mice infected with cysts of *E. granulosus* (Infected Group) and 11 control samples from mice without infection (Uninfected Group). A total of 1,383,569 16S rRNA genes were identified by high-throughput DNA sequencing analysis after filtering through quality control filters. The gene numbers ranged from 24,097 (from one in Uninfected Group) to 86,478 genes. To normalize the data to avoid statistical bias, 24,097 genes from the mice with the lowest gene number were used as a baseline for normalization of all the sequences. OTU (97% similarity) analysis was used to estimate richness, evenness and diversity of the bacterial communities. A total of 13,353 OTUs were obtained including 9,118 OTUs from mice infected with *E. granulosus*, and 8,423 OTUs from the uninfected mice (Table 2). No significant difference was evident between the two groups of mice in terms of OTU numbers (p > 0.05). The Good’s coverage was over 93.5% (93.5–97.8%) for each sample, and over 98% for the two groups, respectively (Table 2), meaning that the sequencing depth was sufficient to undertake microbiota analysis with two groups.

**Core Microbiome in the Gut of Mice**

Ribosomal Database Project analysis showed that 99.7% of the 16S rRNA genes were aligned into nine phyla with the common bacteria *Firmicutes, Bacteroidetes* and *Proteobacteria* being dominant in both infected and uninfected groups. RDP analysis clustered 93.5% of the genes (OTUs) into 58 families and 13 families were identified as the major taxa and core microbiomes co-existing in the two groups. The genes in those families accounted for 91.61 and 94.27% of the microbiome community in the infected group and uninfected group, respectively (Table 3). Among the 13 families, *Lachnospiraceae* was mostly predominant in both groups, accounting for 41.42 and 43.92% of the total microbiome, respectively. *Ruminococcaceae* and

**TABLE 2 |** The diversity evaluation of the microbiomes of mice infected with cystic echinococcosis and uninfected mice.

| Group        | Sample | OTUs | Coverage (%) | Chao     | ACE       | Simpson Even | Shannoneven | Shannon | Simpson |
|--------------|--------|------|--------------|----------|-----------|--------------|-------------|---------|---------|
| Infected     | 14     | 9,118| 98.50        | 26,873.315| 32,509.715| 0.01444      | 0.646977    | 5.899138| 0.007595|
| Uninfected   | 11     | 8,423| 98.25        | 25,099.202| 27,006.635| 0.01675      | 0.660714    | 5.972012| 0.007084|

Infected, mice infected with *E. granulosus*; Uninfected, uninfected mice; OTUs, operational taxonomic units; Chao, Chao index; ACE, ACE index.

**TABLE 3 |** The major families of microbiota in *E. granulosus* infected mice and uninfected mice.

| Family                      | Uninfected (%) | Infected (%) | p-value |
|-----------------------------|----------------|--------------|---------|
| Lachnospiraceae             | 41.42          | 43.92        | > 0.05  |
| Porphyromonadaceae          | 14.21          | 15.75        | > 0.05  |
| Ruminococcaceae             | 11.52          | 18.07        | > 0.05  |
| Rikenellaceae               | 6.61           | 5.33         | > 0.05  |
| Bacteroidaceae              | 5.80           | 5.93         | > 0.05  |
| Helicobacteraceae           | 3.31           | 3.94         | > 0.05  |
| Prevotellaceae              | 2.32           | 2.20         | > 0.05  |
| Desulfovibrionaceae         | 1.94           | 1.94         | > 0.05  |
| Coriobacteriaceae           | 0.14           | 0.20         | > 0.05  |
| Deferribacteraceae          | 1.82           | 1.80         | > 0.05  |
| Lactobacillaceae            | 1.80           | 0.90         | > 0.05  |
| Enterobacteriaceae          | 0.39           | 0.17         | > 0.05  |
| Erysipelotrichaceae         | 0.33           | 0.29         | > 0.05  |

Uninfected, uninfected mice; Infected, mice infected with *E. granulosus*. Echinoscopiosis Impact on Microbiota in Mice.
Porphyromonadaceae were also dominant (>10% of the entire microbiome in both groups).

Among the 58 families, 40.4% of genes (OUTs) have genus reference information and were aligned into 105 classified genera (Figure 2). There were 57 genera co-existing in both groups. However, there were 24 genera present in the Infection Group and another 24 genera in the Uninfected Group. The proportion of all the group unique genera was less than 0.01%, and no significant differences were found between the two groups.

Among the 105 classified genera, 33 were core genera (with each genus comprising >0.1% of total the microbiome), including Bacteroides, Odoribacter, Clostridium XIVA, Helicobacter, Alistipes, Barnesiella and Mucispirillum (Table 4). Among the predominant genera, there were 27 ubiquitous (core) genera which were consistently found in all samples and comprised more than 38% of the total microbiome.

At the OTU level, there were significant differences between the two groups; 101 OTUs were significantly increased and 49 OTU were significantly decreased in the infected group (p < 0.05) (Table 5). Of note, most (59.6%) OTUs were unclassified into genera as no classification information is available for these OTUs.

### Bacterial Composition in Different Groups

LEfSe analysis showed the composition of the bacterial populations in the guts of the infected and uninfected mice was similar, whereas richness, evenness and diversity were only slightly changed (Table 2 and Figure 3). In contrast, AMOVA analysis showed significant difference between the two groups for the microbiota (PAMOVA = 0.029). Species richness (OTU, ACE and Chao index) was higher in the E. granulosus Infected Group, and the evenness (Shannoneven and Simpson even) was lower in this group compared with the uninfected mice. As richness and evenness combined, there was no significant difference in the diversity between the two groups (p > 0.5). Consequently, the results of these analyses indicated that E. granulosus infection did not alter much of the composition of the core bacteria present in the mouse gut significantly, although some rare bacteria in very low abundance were increased.

Among the 13 core families, LEfSe analysis showed no significant difference between the E. granulosus Infected Group and Uninfected Group in terms of microbiome. Among the major abundant genera, three showed significant differences between the groups (Table 4, LDA > 2, p < 0.05). The infected mice significantly increases two genera, including Eisenbergiella (1.9 times) and Parabacteroides (17.5 times) compared with the uninfected mice (p < 0.05).

### Predicted Functional Potential Changes in the Microbiomes of the E. granulosus Infection and Uninfected Groups

We used PICRUSt to predict and compare the microbial functional potential changes between the two groups. A total of 47 Kos were found to be significantly increased in the Infected
**Correlations Between Bacterial Composition and Immunoglobulin Isotypes**

Spearman correlation analysis showed strong correlations of IgG, IgG1 and IgG2a with nine major genera (Table 7 and Figure 4). The numbers of *Enterorhabdus*, *Barnesiella* and *Clostridium* XIVa were positively correlated with IgG1, IgG2a and IgG2b levels, respectively. IgA was positively associated with increased numbers of genera *Clostridium IV*, *Lachnospiraceae Incertae sedis* and *Mucispirillum*. In addition, IgG3 was associated with decreased numbers of genera *Escherichia/Shigella*, *Ruminococcus*, *Ruminococcus/Intestinimonas* and *Ruminococcus*, respectively (Table 7).

**DISCUSSION**

The ecological balance of the microbiota in the gut is crucial for maintaining healthy condition (Cani et al., 2008). Disruption of the balance of the gut microbiota is associated with a range of diseases, including colorectal cancer, autoimmune diseases, metabolic diseases, among others (Sokol et al., 2008; Jiang et al., 2015). In this study, we showed that *E. granulosus* infection increased two genera of gut microbiota: *Eisenbergiella* and genus...
**TABLE 5** | The abundance of OTU (p < 0.01) in the infected mice and uninfected mice determined by LEfSe analysis.

| #OTU     | Taxonomy                                      | Uninfected: mean rel.freq. (%) | Infected: mean rel.freq. (%) | p-values |
|----------|-----------------------------------------------|---------------------------------|------------------------------|----------|
| OTU00452 | Clostridiales; Lachnospiraceae; Clostridium XIVA | 0.0139687                       | 0.04387                      | 0.000351 |
| OTU00048 | Desulfovibrionales; Desulfovibrionaceae; unclassified Desulfovibrionaceae | 0.0067907                       | 0.93419                      | 0.000472 |
| OTU00409 | Clostridiales; Lachnospiraceae; Acetatifactor | 0.0033954                       | 0.040906                     | 0.001051 |
| OTU00832 | Desulfovibrionales; Desulfovibrionaceae; unclassified Desulfovibrionaceae | 0                          | 0.013043                     | 0.001199 |
| OTU01146 | Clostridiales; Lachnospiraceae; unclassified Lachnospiraceae | 0.0075453                       | 0.000296                     | 0.001367 |
| OTU01119 | Clostridiales; Lachnospiraceae; unclassified Lachnospiraceae | 0.003773                        | 0.005336                     | 0.001834 |
| OTU00050 | Bacteroidales; Porphyromonadaceae; Parabacteroides | 0.1011065                       | 0.450856                     | 0.002202 |
| OTU00186 | Clostridia; Clostridiales; unclassified Clostridiales | 0.1844817                       | 0.046538                     | 0.00282  |
| OTU00122 | Clostridiales; Lachnospiraceae; Eisenbergiella | 0.0211267                      | 0.335252                     | 0.002849 |
| OTU00410 | Clostridiales; Ruminococcaceae; Oscillibacter | 0.0045272                      | 0.034681                     | 0.003348 |
| OTU00058 | Campylobacterales; Helicobacteraceae; Helicobacter | 0.5880985                       | 0.596103                     | 0.003788 |
| OTU00720 | Clostridiales; Clostridiales; unclassified Clostridiales | 0.0015091                      | 0.01245                      | 0.00456  |
| OTU01984 | Desulfovibrionales; Desulfovibrionaceae; unclassified Desulfovibrionaceae | 0                          | 0.002668                     | 0.006591 |
| OTU01307 | Clostridiales; Lachnospiraceae; unclassified Lachnospiraceae | 0.0003773                      | 0.004743                     | 0.007635 |
| OTU01051 | Bacteria; Firmicutes; unclassified Firmicutes | 0.0003773                       | 0.007411                     | 0.007741 |
| OTU02591 | Desulfovibrionales; Desulfovibrionaceae; unclassified Desulfovibrionaceae | 0                          | 0.001779                     | 0.008089 |
| OTU01588 | Bacteroidales; Porphyromonadaceae; Odoribacter | 0.0033954                      | 0.00296                      | 0.008438 |
| OTU00715 | Desulfovibrionales; Desulfovibrionaceae; unclassified Desulfovibrionaceae | 0.0120724                      | 0.003261                     | 0.008709 |
| OTU00480 | Clostridiales; Lachnospiraceae; unclassified Lachnospiraceae | 0.0049044                      | 0.049206                     | 0.009747 |

**FIGURE 3** | The microbial diversity variability in mice infected (Infected Group) or uninfected (Uninfected Group) with *E. granulosus*. LEfSe analysis showed the composition of the bacterial populations in the guts of the infected and uninfected mice was similar, whereas richness, evenness and diversity were no changed.

*Parabacteroides*, with most genera remaining unchanged. The two genera are in the family *Lachnospiraceae*. Their increase may impact on human health (Plieskatt et al., 2013), and may be associated with diabetes in mice (Kameyama and Itoh, 2014).

At the OTU level, there were 150 OTUs significantly changed in the infected mice. However, among the OTUs, only 49 OTUs have taxonomic information at genus level with 101 without predicted taxonomic classification information, which limited our further analysis (**Table 5**). Additionally, the LEfSe analysis also showed genera *Eisenbergiella* and *Parabacteroides* increased in the infected mice, suggesting that these two genera of bacteria might be biomarkers for *E. granulosus* infection. In our study, the genus *Eisenbergiella* in the family *Lachnospiraceae* was up-regulated significantly in the infected mice, however, there is very limited biological function information on this genus. Combined with antibody analysis in this study, this genus of bacteria may be associated with a Th2 response.

Another increased genus in Infected Group is *Parabacteroides* whereas species belonging to the genus *Parabacteroides* are saccharolytic (Rajilic-Stojanovic and de Vos, 2014), being...
TABLE 6 | Functional predictions using the PICRUSt base on 16S rRNA gene copy numbers.

| KO accession | Annotation                                                                 | Infected | Uninfected | p-value |
|--------------|-----------------------------------------------------------------------------|----------|------------|---------|
| K00655       | 1-acyl-sn-glycerol-3-phosphate acyltransferase [EC:2.3.1.51]                | 17,607   | 15,826     | 0.01887 |
| K02312       | 2,3-dihydroxybenzoate-AMP ligase [EC:2.7.7.58]                               | 4,214    | 3,074      | 2.43E−02|
| K01826       | 5-carboxymethyl-2-hydroxymuconate isomerase [EC:5.3.3.10]                   | 2        | 0          | 0.015531|
| K01905       | Acetyl-CoA synthetase (ADP-forming) [EC:6.2.1.13]                            | 4,216    | 3,063      | 0.02436 |
| K12554       | Alanine adding enzyme [EC:2.3.3.2.-]                                        | 1        | 0          | 0.020538|
| K03541       | Amylosucrase [EC:2.4.1.4]                                                   | 4,223    | 3,068      | 0.023681|
| K03325       | Arsenite transporter, ACR3 family                                           | 5,022    | 3,773      | 1.16E−02|
| K01305       | Beta-aspartyl-dipeptidase (metallo-type) [EC:3.4.19.-]                      | 4,245    | 3,082      | 0.023369|
| K11754       | Dihydrofolate synthase/folylpolyglutamate synthase [EC:6.3.2.12 6.3.2.17]    | 26,899   | 24,234     | 0.011904|
| K07458       | DNA mismatch endonuclease, patch repair protein [EC: 3.1.-.-]               | 4,883    | 3,692      | 0.01095 |
| K06212       | Formate transporter                                                         | 4,268    | 3,116      | 0.021367|
| K13892       | Glutathione transport system ATP-binding protein                            | 4,226    | 3,072      | 0.024579|
| K04653       | Hydrogenase expression/formation protein HypC                               | 7,620    | 6,473      | 0.017468|
| K04654       | Hydrogenase expression/formation protein HypD                               | 7,624    | 6,475      | 0.016744|
| K04655       | Hydrogenase expression/formation protein HypE                               | 7,696    | 6,567      | 0.019062|
| K04656       | Hydrogenase maturation protein HypF                                         | 7,630    | 6,479      | 0.016758|
| K04652       | Hydrogenase nickel incorporation protein HypB                               | 7,888    | 6,740      | 0.021211|
| K09091       | Hypothetical protein                                                        | 4,225    | 3,076      | 2.37E−02|
| K09703       | Hypothetical protein                                                        | 4,217    | 3,064      | 0.024347|
| K07301       | Inner membrane protein                                                      | 8,454    | 7,324      | 0.000592|
| K03779       | L(+)-tartrate dehydratase alpha subunit [EC:4.2.1.32]                       | 4,264    | 3,106      | 0.024966|
| K00879       | L-fuculokinase [EC:2.7.1.51]                                                | 4,246    | 3,099      | 0.023788|
| K08369       | MFS transporter, putative metabolite:H+ symporter                           | 4,280    | 3,112      | 0.021188|
| K02018       | Molybdate transport system permease protein                                 | 10,849   | 8,353      | 2.09E−02|
| K03637       | Molybdenum cofactor biosynthesis protein C                                  | 5,564    | 4,311      | 0.024072|
| K07474       | Phage terminase small subunit                                               | 4,318    | 3,190      | 0.022446|
| K02759       | PTS system, cellbiose-specific IIA component [EC:2.7.1.69]                  | 4,336    | 3,207      | 0.022898|
| K02760       | PTS system, cellbiose-specific IIB component [EC:2.7.1.69]                  | 4,448    | 3,372      | 0.019282|
| K02777       | PTS system, glucose-specific IIA component [EC:2.7.1.69]                    | 4,677    | 3,516      | 0.016117|
| K10026       | Queuosine biosynthesis protein QueE                                         | 4,760    | 3,668      | 2.47E−02|
| K12996       | Rhamnosyltransferase [EC:2.4.1.-]                                           | 4,219    | 3,080      | 0.024267|
| K03086       | RNA polymerase primary sigma factor                                         | 17,918   | 16,006     | 3.41E−03|
| K05297       | Rubredoxin-NAD+ reductase [EC:1.18.1.1]                                      | 4,352    | 3,216      | 0.020307|
| K03438       | S-adenosyl-methyltransferase [EC:2.1.1.-]; 16S rRNA (cytosine1402-N4)-methyltransferase [EC:2.1.1.199] | 16,924   | 15,121     | 0.014162|
| K07313       | Serine/threonine protein phosphatase 1 [EC:3.1.3.16]                        | 5,067    | 3,888      | 0.010963|
| K02945       | Small subunit ribosomal protein S1                                         | 16,311   | 14,363     | 1.68E−03|
| K05814       | sn-glycerol 3-phosphate transport system permease protein                   | 4,248    | 3,118      | 0.023484|
| K11928       | Sodium/proline symporter                                                    | 4,419    | 3,257      | 0.022096|
| K01695       | Tryptophan synthase alpha chain [EC:4.2.1.20]                               | 9,531    | 8,500      | 0.021878|
| K07665       | Two-component system, OmpR family, copper resistance phosphate regulon response regulator CusR | 7,719   | 6,661      | 0.012516|
| K04784       | Yersiniabactin non-ribosomal peptide synthetase                             | 4,212    | 3,062      | 0.024426|
| Average      |                                                                           | 2,156    | 2,116      |         |

The numbers of Infected and Uninfected were the normalization of copy numbers. Uninfected, uninfected mice; Infected, mice infected with E. granulosus.

producers of short chain fatty acids (SCFAs) including acetate, propionate and butyrate as bacterial fermentation products (Cummings et al., 1987; Correa-Oliveira et al., 2016; Lloyd-Price et al., 2016). SCFAs act as links between the microbiota and the host immune system (Correa-Oliveira et al., 2016). The liver is the major systemic organ for SCFA metabolism and consumption
TABLE 7 | Genera of bacteria correlating highly with serum antibody isotypes in *E. granulosus* infected mice and uninfected mice.

| Genus           | Uninfected (%) | Infected (%) | Factor | Uninfected | Infected | Spearman Coef | p-value   |
|-----------------|----------------|--------------|--------|------------|----------|---------------|-----------|
| Barnesiella     | 2.06           | 2.99         | IgG2a  | 0.2758     | 0.8682   | 0.9           | 0.037386  |
| Ruminococcus    | 0.13           | 0.02         | IgG1   | 0.2801     | 3.4034   | −0.9          | 0.037386  |
| Ruminococcus    | 0.13           | 0.02         | IgG2b  | 0.2434     | 0.8148   | −0.9          | 0.037386  |
| Ruminococcus    | 0.13           | 0.02         | IgG3   | 0.2329     | 1.0469   | −1            | 0         |
| Ruminococcus    | 0.13           | 0.02         | IgM    | 0.3879     | 2.7058   | −0.9          | 0.037386  |
| Clostridium IV  | 0.04           | 0.06         | IgA    | 0.1563     | 0.3758   | 0.9747        | 0.004818  |
| Enterorhabdus   | 0.06           | 0.08         | IgG1   | 0.2801     | 3.4034   | 0.9           | 0.037386  |
| Intestinimonas  | 0.13           | 0.11         | IgG2b  | 0.2434     | 0.8148   | −0.9          | 0.037386  |
| Escherichia/Shigella | 0.39 | 0.17 | IgG    | 0.7014     | 3.9271   | −1            | 0         |
| Lachnospiracea incertae sedis | 0.56 | 0.61 | IgA    | 0.1564     | 0.3758   | 0.9747        | 0.004818  |
| Mucispirillum   | 1.81           | 2.72         | IgA    | 0.1564     | 0.3758   | 0.9747        | 0.004818  |
| Clostridium XIVA| 2.70           | 4.19         | IgG2b  | 0.2434     | 0.8148   | 0.9           | 0.037386  |

Uninfected, uninfected mice; Infected, mice infected with *E. granulosus*.

(†Kim et al., 2014), SCFAs released by the gut and equaled by hepatic uptake (Bloemen et al., 2009). *Parabacteroides* has evolved to contain a gene encoding a major capsid protein (Rosenwald et al., 2014) one of the phage orthologous groups (Kristensen et al., 2013). One report demonstrated that *Parabacteroides* was prevalent in diabetic (Wu et al., 2010). The increasing of genus *Parabacteroides* in hydatid infection may be associated with hepatic alteration.

Functional predictions showed seven pathways of the gut microbiota in the *E. granulosus* infection group were altered compared with the uninfected group. These pathways included biotin, lipid metabolism, and tryptophan metabolism.
The synergistic effect of bacteria leads to the difference of gut flora metabolic pathways due to some or all intestinal bacteria involving in metabolism. Biotin metabolism in the intestine is regulated through transcriptional and post-transcriptional mechanisms. Its balance plays a key role in regulating the absorption and the function of biotin in tissues (Zoetendal et al., 2012). Based on pathway impact analysis, we found that tryptophan metabolism was decreased in the E. granulosus infection group. In mice infected with schistosomes, tryptophan or compounds from tryptophan metabolism were up-regulated and increased in urine which indicate possible problems in tryptophan metabolism in these infected animals (Njagi et al., 1992; Wang et al., 2004).

We showed that the bacterial composition of nine major genera had strong correlations with the levels of IgG, IgG1 and IgG2a antibodies against HCF antigens (Figure 4). The numbers of genera Enterorhabdus, and Clostridium XIVA were positively correlated with IgG1 and IgG2b levels, indicating that these bacteria can be tolerated with those Th2 associated antibodies or Th2 responses may benefit those genera of bacteria. Meanwhile, IgG2a, a Th1 associated antibody, was associated with increased number of genus Barnesiella, indicating Th1 has a role for increasing genus Barnesiella. Our data also showed that Th2 associated antibodies IgG1 and IgG2b and IgG3 decreased numbers of genera Escherichia/Shigella, and Ruminococcus, Ruminococcus/Intestinimonas (Table 7 and Figure 4). Interestingly, Intestinimonas decreased significantly in E. granulosus infection group as a differential genus by LEfSe analysis, perhaps it is associated with some kinds of change, then we concluded that it is highly related to IgG2b by Spearman coefficient correlation analysis and is consistent with the immune background of E. granulosus infection. So IgG2b may play impartment role in inhibition of Intestinimonas. Clostridium has been found to be associated with a number of diseases. It showed that Clostridium may participate in antibiotic-associated diarrhea (Buffie et al., 2015) and damages the human intestine in vitro (Fernandez Miyakawa et al., 2005). Barnesiella is present in the healthy intestinal tract and is influenced by antibiotics, and intestinal colonization with Barnesiella confers resistance to intestinal domination and bloodstream infection with vancomycin-resistant Enterococcus (Ubeda et al., 2013).

In summary, we explored gut microbiota in mice infected with E. granulosus, and found that chronic E. granulosus infection increased 101 OTUs including two genera of gut microbiota in mice. Functional prediction showed seven pathways of gut microbiota were altered, and bacterial composition of major genera had positive correlations with IgG1 and IgG2b in E. granulosus infected mice.

Whereas more than 85% of the genomic sequences between mouse and Homo are conserved, overall gene expression and its regulation are considerably different between the two species (Hugenholtz and de Vos, 2018). Human and mouse seems to be similar at phylum level, Bacteroidetes and Firmicutes are the two major bacterial phyla of the intestinal tract (Rawls et al., 2006). However, we do not know whether E. granulosus infection will affect human intestinal flora in the same way as the mouse and further studies are now required to understand the further possible mechanisms associated with altered colonization resistance after helminth infection and to determine changes in the gut microbiota of patients with CE.

ACCESSION NUMBERS
The sequence data have been submitted to the GeneBank Sequence Read Archive (Accession Number PRJN596089).

AUTHOR CONTRIBUTIONS
WZ, JL, and DM contributed to conception and design of the study. JB organized the database. LH, WQ, and TW finished the animal experiments. HZ, ZZ, and YW performed the statistical analysis. JB and YW wrote the first draft of the manuscript. GG, XZ, and BG wrote sections of the manuscript. All authors contributed to manuscript revision, read and approved the submitted version.

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