Intestinal Protozoan Infections Shape Fecal Bacterial Microbiota in Children from Guinea-Bissau.

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Research

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Abstract

Background: Intestinal parasitic infections, caused by helminths and protozoa, are globally distributed and major causes of worldwide morbidity. The gut microbiota may modulate parasite virulence and host response upon infection. The complex interplay between parasites and the gut microbiota is poorly understood, partly due to sampling difficulties in remote areas with high parasite prevalence and burden.

Results: In a large study of children in Bissau, Guinea-Bissau, we found high prevalence of intestinal parasites, including hookworms, Entamoeba spp. and Giardia lamblia. By high-throughput sequencing of the 16S rRNA gene of fecal samples stored on filter paper from a total of 1,204 children, we demonstrate that the bacterial microbiota is more or less unaltered by helminth infections, whereas it is shaped by the presence of both pathogenic and non-pathogenic protozoa, including Entamoeba spp. and Giardia lamblia. Within-sample (alpha) diversity remains largely unaffected, whereas overall community composition (beta diversity) is significantly affected by infection with both non-pathogenic Entamoeba coli \((R^2=0.0131, P=0.0001)\) and Endolimax nana \((R^2=0.00902, P=0.0001)\), and by pathogenic Entamoeba histolytica \((R^2=0.0164, P=0.0001)\) and Giardia lamblia \((R^2=0.00676, P=0.0001)\). Heavy infection load with multiple parasite species induces more pronounced shifts in microbiota community than mild ones. A total of 31 bacterial genera across all four major gut bacterial phyla associates with protozoan infection, including increased abundance of Prevotella, Campylobacter and two Clostridium clades (IV and WIVb), and decreased abundance of Collinsella (associated with irritable bowel syndrome), Lactobacillus, Ruminococcus, Veillonella and one Clostridium clade (XVIII). Further, we demonstrate that filter papers are usable for storage of samples at room temperature in large-scale studies where immediate freezing is not possible with minor alterations of microbiota composition.

Conclusion: In the present largest-to-date study, we demonstrate that the fecal bacterial microbiota is shaped by intestinal parasitic infection, with most pronounced associations for protozoan species. Our results provide insights into the interplay between the microbiota and intestinal parasites, which are useful to understand infection outcome and design further studies aimed at optimizing treatment strategies.

Background

Intestinal parasitic infections are among the most common infections in humans and contribute to global morbidity and mortality. Infections are distributed worldwide, with highest prevalence in tropical and subtropical regions, predominantly in developing countries [1]. Intestinal parasitic infections are caused by a wide range of both helminth and protozoan species of varying importance and severity. Soil-transmitted helminths (STHs) include roundworm (Ascaris lumbricoides), whipworm (Trichuris trichiura) and hookworms (Necator americanus and Ancylostoma duodenale), and are estimated to affect more than 1.75 billion people worldwide [2, 3], and further cause a loss of almost 4 million disability-adjusted life years (DALYs) [4]. Among the most important intestinal protozoans to humans are Giardia lamblia, which infect more than 250 million people worldwide [5–8], and Entamoeba histolytica, which contributes with approx. 400,000 cases annually and account for 40–110,000 deaths [5, 9].

Despite several initiatives to lower incidence, including the distribution of mass drug treatment to STHs [10] and improvement of water, sanitation and hygiene [11], intestinal parasitic infections remain a severe global health concern. Although a general decline in STH since 1990 has been documented, a rise in other intestinal parasitic infections has been observed, including amebiasis [4], and the infections are still among the most important causes of morbidity in high-prevalent regions. Mass drug treatment campaigns against STHs have highly varying efficacy, especially towards T. trichiura-infection [12, 13]. Additionally, drug resistance is an increasing problem in veterinary medicine, and potentially also in humans, as metronidazole resistance against Entamoeba spp. has been reported [14, 15]. Clearly, there is an urgent need to develop new approaches to control intestinal parasitic infections. One possible way to achieve this is by enhancing understanding of the interaction between intestinal parasites and their host.

The parasite-host relationship is highly complex, and at present only partly understood. It is generally accepted that intestinal parasites have co-evolved with and adapted to their hosts, a feature that is also proposed for prokaryotic microbes of the intestinal tract [14]. To enhance understanding of the interaction between intestinal parasites and their vertebrate host, some focus over the last years has been on how intestinal parasites modulate or manipulate the hosts' gut microbiota, i.e. the commensal microbes (mainly bacteria) within the gastrointestinal tract [15–21]. As the gut microbiota has been demonstrated to be pervasive and essential for human health, including maturation and regulation of the host immune system and protection against pathogens [15–17], and as intestinal parasites share the same niche as the microbiota, it is plausible that a parasite-microbiota-interaction occurs, and that this interaction may determine infection symptomatology, virulence and outcome [18]. Most intestinal parasites secrete immunomodulatory molecules, which may change the local environment within the gut, and thereby induce alterations in the microbiota [19]. Most clinical and preclinical research has focused on how helminths affect the gut microbiota, and the results have not been uniform, as reviewed in [20–22]. The majority of studies undertaken in humans are characterized by a relatively low number of study participants (<100), with variation in both sampling and sequencing technique [20]. The microbiota diversity has both been observed to increase and decrease, or even be unaffected due to intestinal helminth infection. For instance, one study reported a decrease in diversity due to infection with T. trichiura [23], whereas another reported an increase [24]. Further, various bacterial taxa have been associated with infection, but with no clear trend across studies. A recent study covering two geographically separate regions, Liberia and Indonesia, demonstrated that specific members of the gut microbiota could discriminate STH-infected and non-infected individuals. Further, the study demonstrated an increase in Actinobacteria spp. due to T. trichiura infection in a cohort from Liberia, but not in Indonesia [24], which may reflect overall differences in the gut microbiota between geographically separate populations. Additionally, the study found that microbiota diversity was increased in both cohorts due to infection with both T. trichiura, A. lumbricoides and hookworms. In a study conducted in Cameroon, an increased diversity and increase in Bacteroidetes was seen in individuals infected with Entamoeba histolytica [25]. This study further highlighted the importance of lifestyle and standard of living on the gut microbiota composition, as the authors studied both hunter-gatherers, farming- and fishing populations. In a study from Côte d’Ivoire, the relative abundance of Bifidobacteria and Escherichia was demonstrated to increase upon infection with G. lamblia by targeted qPCR [26].
In the present study, we explored alterations in the fecal bacterial microbiota due to intestinal parasitic infections in a large cohort of children from the capital of one of the poorest countries in the world, Guinea-Bissau, Western Africa. From here on the bacterial microbiota studies is referred to simple as the microbiota. Using this cohort, we have recently investigated the prevalence of intestinal parasitic infections in both health-care seeking children (referred to as cohort I) and children from the background population (referred to as cohort II), all aged 2–15 years and found that infections were highly prevalent in both cohorts [27]. In 566 children in cohort I and 708 in cohort II, we found that prevalence of intestinal helminths was 13.8% and 9.6%, respectively (Fisher's exact test between groups, P = 0.021), whereas prevalence of intestinal protozoa was 41.5% and 46.0%, respectively (Fisher's exact test between groups, P = 0.112). Helminth infections were mainly due to hookworms, and protozoan infections were dominated by both pathogenic and non-pathogenic species, including *Entamoeba coli*, *Entamoeba histolytica/dispar* and *Giardia lamblia*. Upon fecal parasitological investigation by microscopy, fecal samples were applied to filter papers, and kept at ambient temperature. By high-quality 16S rRNA gene sequencing (> 10,000 reads per sample) of fecal samples from 1,204 of these children, we demonstrate that the fecal microbiota significantly associates with intestinal parasitic infection, and that the association is stronger for children infected with protozoa compared to helminths. Further, we here demonstrate that the fecal microbiota from samples stored at ambient temperature on filter (fecal occult blood test, FOBT) papers for up to 1,000 days can facilitate largescale microbiome studies in remote areas. We previously validated the use of filter paper in a benchmark study that show high resemblance of fecal microbiotas from filter paper with conventionally stored samples, with satisfying DNA yield and sequencing quality [28]. Thereby, we demonstrate that long-term storage on FOBT papers is an applicable approach for large-scale sample collection in field settings, where immediate freezing of samples is not possible.

To our knowledge, this is the largest study to date investigating the relationship between intestinal parasites and alterations of the fecal microbiota. This exploratory study should enable us to move forward with targeted questions towards understanding the role of the microbiota in intestinal parasitic infections and infection-associated complications. This may lead to new therapeutic strategies, as manipulation of the microbiota by the administration of probiotics might be an effective way to enhance the host's immune system to control the parasite. Furthermore, expanding the knowledge may help to differentiate between beneficial and pathogenic intestinal parasites.

**Results**

**Cohort characteristics and parasite prevalence**

The dataset includes microscopic investigation for intestinal parasites from 1,274 children, included between August 2015 and April 2017 in urban Bissau, Guinea-Bissau. Details on the study design, parasitological examination methods and results are described in detail elsewhere [27]. From the cohort, a total of 1,264 fecal samples were applied to FOBT paper, of which 1,253 underwent 16S rRNA sequencing. A total of 60 samples were excluded due to low DNA yield or quality, low sequence reads or recent antibiotic usage, and the final sample size included in this study was 1,204 samples. A flow diagram for the inclusion and final sample size is provided in Fig. 1.

Characteristics of the study participants, as well as storage time of fecal sample on filter paper at room temperature, is provided in Table 1. Of the included subjects with available sequence data, the median age was 6 years in both cohorts (healthcare seeking children and children from the background population) with 54% boys (Table 1). There were no significant differences between the two cohorts with regards to age- and gender distribution. However, there were significant differences in some parameters between the two cohorts, including chicken husbandry, source of drinking water, inclusion period (rainy vs. dry season) and sample storage time (Table 1).
Table 1

Cohort characteristics and intestinal parasite prevalence. Characteristics of cohort I (n=529) and cohort II (n=675), which were included in the microbiota analysis. Between-group differences are calculated using Wilcoxon rank sum test, Fisher’s exact test or Kruskal–Wallis equality-of-populations rank test, when appropriate.

| Characteristic                  | Cohort I (n = 529) | Cohort II (n = 675) | Total (n = 1,204) | P value |
|--------------------------------|--------------------|--------------------|-------------------|---------|
| **Gender**                     |                    |                    |                   |         |
| Male                           | 289 (54.6%)        | 361 (53.5%)        | 650 (54.0%)       | 0.367   |
| Female                         | 240 (45.4%)        | 314 (46.5%)        | 554 (46.0%)       |         |
| **Age**                        |                    |                    |                   | 0.227   |
| Age, mean (range)              | 6.7 (2–14)         | 6.9 (2–15)         | 6.8 (2–15)        |         |
| Age, median                    | 6                  | 6                  | 6                 |         |
| **Husbandry**                  |                    |                    |                   |         |
| None                           | 298 (56.3%)        | 374 (55.4%)        | 672 (55.8%)       | 0.397   |
| Pigs                           | 43 (8.1%)          | 35 (5.2%)          | 78 (6.5%)         | 0.181   |
| Ducks                          | 5 (0.9%)           | 14 (2.1%)          | 19 (1.6%)         | 0.247   |
| Chicken                        | 46 (8.7%)          | 81 (12%)           | 127 (10.5%)       | 0.002   |
| Dogs                           | 85 (16.1%)         | 100 (14.8%)        | 185 (15.4%)       | 0.253   |
| Other animals                  | 52 (9.8%)          | 71 (10.5%)         | 123 (10.2%)       | 0.487   |
| **Toilet source**              |                    |                    |                   | 0.129   |
| Poor                           | 422 (79.8%)        | 519 (76.9%)        | 941 (78.2%)       |         |
| Good                           | 107 (20.2%)        | 156 (23.1%)        | 263 (21.8%)       |         |
| **Water source**               |                    |                    |                   | <0.001  |
| Poor                           | 319 (60.3%)        | 171 (25.3%)        | 490 (40.7%)       |         |
| Good                           | 210 (39.7%)        | 504 (74.7%)        | 714 (59.3%)       |         |
| **Season of inclusion**        |                    |                    |                   | 0.038   |
| Dry season                     | 255 (48.2%)        | 366 (54.2%)        | 621 (51.6%)       |         |
| Rainy season                   | 274 (51.8%)        | 309 (45.8%)        | 583 (48.4%)       |         |
| **Sample storage time at RT**  |                    |                    |                   | <0.001  |
| 200–300 days                   | 0 (0.0%)           | 27 (4.0%)          | 27 (2.2%)         |         |
| 301–400 days                   | 20 (3.8%)          | 146 (21.6%)        | 166 (13.8%)       |         |
| 401–500 days                   | 83 (15.7%)         | 124 (18.4%)        | 207 (17.2%)       |         |
| 501–600 days                   | 93 (17.6%)         | 9 (1.3%)           | 102 (8.5%)        |         |
| 601–700 days                   | 101 (19.1%)        | 55 (8.1%)          | 156 (13.0%)       |         |
| 701–800 days                   | 63 (11.9%)         | 94 (13.9%)         | 157 (13.0%)       |         |
| 801–900 days                   | 98 (18.5%)         | 149 (22.1%)        | 247 (20.5%)       |         |
| > 900 days                     | 71 (13.4%)         | 71 (10.5%)         | 142 (11.8%)       |         |
| **Parasite prevalence**        |                    |                    |                   |         |
| Parasite positive (overall)    | 272 (51.4%)        | 336 (49.8%)        | 608 (50.5%)       | 0.817   |
| Positive for 1 parasite         | 182 (34.4%)        | 229 (33.9%)        | 411 (34.1%)       |         |
| Positive for 2 parasites        | 74 (14.0%)         | 92 (13.6%)         | 166 (13.8%)       |         |
| Positive for ≥ 3 parasites      | 16 (3.0%)          | 15 (2.2%)          | 31 (2.6%)         |         |
Overall parasite prevalence was statistically indifferent between the two cohorts (Table 1): In cohort I, 51.4% were positive for at least one intestinal parasite, whereas 49.8% were positive in cohort II (Fisher’s exact test, \( P = 0.817 \)). Prevalence of helminths was higher in cohort I than in cohort II (14.0% vs. 9.6%, Fisher’s exact test, \( P = 0.012 \)), whereas prevalence of protozoa was equal between the two (43.1% vs. 45.3%, Fisher’s exact test, \( P = 0.237 \)). Prevalence of hookworm and *Giardia lamblia* differed between cohorts (10.4% vs. 5.8%, \( P = 0.002 \) and 21.2% vs. 25.6%, \( P = 0.041 \), respectively). Infection with multiple species was equally common in the two cohorts; 14.0% of children from cohort I and 13.6% from cohort II were infected with two parasite species, and 3.0% in cohort I and 2.2% in cohort II were infected with three or more parasite species.

The major intestinal helminth species found in the study participants were *Ancylostoma duodenale* (hookworm, \( n = 94 \)) and *Hymenolepis nana* (dwarf tapeworm, \( n = 36 \)). The most common intestinal protozoan species found in the participants were *Entamoeba coli* (\( n = 98 \)), *Entamoeba histolytica/dispar* (\( n = 214 \)), *Giardia lamblia* (\( n = 285 \)) and *Endolimax nana* (\( n = 94 \)). A number of other intestinal parasite species were found at lower prevalence, but were not included in the present study due to lack of statistical power. Prevalence and distribution of the major intestinal parasites in the two cohorts and statistical differences between groups are provided in Table 1.

### Identification of confounding variables

Several host phenotypic and environmental variables have previously been linked to differences in the gut microbiota composition, including age, diet, vitamin supplementation and antibiotic treatment (as reviewed in [29]). Thus, we excluded all individuals with a history of antibiotics use three months prior to inclusion from the analysis, and adjusted for the potential confounding effect of age, history of vitamin A supplementation (binary variable, (i) yes or (ii) no), toilet source (binary variable, either (i) no private toilet/latrine or (ii) access to private latrine/toilet) and tropical season for sample collection (binary variable, (i) rainy or (ii) dry season). Further, as samples were kept at room temperature on FOBT cards prior to DNA extraction, we adjusted for sample storage time (in days). As there were some differences in characteristics and parasite prevalence between the two cohorts (Table 1), all analyses were performed for both cohorts separately, and jointly, the latter adjusted for cohort status. In the following sections, the reported results are from the joined analysis if not stated otherwise.

### Alpha diversity is largely unaffected by intestinal parasite infection

Relative abundance of bacterial phyla across all samples revealed that the composition of the microbiota in all study participants was dominated by Firmicutes and Bacteroidetes, as expected (Fig. 2).

Three different measures of alpha diversity were calculated to explore possible diversity alterations due to intestinal parasitic infections: Shannon entropy; ACE as a measure of species richness; and phylodiversity as a measure of total unique phylogenetic branch length (Table 2). We compared alpha diversity measurements for individuals with each of nine infection variables (either overall parasite positive, helminth positive, protozoa positive or positive for one of the six specific species, i.e. *Ancylostoma duodenale, Hymenolepis nana, Entamoeba coli, Entamoeba histolytica/dispar, Giardia lamblia* or *Endolimax nana*) against non-infected individuals. Effects on all three diversity indices were limited for all infections, as illustrated in Fig. 3 for phylodiversity (Table 2 and Fig. 3). A significant decrease in ACE diversity was seen in participants in cohort I with *Giardia lamblia* infection (\( \beta = -3.42; P = 0.0470 \)), which was not found in cohort II. Increase in phylodiversity index was seen in cohort II with any intestinal parasite and protozoa (\( \beta = 7.00; P = 0.0153 \) and \( \beta = 6.73; P = 0.0266 \), respectively). Further, all three alpha diversity indices were increased in cohort II with *Entamoeba* spp., which were not observed in cohort I. In the pooled dataset containing both cohorts, a significant increase in phylodiversity was seen upon infection with *Entamoeba* spp. (\( \beta = 11.35; \text{P.adj.}=0.0115 \) for *Ent. coli* and \( \beta = 8.15; \text{P.adj.}=0.0115 \) for *Ent. histolytica/dispar*), however only nominally significant in cohort II. Therefore, changes in alpha diversity was predominantly observed in cohort II, and predominantly for phylodiversity. The changes were mainly an increased diversity in infected individuals in agreement with findings reported in [25].

| Dataset                        | Cohort I (n = 529) | Cohort II (n = 675) | Total (n = 1,204) | \( P \) value |
|--------------------------------|--------------------|--------------------|-------------------|---------------|
| Helminth positive              | 74 (14.0%)         | 65 (9.6%)          | 139 (11.5%)       | 0.012         |
| *Ancylostoma duodenale*        | 55 (10.4%)         | 39 (5.8%)          | 94 (7.8%)         | 0.002         |
| *Hymenolepis nana*            | 14 (2.6%)          | 22 (3.3%)          | 36 (3.0%)         | 0.329         |
| Protozoa positive              | 228 (43.1%)        | 306 (45.3%)        | 534 (44.4%)       | 0.237         |
| *Entamoeba coli*               | 50 (9.5%)          | 48 (7.1%)          | 98 (8.1%)         | 0.086         |
| *Entamoeba histolytica/dispar* | 95 (18.0%)         | 119 (17.6%)        | 214 (17.8%)       | 0.470         |
| *Giardia lamblia*              | 112 (21.2%)        | 173 (25.6%)        | 285 (23.7%)       | 0.041         |
| *Endolimax nana*               | 44 (8.3%)          | 50 (7.4%)          | 94 (7.8%)         | 0.316         |
Intestinal parasitic infections have limited effects on alpha diversity. Three different alpha diversity indices were analyzed for association with infection status using robust regression (Methods). The table shows summary statistics: P value for analysis within participants from cohort I, within participants from cohort II and across all participants (Total), Benjamini-Hochberg adjusted P value for analysis across all individuals (Padj), and association coefficient (Beta). Most pronounced effects are seen in cohort II with protozoan infections. P values and Padj values below 0.05 are highlighted.

| Parasite species | Alpha.div | Cohort I (n = 529) | Cohort II (n = 675) | Total (n = 1,204) |
|------------------|-----------|--------------------|--------------------|------------------|
|                  | Beta      | P                   | Beta               | P                | P                |
| Overall          | Parasite pos. (overall) | PhyloDiv | 1.49±10\(^{-00}\) | 6.24±10\(^{-01}\) | 7.00±10\(^{-00}\) | 1.53±10\(^{-02}\) | 4.59±10\(^{-00}\) | 3.21±10\(^{-02}\) | 2.37±10\(^{-01}\) |
|                  | Shannon   | 8.52±10\(^{-04}\) | 9.76±10\(^{-01}\) | 2.42±10\(^{-02}\) | 3.65±10\(^{-01}\) | 1.66±10\(^{-02}\) | 4.00±10\(^{-01}\) | 5.84±10\(^{-01}\) |
|                  | ACE       | -8.87±10\(^{-01}\) | 4.50±10\(^{-01}\) | 1.44±10\(^{-00}\) | 1.38±10\(^{-01}\) | 7.00±10\(^{-01}\) | 3.49±10\(^{-01}\) | 5.58±10\(^{-01}\) |
| Helminths        | Parasite pos. | PhyloDiv | 2.75±10\(^{-00}\) | 5.79±10\(^{-01}\) | 3.09±10\(^{-00}\) | 5.89±10\(^{-01}\) | 2.52±10\(^{-00}\) | 4.92±10\(^{-00}\) | 6.31±10\(^{-01}\) |
|                  | Shannon   | -1.34±10\(^{-02}\) | 8.31±10\(^{-02}\) | 6.59±10\(^{-02}\) | 3.85±10\(^{-02}\) | 3.90±10\(^{-02}\) | 4.11±10\(^{-02}\) | 5.84±10\(^{-02}\) |
|                  | ACE       | -1.66±10\(^{-01}\) | 9.47±10\(^{-01}\) | -6.28±10\(^{-01}\) | 8.07±10\(^{-01}\) | 6.60±10\(^{-01}\) | 9.74±10\(^{-01}\) | 9.74±10\(^{-01}\) |
| Protozoa         | Parasite pos. | PhyloDiv | 6.20±10\(^{-01}\) | 8.61±10\(^{-01}\) | 6.73±10\(^{-00}\) | 2.66±10\(^{-02}\) | 3.99±10\(^{-00}\) | 4.41±10\(^{-00}\) | 2.37±10\(^{-01}\) |
|                  | Shannon   | 2.32±10\(^{-03}\) | 9.42±10\(^{-02}\) | 2.18±10\(^{-02}\) | 4.36±10\(^{-02}\) | 1.37±10\(^{-02}\) | 5.15±10\(^{-02}\) | 6.31±10\(^{-02}\) |
|                  | ACE       | -1.34±10\(^{-03}\) | 3.01±10\(^{-01}\) | 1.70±10\(^{-00}\) | 9.59±10\(^{-02}\) | 6.17±10\(^{-01}\) | 4.38±10\(^{-01}\) | 5.91±10\(^{-01}\) |
| Helminths Ancylostoma duodenale | Parasite pos. | PhyloDiv | 2.46±10\(^{-00}\) | 6.90±10\(^{-02}\) | 5.37±10\(^{-00}\) | 5.04±10\(^{-01}\) | 4.29±10\(^{-00}\) | 3.06±10\(^{-01}\) | 5.58±10\(^{-01}\) |
|                  | Shannon   | -1.96±10\(^{-03}\) | 9.78±10\(^{-02}\) | 7.54±10\(^{-02}\) | 3.75±10\(^{-02}\) | 5.07±10\(^{-02}\) | 3.51±10\(^{-02}\) | 5.58±10\(^{-02}\) |
|                  | ACE       | 6.62±10\(^{-01}\) | 8.13±10\(^{-01}\) | -5.82±10\(^{-01}\) | 8.36±10\(^{-01}\) | -2.34±10\(^{-01}\) | 9.09±10\(^{-01}\) | 9.44±10\(^{-01}\) |
| Hymenolepis nana | Parasite pos. | PhyloDiv | 8.57±10\(^{-00}\) | 4.98±10\(^{-01}\) | -6.08±10\(^{-00}\) | 6.40±10\(^{-01}\) | 2.80±10\(^{-00}\) | 7.49±10\(^{-01}\) | 8.79±10\(^{-01}\) |
|                  | Shannon   | -5.60±10\(^{-02}\) | 6.88±10\(^{-01}\) | -1.81±10\(^{-02}\) | 9.22±10\(^{-01}\) | -2.23±10\(^{-02}\) | 8.38±10\(^{-01}\) | 9.05±10\(^{-01}\) |
|                  | ACE       | -6.06±10\(^{-00}\) | 2.92±10\(^{-01}\) | -1.51±10\(^{-01}\) | 1.64±10\(^{-01}\) | -5.69±10\(^{-00}\) | 1.59±10\(^{-01}\) | 4.77±10\(^{-01}\) |
| Protozoa Entamoeba coli | Parasite pos. | PhyloDiv | 7.14±10\(^{-00}\) | 1.16±10\(^{-01}\) | 1.64±10\(^{-01}\) | 1.37±10\(^{-02}\) | 1.13±10\(^{-01}\) | 8.49±10\(^{-01}\) | 1.15±10\(^{-02}\) |
|                  | Shannon   | 7.44±10\(^{-03}\) | 8.94±10\(^{-01}\) | 1.37±10\(^{-01}\) | 2.26±10\(^{-02}\) | 7.71±10\(^{-01}\) | 5.93±10\(^{-01}\) | 2.37±10\(^{-01}\) |
|                  | ACE       | 1.20±10\(^{-01}\) | 9.58±10\(^{-01}\) | 5.37±10\(^{-00}\) | 7.18±10\(^{-03}\) | 3.02±10\(^{-00}\) | 4.62±10\(^{-02}\) | 2.37±10\(^{-01}\) |
| Entamoeba histolytica/dispar | Parasite pos. | PhyloDiv | 6.78±10\(^{-00}\) | 1.29±10\(^{-01}\) | 9.82±10\(^{-00}\) | 1.42±10\(^{-02}\) | 8.15±10\(^{-00}\) | 8.38±10\(^{-00}\) | 1.15±10\(^{-02}\) |
|                  | Shannon   | 3.45±10\(^{-02}\) | 4.36±10\(^{-01}\) | 7.49±10\(^{-02}\) | 5.18±10\(^{-02}\) | 5.47±10\(^{-01}\) | 6.15±10\(^{-02}\) | 2.37±10\(^{-01}\) |
|                  | ACE       | -1.13±10\(^{-00}\) | 5.29±10\(^{-01}\) | 2.91±10\(^{-00}\) | 3.37±10\(^{-00}\) | 1.36±10\(^{-00}\) | 2.22±10\(^{-01}\) | 5.08±10\(^{-01}\) |
| Giardia lamblia | Parasite pos. | PhyloDiv | -2.70±10\(^{-00}\) | 4.87±10\(^{-01}\) | 2.97±10\(^{-00}\) | 4.20±10\(^{-01}\) | 5.82±10\(^{-01}\) | 7.97±10\(^{-01}\) | 8.97±10\(^{-01}\) |
for individuals infected with helminth infection; for the overall helminth variable no associations were observed only few associations regarding specific taxa of the fecal microbiota and helminth infections; for the overall helminth variable no associations were observed.

Associations between the parasite infection variables and the relative abundance of individual taxa was evaluated for the phylogenetic level phylum to genus (Table 3). For overall variables, the most pronounced association was seen for protozoa ($R^2 = 1.03 \times 10^{-2}; P = 1.00 \times 10^{-4}$) followed by any parasite ($R^2 = 9.73 \times 10^{-3}; P = 1.00 \times 10^{-4}$) while association for any helminth was insignificant ($R^2 = 2.40 \times 10^{-3}; P = 6.61 \times 10^{-2}$). Therefore, the association between overall parasite infection and the microbiota may be driven by protozoan infections, as further supported by the similar appearance of the ordination-based visualization of overall parasite and protozoa infections in Fig. 4.

A large portion of tested bacterial taxa are associated with intestinal protozoan infection

Associations between the parasite infection variables and the relative abundance of individual taxa was evaluated for the phylogenetic level phylum to genus (Table 4). As mentioned, only minor effects on the fecal microbiota beta-diversity were observed in individuals with helminth infections. Accordingly, we observed only few associations regarding specific taxa of the fecal microbiota and helminth infections; for the overall helminth variable no associations remained significant at genus level ($P_{adj} < 0.05$) while the Epsilonproteobacteria-Campylobacterales-Campylobacteraceae branch was increased in abundance for individuals infected with Ancylostoma duodenale. However, Campylobacter abundance was broadly increased across individuals infected with protozoa, as supported by highly significant association with microbiome composition ($adonis, R^2 = 0.012, P = 0.0001$).
thereby causing the association with *Ancylostoma duodenale* to be non-specific. Beyond *Campylobacter*, the only genus associating with helminth infection was *Collinsella* that associated with *Hymenolepis nana* infection (β = -0.48, Padj = 3.22 × 10⁻²).
### Table 4

Alterations of individual genera due to intestinal protozoa infections. A total of 31 genera from the four major phyla of the gut microbiota were either found to be increased or decreased in abundance due to any of the protozoan infections. The table shows coefficients (Beta) and significance (Padj) for each genus. The Campylobacter genus was found at increased abundance due to any of the protozoan infections. The table shows coefficients (Beta) and significance (Padj) for each genus. The associations were evaluated for genera and the higher taxonomic levels are listed to support evaluation.
| Taxa | Protozoa | Ent. coli |
|------|----------|----------|
| Firmicutes | Negativicutes | Acidaminococcales | Acidaminococcaceae | Phascolarctobacterium | 7.92×10⁻² | 2.77×10⁻² |
| Firmicutes | Negativicutes | Veillonellales | Veillonellaceae | Veillonella | -5.49×10⁻² | 1.98×10⁻³ | -9.43×10⁻² |
| Proteobacteria | Betaproteobacteria | Burkholderiales | Sutterellaceae | Sutterella | -2.10×10⁻¹ | 1.92×10⁻³ |
| Proteobacteria | Gammaproteobacteria | Aeromonadales | Succinivibrionaceae | Succinivibrio | 1.77×10⁻³ | 3.02×10⁻² | 5.85×10⁻³ |
| Proteobacteria | Gammaproteobacteria | Enterobacteriales | Enterobacteriaceae | Escherichia Shigella | -1.24×10⁻⁴ | 7.20×10⁻⁴ |
| Proteobacteria | Gammaproteobacteria | Enterobacteriales | Enterobacteriaceae | Klebsiella | -6.19×10⁻² | 1.61×10⁻² |
| Proteobacteria | Gammaproteobacteria | Pasteurellales | Pasteurellaceae | Haemophilus | -9.47×10⁻² | 8.67×10⁻³ | -1.91×10⁻¹ |
| Proteobacteria | Epsilonproteobacteria | Campylobacterales | Campylobacteraceae | Campylobacter | 1.38×10⁻⁰ | 2.66×10⁻⁰ | 1.68×10⁻⁰ |
| | | | | Lachnospiraceae incertae sedis | -2.45×10⁻⁰ |

From the 43 genera analyzed, a total of 31 genera from the four major gut microbiota phyla associated with protozoa infection, either overall protozoa positive, or for individual species (P.adj.<0.05) (Table 4, Fig. 5). Most genera were found to be altered by overall protozoa infection (10 with increased abundance, 16 with decreased abundance), followed by infection with Entamoeba histolytica/dispar (6 with increased abundance, 15 with decreased abundance), Entamoeba coli (5 with increased abundance, 10 with decreased abundance), Endolimax nana (6 with increased abundance, 7 with decreased abundance) and Giardia lamblia (3 with increased abundance, 7 with decreased abundance).

The genus Collinsella within the Actinobacteria phylum was associated with both Entamoeba histolytica/dispar and Giardia lamblia infection (β = -1.82×10⁻¹; P.adj. = 2.91×10⁻³ and β = -1.63×10⁻¹; P.adj. = 4.32×10⁻³, respectively). Within the Bacteroidetes phylum, the genus Prevotella was associated with Giardia lamblia infection (β = 3.80×10⁻¹; P.adj. = 6.00×10⁻³).

Nineteen genera were within the Firmicutes phylum to be associated with protozoa infections. Overall protozoa infection was associated with lower abundance of genera within the Lactobacillales order, including Enterococcus (β = -3.31×10⁻²; P.adj. = 3.45×10⁻²), Lactobacillus (β = -9.95×10⁻²; P.adj. = 2.42×10⁻²) and Streptococcus (β = -1.48×10⁻¹; P.adj. = 2.97×10⁻⁴), and these trends were found for specific protozoa species as well. Genera within the Clostridiales order were found associated with both increased and decreased abundance, depending on protozoan species. For example, decreased abundance of Biautia was observed in individuals with Entamoeba coli (β = -1.01×10⁻¹; P.adj. = 3.45×10⁻²) and Entamoeba histolytica/dispar (β = -7.01×10⁻²; P.adj. = 4.90×10⁻²), whereas abundance of the genus Clostridium IV was increased by Entamoeba histolytica/dispar and Endolimax nana, but not by Entamoeba coli or Giardia lamblia. Decreased abundance of the closely related genus Clostridium XVIII was associated with all protozoa parasites except Giardia lamblia.

Of the four tested genera within the Gammaproteobacteria class, three were found to be less abundant in individuals with protozoan infections, namely Escherichia/Shigella, Klebsiella and Haemophilus; while the fourth, Succinivibrio, was increased.

One genus within the Epsilonproteobacteria class, Campylobacter, was the only genus found with increased abundance for all of the aforementioned protozoan infection variables (β = 1.38×10⁻¹; P = 2.66×10⁻⁴ for overall protozoa infection, β = 1.68×10⁻¹; P.adj. = 5.52×10⁻³ for Ent. coli infection, β = 1.49×10⁻¹; P.adj. = 1.58×10⁻³ for Ent. histolytica/dispar infection, β = 1.66×10⁻¹; P.adj. = 2.66×10⁻⁴ for Giardia lamblia infection, β = 1.98×10⁻¹; P.adj. = 1.45×10⁻³ for Endolimax nana infection).

Additionally, the Lachnospiraceae incertae sedis family (at present taxonomically organized within the Clostridia class) was found at lower abundance by infection with any Entamoeba spp. (β = -2.45×10⁻¹; P.adj. = 9.43×10⁻³ for Ent. col and β = -1.64×10⁻¹; P.adj. = 2.35×10⁻² for Ent. histolytica/dispar).

In summary, a large portion of the analyzed genera associated with protozoa infection, while only few taxa were significantly associated with helminth infection. All major phyla normally found in the human gut microbiota were represented across associated genera.

**Storage time associates with small but significant differences in microbiota composition**

Due to challenges associated with collecting fecal samples in Guinea-Bissau, including unstable electricity supply and limited infrastructure to secure adequate storage at freezing temperatures, fecal samples were collected on FOBT paper and stored at ambient temperature for a longer time period. Existing studies of this procedure has reported a general good performance of the FOBT papers [30–34]. We have recently demonstrated that the storage method used...
in this study provide microbiota results that are very similar to conventionally stored samples kept at -80°C [28]. Within the present study, both overall DNA and sequencing quality from FOBT paper samples was satisfactory, and microbiota data resembled that of a microbiota from a conventionally stored sample. However, to limit potential confounding effects of storage time, days of storage was included as a covariate in the analyses.

We explored changes in phylogenetic diversity, beta diversity and compositional alterations of selected bacterial taxa with increasing storage time. We observed a relative increase in Bacteroidetes abundance and a corresponding decrease in Firmicutes with increasing storage time (Fig. 6A). The decrease in Bacteroidetes seemed to be driven by a decreased abundance of Prevotellaceae at family level (Fig. 6B), and by Prevotella at genus level (Fig. 6C). Alpha and beta diversity associated with storage time (Spearman rho ~ 0.097, P = 0.00077, and adonis R² = 0.048, P = 0.007, respectively). Furthermore, correlation analysis of the relative abundance of selected taxa demonstrated a general significant association with storage time, however with limited average change in abundance (strongest coefficient (rho) = 0.23 observed in the Firmicutes phylum, P-values ranging from 4.0⋅10⁻⁹ to 6.80⋅10⁻¹ for the investigated taxa) (Supplementary fig. S1).

Discussion

In the present study, we analyzed fecal microbiota composition from 1,204 children from Bissau, Guinea-Bissau, with an overall high prevalence of intestinal parasitic infections, predominantly caused by protozoans including Entamoeba spp. and Giardia lamblia. We demonstrated that microbial alpha diversity was largely unaffected by helminth infections, and that protozoan infections had moderate effects on alpha diversity. We demonstrated that beta diversity associates with infection status for both pathogenic and non-pathogenic protozoa, and that the abundance of a total of 32 bacterial genera were altered due to parasite infections. We proved that the microbiota from fecal samples stored at room temperature on FOBT papers resembled that from conventionally stored samples, and that FOBT papers were useful in fieldwork with lack of freezing capacity.

We found a total of 31 genera from four different phyla, out of 43 genera analyzed, to significantly associate with intestinal protozoan infection. These include a decrease of the Collinsella genus in individuals infected with Entamoeba histolytica/dispar and Giardia lamblia. To our knowledge, no previous studies have associated this genus with intestinal parasitic infections, however, the Actinobacteria phylum has recently been demonstrated to be increased upon infection with Trichurus trichiura in humans [24]. Collinsella spp. have been demonstrated to regulate levels of circulating insulin in pregnant women [35], and a reduced abundance has been associated with symptom severity in patients with irritable bowel syndrome [36]. A common and debilitating feature of Giardia infection is the post-giardiasis syndrome after complete elimination of the parasite, with a symptomatology very much alike irritable bowel syndrome [37, 38]. One possible explanation for these long-lasting post-infectious symptoms could be an altered microbiota, including decreased abundance of Collinsella spp. However, as Giardia lamblia resides in the small intestine only [39], an effect seen in the fecal microbiota may be immunological derived, rather than by local interactions. In mice, Giardia infection has been reported to increase the abundance of Proteobacteria in the fore- and hindgut [40]. This is contradictory to our observations, where Proteobacteria itself was not associated and two of three genera in the clade were decreased upon Giardia infection.

We found a decreased abundance of the Bacteroides genus due to infection with Entamoeba histolytica/dispar. There are conflicting results regarding gut microbiota alterations due to Entamoeba histolytica infection, as both increased [25] and decreased [41] abundance of Bacteroides has been previously observed. The two studies in question are conducted in Zimbabwe and India, respectively, and the conflicting findings could be due to geographical changes in gut microbiota. Further, there are differences in the applied technique, as one is based on sequencing, and the other on targeted PCR. We further found a decrease in Lactobacillus spp. due to infection with Entamoeba histolytica/dispar, which is consistent with previous findings [41].

We found limited (single taxa and beta-diversity) or no (alpha-diversity) effects on gut microbiota composition due to helminth infections. This is contradictory to previous studies on the subject, in which several bacterial taxa have been associated with infection. Regarding hookworm infection, previous studies have demonstrated an increase in Bacteroidetes and a decrease in both Lachnospiraceae and Firmicutes [24, 42]. The number of hookworm-infected individuals in these studies vary between 8 and 55, compared to a total of 94 in the present study, and the differences could thus be explained by a lack of power in the previous studies, as these hold an increased risk of false positive findings. Furthermore, helminth prevalence is minor in the present study, compared to protozoa, and may partly explain why less pronounced effects due to helminth infections are observed. Further, regional and geographical differences in gut microbiota composition is another plausible explanation for the lack of uniform results. To our knowledge, no other studies have investigated gut microbiota alterations due to infection with Hymenolepis nana.

Historically, all protozoa and helminths were considered parasitic, and assumed to be pathogenic. As reflected by the number of prevalent cases and related morbidity worldwide, this is indeed true for some species. A distinctive feature of many intestinal parasitic infections is that they cause significant morbidity, and less pronounced mortality. For instance, STI infections, which especially affect children, may cause nutritional deficiency, which may lead to anemia and ultimately reduced growth and cognitive development [2, 3, 43]. With regards to some intestinal parasites, infection can be life-threatening and even fatal, as seen in hyperinfection syndrome of Strongyloides infection, bowel obstruction in Ascaris infection, or invasive amebiasis by Entamoeba histolytica infection [44–46]. Although some intestinal parasites may cause pronounced pathology in humans, evaluation of the existing literature indicates that many common eukaryotic species within the human gut, originally identified as pathogenic parasites, are actually commensals or even beneficial, at least in part, and could be regarded as pathobionts, only causing disease in certain contexts [22, 47, 48]. Some even extend this to state that eukaryotic members of the microbiota (termed the eukaryome or parasitome) are crucial in maintaining gut homeostasis and shaping host immunity [49], and that consequently, absence of e.g. helminths may result in a dysfunctional immune system [14], which partially explain the rise in autoimmune diseases seen in the industrialized world [50]. The most prevalent intestinal parasites found in the present study are pathogenic, with the exception of the amebic species Entamoeba coli and Endolimax nana, which are generally accepted as being non-pathogenic. We demonstrate that there are only minor differences between the two cohorts (children seeking medical attention and children found in the background population, respectively) with regard to parasite prevalence, and this difference is due to helminth infections, which are dominant in cohort I. By so, our findings support that the presence of intestinal parasites does not necessarily cause individuals to seek medical attention, even though the microbiota of these individuals is altered.
One major limitation of our study is the somewhat old-fashioned and rough methods for detecting intestinal parasites. In the industrialized world, conventional light microscopy has largely been replaced by molecular diagnostics including qPCR, which has proven to be superior to microscopy with increased sensitivity and specificity [51]. However, in developing countries and field settings, laboratory access is sparse, and microscopy remains a cheap, fast and reproducible method for parasite examination, with acceptable sensitivity and specificity [52]. One could argue that an increased sensitivity by qPCR would result in overdagnosis and detect intestinal parasites at much lower abundance, thereby limit the effects seen in the fecal microbiota. As the parasites are detected by relatively insensitive light microscopy, it is safe to assume that the parasite burden in positive samples is clinically relevant, which increases the value of our findings regarding fecal microbiota alterations, as these are not driven by clinically irrelevant infections.

Another methodological limitation by microscopy diagnosis is the failure to differentiate between pathogenic and potentially fatal Entamoeba histolytica and non-pathogenic Entamoeba dispar, as the two are indistinguishable by microscopy [53, 54]. The high prevalence of Entamoeba histolytica/dispar found in our study may very likely resemble Entamoeba dispar, and the described associations with fecal microbiota may not be due to Entamoeba histolytica. Microbiota alterations due to infection with Entamoeba histolytica has previously been investigated in Cameroon, where an infection-dependent increase in Bacteroidetes was reported [25].

Due to the sequencing approach to investigate the fecal microbiota, we were unable to detect alterations on strain-level. Furthermore, as the two genera Escherichia and Shigella (within the Gammaproteobacteria class) have very similar 16S rRNA gene sequences, we were unable to differentiate between the two. Both genera are related to gastrointestinal pathology, and differentiation between the two by other approaches would possibly yield interesting aspects.

The use of samples stored on FOBT paper at room temperature could be considered both as a strength and a weakness of the present study. Optimal sample storage has been reported to be among the most important issues in microbiota research, and compositional changes might occur over a relatively short time. General recommendations include immediate freezing within 15 minutes [55] to 24 hours [56] from defecation, and storage at -80 °C is regarded as gold standard [57]. However, several studies, including our own benchmark study, have demonstrated that the use of FOBT filter papers is a valid approach, which does not induce major compositional shifts [33]. However, all analysis in the current study were adjusted for storage time, in order to further ensure biologically reliable results. Amir et al. have found that specific bacterial taxa grow at room temperature, which could induce confusing results. Especially genera within the Gammaproteobacteria class are reported to bloom [58]. However, as we previously demonstrated that this class was unaffected by room temperature storage for five weeks on FOBT papers [28], we judged that adjusting for storage time accommodated adequately for any potential confounding effect of FOBT storage (Table 4, Fig. 5). Although we demonstrated some significant alterations due to room temperature storage, these were minor, and do not argue against the use of FOBT cards in fieldwork without electricity. What seems to be important is a uniform sample collection, and that analyses should be adjusted for storage time.

**Conclusion**

In the present study, we demonstrate that microbiota assemblages are significantly associated with intestinal protozoa infections, whereas limited or no effects are seen due to helminth infections. We find that specific taxa associated with different protozoan infections, which can lead to improvement of treatment strategies by probiotic supplementation, and further enhance our understanding of the interplay between host microbiota and intestinal parasitic infections.

**Methods**

**Sample collection and storage**

Stool samples were collected as a part of a prospective no-intervention two-cohort study, investigating the prevalence and potential risk factors for intestinal parasite infections in children from Bissau, Guinea-Bissau, Western Africa. The study area and sample collection procedure has been described in detail previously [27]. In brief, children aged 2–15 years were included between August 2015 and April 2017 at local health centers (cohort I) or at their private address (cohort II). Upon inclusion, participants delivered fresh stool samples in designated sterile containers, which were kept in a refrigerator prior to microscopic analysis for intestinal parasites. Microscopic parasitological analyses were performed following the local routine, and infection load was determined by the number of different species identified. Upon microscopic investigation, the fecal sample was manually homogenized within the container, and approximately 0.5 mL of the sample was applied to the two filter paper windows of a fecal occult blood test (FOBT) filter card (Hemoccult®, Beckman Coulter) with a clean wooden spatula. The sample was air-dried under laminar airflow, protected from sunlight, for 1–6 hours, after which the sample was packed in an individual airtight zip lock bag with desiccant (Whatman® desiccant packs, Sigma-Aldrich). Samples were subsequently stored in the dark at ambient temperature, which is approx. 25 °C on average in Guinea-Bissau [59], prior to airplane shipment to laboratory facilities in Germany for DNA extraction and 16S rRNA sequencing. Sample storage time was calculated from day of inclusion to the day of DNA extraction. All samples were stored between 209 and 993 days at room temperature prior to DNA extraction.

The storage on filter paper was chosen due to lack of freezing capacity and further lack of a possibility to transport samples from Guinea-Bissau to central laboratory facilities at stable and constant freezing temperatures. We have recently demonstrated that this particular storage method is applicable in microbiota research, as the fecal microbiota from samples stored on FOBT filter papers at room temperature for up to five months is comparable to that of a sample frozen and kept at -80°C immediately after collection, with regards to diversity and cumulative abundances [28].

A total of 1,274 fecal samples were collected for microscopic investigation, all from participants with complete questionnaire data. Of these, samples from 1,264 participants were applied to filter paper and underwent DNA extraction as described below. A flow diagram of the study is provided in Fig. 1.

**DNA extraction**
The actual filter from the FOBT cards was cut free from the card using scissors and handled using tweezers. Instruments were cleansed thoroughly between each sample using absolute ethanol (EMSURE®) to avoid cross-contamination between samples. Bacterial DNA extraction from FOBT papers was performed with the QIAamp DNA Stool Mini Kit (QIAGEN) on a QIACube platform (QIAGEN), according to manufacturer's instructions with minor modifications. In brief, the FOBT paper was placed in PowerBead Tubes with Garnet beads (0.70 mm) (QIAGEN) with ASL lysis buffer (QIAGEN). Samples were homogenized by bead beating at 40–50 MHz for 45 seconds on a SpeedMill PLUS instrument (Analytik Jena AG), spun down and the supernatant stabilized with InhibitEX tablets (QIAGEN), containing PCR inhibitor absorption matrix. Subsequent DNA extraction was automated on the QIAcube, following standard programs. Extracted DNA was stored at -80°C prior to PCR amplification. Blank non-template extraction controls were included to investigate potential contamination.

**Bacterial DNA amplification and pooling**

The two hypervariable regions V1 and V2 of the 16S rRNA gene were amplified using the forward 27F primer and reverse 338R primers and dual MID indexing, as described by Koziuch et al. [60]. Bacterial DNA was dually barcoded by unique forward and reverse primers, as described by Caporaso et al. [61], enabling subsequent multiplexing of the PCR product. PCR products were evaluated by gel analysis and normalized using the SequiPrep Normalization plate Kit (Invitrogen), according to manufacturer's instructions. The pooled PCR products were measured fluorometrically using Qubit 4 Fluorometer (Invitrogen), to test DNA concentration.

**16S rRNA gene sequencing and data processing**

Sequencing was performed on the Illumina MiSeq platform, using the MiSeq Reagent Kit v3 according to manufacturer's instructions. MiSeq FastQ files were trimmed using sickle [62] in PE (paired-end) mode with a sliding window of 0.1 readlength. Trimming was performed when average quality within the window was below 20, and reads were all > 100 bp after trimming. Reads were stitched together using VSEARCH [63] with a length between 280 and 350 bp. Further, VSEARCH filtered reads with more than 1 expected error. Further quality filtering was performed using the FastX-Toolkit::fastq_quality_filter [64] to exclude sequences with > 5% nucleotides with a quality score below 30. Files were subsequently converted to FASTA format, and chimeras were removed in VSEARCH, using the gold.fa database. The remaining reads were classified using the UTAX algorithm, where reads classified as chloroplasts or not classified at domain level were removed.

OTU tables were generated in UPARSE [65], implemented in VSEARCH. After removal of replicates and singletons, reads were clustered based on 97% similarity. Chimeras were once again filtered using VSEARCH in de-novo mode. To generate OTU abundance tables, all reads per sample were mapped to OTU tables using VSEARCH. Using the SINTAX classifier [66] at lowest possible level with minimum 80% bootstrap confidence, one representative sequence for each OTU was annotated. OTUs with identical annotations were grouped into taxonomic bins. Samples from individuals with self-reported antibiotic usage 3 months prior to inclusion were not included in the analysis (n = 31).

**Statistical analysis**

Between-group differences in baseline characteristics and parasite prevalence were calculated using Wilcoxon rank-sum test, Fisher's exact test and Kruskal-Wallis rank test in STATA 15.1 (StataCorp, College Station, TX, USA). P-values < 0.05 were considered significant in these analyses.

Statistical analysis of the microbiota data was performed using the R programming environment v3.2 [67]. All adjusted p-values were obtained using the Padjust function in R package stats and the Benjamini-Hochberg method (method = “BH”). First, data was filtered by excluding samples with less than 10,000 reads and samples possibly affected by overgrowth of facultative anaerobic taxa, which were predominantly found in the Proteobacteria and Firmicutes phyla (the latter in the branch of *Streptococcus*). These samples were identified if they fell above the third quartile plus three times inter quartile range (IQR) of phylum abundance (n = 18). Microbial count data on the remaining 1,204 samples was transformed to adjust for deviating sequencing depth by dividing the counts by sample sum and multiplied by one hundred to obtain relative abundances between zero and 100.

As the life cycle, infection route and severity of infection vary considerably between different species of intestinal parasites, we chose to analyze different aspects of the fecal microbiota separately. First, we performed analyses for any parasite, any helminth and any protozoa, and subsequently performed analysis for the most abundant individual pathogenic parasite species (*Ancylostoma duodenale, Hymenolepis nana, Entamoeba coli, Entamoeba histolytica/dispar, Giardia lamblia* and *Endolimax nana*). For each analysis, the infected group was compared with the subset of individuals with no detectable parasitic infection (n = 596).

All analyses were adjusted for storage time at room temperature. Alterations associated with helminth infections were adjusted for co-infection with protozoa, and vice versa. To further control for potential confounding factors, all analyses were adjusted for age, usage of vitamin A, toilet source and tropical season of sample collection. As described above, participants were enrolled in the study on a two-cohort basis. While there was no overall difference in infection load between the two cohorts, prevalence of some species differed significantly, and we thus adjusted for cohort status in the joint analysis and performed a separate supportive analysis within each group of the two cohorts. Results from the joined analysis is provided in the main text while Supplemental table S1 show results from all three analyses.

**Association with alpha- and beta-diversity**

Associations between alpha-diversity and each of the nine infection states (overall parasite positive; overall helminth positive; overall protozoa positive; *Ancylostoma duodenale* positive; *Hymenolepis nana* positive, *Entamoeba coli* positive, *Entamoeba histolytica/dispar* positive, *Giardia lamblia* positive, *Endolimax nana* positive) was evaluated using a robust regression (lmRob function in R package robust [68]) and the covariates given above. The alpha diversity measures considered were Shannon entropy (diversity function with index = “shannon” in R package vegan [69]), the measure of species richness ACE (estimateR function using result row four in R package vegan), and phylodiversity as a measure of total unique phylogenetic branch length (calculated using mothur's phylodiversity function [70] with the phylogenetic tree built using FastTree with -nt and -gtr and the 16S OTU table as input). Evaluation of the association between the parasite infection status and microbial community structure was performed using adonis function in R package vegan [69] with Bray-
Curtis dissimilarity and 9999 permutations (remaining settings as default). Furthermore, association with infection load was evaluated with infection load ranging from 0 to 4 by the number of identified species within each sample. Covariates were as listed above.

**Analysis of single taxa**

The relative abundance of single bacteria was evaluated from the taxonomic level of phylum to genus, and was filtered to keep most abundant taxa as follows; filtered to require a mean abundance across all samples of at least 0.05, and an abundance of 0.05 in at least one sample. Further, taxa with ≥40% zeroes across samples were removed. After filtering, 43 genera, 23 families, 15 orders, 11 classes and 5 phyla remained. Associations between the selected taxa and parasite infection status were evaluated using a linear regression with square root transformed taxa abundance and the covariates as listed above.

**Additional R-packages used**

- reshape2 v1.4 [71], grid v3.5, gridExtra v2.3 [72], gridBase v0.4 [73], plyr v1.8 [74], ggplot2 v3.1 [75], extrafont v0.17 [76], metafor v2.0 [77], plotly v4.9 [78], data.table v1.12 [79], ggrepel v0.8 [80], MASS v7.3 [81], robust v0.4 [68].

**Ethical statement**

The participant enrollment and microscopic investigation of fecal samples was approved by the Ethical Committee of Guinea-Bissau (Comité Nacional de Ética na Saúde) (ref. no. 0029/CNES/INASA/2015). Participants or parents/guardians to participants gave oral and written consent to participate. Subsequent microbiota analysis was approved by the Ethical Committee of Guinea-Bissau (ref. no. 062/CNES/INASA/2017) and the Regional ethics committee of Region of Southern Denmark (ref. no. S-20160138). The study was conducted in adherence to the Declaration of Helsinki.

**List Of Abbreviations**

- 16S rRNA  16S ribosomal ribonucleic acid
- DALY  Disability-adjusted life years
- FOBT  Fecal occult blood test
- OTU  Operational taxonomic unit
- qPCR  Quantitative polymerase chain reaction
- STH  Soil-transmitted helminth

**Declarations**

**Consent for publication**

Not applicable.

**Data availability**

An anonymized dataset, including sequence data for the 16S rRNA gene, supporting the conclusions of this article, is available at the NCBI SRA repository (Accession no. PRJNA642721).

**Competing interests**

All authors declare that they have no competing interests.

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**Author contributions**

Study design: PK UH AF SH LT

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Funding acquisition: SH PK UH
Laboratory/technical assistance: SH CB
Bioinformatics: LT MR
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Dataset repository:

A complete, anonymized dataset is available at the NCBI SRA repository (Accession no. PRJNA642721).

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Figures
Flow-diagram for inclusion and fecal sampling. The dataset includes microscopic investigation for intestinal parasites from 1,274 children aged 2-15 years from urban Bissau, Guinea-Bissau. Details on the cohort including microscopy method is described elsewhere. A total of 1,253 samples underwent 16S rRNA gene sequencing, and 49 were excluded subsequently, yielding a final study size of 1,204 samples.

Figure 1
Figure 2

Stacked barplot of taxonomic profile of the gut microbiota across participants. Illustration of the relative abundance (y-axis) of phyla across the 1,204 samples (x-axis) in the study, irrespective of infection status. The 10 most abundant phyla are colored and listed in the legend. For all samples, the two dominant phyla are Bacteroidetes and Firmicutes, followed by Proteobacteria and Actinobacteria, in line with that of a normal human gut microbiota.
Phylodiversity is altered by infection with Entamoeba spp. Illustration of differences in phylodiversity levels between samples with different infection status. Phylodiversity, as a measure of alpha diversity, remains unaltered by infection with helminth species, Giardia lamblia and Endolimax nana. Infection with Entamoeba spp. significantly increases phylodiversity. Each boxplot shows the non-infected (with any tested parasite (orange) versus samples infected with the parasite of interest as listed in plot title (grey). The adjusted significance level from the robust regression test is given in parentheses.
Figure 4

Gut microbiota is shaped by protozoan infections and infection load. (A) Increasing infection load, determined by the number of different parasitic species identified by microscopy in each sample, induces increasing shifts in the microbiota, suggesting that multispecies infections have more pronounced effects on microbiota than single species infections. (B) Parasite infection (any positive finding, regardless of species) imposes a visual shift in microbiota structure. (C) Overall protozoan infection imposes a shift in gut microbiota composition very similar to the shift observed for overall parasite infection, indicating that protozoa infection, and not helminth infection, has the strongest effect on microbiota composition, an observation supported by results from the adonis-based analysis. (D) Helminth infection has no apparent effect on microbiota structure. The figure is composed of four ordination plots built using genera relative abundance data and principal coordinates analysis (capscale function in R package vegan with Bray-Curtis dissimilarity and automatic data transformation (metaMDS = T)). Plots are made with R package ggplot2 and ellipses drawn using function stat_ellipse with default parameters. Each dot shows a sample, all of which are colored by infection status. (A) All 1,204 samples colored by infection load from no infection (0) to infected with four different parasite species (4). (B-D) Plots showing relationship between the microbiota community of samples either (B) non-infected (red) or infected with a parasite (blue), (C) non-infected (red) or infected with a protozoa (blue) and (D) non-infected (red) or infected with a helminth (blue)
Figure 5

Specific taxa are altered by different parasitic species. A total of 32 genera are significantly associated with intestinal parasitic infection, either overall infections or specific parasite species. Most associations are seen with protozoan infections, whereas helminth infections have limited effects on taxa alterations. The figure illustrates significant associations between genera and different parasite infections (linear regression, P.adj<0.05). Summary statistics for the shown associations are given in Table 4. The color illustrates coefficient (blue for low values, green for high values), whereas size illustrates the significance (calculated by -log(p-value)) (the larger bubble, the higher significance). The plot is made using ggplot2 package in R.
Figure 6

Compositional alterations by increasing storage time. The illustration depicts mean cumulative abundance of taxa on (A) phylum, (B) family and (C) genera level for the seven different time periods for room temperature storage (each spanning 100 days). (A) A relative decrease in Bacteroidetes and a corresponding increase in Firmicutes is observed. The decrease in Bacteroidetes appear to be driven by Prevotellaceae at the family level (B), and by Prevotella at the genus level (C).

Supplementary Files
This is a list of supplementary files associated with this preprint. Click to download.

- TableS1.docx
- Fig.S1plotstorage.time.pdf