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RESEARCH PAPER

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Restitution of gut microbiota in Ugandan children administered with probiotics (*Lactobacillus rhamnosus* GG and *Bifidobacterium animalis* subsp. *lactis* BB-12) during treatment for severe acute malnutrition

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ABSTRACT

Severe acute malnutrition (SAM) is a major challenge in low-income countries and gut microbiota (GM) dysbiosis may play a role in its etiology. Here, we determined the GM evolution during rehabilitation from SAM and the impact of probiotics (*Lactobacillus rhamnosus* GG and *Bifidobacterium animalis* subsp. *lactis* BB-12) supplementation. The GM (16S rRNA gene amplicon sequencing) of children admitted to hospital with SAM showed distinct composition over admission (e.g. *Klebsiella* spp., and Enterobacteriaceae spp.), discharge (e.g. Clostridiaceae spp., *Veilonella dispar*) and follow-up (e.g. *Lactobacillus ruminis*, *Blautia* spp., *Faecalibacterium prausnitzii*), reaching similar β - and α -diversity as healthy individuals. Children with diarrhea had reduced distribution of Bacteroidaceae, Lachnospiraceae, increased Enterobacteriaceae and Moraxellaceae, and lower α -diversity. Children suffering from edematous SAM had diminished proportion of Prevotellaceae, Lachnospiraceae, Ruminococcaceae and a higher α -diversity when compared to non-edematous SAM. Supplementation of probiotics did not influence β -diversity upon discharge or follow-up, but it increased ($p < .05$) the number of observed species [SE: > 4.5]. Children where the probiotic species were detected had lower cumulative incidence ($p < .001$) of diarrhea during the follow-up period compared to children receiving placebo and children receiving probiotics, but where the probiotics were not detected. The GM of children with non-edematous and edematous SAM differ in composition, which might have implications for future GM targeted treatments. Probiotics treatment reduced the cumulative incidence of diarrhea during the outpatient phase, with the strongest effect in children where the administered probiotics could be detected in the GM.

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Introduction

Malnutrition remains a global challenge with 45% of childhood deaths being attributed to undernutrition.¹ In addition, undernutrition in early life is associated with long-term sequelae, including stunting, less schooling, and reduced economic productivity later in life.² Recent research indicates that gut microbiota (GM) dysbiosis may be involved in development or maintenance of acute malnutrition.^{3–6}

Malnutrition seems to be associated with reduced GM diversity and maturity as determined by

metagenomics and culturomics approaches.^{3–5,7–9}

In Bangladesh, a birth cohort of children from urban slum was followed until 2 years of age with frequent analyses of GM composition.⁴ Based on age-discriminatory bacterial taxa microbiota maturity scores were developed. Children with severe acute malnutrition (SAM) showed significant GM immaturity compared to well-nourished peers. Nutritional interventions, including Ready-to-Use Therapeutic Food (RUTF), only partially and temporarily improved the GM maturity. Using the same GM maturity models on samples from other

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Bangladeshi children, reduced maturity was found in stool samples during and 1 month after diarrhea episodes.⁴ Findings of reduced GM maturity or diversity in malnourished children, and temporary improvement of GM during RUTF treatment has also been observed in Malawian children.³ In Uganda, a cross-sectional study found differences in the GM composition of children with non-edematous and edematous SAM and reported lower numbers of observed species in the GM of children with non-edematous compared to edematous SAM.¹⁰

It has been suggested that disruption of normal development of the GM may be causally related to development of malnutrition.^{3,5,7-9} This was first shown in the GM from a Malawian cohort of twin pairs who became discordant for kwashiorkor (SAM with edema).³ When the GM of twin pairs discordant for kwashiorkor was transferred to germ-free mice, only mice receiving GM from a child with kwashiorkor became malnourished when fed a diet similar to the diet of Malawian children. Likewise, children with SAM in Niger and Senegal were reported to have GM composition depleted in oxygen-sensitive bacteria and being enriched in putatively pathogenic Proteobacteria, Fusobacteria, and *Streptococcus gallolyticus*.^{8,9} Moreover, animal studies have confirmed that transfer of GM originating from undernourished children to germ-free mice impair growth.⁷

If early life GM dysbiosis can contribute to malnutrition, microbial interventions may be able to support repairing or normalization of the GM. Several bacterial species (e.g. species of *Lactobacillus*, *Bifidobacterium adolescentis*, and *Bacteroides salyersiae*) have been previously proposed as probiotic candidates as an alternative to fecal transplantation to address children suffering from SAM.⁹ Meta-analyses have found that probiotics reduce the duration of acute diarrhea by 1 day and reduce the risk of acute diarrhea lasting 4 days or more.¹¹ However, most studies were performed in high-income countries and in well-nourished children and the knowledge of the impact of probiotic treatment in children with SAM is scarce.¹² The PRONUT study investigated the effect of a synbiotic mixture, a combination of four lactic acid bacteria in a total dose of 10^{11} colony-forming

units in total per day and four fermentable fiber sources. No effect on the primary outcome (nutritional cure) was observed, but a near-significant effect (relative risk = 0.65, $p = .06$) on overall mortality was found.¹² The ProbiSAM study investigated the effect of administering a combination of *Lactobacillus rhamnosus* GG (LGG) and *Bifidobacterium animalis* subsp. *lactis* BB-12 (BB-12) to children hospitalized with SAM.¹³ It was found that administration of the probiotics did not influence days with diarrhea during in-patient treatment, but a significant 26% reduction in days with diarrhea during the outpatient treatment (8–12 weeks after hospitalization) was observed in the probiotic group.

In the present study, we investigate GM development during rehabilitation from SAM, whether probiotics supplementation has any effects on the GM during inpatient and outpatient treatment of SAM, and whether GM composition and development is linked to the observed reduction in days with diarrhea when administered probiotics during the randomized controlled ProbiSAM trial.

Results

Cohort overview and sequencing

The study enrolled 400 children with a mean (\pm SD) age of 17.0 (\pm 8.5) months. Males constituted 58% of the population, 14% were HIV seropositive, 66% suffered from edematous malnutrition and 61% had diarrhea at admission. Fecal samples were collected at hospital admission, at discharge and 8 weeks post-discharge (follow-up). The proportion of individuals from which a fecal sample was obtained at admission to hospital, discharge, and after 8 weeks post-discharge varied between 60% and 80% at the different time points (Figure 1). Control children included 22 apparently healthy children aged 6–59 months living in communities similar to the children admitted with SAM (Figure 1).

Sequencing of DNA extracted from fecal samples generated 34.5 million reads derived from the 16S rRNA gene V3-V4-region with an average of 47,996 (max: 532,852, min: 10,289) sequences per subject (Figure 1b). The analysis of amplicon-sequencing

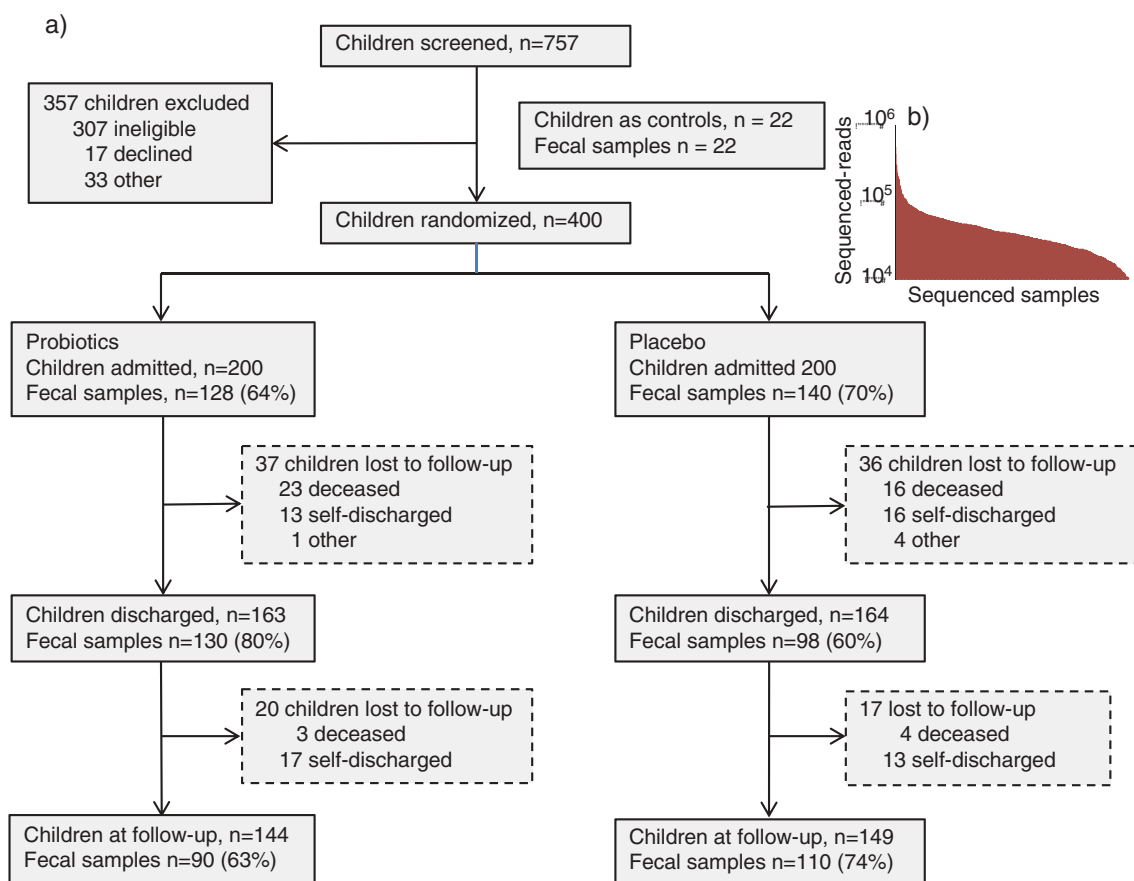


Figure 1. Trial profile.

(a) Dotted boxes: patients lost to follow-up (b) Distribution of high-quality amplicon reads for the total number of samples subjected to high-throughput sequencing.

data generated 44,808 OTU phylotypes (representative sequences clustered at 97% sequence similarity) that were summarized over 365 bacterial species.

GM variation during treatment of SAM

Constrained and un-constrained analyses of Bray-Curtis dissimilarity metrics (Figure 2a-b) displayed GM compositional signatures associated with changes from admission to discharge and follow-up and explaining up to 13% (adonis $p < .001$) of the total variance. At follow-up, the GM of children treated for SAM was indistinguishable from that of healthy individuals from the same-community setting (Figure 2a and b). Based on Constrained Analysis of Principal coordinates (CAP), the most discriminatory GM members (Figure 2b) at admission were associated with Enterobacteriaceae

members (*Klebsiella* and Enterobacteriaceae other). At discharge, the relative abundance of Clostridiaceae members (*Clostridium* uncl., *Veilonella dispar* (Clostridium cluster IX¹⁴) and other Clostridiaceae) had increased and at follow-up, the GM composition was enriched with Lactobacillaceae (*Lactobacillus ruminis*), Campylobacteraceae (*Campylobacter* uncl.), and several members of the *Clostridium* cluster IV and XIVa¹⁵ (such as *Blautia*, Lachnospiraceae, Ruminococcaceae and *Faecalibacterium prausnitzii*).

In relation to mean α -diversity (Figure 2c), the lowest number of observed species was determined at admission (41.4 ± 17.5), followed by an increase as treatment progressed. Upon discharge and follow-up, the mean α -diversity was 42.7 ± 12.7 and 49.3 ± 12.2 ($p < .01$ relative to both admission and discharge), respectively. As also observed with

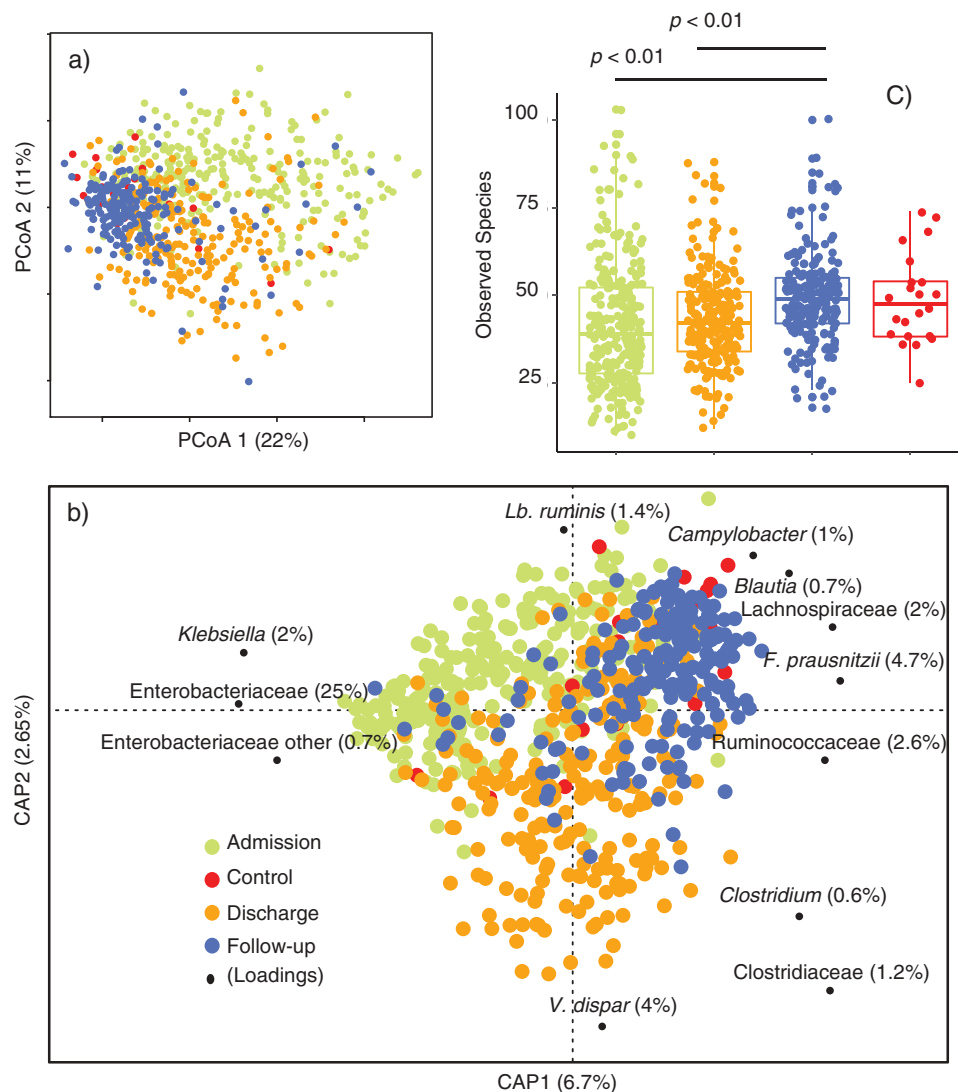


Figure 2. Changes in β - and α -diversity over different stages of GM restitution.

Principal Coordinates Analysis (PCoA) plot based on Bray–Curtis distance of 16S rRNA gene (V3–V4 region) amplicons (phylotypes summarized to species level) determined in the stool samples at different stages of the cohort and healthy subjects (a). Constrained Analysis of Principal Coordinates (CAP) bi-plot of GM composition based on Bray–Curtis distance displaying the most influential species (including their average abundance) discriminating stages (admission, discharge, and follow-up), and healthy subjects during GM restitution (b). α -diversity boxplot depicting the number of observed species (phylotypes summarized to species level) through stages (admission, discharge, and follow-up) and healthy subjects. Changes in α -diversity were analyzed through non-parametric Monte Carlo test (permutations 999) (c).

respect to β -diversity (Figure 2a and b), no significant differences between healthy subjects and at follow-up for children treated for SAM were observed regarding the number of observed species (42.7 ± 12.7) (Figure 2c).

Diarrhea and edema on admission

At admission, significant differences in the number of observed species were observed between

children with and without diarrhea (39 ± 16 vs. 46 ± 19 , $p = .013$), with diarrhea being associated with a lower number of observed species. Similarly, non-edematous SAM was associated with a lower number of observed species relative to edematous SAM (36 ± 15 vs. 45 ± 18 , $p < .001$, those with any grade of edema) (Figure 3a-b). Likewise, Bray–Curtis dissimilarity analysis demonstrated significant compositional differences between children admitted with vs. without

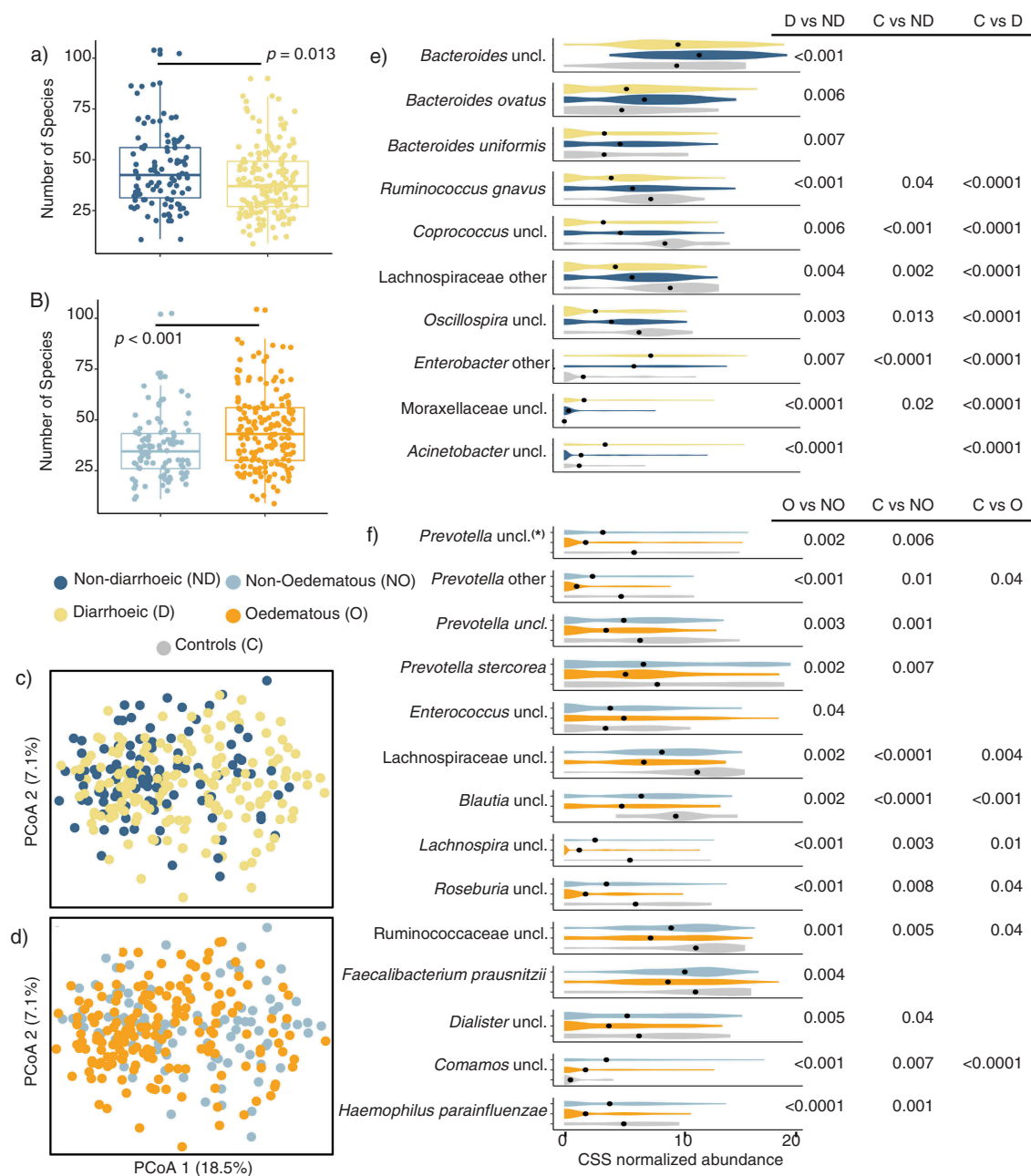


Figure 3. GM composition associates with incidence of diarrhea and edema at admission.

Principal Coordinates Analysis (PCoA) plots based on Bray–Curtis distance on admission discriminating between non-diarrheic and diarrheic children (a), as well as non-edematous from edematous children (b). Relative abundance of the discriminatory bacteria with respect to non-diarrheic/diarrheic (c), non-edematous/edematous (d) children and healthy subjects. p -values correspond to student t -test with False Discovery Rate (FDR) correction on top discriminant bacteria selected through CAP analysis (with minimum absolute index of 0.15 over the canonical axis) among non-diarrheic/diarrheic (c) and non-edematous/edematous (d) children. α -diversity boxplots displaying the number of observed bacterial species with respect to non-diarrheic/diarrheic (c) and non-edematous/edematous (d) children. Differences in α -diversity were analyzed through non-parametric Monte Carlo test (permutations 999).

diarrhea (adonis, $p = .001$, $R^2 = 0.018$), and edematous vs. non-edematous SAM (adonis, $p = .001$, $R^2 = 0.019$) (Figure 3c-d). No significant interaction effects of diarrhea and edema on α - (two-

way ANOVA $p = .98$) nor β -diversity (CAP $p = .24$) were observed indicating no mutual dependence between the two conditions with respect to GM composition.

The GM of children with diarrhea at admission was characterized by lower relative abundance of Bacteroidaceae (*Bacteroides* spp., *B. ovatus* and *B. uniformis*), Lachnospiraceae (*R. gnavus*, *Coprococcus* and *Oscillospira* uncl.), higher Moraxellaceae (Moraxellaceae uncl. and *Acinetobacter* uncl.), and Enterobacteriaceae (*Enterobacter*) relative to children admitted without diarrhea and/or the healthy children (Figure 3e). At admission the GM of children admitted with non-edematous SAM had lower abundance of Prevotellaceae members (*Prevotella* spp. including *Prevotella stercorea*), Lachnospiraceae (*Blautia*, *Lachnospira*, and *Roseburia* spp.), Ruminococcaceae, Clostridiaceae (*F. prausnitzii*), Veillonellaceae (*Dialister* spp.), Comamonadaceae (*Comamos* uncl.), as well as Pasteurellaceae (*H. parainfluenzae*) (Figure 3d) compared to the children admitted with edematous SAM and healthy subjects. Only the abundance of *Enterococcus* uncl. (figure 3f) was significantly higher in children with non-edematous SM compared to edematous SAM.

Probiotics establishment increases α -diversity and reduces days of diarrhea

Probiotic administration did not influence (adonis, $p > .05$) the β -diversity profiles at discharge (Figure 4a) nor follow-up (Figure 4b), and on average the proportion of reads mapping the probiotic strains was 0.33% at discharge and 0.02% at follow-up (as determined by amplicons with >97% similarity to the relevant 16S rRNA gene fragment of *Lb. rhamnosus* and *B. animalis*). Furthermore, in the probiotic group recovery of the two probiotic strains from fecal samples was not evenly distributed among the children (Figure 4c). Children in which at least one probiotic strain was detected were defined as individuals with high level of probiotic response (responders), whereas children where the probiotic strains could not be detected were defined as individuals with low level of probiotic response (non-responders) (Figure 4c and d). Interestingly, at discharge, the average number of observed species in the children defined as responders was 4.8 ($p \leq 0.05$) higher than in the children that received placebo. Likewise, at follow-up, the number of species in responders was 6.4 ($p \leq 0.01$) and 4.5 ($p \leq 0.03$) higher than in the

children defined as non-responders or receiving placebo, respectively (Figure 4e and f), after correcting for the confounding effect of age. There were no significant associations between age, sex, presence of diarrhea or edema at admission, GM composition at admission, HIV status, weight-for-length or duration of hospitalization and being a responder/non-responder.

Throughout the trial, diarrhea incidence (minimum of 1 day) during outpatient treatment was assessed over a median (IQR) period of 56 (56:58) days. Cumulative incidence function (CIF) analysis on outpatient data found that the probability of having more days with diarrhea was significantly higher ($p < .001$) among placebo subjects as compared to the probiotic group (Figure 4g). Interestingly, differences between responders and non-responders ($p < .001$) were also found, where the probability of having more days with diarrhea was significantly reduced (~two fold) in the responders as compared to non-responders (Figure 4g).

Discussion

GM changes during rehabilitation from SAM

Here we show that children with SAM have significantly reduced number of observed species and major compositional differences (β -diversity) compared to healthy subjects. The number of observed species increased during the course of treatment with the lowest number found at admission increasing until it no longer differed from the healthy individuals at follow-up. In line with this, malnutrition in children has previously been associated with reduced diversity and maturity of the gut microbiota.^{3-5,7}

Distinct GM compositional signatures were observed during the treatment of the children admitted with SAM (i.e. admission, discharge and follow-up). Between admission and discharge, a beneficial shift with less Enterobacteriaceae and increasing *Veillonella* and Clostridiales abundance were seen. A development that continued between discharge and follow-up, where Clostridiales members (*Faecalibacterium*, *Blautia*, and other unclassified members of Ruminococcaceae and Lachnospiraceae) became more abundant, similar to the GM of the healthy subjects. This is also in

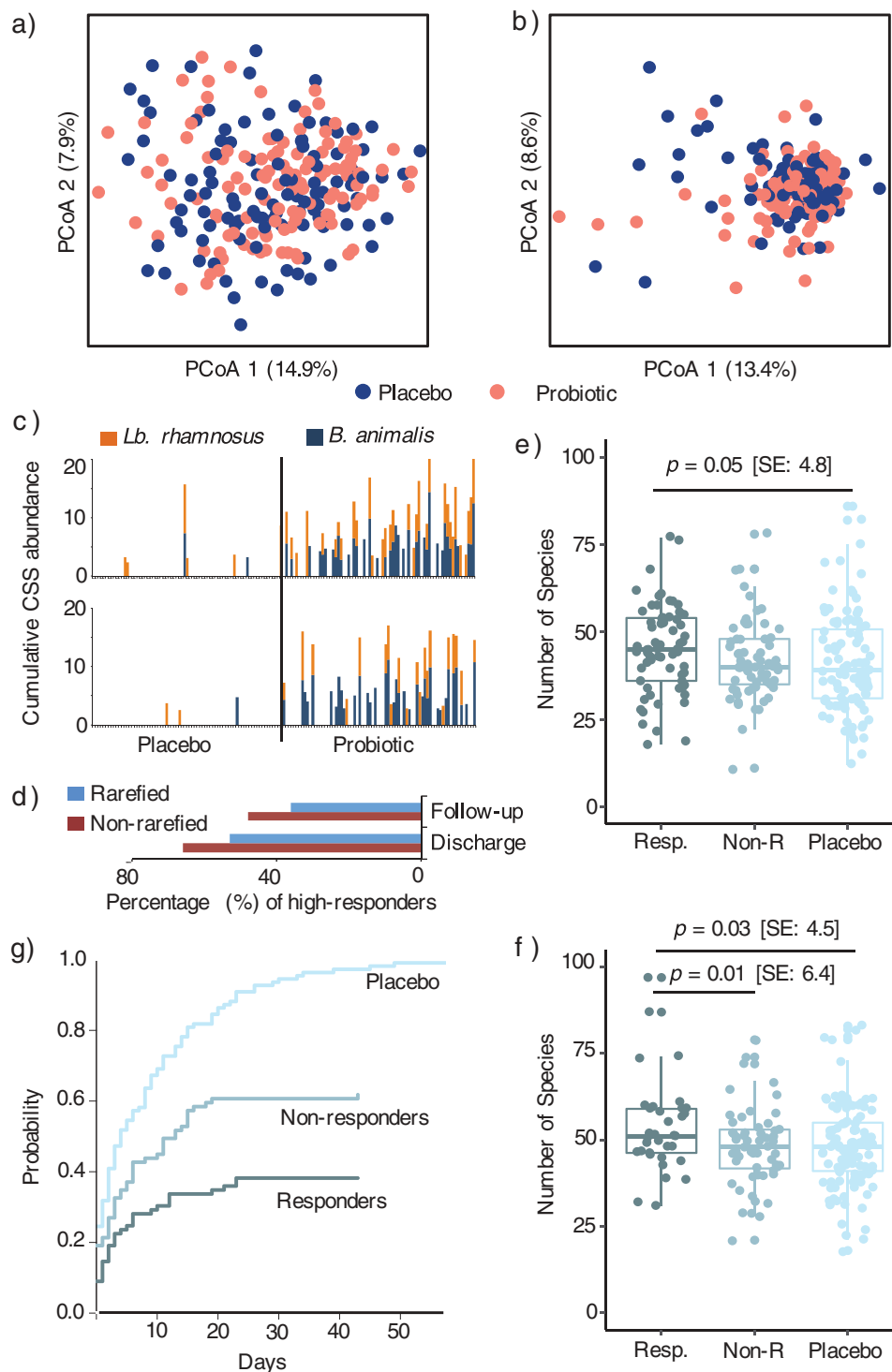


Figure 4. Probiotic administration and establishment in relation to α -diversity and cumulative incidence of diarrhea.

Principal Coordinates Analysis (PCoA) plots based on Bray–Curtis distance on discharge (a) and follow-up (b) describing GM composition in children allocated within placebo and probiotic groups. Normalized abundance of *Lb. rhamnosus*/*B. animalis* and their distribution across children subjected to probiotic administration at discharge (upper panel) and follow-up (lower-panel). For the upper- and lower-panels, every position along the panels represent the same subject evaluated at discharge and follow-up (c). Changes in the distribution of responders to non-responders as a function of rarefied (10,000 reads per sample) and non-rarefied samples (d). α -diversity boxplots displaying the distribution of observed bacterial species within probiotic-responders, probiotic-non-responders and placebo at discharge (e) and follow-up (f). Differences in α -diversity were analyzed through mixed linear models corrected for the age effect. Plots describe the level of significance and the size effect respectively associated. Cumulative incidence function (CIF) of diarrhea determined for placebo, probiotic responders and probiotic non-responders during the follow-up stage. Differences in cumulative incidence rates were assessed with the Gray's test (g).

agreement with previous observations where several obligate anaerobes related to Clostridiales, such as *C. butyricum*, *R. bromii*, and *R. intestinalis*, were reported to be enriched in healthy children as compared to those suffering from kwashiorkor in Western Africa.⁹ Similarly, *B. breve* has also found to be enriched in kwashiorkor cases,⁹ but in the present study, no significant ($p = .36$) differences were found among the healthy controls and children with SAM.

The observed normalization of the GM is likely due to improved nutrition³ in response to the therapeutic foods given during in- and outpatient treatment that contain high-quality protein and high amounts of micronutrients to replenish micronutrient deficiencies and support catch-up growth of the children. The micronutrients may also support development of a GM with a composition beneficial for the child.¹⁶ A study from Bangladesh in 64 children with SAM⁴ found their GM maturity index to be reduced compared to healthy subjects. Treatment with therapeutic foods improved the GM maturity. However, it did not reach the level of healthy children and eventually regressed after cessation of therapeutic food treatment.⁴ Similarly, in a detailed study of 13 Malawian twin pairs discordant for edematous malnutrition (Kwashiorkor) it was found that treatment with ready-to-use therapeutic food (RUTF) resulted in a transient maturation of the GM.³ These changes in GM in children with SAM treated with RUTF could provide a protective barrier to readmission from SAM in the future, but it remains to be investigated how stable the changes are.

Diarrhea, edema, and GM

Diarrhea is a well-known morbidity factor and indicator of poor prognosis in children with SAM.^{17–19} We found the presence of diarrhea at hospital admission to be associated with reduced number of observed species, as well as reduced relative abundance of species belonging to Bacteroidaceae and Lachnospiraceae and higher relative abundance of Moraxellaceae and Enterobacteriaceae. Others have reported transiently reduced GM maturity in children with diarrhea in Bangladesh⁴ and an association with moderate to severe diarrhea and lower GM

diversity in children below 5 years in four low-income countries.²⁰

Distinct features of the GM were associated with non-edematous compared to edematous SAM at admission, with non-edematous children having significantly fewer observed species. A similar finding has been made in an earlier cohort of children with SAM from the same hospital.¹⁰ Several factors could explain the lowered number of observed species seen in relation to non-edematous SAM. It is hypothesized that longer starvation of the GM in non-edematous SAM may lead to a lower GM diversity. The non-edematous children also tend to have more infections when they are admitted to hospital.^{21,22} They may, therefore, have been treated with more antibiotics from health clinics before hospital admission leading to a lower number of observed species. The abundance of taxa normally associated with Sub-Saharan African children not suffering from disease such as Prevotellaceae members²³ was also significantly higher in children admitted with edematous SAM compared to non-edematous SAM and closer to the abundance observed in the healthy subjects.

Responders vs non-responders

The probiotic strains LGG and BB-12 have in previous studies been found to colonize the gut transiently after oral administration. Recovery of both strains depends on the dose administered and vary between individuals.^{24,25} We detected each strain in approximately half of the fecal samples from children randomized to receiving the probiotics. The observed recovery is in line with recovery obtained with BB-12²⁴ and slightly lower recovery than previously observed for LGG²⁵ in previous studies.

There were no significant differences in β -diversity among individuals considered as probiotic high- and non-responders. However, the mean number of observed species was 4.8 higher among responders compared to placebo at discharge. At follow-up, the number of observed species in responders was 6.4 and 4.5 higher compared to non-responders and placebo, respectively. This indicates that in the case of SAM, LGG, and BB-12 may influence GM α -

diversity. Responders had lower cumulative incidence of diarrhea during outpatient treatment compared to probiotic non-responders, which could indicate a better gut colonization and stronger interaction with the immune system.²⁶ Yet, we were unable to identify malnutrition-related factors associated with being a responder/non-responder. As described above, non-responders are also reported in healthy volunteers for both LGG and BB-12 and being a non-responder, may therefore, be very common and not related to malnutrition.^{24,25}

Strengths and limitations

A particular strength of the present study is the large sample size compared to other studies investigating the effect of re-feeding and probiotic administration to children suffering from SAM. However, in this context, it is a limitation, that the number of included children without disease (healthy subjects) is relatively small. At the different sampling time points we obtained samples from 60-80% of the enrolled children. Possibly the mothers/caregivers with the most ill children were those with lowest compliance which might skew the sample set toward the most ill children being under-represented. However, it should be noted that most missing samples are due to failure to pass stool/collect stool within the specified timeframe.

Conclusion

GM diversity and composition change over the course of rehabilitation from SAM and approach the GM of apparently healthy subjects as treatment progresses. Further, our study supports that non-edematous and edematous SAM are associated with GM compositional differences, which might have implications for future GM targeted treatments. Finally, using probiotics alongside the standard treatment protocol for SAM reduces the incidence of days with diarrhea after discharge. This may be partly mediated by the observed increase in the number of observed species seen in the children, where the administered probiotics

could be detected at discharge and follow-up (“Responders”). Although the effect of probiotics on the GM was modest and previous studies have shown that nutritional interventions may only lead to transient improvements of the GM, the results indicate a potential direction for future research and management of SAM.

Materials and methods

Ethics statement

Before study initiation, ethical approval was obtained from the School of Medicine Research and Ethics Committee at Makerere University, Kampala, and a consultative approval was provided by The National Committee of Health Research Ethics in Denmark. Written informed consent was obtained from all caregivers on behalf of their children. In addition, clearance to conduct the study was given by the Uganda National Council of Science and Technology and the Ugandan National Drug Authority. Further details have been described elsewhere [11]. The study was registered at www.isrctn.com, ISRCTN16454889.

Study design, patients, and study procedures

The study is a prospective study nested in a randomized, double-blind, placebo-controlled trial assessing the effect of probiotics on diarrhea among children with SAM. The trial was conducted at Mwanamugimu Nutrition Unit, Mulago National Referral Hospital, Kampala, Uganda between March 2014 and September 2015. Children admitted to Mulago Hospital with SAM generally have multiple medical complications and the case fatality rates are high (approx. 20%). Children between 6 and 59 months with SAM were eligible for the probiotic trial. SAM was defined as mid-upper-arm circumference (MUAC) < 11.5 cm, weight-for-height z-score (WHZ) < -3 SD or bipedal pitting edema. Caregivers also had to provide written informed consent and be willing to return for follow-up. Children were excluded if they were in shock, had severe respiratory distress, an admission weight below 4.0 kg, obvious congenital anomalies or if they had been admitted with SAM the previous 6

months. Controls included apparently healthy 22 children aged 6–59 months with WHZ > -1 and living in communities similar to the children admitted with SAM.

Standard treatment was provided to all children according to the Integrated Management of Acute Malnutrition guidelines for Uganda²⁷ with adaptation from the WHO guidelines.²⁸ In addition, one daily dose of a combination of two probiotic strains BB-12 and LGG or placebo was given. The total probiotic dose was 10 billion colony-forming units [CFU] per day with half of each strain (Chr. Hansen A/S, Hørsholm, Denmark). The probiotic/placebo supplement was administered from hospital admission to discharge and throughout an outpatient treatment period of 8–12 weeks, depending on the nutritional recovery rate of each child. More detailed information about the study is reported elsewhere.¹³

Sample collection, processing, and DNA extraction

Fecal samples were collected at admission, discharge, and after 8 weeks of outpatient treatment. Admission samples were collected from the time of admission to day 3 of hospitalization; discharge and 8-week follow-up samples were collected on the day of discharge/follow-up or the day before. During hospitalization, stool was collected in plastic bags and stored for maximum 1 h before the contents of the stool bags were emptied into 2-ml DNase free cryotubes. The cryotubes were immediately frozen in liquid nitrogen. Outpatient samples were collected in the children's home up to one day before a follow-up visit. Caregivers transferred stool to a lidded 10-ml plastic vial containing 5-ml RNA-later (Qiagen GmbH, Hilden, Germany). Caregivers were asked to fill stool up to a mark ensuring an approximate 5:1 ratio of RNA-later and stool. When samples were received, the 10-ml plastic vial was centrifuged at 1,300–2,200 × g for 10 min. RNA-later was discarded and the stool was re-suspended in 2-ml TE buffer. One ml was transferred to a 2-ml DNase free cryotube and immediately frozen in liquid nitrogen and stored at -80°C. Samples were shipped on dry ice to Denmark for further processing and analysis.

Fecal samples were centrifuged at 13,000 × g for 10 min at room temperature and ~200 mg of the fecal pellet was used for DNA extraction using the PowerSoil® DNA Isolation Kit (MOBIO Laboratories, Carlsbad, CA, USA), following the instructions of the manufacturer, but with minor modifications. Briefly, prior DNA extraction, samples were placed into the PowerBead tubes and heat treated at 65°C for 10 min and then at 95°C for 10 min. Subsequently, solution C1 was added and bead-beating performed in FastPrep (MP Biomedicals, Santa Ana, CA, USA) using 3 cycles of 15 s each, at a speed of 6.5 m.⁻¹ The remaining DNA extraction procedure followed the manufacturer's instructions.

High-throughput 16S rRNA gene amplicon sequencing

GM composition was determined by high-throughput 16S rRNA gene amplicon sequencing. The primers designed with adapters Nextera Index Kit® (Illumina, CA, USA) targeted the V3-V4 region (~466 bp) and the amplicon library preparation, purification and sequencing were performed as previously described.²⁹ Briefly, the amplification profile (1st PCR) followed: Denaturation at 95°C for 2 min; 33 cycles of 95°C for 15 s, 55°C for 15s and 68°C for 40 s; followed by final elongation at 68°C for 5 min, while barcoding (2nd PCR) was performed at 98°C for 1 min; 12 cycles of 98°C for 10 s, 55°C for 20 s and 72°C for 20 s; elongation at 72°C for 5 min. The amplified fragments with adapters and tags were purified using AMPure XP beads (Beckman Coulter Genomic, CA, USA). Prior to library pooling clean constructs were quantified using a Qubit Fluorometer (Invitrogen, Carlsbad, CA, USA) and mixed in approximately equal concentrations to ensure even representation of reads per sample followed 250 bp pair-ended MiSeq (Illumina, CA, USA) sequencing.

Processing of HTS data

The raw dataset containing pair-ended reads with corresponding quality scores were merged and trimmed using settings previously described.²⁹ De-replicating, purging from chimeric reads and constructing de novo Operational Taxonomic Units (OTU, with 97% similarity) was conducted using

the UPARSE pipeline.³⁰ The green genes (v13.8) 16S rRNA gene collection was used as a reference database.³¹

Statistical analyses

For abundance-based analyses (β -diversity), contingency tables (based on OTUs/phylotypes summarized to species level) were normalized with cumulative sum scaling (CSS³²). The influence of explanatory variables over GM composition was evaluated through the Constrained Analysis of Principal Coordinates (CAP) on Bray–Curtis distances, while using the ANOVA-like permutation tests (1,000) to determine the significance of each effect.³³ Through CAP, top discriminant GM features were selected based on a minimum absolute index of 0.15 along the canonical axes; (i) mean differences between such discriminant features was performed with two-tailed Student's *t*-test and corrected for Type I error with False Discovery Rate (FDR).

For mean species diversity (α -diversity) samples were randomly rarefied to 10,000 sequences each, and the number of observed species determined as a function of sequence depth. Differences in α -diversity were determined by either (i) non-parametric using 999 Monte Carlo permutations or (ii) by mixed linear models (MLM) in order to correct for the co-founding effect of age.

Days with diarrhea during outpatient treatment were evaluated using cumulative incidence function (CIF)³⁴ within placebo, probiotic responders and probiotic non-responders subjects, whereas differences in cumulative incidence rates were assessed with the Gray's test.³⁵

Data availability

The accession number of the sequencing-data reported in this paper is ENA: PRJEB29297. Sequencing metadata are available on request.

Disclosure of potential conflicts of interest

Chr. Hansen A/S sponsored parts of the gut microbiome analysis. The company did not play any role in data analysis, data interpretation or manuscript writing.

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