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Bifidobacterium animalis subsp. lactis fermented milk product reduces inflammation by altering a niche for colitogenic microbes

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Intestinal health requires the coexistence of eukaryotic self with the gut microbiota and dysregulated host-microbial interactions can result in intestinal inflammation. Here, we show that colitis improved in T-bet\textsuperscript{−/−}Rag2\textsuperscript{−/−} mice that consumed a fermented milk product containing Bifidobacterium animalis subsp. lactis DN-173 010 strain. A decrease in cecal pH and alterations in short chain fatty acid profiles occurred with consumption, and there were concomitant increases in the abundance of select lactate-consuming and butyrate-producing bacteria. These metabolic shifts created a non-permissive environment for the Enterobacteriaceae recently identified as colitogenic in a T-bet\textsuperscript{−/−}Rag2\textsuperscript{−/−} ulcerative colitis mouse model. In addition, 16S rRNA-based analysis of the T-bet\textsuperscript{−/−}Rag2\textsuperscript{−/−} fecal microbiota suggest that the structure of the endogenous gut microbiota played a key role in shaping the host response to the bacterial strains studied herein. We have identified features of the gut microbiota, at the membership and functional level, associated with response to this B. lactis-containing fermented milk product, and therefore this model provides a framework for evaluating and optimizing probiotic-based functional foods.

Enterobacteriaceae | intestinal inflammation | microbiota | probiotics | colitis

Intestinal health requires the coexistence of eukaryotic self with the gut microbiota (1). This balance is maintained by the intestinal epithelium, mucosal immune system, and gut microbes. In inflammatory bowel disease (IBD), interactions between a host’s immune system and gut microbiota are dysregulated. Studies of the fecal microbiota of IBD patients reveal distinct differences in community membership between healthy and affected individuals (2, 3). An association between gut microbiome profiles and diseases such as IBD (2–4), obesity (5), type 1 diabetes (6), metabolic syndrome (7), and irritable bowel syndrome (8, 9) is emerging from studies of murine disease models and human study populations. The relationship between microbial communities and disease states raises questions as to how microbial communities can be restructured to prevent and treat disease.

Specific probiotics (live microbes that can provide a health benefit to the host) (10), prebiotics (selectively fermentable substances that confer benefits to the host) (11), and antibiotics all represent modalities to alter the composition and function of the gut microbiota (12, 13). The role of antibiotics in the management of IBD remains controversial, with disparate results across different trial populations (14). Furthermore, recent metagenomic studies suggest that antibiotics have sweeping effects on the intestinal microbiota (15, 16). The distal gut microbiome also appears to be a reservoir for antibiotic resistant microbes (17). Thus the use of antibiotics to alter microbial communities is not as predictable or as specific as desired, and could promote the selection and/or outgrowth of antibiotic-resistant strains. Some probiotics may represent an alternative for modulating the gut microbiome. The beneficial effects of specific probiotics have been ascribed to their ability to alter the intestinal microbiota, support colonization resistance against pathogens, and influence host immune responses (18). Defining how consumption of beneficial microbes ameliorates intestinal inflammation in murine models with defined genetics and pathophysiology may facilitate an understanding of who will benefit from probiotic interventions and how probiotics can be optimized for maximal benefit.

The microbiota colonizing the gastrointestinal tract has coevolved with the host to be mutualistic. However, many gut microbes have the capacity to cause or promote disease. Both host and microbial factors influence bacterial opportunism and deficiencies in beneficial microbes may also contribute to chronic intestinal inflammation. Decreased counts of Faecalibacterium prausnitzii have been observed in fecal samples from IBD patients (19, 20), and low counts of these bacteria appear to predict post-surgical recurrence (21). A decreased relative abundance of Bifidobacterium in the gut microbiota of IBD patients has also been observed (22).

Using both culture-dependent and independent methods, we have recently characterized the fecal microbial communities in a murine model of IBD driven by deficiency of T-bet, a T-box family transcription factor, in the innate immune system; and identified culturable colitogenic bacteria that work in concert with gut microbial communities to drive intestinal inflammation (23). In this study, we found that intestinal inflammation improved in the majority of mice that consumed a fermented milk product containing Bifidobacterium animalis subsp. lactis DN-173 010 (B. lactis), Streptococcus thermophilus, two strains of Lactobacillus delbrueckii subsp. bulgaricus, and Lactococcus lactis subsp. cremoris. Marked shifts in cecal pH (decreased pH) and short chain fatty acid (SCFA) profiles (increased acetic acid, propionic acid, and butyric acid and decreased lactic acid) occurred with consumption, and there were concomitant increases in the abundance of select lactate-consuming and butyrate-producing bacteria. These metabolic shifts created...
a nonpermissive environment for the *Enterobacteriaceae* recently identified as colitogenic in a *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> ulcerative colitis mouse model (23). In addition, 16S rRNA-based analysis of the *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> fecal microbiota suggest that the structure of the endogenous gut microbiota plays a key role in shaping host response to probiotics. We have identified features of the gut microbiota associated with response to a *B. lactis*-containing fermented milk product, and therefore this model provides a framework for evaluating and optimizing probiotic-based functional foods.

**Results**

**B. lactis-Containing Fermented Milk Product Improves *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> Colitis and Requires Live *B. lactis*. *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice consumed a *B. lactis*-containing fermented milk product (BFMP) or nonfermented milk product (MP) starting at 4 wk of age for 4 wk; a sham feeding and handling control was performed using sterile water (sham). Mice were provided with additional BFMP, MP, or water (100 mg/per mouse provided in its cage). Colitis scores were decreased in the BFMP group (1.67 ± 1.53) compared with MP (4.11 ± 1.53; *P* < 0.0001) and sham (5.43 ± 0.98; *P* = 0.0001) (Mann–Whitney test) (Fig. L4). 12-wk-old *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice consumed the product for 4 wk, as in Fig. L4, to determine whether similar effects would be observed in more severe disease. Colitis scores were lower in BFMP group (1.77 ± 1.96) as compared with MP (4.36 ± 2.66; *P* = 0.015) and sham (8.38 ± 1.04; *P* < 0.001) (Mann–Whitney test) (Fig. 1B). The BFMP reduced colitis scores in the majority of mice both at early stages and at late stages of disease when intestinal inflammation was more severe.

To determine whether these observations were dependent upon live bacteria in the BFMP, we generated BFMP without live bacteria by irradiation (4.4 × 10^6 Gy). The product was only orally instilled in these experiments—ensuring that it was free of live bacteria and unlikely to support the growth of environmental bacteria as might occur if placed in the cage. Colitis scores in the live BFMP were lower than those of the irradiated product (3.0 ± 1.75 vs. 4.44 ± 1.13, *P* = 0.03) or sham (6.0 ± 1.22, *P* = 0.018) (Mann–Whitney test) (Fig. 1C).

**Consumption of *B. lactis*-Containing Fermented Milk Product Creates a Nonpermissive Environment for Colitogenic *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> *Enterobacteriaceae*.** Select *Enterobacteriaceae* instigate inflammation in *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice (23). BFMP consumption decreased fecal levels of *Enterobacteriaceae* sevenfold (*P* = 0.0132, Mann–Whitney test) as compared with MP by RT-quantitative PCR (Fig. 2A). We recently identified *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup>-derived strains of *Klebsiella pneumoniae* and *Proteus mirabilis* as specific *Enterobacteriaceae* that were capable of inducing colonic inflammation in concert with the endogenous gut microbiota (23). Quantitative culture-based fecal counts of both *K. pneumoniae* and *P. mirabilis* markedly decreased in response to BFMP consumption (*P* = 0.0031 *K. pneumoniae*, *P* = 0.02 *P. mirabilis*, Mann–Whitney test) (Fig. 2B). For seven mice, *K. pneumoniae* was below our limit of detection as was *P. mirabilis* for four mice (Fig. 2B).

Because acidic pH can inhibit the growth of the *Enterobacteriaceae*, we measured theecal pH. Indeed, the ecalf pH of *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice was markedly lower 8 h after BFMP consumption (*P* < 0.0001, one-way ANOVA) (Fig. 2C). Survival of lactic acid producing bacteria from the BFMP could account for the decreased pH. Fecal levels of the four lactic acid producing species present in the BFMP: *S. thermophilus*, *B. lactis*, and *B. lactis* were measured by qPCR and RT-qPCR. The BFMP species were detected in most of the postconsumption BFMP group and all changes observed in response to BFMP were significant (*B. lactis*, *P* = 0.006, *S. thermophilus*, *P* = 0.0139, Wilcoxon signed rank test; *B. lactis*, *P* = 0.0001, *L. lactis*, *P* = 0.0002, Mann–Whitney test) (Fig. 2D).

**Altered SCFA Profiles and Increased Levels of Specific Lactate-Consuming and Butyrate-Producing Bacteria Define the Nonpermissive Environment for the Colitogenic *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> *Enterobacteriaceae*.** The detection of lactic acid producing bacteria suggested that lactic acid may be responsible for the lowecal pH observed, so we measured SCFA levels. Acetic, propionic, and butyric acid levels increased (*P* < 0.0001, one-way ANOVA), whereas lactic acid decreased (Fig. 3A). This observation was counterintuitive, as the BFMP strains are lactic acid producers. However, the decreased lactic acid and increased butyric acid could come from increased endogenous lactic acid-consuming and butyrate-producing bacteria. qPCR and RT-qPCR of fecal material of *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice pre- and postconsumption revealed increases in the lactate-consuming *Desulfovibrio* spp (*P* = 0.0089) and of the lactate-consuming, butyrate-producing *Anaerostipes cacaee* subgroup (*P* = 0.02) and *Eubacterium hallii* (*P* = 0.0044) (Mann–Whitney test) in response to the BFMP as compared with the MP (Fig. 3B). These shifts were specific, as there were not significant shifts in other butyrate producers, such as the *Blautia cocoides* subgroup, *Clostridium leptum* subgroup, or the *Roseburia* subgroup (Fig. 3B).

Bacteriostatic and bactericidal activities have been ascribed to volatile SCFAs (24, 25), and thus theecal conditions of *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice that were capable of inducing colonic inflammation in concert with the endogenous gut microbiota as well as probiotics, such as the *B. lactis* strains present in the BFMP. Additionally, the cecal conditions of *T-bet*<sup>−/−</sup>*Rag2*<sup>−/−</sup> mice improved by the BFMP are shown to be nonpermissive for the *Enterobacteriaceae* as the levels of both *K. pneumoniae* and *P. mirabilis* are markedly decreased in response to the BFMP.

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respectively). Media with the mean cecal pH of these two groups but without the SCFAs were also prepared. K. pneumoniae growth was inhibited by media with the pH of the BFMG group (pH 4.50)

**Fig. 2.** Consumption of B. lactis-containing fermented milk product creates a nonpermissive environment for colitogenic T-bet−/−Rag2−/− Enterobacteriaceae. (A) qPCR using primers targeting Enterobacteriaceae. Pre- and postconsumption levels for MP (●) or BFMG (○) (y axis), expressed as log_{10}(equivalent cells/g feces). Each circle indicates data from one mouse; horizontal bars show the mean. *P* values, Mann–Whitney test. (B) Quantitative fecal culture pre- and post-BFMG and MP (as in A). Colony counts for K. pneumoniae (○) and P. mirabilis (□); detection limit is shown. Each symbol shows data from one mouse; bars show the mean; *P* values, where significant, are shown. (C) Cecal pH for T-bet−/−Rag2−/− mice that consumed BFMG, MP, or sham and for sham Rag2−/− mice. Mean values from three independent experiments are shown (n = 5 mice/sample). Error bars represent SD. (D) qPCR and RT-qPCR of fecal samples pre- and postconsumption of MP or BFMG. Data expressed as log_{10}(equivalent cells/g feces).

**Fig. 3.** Altered SCFA profiles and increased levels of specific lactate-consuming, butyrate-producing bacteria define a nonpermissive environment for colitogenic T-bet−/−Rag2−/− Enterobacteriaceae. (A) Volatile and nonvolatile SCFA levels of cecal contents for BFMG, MP, and sham T-bet−/−Rag2−/− mice. Bars show mean values from three independent experiments (n = 5 mice/sample). Error bars indicate standard deviation. (B) qPCR and RT-qPCR using primers targeting the indicated bacteria genera or species. Pre- and postconsumption levels for MP (●) or BFMG (○) shown along the y axis (log_{10}(equivalent cells/g feces)). Each circle indicates data from a single mouse. Horizontal bars show the mean. *P* values were calculated by Mann–Whitney test or Wilcoxon signed-rank test. (C) Growth kinetics of T-bet−/−Rag2−/−-derived strains of K. pneumoniae and P. mirabilis over a 3-h time course in media with pH and SCFA concentration adjusted to model cecal contents of T-bet−/−Rag2−/− mice that consumed BFMG or MP.
(Fig. 3C), and these slowed growth kinetics were observed up to pH 5.8. In contrast, *P. mirabilis* growth appeared selectively inhibited by media that contained both the lower pH and VFA and non-VFA of the BFMP group (Fig. 3C). Growth inhibition for *P. mirabilis* was significant at the 3 h time point (*P = 0.05*, one-way ANOVA), and decreased growth at 3 h was also significant for *K. pneumoniae* when both BFMP group pH media were compared with the higher-pH MP group media (*P = 0.005*, Mann–Whitney test). Thus, both in vivo and in vitro experiments suggest that consumption of this BFMP creates a nonpermissive environment (low pH and altered SCFA profiles) for colitogenic *T-bet*+/−* Rag2*+/− Entero bacteriaceae.

**Structure of the Gut Microbiota Influences Response to *B. lactis*-Containing Fermented Milk Product.** To identify additional features of the microbiota that could explain why *T-bet*+/−* Rag2*+/− mice respond to the BFMP, we performed a 16S rRNA gene-based survey of the fecal microbiota of *T-bet*+/−*Rag2*+/− and *Rag2*+/− mice before BFMP consumption (*n = 10* group, age 4 wk). Multiplex pyrosequencing of amplicons generated from the V5 and V6 region of the 16S rRNA gene was performed (*n = 20* samples; 7379 ± 2579 reads/sample). Compared with *Rag2*+/− controls, *T-bet*+/−*Rag2*+/− samples had a significantly lower proportional representation of operational taxonomic units (OTUs) belonging to the families *Bifidobacteriaceae* (P = 0.022), *Porphyromonadaceae* (P = 0.011), *Prevotellaceae* (P = 0.011), and *Staphylococcaceae* (P = 0.0037) (Mann–Whitney test with Bonferroni correction) (Fig. 4A). In contrast, the proportional representation of the *Lachnospiraceae* (P = 0.05, Mann–Whitney test with Bonferroni correction) was higher in *T-bet*+/−*Rag2*+/− samples vs *Rag2*+/− (Fig. 4A). The reduced relative abundance of the *Bifidobacteriaceae* was particularly striking, as it suggests a deficiency in *T-bet*+/−*Rag2*+/− mice. The BFMP may replete this deficiency, leading to decreased intestinal inflammation.

To validate and quantify the reduced proportional representation of bifidobacteria, we measured fecal *Bifidobacterium* levels from *T-bet*+/−*Rag2*+/− mice (*n = 20*) at 4 wk of age (the onset of colitis detectable by histology) and healthy age-matched *Rag2*+/− mice (*n = 11*) by qPCR. There was a 316-fold mean reduction in *Bifidobacterium* genus levels in *T-bet*+/−*Rag2*+/− relative to *Rag2*+/− mice (P = 0.0005, Mann–Whitney test) (Fig. 4B). Thus *T-bet*+/−*Rag2*+/− mice were indeed relatively deficient in bifidobacteria.

We asked whether there was an association between repleting these decreased bifidobacterial levels, as measured by recovery of live *B. lactis* from the feces of mice that consumed the BFMP, and colitis score. *B. lactis* was recovered from mice with colitis scores of 0–3 but was below detection (10^2.5 CFU/g feces) in mice with higher scores and was not detected in controls (Fig. 4C). This observation raised the question as to whether subsets of mice could be identified, before product consumption, that would or would not respond to the BFMP. A hierarchical cluster analysis using the family level OTU assignments (Fig. 4D) revealed that *T-bet*+/−*Rag2*+/− with colitis score ≥2 (F, G, I, and J) clustered together in two subsets. Both samples I and J had markedly elevated proportional representation of the *Lactobacillaceae* (27.9% and 31.9%, respectively, approximately two standard deviations above the mean) relative to the other *T-bet*+/−*Rag2*+/− mice. The F and G subset were also notable for low levels of the strains following 4 wk of BFMP (Fig. 4E).

In aggregate, these data suggest that the structure of the gut microbiota may influence and predict response to the BFMP.

**Discussion**

Here, we have used a murine model of colitis to gain insight into how a defined probiotic-containing food reduces intestinal inflammation. In this study, we focused on the microbiota and its response to host consumption of BFMP containing *B. lactis*, *L. lactis*, *L. bulgaricus*, and *S. thermophilus*. We found that *T-bet*+/−*Rag2*+/− mice have a relative deficiency of *Bifidobacterium*, that

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**Fig. 4.** Structure of gut microbiota influences response to *B. lactis*-containing fermented milk product. (A) Distribution of family-level phylogenotypes in *T-bet*+/−*Rag2*+/− and *Rag2*+/− fecal microbiota at 4 wk of age (pre-consumption). Percent relative abundance is plotted. (B) *Bifidobacterium* levels (qPCR) from 4-wk-old *T-bet*+/−*Rag2*+/− (*n = 20*) (C) and *Rag2*+/− (*n = 10*) (D) fecal samples. (C) *B. lactis* counts (log10~CFU/g feces) (y axis) of sufficient quantity from mice in Fig. 1. Mice grouped by product administered and colitis score (x axis). Each circle represents a single mouse; horizontal bars show the mean. (D) Family-level hierarchical clustering analysis of *T-bet*+/−*Rag2*+/− fecal microbial communities before product consumption. Each sample is labeled corresponding to its relative abundance data in A, and colitis scores postconsumption are heat map color coded based on tertile distribution. (E) Postconsumption recovery levels of BFMP strains are heat map color coded based on tertile distribution for the corresponding sample in A and D.
supplementation with a BFMP reduced intestinal inflammation, and that recovery of live \textit{B. lactis} from mice correlated with optimal response. Consumption of this BFMP resulted in an increase of certain lactate-consuming and butyrate-producing bacteria, decrease in cecal pH, and increases in select cecal SCFAs. These conditions proved inhospitable to \textit{T-bet}−Rag2−− mice whose disease is in part driven by colitogenic members of the \textit{Enterobacteriaceae}. In addition, the data from our 16S rRNA sequencing and qPCR-based experiments suggest that the structure of a host’s gut microbiota influences response to probiotics. Thus, the development of gut microbe-based biomarkers for both identification of those individuals likely to benefit from probiotics and for monitoring response to probiotics seems an attainable goal. The BFMP used in these studies is a complex mixture of live bacterial strains, bacterial products, and fermentation products that exert effects on gut microbes as well as host cells. Characterization of both gut microbiomes and host responses to functional foods will benefit from emerging metagenomic and metatranscriptomic techniques, as further light is shed on how chronic inflammation resolves in response to consumption of these products.

### Materials and Methods

**Animal Husbandry.** \textit{T-bet}−Rag2−−, mice, their husbandry, and chow have been described (23, 43). Animal studies and experiments were approved and carried out according to Harvard University’s Standing Committee on Animals as well as National Institutes of Health guidelines.

**Study Product.** The test product was a fermented milk (Activia; Danone); details are given in SI Materials and Methods for product strain details. The control product was a milk-based nonfermented dairy product without bacteria and with 1.6% lactose per serving. Control product details are provided in SI Materials and Methods. Products were provided by Danone Research. A 100-μg quantity was orally instilled daily at the same time daily and, if indicated, 100 mg per mouse in its cage was provided for additional consumption.

**Production Sterilization.** The BFMP was sterilized with 4.4 × 104 Gy in a gamma irradiator. To confirm product sterility after irradiation, the product was cultured under aerobic and anaerobic conditions.

**Histology.** Colonos were prepared for histology, and histopathology was evaluated in a blinded fashion (with respect to genotype and experimental protocol) by J.N.G. as previously described (43).

**Fecal Collection, Storage, and RNA and DNA Extraction.** Fecal pellets were collected into tubes and weighed. RNA later (Ambion) was added to the pellets, samples were homogenized and then stored at −80 °C until RNA and DNA were extracted as described previously (44).

**Quantification of Bacteria Equivalents by qPCR and RT-qPCR.** Determination of bacterial concentration was based on the quantification of either DNA or RNA molecules using primers targeting 16S rRNA sequences (Tables S1–S3). Additional details are provided in SI Methods and Materials. The qPCR and RT-qPCR reactions were performed according to previously published work (44, 45). Primers and their annealing temperature are found in Tables S1–S3.

**Fecal Collection and Culture of Gram-Negative Aerobes.** From each mouse, four to six pellets were collected. Pellets were resuspended in sterile PBS, and 10-fold serial dilutions were generated, plated on MacConkey’s medium, and incubated in ambient air at 37 °C overnight. Biochemical assays with the API-20E panel (bioMérieux) confirmed that colony morphology correlated with identification as \textit{K. pneumoniae} and \textit{P. mirabilis}.

**Cecal pH and SCFA Measurement.** Cecal contents were collected after the animals were killed. An aliquot was removed for pH measurement and the remaining sample was flash frozen in N2(0). SCFA analyses were performed through a panoply of structural and metabolic microbial features, and thus further research is needed to determine the contribution of the specific microbial components in this BFMP to the reduction of intestinal inflammation observed.

Understanding both the microbial community and host response differences underpinning the spectrum of responses to probiotic-based treatments in inflammation will be essential for realizing the full potential of these treatments. Our results demonstrate that this BFMP is effective in \textit{T-bet}−Rag2−− mice whose disease is in part driven by colitogenic members of the \textit{Enterobacteriaceae}. In addition, the data from our 16S rRNA sequencing and qPCR-based experiments suggest that the structure of a host’s gut microbiota influences response to probiotics. Thus, the development of gut microbe-based biomarkers for both identification of those individuals likely to benefit from probiotics and for monitoring response to probiotics seems an attainable goal. The BFMP used in these studies is a complex mixture of live bacterial strains, bacterial products, and fermentation products that exert effects on gut microbes as well as host cells. Characterization of both gut microbiomes and host responses to functional foods will benefit from emerging metagenomic and metatranscriptomic techniques, as further light is shed on how chronic inflammation resolves in response to consumption of these products.
using a Shimadzu GC-14A gas chromatograph (Shimadzu Scientific Instruments) and a SPB-1000 capillary column (Supelco). VFA and non-VFA standard mixtures were used to identify and quantify acids present in each sample. Conjugate base concentrations were weighted correct and converted to conjugate acid concentrations using the cecal pH and Henderson-Hasselbalch equation. Additional details on VFA and non-VFA extractions are provided in SI Materials and Methods.

Growth Kinetics in Modeled Media. Brain-heart infusion medium was prepared to generate the following media. Cecal pH of BMF mice (BHI pH 4.50), MP mice (BHI pH 6.07), cecal pH of BMF mice with VFA and non-VFA (BHI pH 4.50, 99.2 mM acetic acid, 31.05 mM propionic acid, 43.87 mM butyric acid, and 2.08 mM lactic acid), and cecal pH of MP mice with VFA and non-VFA (BHI pH 6.07, 2.35 mM acetic acid, 1.34 mM propionic acid, 0.37 mM butyric acid, and 12.9 mM lactic acid) were maintained at constant pH levels for the experiment. Statistical tests used to determine the significance of differences between treatments were performed using the complete linkage hierarchical clustering (hclust from R http://www.r-project.org/.

B. lactis Quantitative Culture. Quantitative fecal culture was performed using Roga media supplemented with 1% irradiated sterile fermented milk product. Plates were incubated for 5 d (anaerobic conditions). Gram stains were performed and morphotypes consistent with Bifidobacterium as well as control were screened by PCR (46). B. lactis counts were then calculated from the quantitative fecal culture and PCR results and weight corrected based on fecal dry weight.

Statistical Analysis. Error bars represent standard deviation. Statistical tests used to follow the stated P value. Prism (GraphPad) was used for statistical calculations. The Bonferroni correction was applied as noted for multiple comparison correction.

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Metagenomic Sequencing and Analysis. The VS and V6 regions of the 16S rna gene were targeted for amplification and multiplex pyrosequencing with error-correcting barcodes. Sequencing was performed using a Roche FLX Genome Sequencer at DNAStruction. Data were preprocessed to remove sequences with low-quality scores (7,579 and 2,379 high-quality reads per sample, mean read length: 278). Taxonomy was assigned using RDP classifier v.2.01 and similar sequences were binned into operational taxonomic units (OTUs) using cd-hit with minimum pairwise identity of 97%. Sample clustering was performed using the complete linkage hierarchical clustering (hclust from R http://www.r-project.org/.


