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**Title**
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**Permalink**
https://escholarship.org/uc/item/4wj9g27d

**Journal**
The ISME journal, 7(6)

**ISSN**
1751-7362

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**Publication Date**
2013-06-01

**DOI**
10.1038/ismej.2013.15

Peer reviewed

## ORIGINAL ARTICLE

# Biphasic assembly of the murine intestinal microbiota during early development

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The birth canal provides mammals with a primary maternal inoculum, which develops into distinctive body site-specific microbial communities post-natally. We characterized the distal gut microbiota from birth to weaning in mice. One-day-old mice had colonic microbiota that resembled maternal vaginal communities, but at days 3 and 9 of age there was a substantial loss of intestinal bacterial diversity and dominance of *Lactobacillus*. By weaning (21 days), diverse intestinal bacteria had established, including strict anaerobes. Our results are consistent with vertical transmission of maternal microbiota and demonstrate a nonlinear ecological succession involving an early drop in bacterial diversity and shift in dominance from *Streptococcus* to *Lactobacillus*, followed by an increase in diversity of anaerobes, after the introduction of solid food. Mammalian newborns are born highly susceptible to colonization, and lactation may control microbiome assembly during early development.

## Introduction

Mammals are thought to develop in a bacteria-free environment within the mother's womb. They are born through a birth canal densely populated by lactic acid bacteria (Harrison *et al.*, 1953; Dominguez-Bello *et al.*, 2010; Ravel *et al.*, 2011) and, during a substantial period of their early development, feed exclusively on maternal milk. These conserved traits may be important for the nutrition and protection of the newborn (Sela and Mills, 2010). In this work, we characterized the mouse intestinal microbiota from the early stages of development until weaning.

## Materials and methods

Six Friend leukemia virus B (FVB) female mice, 4–5 weeks old, were bred with males, and the intestinal contents of litters and mothers were sampled until weaning (Supplementary Figure S1). This work was approved by the University of Puerto Rico IACUC (604-2008). Owing to the low fecal yield in newborn mice, colon contents were collected at these stages. We confirmed that colon contents were good proxies for feces (Supplementary Figures S2–S5), as reported (Peterson *et al.*, 2008; Turnbaugh *et al.*, 2009).

DNA was extracted from 81 samples using MoBio PowerSoil Kits (MoBio Laboratories, Carlsbad, CA, USA) as recommended by the manufacturer, including bead beating. Sample DNA was PCR-amplified from the variable V2 region of the 16S *rRNA* gene, then sequenced using 454 pyrosequencing (Life Sciences Genome Sequencer FLX instrument, Roche, Branford, CT, USA) as described (Andersson *et al.*, 2008; Fierer *et al.*, 2008), (Hamady *et al.*, 2008). Sequences were processed using Quantitative Insights into Microbial Ecology (QIIME) 1.4 (Caporaso *et al.*, 2010). Rarefaction analysis was performed based on the number of operational taxonomic units (OTUs) and the amount of phylogenetic branch length observed in each sample (Hamady *et al.*, 2010). Good's coverage estimator was also calculated. Beta diversity was

estimated using the UniFrac metric (Lozupone and Knight, 2005), and hierarchical clustering was performed using the unweighted pair group method with arithmetic mean (UPGMA). Analyses of variance (ANOVA) were made using the statistical program R, version 2.14.0 (2011-10-31), using the R library.

## Results

We obtained 121 893 V2 region *16S rRNA* gene sequences from 81 samples, with a mean of $1505 \pm 274$ sequences per sample. Sequences were clustered into 3822 OTUs. Unknown bacteria represented a total of 3.9% of the 121 893 sequences. The global estimated coverage was 98.8%, the coverage values for the offspring at each age were as follows: day 1 = 96.2%; day 3 = 98.5%; day 9 = 99.3%; day 21 = 97.3%; adults (older than 21 days) = 96.9%.

Adult mothers harbored $61 \pm 21.7$ fecal OTUs, with dominance of phyla Firmicutes (52.3%), Bacteriodetes (42.1%) and unclassified OTUs (4.8%) (Supplementary Figure S6; Supplementary Table S1). Bacterial communities of the adults did not vary significantly with age (Supplementary Figure S7), gender (Supplementary Figure S8) or physiological stage associated to parturition (ANOVA, $P = 0.124$; Supplementary Figure S9).

At day 1 after delivery, the vaginas of the six mothers harbored $\sim 38 \pm 11.5$ OTUs, with a dominance of Proteobacteria (80%) and Firmicutes (17%) (Supplementary Figure 1; Supplementary Table S2). Colonic bacterial communities in newborn mice at early ages were closer to maternal communities in vaginas than to those in feces (ANOVA, $P = 0.000$), but by age 21 days, they clustered closer to maternal feces (ANOVA, $P < 0.001$), as shown by Principal Coordinates Analysis (Supplementary Figure S10) and UPGMA clustering (Supplementary Figure S11).

The colonic bacterial diversity decreased considerably at age 3 and 9 days (Figure 1, Supplementary Figures S12–S15, Supplementary Table S3), from nearly complete dominance of *Streptococcus* to *Lactobacillus* (Figure 2, Supplementary Figure S16, Supplementary Table S2). By the time of weaning (21 days), the fecal diversity had increased to similar levels to those in the mothers (Figure 1, Supplementary Figures S13–S15, Supplementary Table S2), and with nearly complete disappearance of the *Streptococcus* species (Figure 2). Procrustes analysis shows that these results are robust to the sequence coverage obtained here (Supplementary Materials), and the decrease in diversity during the mid-strict lactation period is supported by other diversity metrics, including phylogenetic diversity (Supplementary Figure S13), Chao1 (Supplementary Figure S14) and Shannon entropy (Supplementary Figure S15).
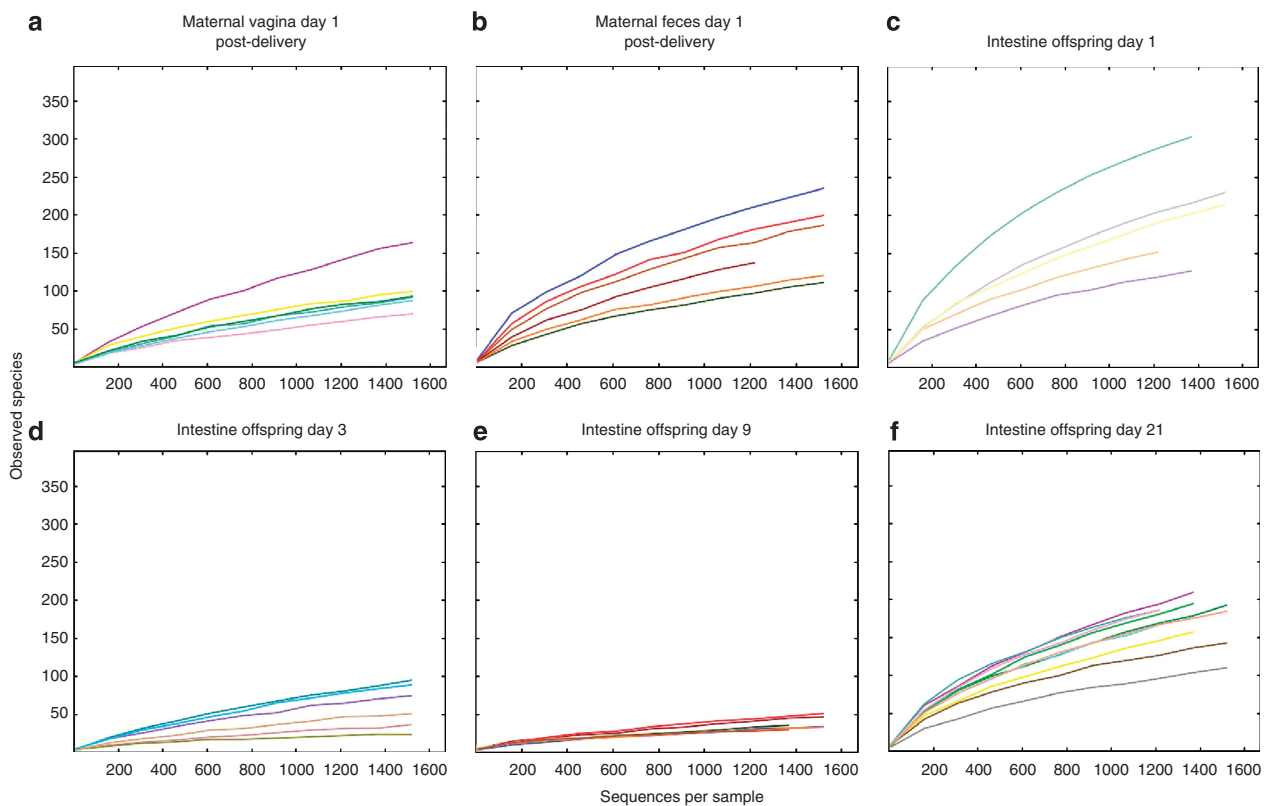


**Figure 1** Rarefaction curves of observed species from mothers and offspring mice. (**a**) Maternal vagina, day 1 post delivery; (**b**), maternal feces, day 1 post delivery; (**c**), offspring intestine, day 1 of life; (**d**), offspring intestine, day 3 of life; (**e**), offspring intestine, day 9 of life; (**f**), offspring intestine, day 21 of life.
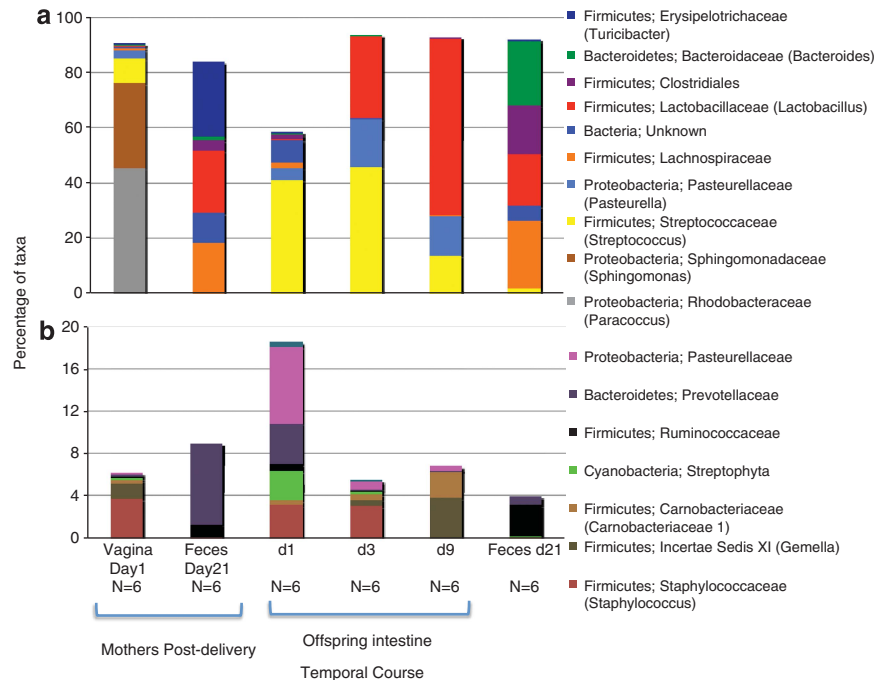
**Figure 2** Proportions of colonic bacterial families in maternal vagina and feces, and during offspring development. (**a**) dominating taxa. (**b**) low abundance taxa.

## Discussion

Consistent with prior studies in humans (Matsumiya *et al.,* 2002; Dominguez-Bello *et al.,* 2010), bacterial communities in the colon of newborn mice resemble maternal vaginal communities. Site-specific selective factors exert pressure during development, and divergence of communities occurs in each body location. Previous work based on cultivable bacteria in mice (Schadler, 1973) has shown an initial colonization by *Lactobacilli*, followed by coliforms, and finally by obligate anaerobes. In the present study, there was an initial bacterial bloom of *Streptococcus* immediately after birth, which decreased after day 3 to be replaced by *Lactobacillus* species that are facultative anaerobes that ferment milk lactose and casein, and produce lactic acid (Kunji *et al.,* 1996; Jiang and Savaiano, 1997; Angelakis *et al.,* 2012). Lactate production acidifies (pH<5.5) the intestinal contents and inhibits the growth of anaerobes (Soergel, 1994; Jiang and Savaiano, 1997), including Lachnospiracea, Clostridiales and Bacteroidales, whose abundance was increased by the time of weaning. The introduction of solids to the milk diet increases the diversity of substrates for intestinal bacteria, and the strictly anaerobic colonizers become established with new pathways of fermentation, leading to the production of short-chain fatty acids, hydrogen, methane and $CO_2$ (Ruppin *et al.,* 1980). The importance of select groups of bacteria including SFB, *Clostridium, Bacteroides, Bifidobacterium* and *Lactobacillus,* in their role on the host mucosal immune system, has been recently reviewed (Reading and Kasper, 2011).

Contrary to the proposed developmental choreography with steady age-associated increase in microbiota alpha diversity (Koenig *et al.,* 2011), the results presented here provide evidence of a biphasic progression toward the adult colonic microbiota, with an early reduction of diversity during suckling with dominance by lactate producers, and a second phase with increased diversity by anaerobes, coinciding with the introduction of solid food.

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