Longitudinal Long-Read Microbiome Profiling in a Canine Model Reveals How Age, Diet, and Birth Mode Shape Gut Community Dynamics

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Running title: Longitudinal Insights into Canine Gut Communities

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ABSTRACT

The gut microbiome undergoes dynamic age-related changes, shaped by diet and maternal factors. Here, we present a species-level, long-read 16S rRNA survey of the developing gut microbiome in a translational canine model, profiling 89 purebred Hungarian Pumis across early-life and reproductive stages. We collected 456 fecal samples longitudinally: 60 puppies followed from birth to 81 weeks, their mothers sampled during pregnancy and lactation, and adult controls from six kennels. We recorded detailed dietary metadata and reproductive status throughout the study. Age was the strongest determinant of alpha diversity, with a rapid increase during weaning and stabilization by six months of age. Beta diversity analyses revealed structured compositional transitions from early developmental phases to adulthood, including a shift toward more uniform, adult-like communities. Within-kennel

variation was modest, consistent with shared environmental exposures. Mixed-effects models showed robust associations between specific taxa and age, diet, and kennel, while SparCC-inferred co-occurrence networks indicated increasing ecological complexity with age. We also demonstrated that the delivery mode - vaginal versus cesarean - impacted early-life microbiome composition: *Lactobacillus spp*. was significantly more abundant in cesarean-born puppies than in vaginally delivered littermates during the 8–10-week window. We also observed reproducible maternal microbiome shifts during pregnancy and lactation, with potential implications for vertical microbial transfer. Taken together, our results show that domestic dogs follow a reproducible, age-structured trajectory of microbial maturation that parallels human development, including delivery-mode effects and diet-responsive taxa.

IMPORTANCE

Microbiome research is among the fastest-moving areas in biomedicine, driven by major global efforts to understand how microbial communities shape human health and disease. Dogs provide an ideal translational model because their gut microbiota more closely resembles that of humans than that of other studied animals; moreover, breeds show high within-breed genetic homogeneity, diets can be tightly regulated, and longitudinal sampling across the lifespan is feasible. Mapping shifts driven by diet and maternal factors - from early-life events through later life, including senior stages - is essential to leverage microbial plasticity for prevention, with implications for inflammation, metabolic disease, and neurodegeneration. Here, we advance this goal by providing a longitudinal, high-resolution dataset and demonstrating that full-length 16S rRNA sequencing is a powerful tool for resolving fine-scale patterns of gut colonization and maturation.

INTRODUCTION

The gut microbiota - comprising bacteria, archaea, viruses, fungi, and protozoa - plays a critical role in host immunity and metabolism via the production of metabolites that influence both intestinal and systemic health(1, 2). The collective genomes of these microorganisms constitute the gut metagenome. How these communities assemble and stabilize in early life remains a major focus of microbiome research. Dogs (*Canis lupus familiaris*) are valuable comparative and translational models because they share human environments and exhibit human-like dietary patterns, physiology, and disease risk profiles (3). Notably, compared with pigs or mice, canine gut microbiomes are more similar to those of humans in species-level composition and diet-responsive dynamics (4), supporting their relevance for cross-species investigations of host–microbiome interactions.

Canine gut microbiome composition varies with age, diet, reproductive status, and genetics (5). In older age, diversity decreases, taxon abundances shift (6–9), and dysbiosis emerges (10); this trajectory coincides with immunosenescence and chronic low-grade inflammation ("inflammaging") (11–13). Although the human infant microbiome is well characterized, comparable longitudinal canine data remain limited (14, 15). Previously, the mammalian fetal gut was considered sterile (16), but accumulating molecular evidence has challenged the "sterile womb" hypothesis, with reports of microbial DNA in the placenta, amniotic fluid, and meconium (17, 18). In dogs, microbial DNA has been detected in meconium and amniotic fluid following both vaginal and cesarean deliveries (19, 20). Vertical transmission occurs via birth, nursing, and close maternal contact (21). In humans, delivery mode has a major impact on early colonization dynamics and microbial diversity (22–25), but corresponding canine studies remain scarce.

Diet is another key modulator of gut microbiota composition. Although classified as facultative carnivores, domestic dogs can adapt to a wide range of dietary patterns. Most commercial products are

68 high in carbohydrates and are extruded into dry kibble, while raw and high-protein formulations offer
69 alternative macronutrient profiles (26). Dietary interventions modulate microbiota composition and
70 metabolic outputs, with downstream effects on inflammation and health status (4, 27). Several studies
71 have shown that the canine microbiome responses to diet resemble those observed in humans,
72 particularly with respect to macronutrient-driven shifts (4, 28, 29). For example, both protein and fiber
73 content influence gut taxonomic and functional profiles (29). Despite these insights, key knowledge gaps
74 remain in our understanding of microbial development, vertical transmission dynamics, and diet–
75 microbiome interactions during early life (15, 30, 31). In particular, the temporal structure of microbiome
76 maturation and the impact of birth mode or maternal status in dogs are not well characterized. Moreover,
77 although diet is a well-known modulator of adult gut communities, its role across different stages of
78 postnatal development is less understood (29, 32, 33).

79 Breed-specific microbiota variation has been reported across mammalian species (34–37). In dogs,
80 *Firmicutes* and *Bacteroidetes* consistently dominate the gut microbiota across breeds and physiological
81 contexts (34, 38–40). However, few studies have investigated how breed-specific or kennel-specific
82 factors interact with developmental and dietary influences in shaping microbial succession. Longitudinal
83 sampling is essential to capture temporal microbial dynamics and disentangle age-related effects from
84 environmental variation (41). Full-length 16S rRNA gene sequencing, enabled by long-read platforms
85 such as Oxford Nanopore Technologies (ONT) and Pacific Biosciences, provides species-level
86 resolution and enables high-resolution taxonomic mapping across diverse hosts (42–46).

87 In this study, we present a longitudinal canine microbiome dataset with species-level resolution,
88 capturing the developmental trajectory of gut microbial communities from birth to 81 weeks of age in
89 purebred Hungarian Pumi dogs. We also examine the effects of delivery mode, maternal status, and diet
90 on community structure, and explore community co-occurrence patterns across early life. This work aims

to advance our understanding of postnatal microbiome assembly in a naturally reared, translational model species.

RESULTS

Study Design and Analytical Pipeline

In this study, we profiled 456 fecal samples from 89 purebred Hungarian Pumi dogs - 55 puppies followed from birth to 81 weeks, dams across pregnancy/lactation, and time-matched adult controls - with a small cesarean-delivered subgroup (5 puppies from one litter) analyzed separately. DNA was extracted (ZymoBIOMICS 96 MagBead); full-length 16S rRNA amplicons (V1–V9) were sequenced on MinION (R9) and basecalled with Dorado. Taxa were profiled with EMU and diversity computed in phyloseq from unfiltered counts. Downstream R analyses applied prevalence/abundance filters (>5%; ≥5 samples), mixed-effects models (lme4) for age, sex, kennel, kennel:litter, and diet (age modeled categorically and with splines:diets categorized based on FEDIAF guidelines), and comparative tests (Wilcoxon, t-test, Mann–Whitney; $\alpha = 0.05$). Community structure was summarized by hierarchical clustering and heatmaps (ComplexHeatmap), taxonomic trees in iTOL, and SparCC co-occurrence networks (CONs; SpiecEasi) visualized in igraph/Cytoscape (**Fig. 1**).

Sequencing output: The MinION sequencing runs yielded a total of 121.30 million pass ($Q \geq 10$) reads (mean ± SD: 266017.09 ± 127742.50 per sample) with an average read length of 1521.81 ± 50.07 bp and a mean Q-score of 20.40 ± 1.05. This corresponded to approximately 185.01 GB of raw data.

Microbial Diversity Increases during Puppyhood but Stabilizes in Adulthood

Ordination of the sampled cohort by non-metric multidimensional scaling (NMDS) reveals a clear age-associated gradient (**Fig. 2A**): neonatal and early-life samples are widely dispersed, indicating high inter-individual variability, whereas profiles from later time points cluster more tightly, consistent with

113 progressive convergence toward an adult-like community. In line with this, alpha diversity - measured
114 by the Shannon index - increases markedly during the first weeks of life, with a pronounced rise around
115 the weaning window (3.5–6 weeks), followed by a gradual tapering and an apparent plateau by
116 approximately 6 months of age (**Fig. 2B**). This trajectory is evident across kennels and age strata (age
117 groups 1–6 as defined in **Fig. 2B**), indicating that the maturation pattern is robust to kennel-level
118 differences in husbandry and environment.

119 Effects of Environment and Host Sex on Community Composition are Modest

120 To quantify the relative contribution of host and environmental variables, we estimated effect sizes for
121 alpha diversity using a cumulative link model adjusted for covariates and subject ID. Age emerged as
122 the dominant predictor ($p < 0.001$), with kennel exerting a smaller but statistically significant effect and
123 sex showing no detectable contribution (**Fig. 2C**). Comparisons between litters within the same kennel
124 revealed generally modest differences in overall Shannon diversity, with only a single litter pair reaching
125 statistical significance (Wilcoxon test, $p < 0.05$; **Fig. 2D**). Finally, within a representative litter, beta
126 diversity distances relative to the dam show early divergence at ≤10–20 days, followed by increasing
127 similarity by 40–60 days that remains stable at later intervals (**Fig. 2E)**. Together, these patterns indicate
128 that age-linked maturation - and the diet transitions that accompany it - are the primary drivers of
129 community structure, while environmental variation at the kennel or litter level exerts comparatively
130 smaller effects.

131 Developmental Stage Shapes Taxonomic Composition

132 Analysis of the most abundant genera across litters and kennels identified clear compositional shifts over
133 time (**Fig. 3A**). During the weaning transition (3.5–8 weeks), early-life taxa — *Escherichia*, *Megamonas*,
134 *Lactobacillus*, and *Ruminococcus* — were dominant, whereas the post-weaning period was characterized
135 by the higher relative abundance of *Blautia*, *Peptacetobacter*, and *Fusobacterium*. At early weaning (6–

136 8 weeks), *Lactobacillus* exhibited kennel-specific presence and diet-linked abundance patterns (**Fig. S1**).
137 Notably, *Blautia* was consistently abundant across all life stages and showed a steady rise over time (**Fig.**
138 **3A**), suggesting its establishment as a core post-weaning genus.

139 To identify the dominant members of the gut microbiome in the cohort, we examined the most prevalent
140 and abundant species. As shown in **Fig. 3B** *Blautia producta*, *Blautia hansenii, Faecalimonas*
141 *umbilicata*, and *Megamonas rupellensis* were among the most prevalent species, while *Blautia*
142 *glucerasea* and *Peptacetobacter hiranonis* exhibited the highest relative abundance. A taxonomic tree
143 highlighting the top 350 taxa revealed distinct associations with age and kennel of origin (**Fig. 3C**).
144 Species-level resolution enabled by full-length 16S sequencing captured fine-scale taxonomic patterns,
145 including taxa showing consistent shifts across multiple litters and kennels. At the genus level,
146 hierarchical clustering of the top 40 taxa showed that age and diet were more influential than sex in
147 shaping microbial community structure (**Fig. S2**). To further explore host and environmental influences
148 on taxonomic profiles, we performed mixed-effects modeling across four key factors: age, kennel, diet,
149 and sex. Each model revealed distinct taxa significantly associated with each factor (**Fig. S3–S6**).

150 Species-level Core Microbiome Across Age

151 Using a prevalence-based definition, we identified a species-level core within each age group and present
152 three complementary analyses. First, focusing on the strict 90%-prevalence criterion, a set of 13 species
153 was present in all age groups; their mean relative abundances varied systematically with age (**Fig. S7A**).
154 This ubiquitously detected set was dominated by short-chain fatty acid (SCFA)–associated *Clostridia*
155 (e.g., *Blautia* spp., *Faecalimonas umbilicata*) (47) and the bile-acid–transformer *Peptacetobacter*
156 *hiranonis*, which has been associated with maintenance of intestinal barrier integrity, colonization
157 resistance against pathogens such as *Clostridioides difficile*, and broader host metabolic homeostasis

158 (48). Some taxa (e.g., *Megamonas rupellensis*) showed prominence in both early and late life, aligning
159 with shifts in diet and overall community complexity.

160 Second, pairwise Jaccard similarities between 90%-prevalence core sets (**Fig. S7B**) were highest between
161 adjacent late-life stages (e.g., 3–7 y *vs*. 7–10 y), indicating a stable, adult-like core with gradual turnover,
162 whereas overlaps between pre-weaning and adult groups were minimal, highlighting weaning as a major
163 ecological breakpoint.

164 Third, to visualize broader intersections, we applied a more inclusive 80%-prevalence threshold in an
165 UpSet analysis (**Fig. S7C**). The largest intersection corresponded to the same 13 universally present
166 species, while multiple intermediate intersections spanned 2–5 adjacent age groups, indicating
167 progressive rather than abrupt replacement during maturation.

168 *Functional context.* The persistent core is enriched for SCFA producers and bile acid modulators, which
169 contribute to mucosal integrity and immune-metabolic balance across life stages; early-life deviations
170 and late-life shifts likely reflect changing dietary substrates, host physiology, and age-related reductions
171 in community redundancy.

172 Developmental Rewiring of the Microbial Network

173 SparCC-based CONs revealed striking differences in microbial interactions across age groups (**Fig. 4**).
174 In pups younger than 3.5 weeks, the network was sparsely connected with few co-occurring taxa. At 6–
175 8 weeks, network density and modularity increased, indicating more structured microbial communities.
176 By 1–2 years of age, the adult microbiome exhibited robust and highly connected networks, with strong
177 positive associations (red edges) among core taxa such as *Bacteroides*, *Clostridium*, and *Sutterella*.
178 These transitions are consistent with compositional and functional maturation of the gut microbiota.

179

Birth Mode Influences Microbiome Composition During Early Postnatal Development

A focused comparison of two litters from the same dam and kennel (Le Petit Lapin) - one delivered by cesarean section and one by vaginal delivery - matched for age (8–10 weeks) and diet revealed differences in beta diversity (NMDS ordination) between the groups (**Fig. 5A**). However, Shannon alpha diversity did not differ significantly (**Fig. 5B)**. Notably, C-section pups showed higher relative abundances of *Lactobacillus* (p.adj = 0.008) and *Prevotella* (p.adj = 0.038), and a lower abundance of *Romboutsia* (p.adj = 0.045) than vaginally delivered pups (**Fig. 5C–D**). These findings suggest that delivery mode can leave a lasting imprint on early gut colonization under otherwise controlled conditions. Contrary to many human studies - where C-section is associated with reduced *Bacteroides*/*Prevotella* and lower *Lactobacillus* early in life - our C-section pups showed higher *Lactobacillus* and *Prevotella* at 8–10 weeks. This discrepancy may reflect species-specific ecology, kennel environment, weaning diet, and the later sampling window. Given the single-litter C-section cohort, these results should be interpreted cautiously.

Maternal Gut Microbiome Shifts with Reproductive Stage

The gut microbiota of dams displayed transient reductions in alpha diversity during late pregnancy and early lactation, followed by partial restoration post-weaning (**Fig. 6A**). Despite temporal proximity, microbiome profiles on the prepartum and postpartum days exhibited distinct compositional signatures (**Fig. 6B**). The transition from prepartum to postpartum was marked by clear taxonomic turnover, with the disappearance of *Blautia* and *Fusobacterium*, and the emergence of *Clostridium* and *Streptococcus* **(Fig. 6B)**. These findings highlight the dynamic microbiome shifts associated with reproductive status and their potential role in seeding the neonatal microbiota.

DISCUSSION

This longitudinal, species-resolved survey of purebred Hungarian Pumi dogs shows that gut microbiome maturation is a strongly age-linked process that proceeds through an early, variable phase toward a more homogeneous adult community. Alpha diversity rose steeply around weaning and plateaued by ~6 months, while ordination revealed an age-graded convergence in beta diversity. These patterns were consistent across kennels and litters, indicating that developmental dynamics are largely robust to kennel-level environmental heterogeneity.

Taxonomic analyses reinforced this trajectory. Early samples (0–8 weeks) were enriched for *Escherichia*, *Megamonas*, *Lactobacillus*, and *Ruminococcus*, whereas later stages showed dominance of *Blautia*, *Peptacetobacter*, and *Fusobacterium*, with *Blautia* increasing progressively across life stages. The short weaning window (6–8 weeks) emerged as a notable inflection point, including kennel-specific and diet-linked *Lactobacillus* patterns, consistent with a transition from milk-adapted to solid-food-adapted communities. Together, these findings emphasize weaning as a critical ecological breakpoint for canine gut community assembly.

Comparable patterns have been described in humans and other mammals, where an early-life dominance of *Enterobacteriaceae* and *Lactobacillaceae* is progressively replaced by fiber-associated *Clostridia* (e.g., *Blautia*, *Ruminococcus*, *Faecalibacterium*) following the introduction of solid food (49). In carnivorous mammals, including dogs and cats, *Fusobacterium* often becomes a prominent post-weaning taxon, reflecting adaptation to meat-rich diets (15). The continuous increase of *Blautia* in our cohort parallels its rise in post-weaning communities across species and is consistent with its proposed role as a core genus during gut maturation (50). In contrast to humans, where *Bifidobacterium* can remain abundant during weaning (51), our dogs showed low and inconsistent *Bifidobacterium* levels throughout development. This pattern consistent with previous canine studies reporting a limited prevalence of this

genus (15). Together, these similarities and differences underscore both the translational relevance of canine models and the species-specific features of microbiome development.

Modeling of covariates demonstrated that age is the dominant predictor of alpha diversity, with kennel exerting a smaller yet detectable effect and host sex showing no measurable impact in this cohort. Comparisons between litters within the same kennel also revealed generally modest differences. These results suggest that, under typical husbandry and diet practices, intrinsic developmental programs and age-coupled diet transitions outweigh contextual environmental variation in shaping community structure. Practically, this implies that longitudinal age matching and careful accounting for weaning status are essential for study design and interpretation in canine microbiome research.

Network analyses indicated that microbe–microbe association structure also matures with age. SparCC-based association networks were sparse in neonates, became denser and more modular at 6–8 weeks, and were robust and highly connected in adults, with strong positive correlations among core taxa (e.g., *Bacteroides*, *Clostridium*, *Fusobacterium*). While correlation does not establish interaction, the observed densification and modularity are consistent with increasing niche differentiation and cross-feeding potential as diets diversify and host physiology stabilizes.

In our study, the focused comparison of two litters from the same dam and kennel suggests that birth mode can leave a detectable imprint on community composition beyond the immediate neonatal period. At 8–10 weeks, C-section pups exhibited higher *Lactobacillus* and *Prevotella* and lower *Romboutsia* than vaginally delivered pups, without a difference in Shannon diversity. This pattern diverges from many human reports and may reflect species-specific postnatal ecology (intense dam–pup contact, coprophagy), kennel environment, weaning practice, or the later sampling window. Given that our cesarean cohort comprised a single litter, these observations should be interpreted cautiously and validated in larger, multicenter cohorts that explicitly track perinatal practices. Zakošek Pipan and

249 colleagues (2020) reported that vaginally delivered puppies exhibited meconium microbiota primarily of
250 maternal vaginal origin, whereas cesarean-delivered puppies showed lower diversity with mixed
251 resemblance to maternal vaginal and oral bacteria (52). Another study by Rota et al. (2021) reported that
252 in cesarean-delivered puppies, amniotic fluid and meconium were found to contain *Acinetobacter*,
253 *Staphylococcus spp*., and *Bacillus spp*. as primary microbiota (20). In humans, Pahirah et al. (2024)
254 reported a higher relative abundance of *Lactobacillus* in cesarean-delivered neonates compared to those
255 delivered vaginally, suggesting that delivery mode–associated microbial shifts at the genus level may be
256 conserved across species (53).

257 We also observed stage-linked shifts in dams, with transient reductions in alpha diversity during late
258 pregnancy and early lactation and compositional turnover across the pre- to postpartum transition. These
259 maternal dynamics likely influence early seeding and may interact with birth mode and nursing behaviors
260 to shape early pup communities. Future work that jointly profiles dams and pups at higher temporal
261 resolution, including milk and environmental sources, will help disentangle vertical and horizontal
262 transmission routes.

263 Methodologically, this study leverages full-length 16S rRNA sequencing to achieve species-level
264 resolution across a dense longitudinal design, minimizing batch effects and enabling fine-scale
265 taxonomic and association analyses. Nonetheless, several limitations merit note. First, while kennel
266 effects were modest, our cohort is breed-specific; generalization beyond Pumis requires replication
267 across breeds and mixed-breed populations. Second, the observational design cannot establish causality,
268 and co-occurrence networks derived from compositional data capture correlations rather than direct
269 interactions. Third, amplicon sequencing limits functional inference; integrating shotgun metagenomics,
270 metatranscriptomics, and metabolomics would clarify metabolic capacity and activity across
271 developmental stages. Finally, the cesarean comparison was underpowered (single litter), and residual
272 confounding (e.g., subtle husbandry differences) cannot be fully excluded.

Taken together, our results position age - and the diet transitions that accompany it - as the primary drivers of canine gut microbiome assembly, with kennel/litter context and host sex exerting comparatively smaller effects. However, because age and diet co-vary in early life, their effects cannot be fully disentangled; by contrast, adult diets were stable (BARF confined to a single kennel), limiting adult diet–related confounding. We identify weaning as a key window of rapid ecological change, document progressive remodeling of association network topology with development, and provide preliminary evidence that birth mode can shape early postnatal composition under otherwise controlled conditions. These insights support the use of dogs as a translational model for early-life microbiome development and highlight practical levers - age matching, weaning status, and perinatal factors - to improve study design and potential intervention strategies. Future studies that expand cesarean cohorts, incorporate functional multi-omics, and prospectively track dam–pup–environment triads will be critical to establishing mechanisms and guiding targeted microbiome modulation in veterinary and comparative medicine.

MATERIALS AND METHODS

Experimental Animals and Feeding Protocols

This study included a main cohort of 84 healthy purebred Hungarian Pumi dogs, comprising 55 puppies from 12 naturally delivered litters, their mothers, and additional adult male and female counterparts from six kennels. Kennel abbreviations used throughout: BCS = Bükki Cserfes; DMD = Duna-menti Dumás; SRT = Serteperti; PPA = Pattogó Parázs; RRT = Rezerta-Réti; LPL = Le Petit Lapin. Puppies were followed longitudinally from birth until adulthood, up to 81 weeks of age. The average age of the mothers was 4.3 ± 1.6 years (mean ± SD), while the control adult dogs had a mean age of 5.9 ± 3.6 years (mean ± SD) (see **Table 1**). Additionally, a subgroup of five puppies (10 samples) born by cesarean section from a single litter in the Le Petit Lapin kennel was included, delivered by the same dam that

296 previously gave birth vaginally; C-section samples were analyzed separately, and excluded from the
297 primary cohort analysis presented in **Table 1** (see **Table 2** for details).

298 Puppies were exclusively breastfed until ~8 weeks, with solid food introduced at ~3.5 weeks alongside
299 continued nursing. All dogs were primarily fed dry commercial diets; however, a subset of adult dogs
300 from one kennel (Duna-menti Dumás) received a raw food diet (BARF). Diet composition and
301 classification followed 2024 FEDIAF guidelines (54)**;** details are provided in **Table S1**. None of the
302 animals received antibiotics during sampling.

303 Study Samples

304 Stool samples were collected at multiple time points from dams during gestation, lactation, and post-
305 lactation; from their offspring from birth through adulthood (up to 81 weeks); and from adult control. In
306 total, 463 samples were collected and sequenced, of which 456 were included in the analyses presented
307 here. The remaining 7 samples (see **Table S2** for accession numbers and exclusion criteria) are available
308 in ENA (BioProject accession **PRJEB82125**) but were excluded from our analyses. These 7 samples
309 included one sample from a single puppy, not part of the main cohort, which accounts for the difference
310 between the 90 dogs archived in ENA and the 89 dogs included in this study. All samples were obtained
311 immediately after defecation using DNA/RNA Shield-Fecal Collection Tubes (Zymo Research) and
312 stored at −80 °C until extraction.

313 DNA Extraction

314 Genomic DNA was extracted from stool samples using the ZymoBIOMICS™ 96 MagBead DNA Kit-
315 Dx (Zymo Research, # Cat. No. D4308-E) following the manufacturer's protocol under sterile conditions
316 in a Class II biosafety cabinet. Samples were homogenized before aliquoting, and bead-beating plus
317 magnetic bead–based purification steps were performed according to the manufacturer's instructions.

318 DNA yield (**Table S3**) and quality were assessed by Qubit® 4 Fluorometer and Agilent 4150 TapeStation.
319 gDNA was stored at −20 °C until library preparation.

320 16S Library Preparation and Sequencing

321 Full-length 16S libraries (V1–V9) were prepared from high-quality gDNA (DINs ≥6) using Oxford
322 Nanopore Technologies' Rapid Sequencing Amplicons–16S Barcoding Kit (SQK-16S024) with
323 AMPure XP bead cleanup, following the manufacturer's protocol. Library concentration (**Table S3**) and
324 fragment size (~1500 bp) were verified using Qubit® 4 and Agilent 4150 TapeStation before pooling,
325 respectively. Libraries were pooled equimolarly and sequenced on an ONT MinION (Version R9; 25
326 flow cells total). Sequencing raw data were basecalled with Dorado v0.8.0 in super accurate mode
327 (minimum Q score = 10) before downstream analysis. See **Table S4** for details of the libraries pooled
328 per flow cell and sequencing read statistics.

329 Bioinformatic Analysis

330 EMU v3.4.5 (55) was used to generate read count tables per taxon and calculate relative abundances.
331 Diversity metrics were computed from unfiltered counts using phyloseq v3.2 in R v4.2 (56). Taxa were
332 filtered for >5% relative abundance and presence in ≥5 samples. Mixed-effects models **(**lme4**)** assessed
333 the effects of age, sex, kennel, kennel:litter, and major diet categories; age was modeled both
334 categorically and with splines to account for diet–age confounding. A taxonomic tree of the top 350 taxa
335 was visualized using iTOL (57), and genus-level hierarchical clustering and heatmaps were produced
336 with ComplexHeatmap (58). Microbial co-occurrence networks were inferred with SparCC **(**SpiecEasi**)**
337 and visualized using igraph and Cytoscape (59). Comparative statistics (Wilcoxon rank-sum, Student's
338 t-test, Mann–Whitney U-test) were applied as indicated in figure legends, with $p < 0.05$ considered
339 significant unless stated otherwise.

340 **Core Microbiome Definition and Visualization:** Core sets were defined at the species level using
341 prevalence thresholds within each age group. A strict cutoff (≥90% prevalence) was applied for the
342 heatmap and Jaccard similarity analyses, while a more inclusive cutoff (≥80% prevalence) was used for
343 the UpSet plot, consistent with established practices in microbiome research (60). Heatmaps and Jaccard
344 indices were computed from prevalence tables, and intersections among core sets were visualized using
345 an UpSet plot. All visualizations were generated in Python (v. 3.12.3) using the pandas, numpy, and
346 matplotlib libraries. The UpSet representation was created with custom plotting code based on
347 matplotlib.

348 DNA isolation, library preparation, and the bioinformatic pipeline followed current best-practice
349 guidelines as described in recent benchmarking studies (61, 62). The detailed methods description is
350 provided in **Supplementary File S1.**

351 ACKNOWLEDGEMENTS

352 This work was supported by the Momentum program I of the Hungarian Academy of Sciences LP2020-
353 8/2020 (D.T.), and by the National Research, Development and Innovation Office (NRDIO) FK 142676
354 (D.T.). The APC fee was covered by the University of Szeged Open Access Fund: 8004. M.A., A.T., and
355 N.J.Y. are awardees of the Stipendium Hungaricum Scholarship for full-time Ph.D. studies, under the
356 registry numbers SHE-29026-004/2021, SHE-102901-004/2022, and SHE-33967-004/2023,
357 respectively. The project was also funded by the NRDIO grant K 142674 to Z.B.

358 The authors declare no conflicts of interest.

359 We would like to express our sincere gratitude to the following kennel owners for providing samples
360 from their dogs (**Supplementary Table S3**):

Ildikó Abonyi, owner of the Duna-menti Dumás Kennel [Baja, Hungary, https://www.pumikennel.eu/];
Dr. Csaba Dobó-Nagy, owner of the Pattogó Parázs Kennel [Gödöllő, Hungary, https://pumi.pedigreedatabaseonline.com/hu/Pattog%C3%B3-Par%C3%A1zs-kennel-Csaba-Dobo-Nagy/breeder/638];
Viktória Hordósné Vasas, owner of the Bükki Cserfes Pumi Kennel [Bükkzsérc, Hungary, https://www.xn--bkki-cserfes-pumi-kennel-vsc.hu/];
Gabriella Kassai, owner of the Serteperti Pumi Kennel [Kiskunmajsa (formerly Dány), Hungary, https://serteperti.hu/];
Bianka Máthé-Szentes owner of the Le Petit Lapin Pumi Kennel [Zomba, Hungary, https://www.facebook.com/lepetitlapinkennel/];
Melinda Takácsné Horváth, owner of the Rezerta-Réti Pumi Kennel [Ajka-Padragkút, Hungary, https://www.facebook.com/p/Rezerta-R%C3%A9ti-PUMI-Kennel-100057382122448/].

We would also like to thank the following dog owners for kindly providing samples: Kimberley Birkett (Beni; Finland), Eszter Dandé (Borcsa; Hungary), Dénes Gergely (Magor; Hungary), Tünde Gyöngyös (Briós; Switzerland), Ildikó Hernádi (Csúzli; Hungary), Yvett Király (Csepke; Hungary), Júlia Lang (Blöff; Hungary), Minna Silander (Pusta; Finland) and Éva Szőke (Rozi; Hungary).

Finally, we are especially grateful to Tünde Baloghné Jancsovics, board member of the Hungarian Pumi Club (https://pumiklub.eu/english), for establishing the crucial contact with breeders and facilitating the collaboration.

We gratefully acknowledge Balázs Kakuk [Department of Medical Biology (DMB), University of Szeged (USZ)] for his work on basecalling and EMU output generation. We also thank Zsolt Csabai (DMB, USZ) and Ákos Dörmő (DMB, USZ) for laboratory support and helpful discussions.

Author Contributions (CRediT)

- **Conceptualization:** D.T., Z.B.
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- **Data curation:** M.A., T.J., G.G.
- **Formal analysis:** P.O., D.T., M.A.
- **Validation:** D.T., M.A.
- **Project administration:** D.T.
- **Supervision:** D.T.
- **Writing – original draft:** D.T., M.A., A.T., P.O., Z.B.
- **Writing – review & editing:** M.A., A.T., N.J.Y., T.J., G.G., P.O., D.T., Z.B.

All authors read and approved the final paper.

ETHICS APPROVAL

In accordance with institutional and national regulations, formal animal ethics approval was not required for this work. The study involved only noninvasive fecal sampling during routine husbandry without any changes to housing, diet, or veterinary care; samples were provided voluntarily by breeders. All procedures complied with national animal welfare regulations.

DATA AVAILABILITY STATEMENT

The sequencing raw data were basecalled using Dorado in super-accurate mode ($Q \geq 10$), and the resulting FASTQ files have been deposited in the European Nucleotide Archive (ENA) under the

405 BioProject accession number PRJEB82125. The repository also contains sample-associated metadata,
406 and the metadata used for analyses are provided in **Table S5-S6**.

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591

592 # LEGENDS

593 **Fig. 1. Graphical Abstract of the Study Workflow.**

594 Fecal samples from purebred Hungarian Pumi dogs were processed for full-length 16S rRNA gene
595 sequencing on the Oxford Nanopore MinION platform, followed by Dorado basecalling and taxonomic
596 profiling with the Emu 16S database.

597

598 **Fig. 2. Longitudinal Diversity Patterns and Within-Litter Variation in Canine Gut Microbiome.**

599 **(A)** NMDS plot revealing a developmental gradient in the gut microbiota, with increasing homogeneity
600 in adulthood. **(B)** Shannon diversity index across age groups, showing microbial expansion during early
601 development. Age groups: 1 (<3.5 wks), 2 (3.5–6 wks), 3 (6–8 wks), 4 (10–26 wks), 5 (26–52 wks), 6
602 (>52 wks)**. (C)** Relative effect sizes of cohort variables on alpha diversity (Shannon index). Analysis was
603 based on a cumulative link model adjusted for covariates and subject ID (***$p < 0.001$). **(D)** Within-
604 kennel litter comparisons of Shannon diversity, showing mostly modest differences with one significant
605 exception (Wilcoxon test, *$p < 0.05$). **(E)** Beta-diversity distances within a single litter. Samples from
606 the mother and pups are shown in red and blue, respectively. Dotted lines connect longitudinal samples
607 from the same subject taken within the same age period.

608

609 **Fig. 3. Key Taxa of the Canine Gut Microbiome: Abundance–Prevalence Patterns and Age/Kennel**
610 **Associations.**

611 **(A)** Relative abundances of the top genera, stratified by kennel, litter, and life phase. **(B)** Abundance–
612 prevalence scatter plot of the most common species. Dot size corresponds to prevalence; color represents
613 the genus (shown for the top 25 species). **(C)** Taxonomic tree of the top 350 prevalent and abundant taxa,
614 annotated with mean effect sizes from mixed-effect models for age and kennel associations (contrasting
615 color scales). Key species with the strongest associations are highlighted.

616
617 **Fig. 4. Co-occurrence Relationships between Age Categories Reflect Dietary Shifts.**

618 **(A)** Co-occurrence network (SparCC) at up to 3.5 weeks. **(B)** Network at 6-8 weeks. **(C)** Network at 1-
619 2 years. Node size is proportional to genus prevalence. Nodes are colored by Louvain cluster; edges are
620 colored by correlation (blue = negative, red = positive).

621

622 **Fig. 5. Comparison of Cesarean-section vs. Vaginally Delivered Pup Microbiomes.**

623 **(A)** Non-metric Multidimensional Scaling based on delivery mode. **(B)** Shannon diversity between
624 delivery modes. **(C)** Stacked barplots of the relative abundances of the top 10 genera of puppies grouped
625 by delivery mode. **(D)** Boxplots of differential relative abundance according to the mode of delivery,
626 showing the top four significantly different genera (Wilcoxon rank-sum test, B-H adjusted p-values).

627
628 **Fig. 6. Gut Microbiome Dynamics in Mother Dogs Across Reproductive States.**

629 **(A)** Distribution of alpha diversities within mother dogs throughout normal state, pregnancy and
630 lactation. **(B)** Stacked barplots of relative abundances in the pre- and postpartum days shows a temporary
631 shift in the microbiome.

632

633 **Table 1. Study Population Summary by Kennel, Including Litter Delivery Type, Puppy Sex**

634 **Distribution, Maternal Age, Adult Counterparts, and Total Sample Counts.**

Kennel ID	Number of litters	Number of puppies	Puppies ♀	Puppies ♂	Number of unique mother (s)	Mothers' age years (mean ± SD)	Adult female counterparts	Adult male counter-parts	Adults' age years (mean ± SD)	Samples per kennel
BCS	2	7	3	4	2	3.9 ± 1.6	2	0	3.2 ± 2.5	61
DMD	4	20	13	7	2[a]	4.5 ± 3.2	0	1	6.8	114
SRT	2	4	1	3	1[b]	4.2	3	4	6.2 ± 1.9	80
PPA	1	7	2	5	1	6.0	4	3	7.9 ± 4.6	46
RRT	2	12	5	7	2	4.0 ± 1.8	1	1	1.8 ± 0.1	100
LPL	1	5	3	2	1[c]	4.1	0	1	1.8	45
Total	**12**	**55**	**27**	**28**	**9**	**4.3 ± 1.6**	**10**	**10**	**5.9 ± 3.6**	**446**

635

636 **a.** Indicates that both mothers gave birth twice during the study period. **b.** Refers to a case where the

637 second mother was not sampled during gestation/lactation and is therefore excluded from the maternal

638 age calculation**.** **c.** Denotes that this mother later delivered an additional litter by cesarean section, which

639 is summarized separately in **Table 1B**. The Mean age of adult control dogs was calculated at the time of

640 first sample collection. For mother dogs, age was calculated at the time of parturition. Kennel IDs: BCS

641 = Bükki Cserfes; DMD = Duna-menti Dumás; LPL = Le Petit Lapin; PPA = Pattogó Parázs; RRT =

642 Rezerta-Reti; SRT = Serteperti

643
644
645

646 **Table 2. Summary of the cesarean section (C-section) delivered puppy litter from Kennel Le Petit**

647 **Lapin (LPL).**

648

Kennel ID	Number of litters	Number of puppies	Puppies ♀	Puppies ♂	Mother ID	Samples per Puppy	Total samples
LPL	1	5	0	5	Same as LPL 1[c]	2	10

649

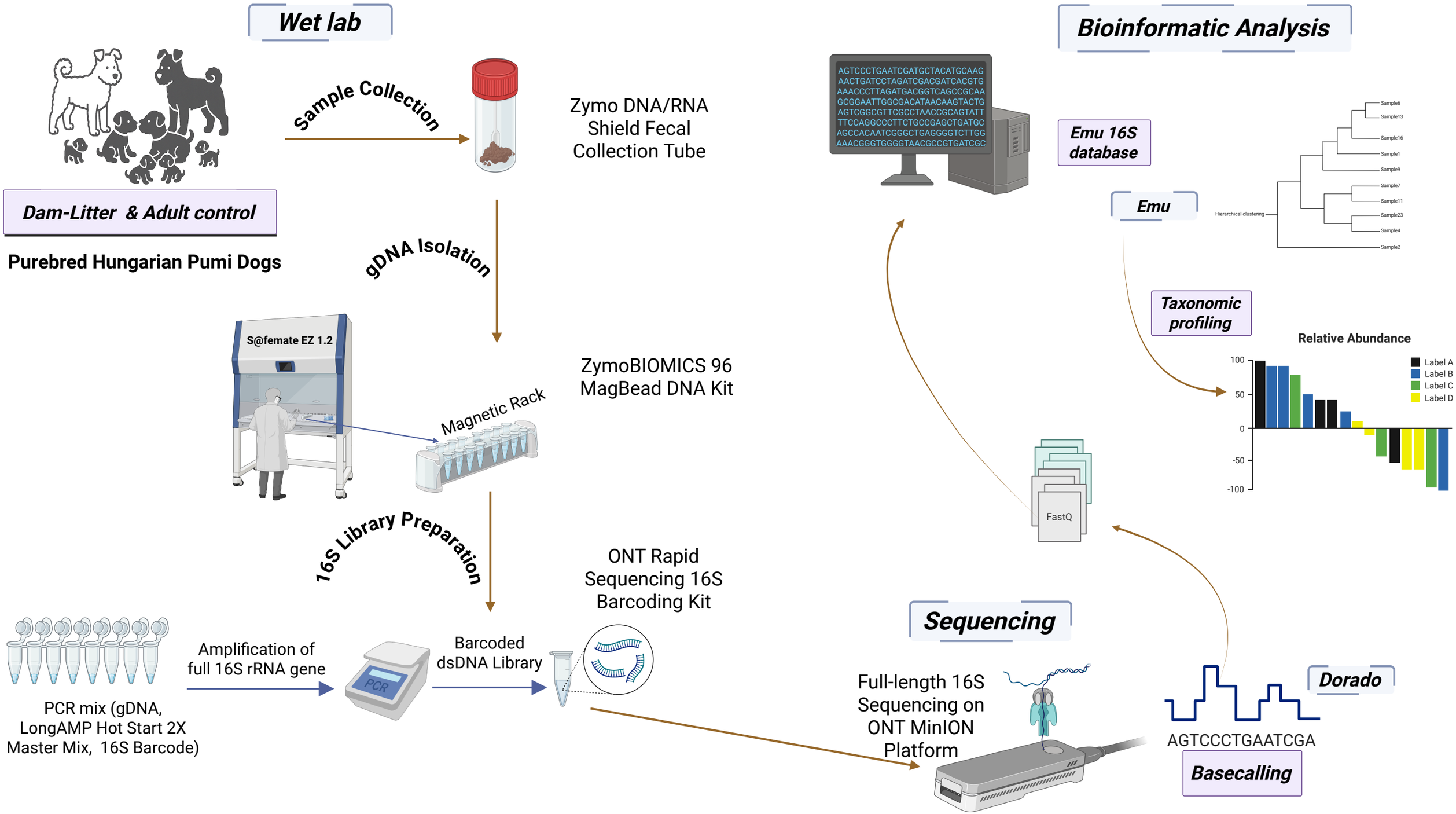
650 This litter was delivered via cesarean section by the same mother who also delivered a vaginal litter

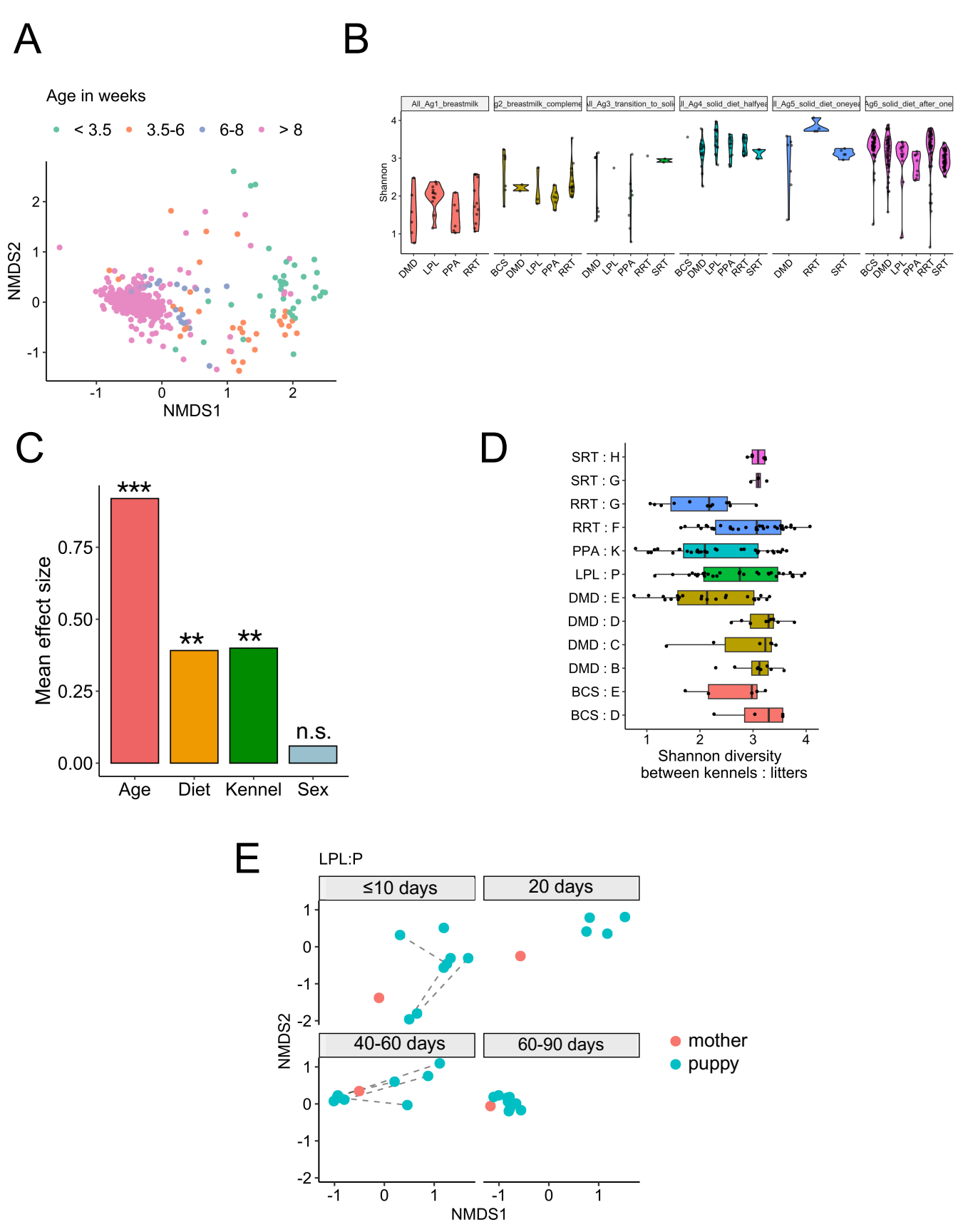
651 included in Table 1 (see footnote c). These samples were analyzed separately and were excluded from

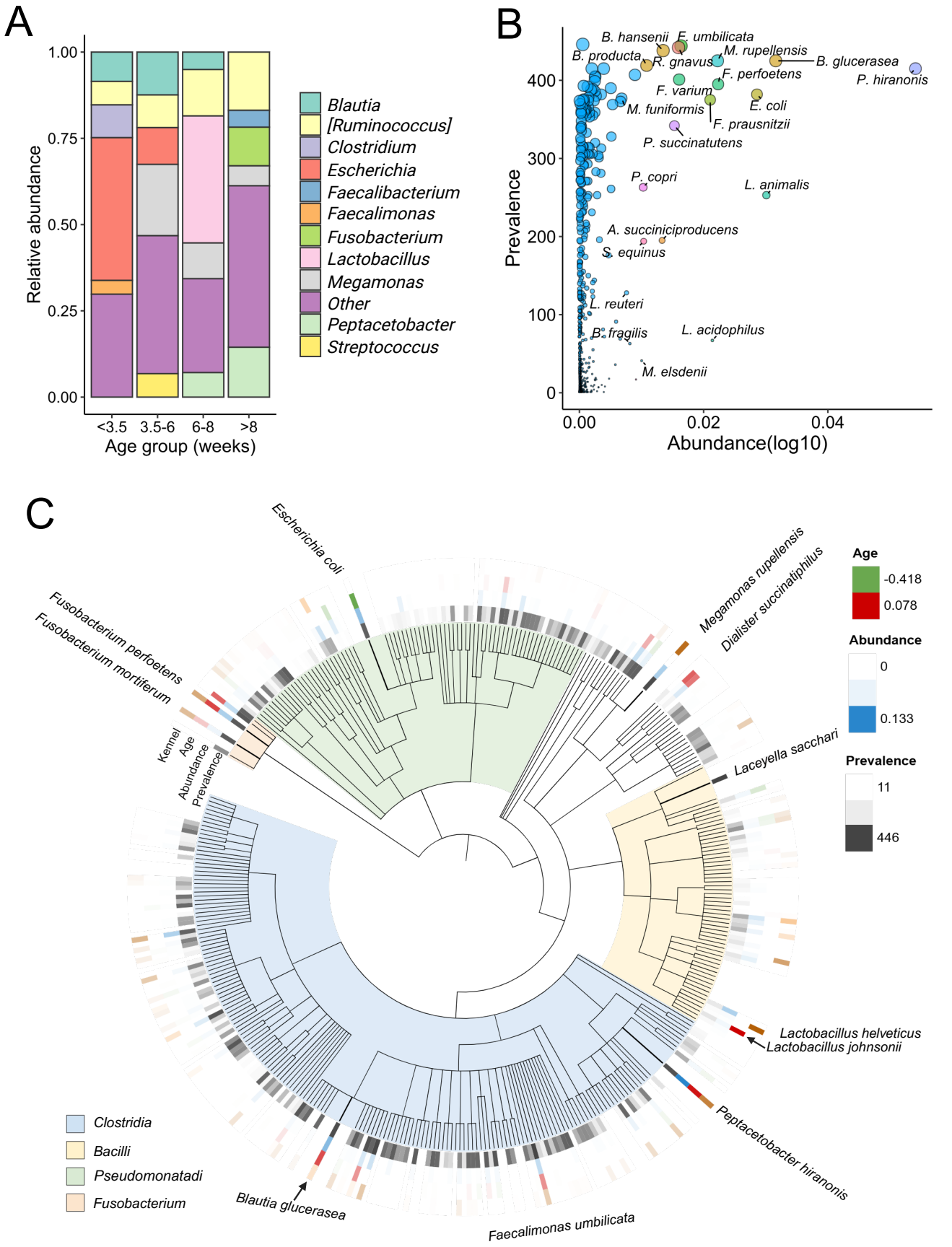
652 the main cohort analysis presented in Table 1.

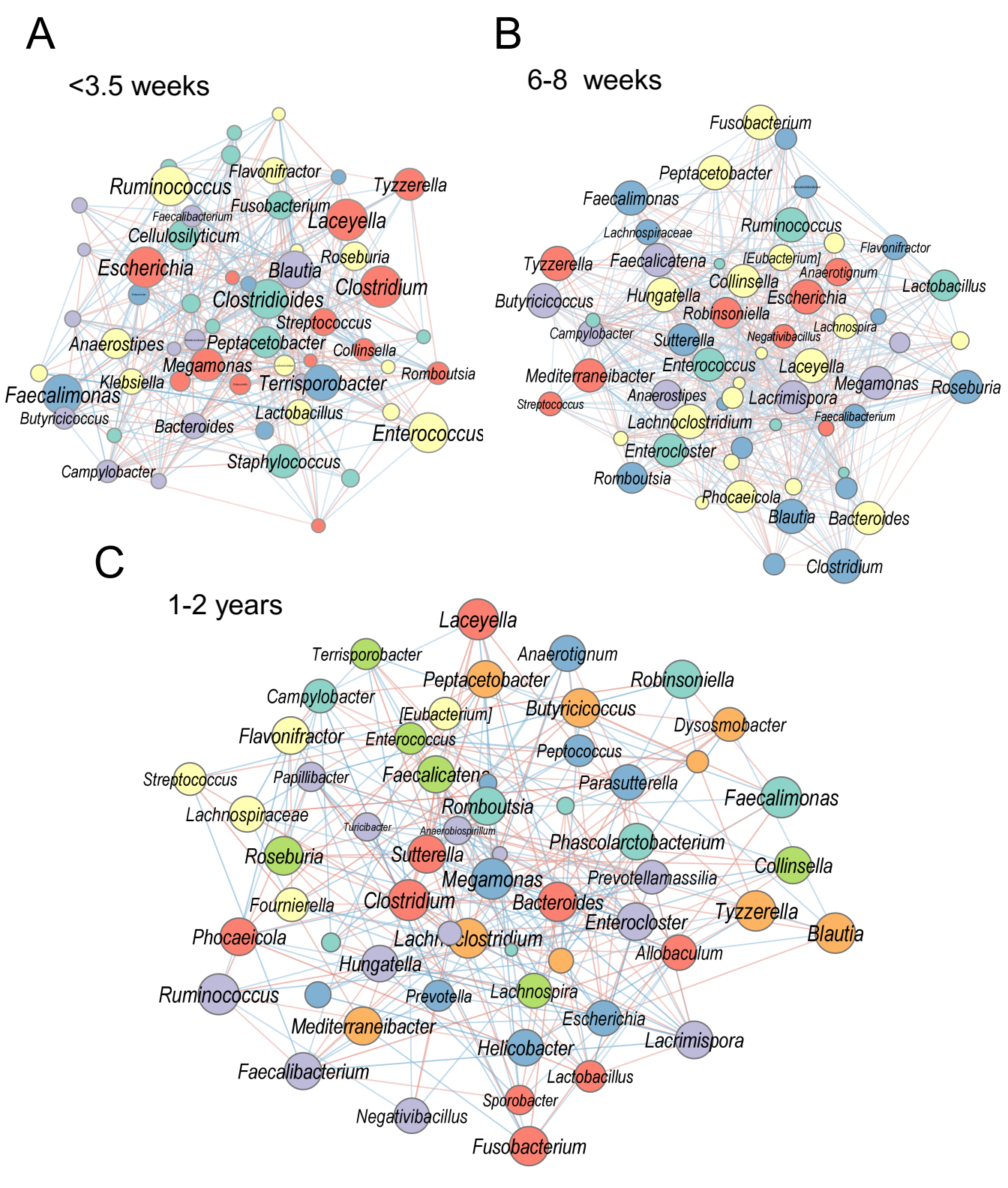
653

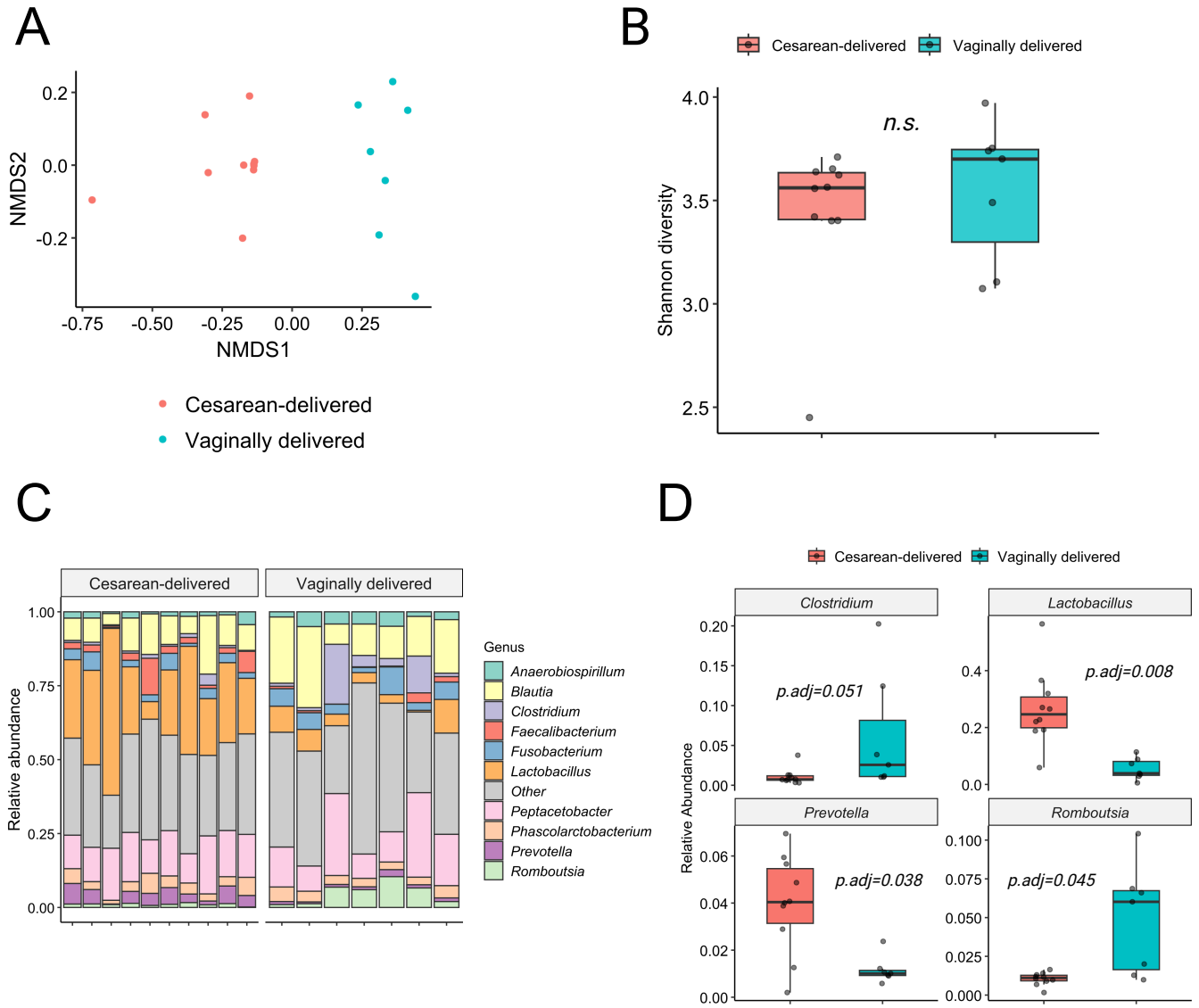
654

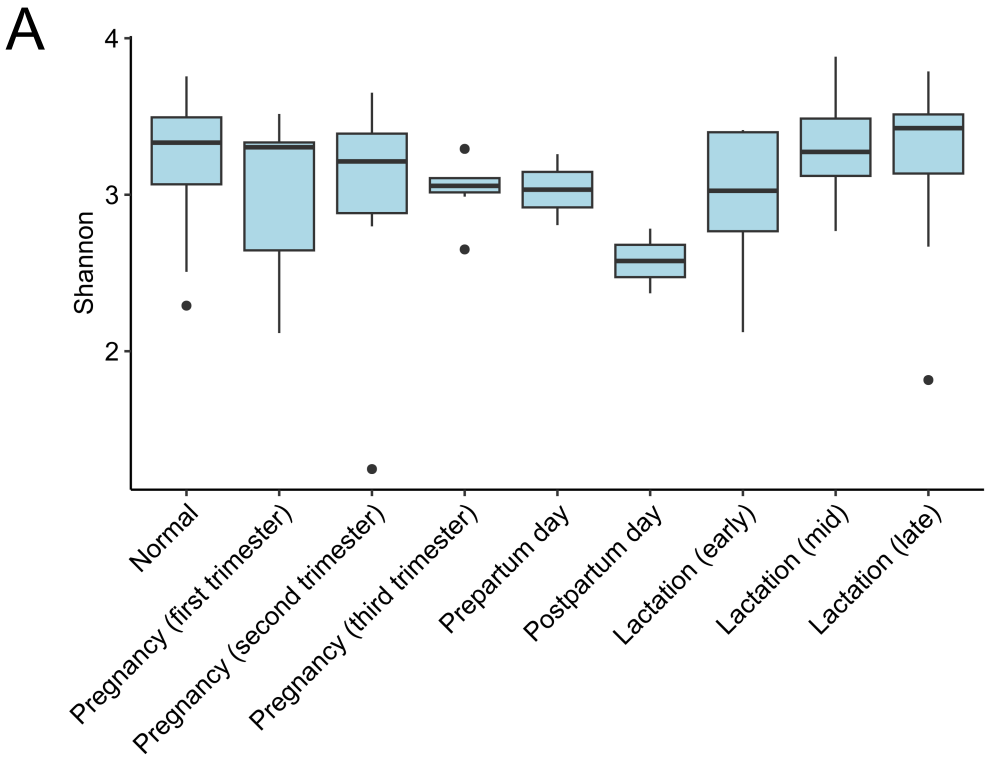

Wet lab
Dam-Litter & Adult control
Purebred Hungarian Pumi Dogs
Sample Collection
Zymo DNA/RNA Shield Fecal Collection Tube
gDNA Isolation
S@femate EZ 1.2
Magnetic Rack
ZymoBIOMICS 96 MagBead DNA Kit
16S Library Preparation
ONT Rapid Sequencing 16S Barcoding Kit
PCR mix (gDNA, LongAMP Hot Start 2X Master Mix, 16S Barcode)
Amplification of full 16S rRNA gene
PCR
Barcoded dsDNA Library
Sequencing
Full-length 16S Sequencing on ONT MinION Platform
Dorado
AGTCCCTGAATCGA
Basecalling
FastQ
Bioinformatic Analysis
Emu 16S database
Emu
Hierarchical clustering
Taxonomic profiling
Relative Abundance
Label A
Label B
Label C
Label D

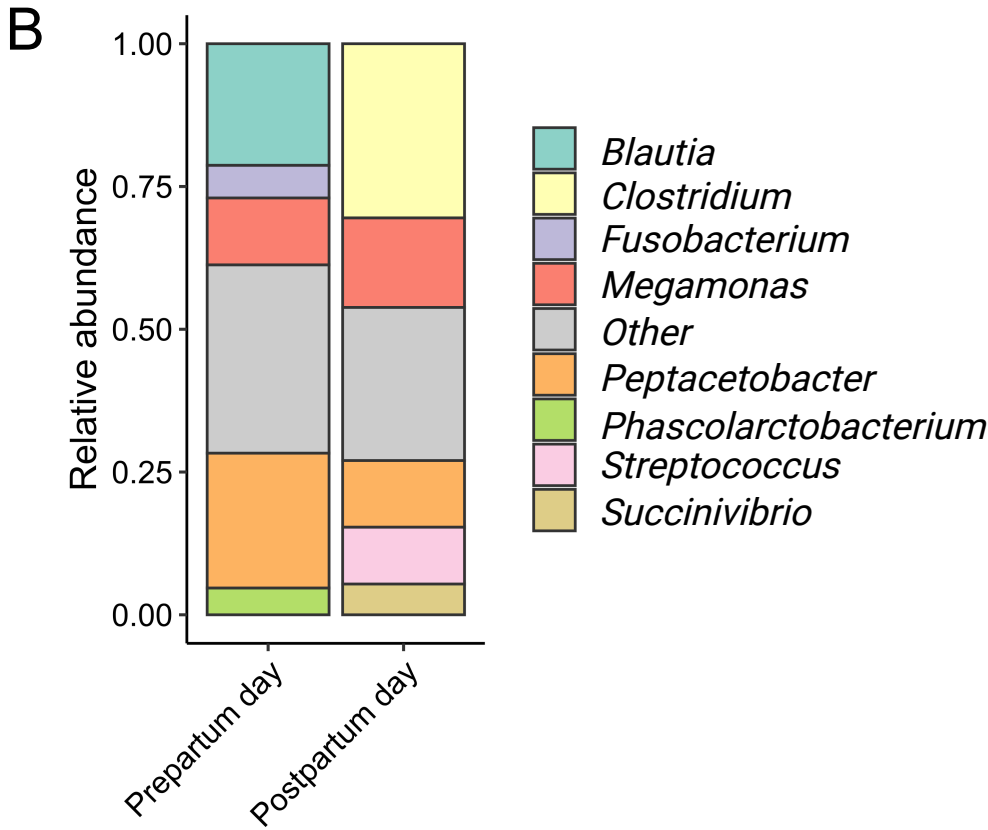

A
Age in weeks
< 3.5
3.5-6
6-8
> 8
NMDS2
NMDS1
B
All_Ag1_breastmilk
g2_breastmilk_compleme
ll_Ag3_transition_to_soli
ll_Ag4_solid_diet_halfyea
ll_Ag5_solid_diet_oneyea
Ag6_solid_diet_after_one
Shannon
DMD
LPL
PPA
RRT
BCS
SRT
C
Mean effect size
***
**
**
n.s.
Age
Diet
Kennel
Sex
D
SRT : H
SRT : G
RRT : G
RRT : F
PPA : K
LPL : P
DMD : E
DMD : D
DMD : C
DMD : B
BCS : E
BCS : D
Shannon diversity
between kennels : litters
E
LPL:P
≤10 days
20 days
40-60 days
60-90 days
NMDS2
NMDS1
mother
puppy


A
Relative abundance
1.00
0.75
0.50
0.25
0.00
<3.5
3.5-6
6-8
>8
Age group (weeks)
Blautia
[Ruminococcus]
Clostridium
Escherichia
Faecalibacterium
Faecalimonas
Fusobacterium
Lactobacillus
Megamonas
Other
Peptacetobacter
Streptococcus
B
Prevalence
Abundance(log10)
B. hansenii
F. umbilicata
M. rupellensis
B. producta
R. gnavus
B. glucerasea
P. hiranonis
F. perfoetens
F. varium
E. coli
M. funiformis
F. prausnitzii
P. succinatutens
P. copri
L. animalis
A. succiniciproducens
S. equinus
L. reuteri
B. fragilis
L. acidophilus
M. elsdenii
C
Escherichia coli
Fusobacterium perfoetens
Fusobacterium mortiferum
Kennel
Age
Abundance
Prevalence
Megamonas rupellensis
Dialister succinatiphilus
Laceyella sacchari
Lactobacillus helveticus
Lactobacillus johnsonii
Peptacetobacter hiranonis
Faecalimonas umbilicata
Blautia glucerasea
Clostridia
Bacilli
Pseudomonatadi
Fusobacterium
Age
-0.418
0.078
Abundance
0
0.133
Prevalence
11
446


A
<3.5 weeks
Flavonifractor
Ruminococcus
Tyzzerella
Fusobacterium
Faecalibacterium
Laceyella
Cellulosilyticum
Roseburia
Escherichia
Blautia
Clostridium
Clostridioides
Streptococcus
Anaerostipes
Peptacetobacter
Collinsella
Megamonas
Romboutsia
Klebsiella
Terrisporobacter
Faecalimonas
Butyricicoccus
Lactobacillus
Bacteroides
Enterococcus
Staphylococcus
Campylobacter
B
6-8 weeks
Fusobacterium
Peptacetobacter
Faecalimonas
Lachnospiraceae
Ruminococcus
Flavonifractor
Tyzzerella
Faecalicatena
[Eubacterium]
Anaerotignum
Collinsella
Lactobacillus
Butyricicoccus
Hungatella
Escherichia
Robinsoniella
Lachnospira
Campylobacter
Sutterella
Negativibacillus
Enterococcus
Laceyella
Mediterraneibacter
Megamonas
Roseburia
Anaerostipes
Lacrimispora
Streptococcus
Lachnoclostridium
Faecalibacterium
Enterocloster
Romboutsia
Phocaeicola
Blautia
Bacteroides
Clostridium
C
1-2 years
Laceyella
Terrisporobacter
Anaerotignum
Peptacetobacter
Robinsoniella
Campylobacter
Butyricicoccus
[Eubacterium]
Dysosmobacter
Flavonifractor
Enterococcus
Peptococcus
Streptococcus
Papillibacter
Faecalicatena
Parasutterella
Faecalimonas
Lachnospiraceae
Romboutsia
Turicibacter
Anaerobiospirillum
Phascolarctobacterium
Roseburia
Sutterella
Collinsella
Megamonas
Prevotellamassilia
Fournierella
Clostridium
Bacteroides
Tyzzerella
Enterocloster
Blautia
Phocaeicola
Lachnoclostridium
Allobaculum
Hungatella
Ruminococcus
Prevotella
Lachnospira
Escherichia
Mediterraneibacter
Lacrimispora
Helicobacter
Faecalibacterium
Lactobacillus
Sporobacter
Negativibacillus
Fusobacterium


A
NMDS2
NMDS1
0.2
0.0
-0.2
-0.75
-0.50
-0.25
0.00
0.25
Cesarean-delivered
Vaginally delivered
B
Cesarean-delivered
Vaginally delivered
Shannon diversity
4.0
3.5
3.0
2.5
n.s.
C
Cesarean-delivered
Vaginally delivered
Relative abundance
1.00
0.75
0.50
0.25
0.00
Genus
Anaerobiospirillum
Blautia
Clostridium
Faecalibacterium
Fusobacterium
Lactobacillus
Other
Peptacetobacter
Phascolarctobacterium
Prevotella
Romboutsia
D
Cesarean-delivered
Vaginally delivered
Clostridium
Lactobacillus
Prevotella
Romboutsia
Relative Abundance
p.adj=0.051
p.adj=0.008
p.adj=0.038
p.adj=0.045


A
Shannon
4
3
2
Normal
Pregnancy (first trimester)
Pregnancy (second trimester)
Pregnancy (third trimester)
Prepartum day
Postpartum day
Lactation (early)
Lactation (mid)
Lactation (late)


B
Relative abundance
1.00
0.75
0.50
0.25
0.00
Prepartum day
Postpartum day
Blautia
Clostridium
Fusobacterium
Megamonas
Other
Peptacetobacter
Phascolarctobacterium
Streptococcus
Succinivibrio