Evolution of the murine gut resistome following broad-spectrum antibiotic treatment

Laura de Nies1,3, Susheel Bhanu Busi1,3, Mina Tsenkova2, Rashi Halder1, Elisabeth Letellier2✉ & Paul Wilmes1,2✉

The emergence and spread of antimicrobial resistance (AMR) represent an ever-growing healthcare challenge worldwide. Nevertheless, the mechanisms and timescales shaping this resistome remain elusive. Using an antibiotic cocktail administered to a murine model along with a longitudinal sampling strategy, we identify the mechanisms by which gut commensals acquire antimicrobial resistance genes (ARGs) after a single antibiotic course. While most of the resident bacterial populations are depleted due to the treatment, *Akkermansia muciniphila* and members of the Enterobacteriaceae, Enterococcaceae, and Lactobacillaceae families acquire resistance and remain recalcitrant. We identify specific genes conferring resistance against the antibiotics in the corresponding metagenome-assembled genomes (MAGs) and trace their origins within each genome. Here we show that, while mobile genetic elements (MGEs), including bacteriophages and plasmids, contribute to the spread of ARGs, integrons represent key factors mediating AMR in the antibiotic-treated mice. Our findings suggest that a single course of antibiotics alone may act as the selective sweep driving ARG acquisition and incidence in gut commensals over a single mammalian lifespan.

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Prior to the advent of antibiotics, bacterial infections were the leading cause of disease and mortality in humans. Antibiotic usage is now commonplace in treating infections, as well as ensuring the safety of surgical procedures and organ transplantation. In addition, they are extensively used in animal husbandry and also in animal models for studying the gut microbiome. Concomitantly, their prevalence and administration are widespread in developing and developed countries alike, whereby antibiotics are easily accessible. However, many bacterial taxa have evolved antimicrobial resistance (AMR) to several classes of antibiotics, and multidrug-resistant bacteria have now emerged, preventing the comprehensive treatment of infections and resulting in a growing number of deaths. Due to the overall rise in resistance, as well as a lack in the development of new antibiotics, AMR has emerged as a growing global threat to human health, whereby antibiotics are easily accessible. 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Fig. 1 Experimental design and metagenome-assembled genome profiles. a Representative illustration demonstrating the project overview, including dates and collections of treatment, and fecal sample collection. Eight single-housed mice per group were treated with an antibiotic cocktail (treatment) or water (control), longitudinally. Fecal samples collected at days 0, 7, and 21 were used for microbiome and antimicrobial resistance profiling. b Genus level representation of the MAGs recovered from control and treatment groups – pre and post (day 0, day 7, and day 21 respectively) antibiotic administration. Source data are provided as a Source Data file.
Fig. 2 Resistome in antibiotic-treated mice. a Overall ARG relative abundance levels are shown for each group (n = 8 biological replicates). *represents significance with an adjusted p-value less than 0.05 as assessed using a Wilcoxon rank-sum test. The center line denotes the median value (50th percentile), while the outer lines of the box represent the 25th to 75th percentiles. The black whiskers mark the 5th and 95th percentiles. 

b Significantly differentially abundant AMR categories found to be enriched in the mice treated with antibiotics compared across different timepoints, i.e., day 0, day 7, and day 21, *adjusted p-value < 0.05 (Wilcoxon rank-sum test). n = 8, biological replicates per group. The center line denotes the median value (50th percentile), while the outer lines of the box represent the 25th–75th percentiles. The black whiskers mark the 5th and 95th percentiles. 

c Barplots showing the relative abundance of MAGs (Family level) associated with ARGs in each sample. 

d Relative abundance of ARGs associated with Akkermansiaceae, Enterococcaceae, and Lactobacillaceae in the control and treated mice (n = 8 biological replicates per group). *adjusted p-value < 0.05 (Wilcoxon rank-sum test). The center line denotes the median value (50th percentile), while the outer lines of the box represent the 25th–75th percentiles. The black whiskers mark the 5th and 95th percentiles. Significance for all analyses was assessed using a two-sided Wilcoxon rank sum test, where, p-values are indicated by *, i.e., * < 0.05, ** < 0.01, *** < 0.001. Source data are provided as a Source Data file.
Antibiotic-induced changes in taxonomic composition. Since the metagenomes revealed an enrichment in different ARG categories, we investigated taxa harboring these ARGs. We linked ARGs to individual genomes by identifying contigs encoding ARGs and their corresponding assignment to MAGs including taxonomic classification of the MAGs using GTDBtk. Based on the MAGs, we subsequently compared taxa contributing to AMR between the groups, including mice treated with antibiotics and those without. Interestingly, we did not recover any MAGs at day 7, likely due to the antibiotic-mediated depletion of the microbiota. However, in contrast to the MAGs, based on operational taxonomic units (OTUs), at day 7 we detected taxa that consisted predominantly of Bacteroidales spp. (Supplementary Fig. 2d). While taxa contributing to AMR within the control group remained constant, a shift in ARG-encoding taxa was observed within the treatment group after recovery, at day 21 (Fig. 2c). Alongside the increase in the abundance of several taxa (Fig. 2c), we found that the abundance of overall ARGs was increased in taxa belonging to the Akkermansiae, Enterococccaceae, and Lactobacillaceae families across all treated mice, as well as compared to the control group, at day 21 (Fig. 2d). Given the enrichment in ARGs at day 21, as opposed to days 0 and 7, and the specific lack of Akkermansiae, Enterococccaceae, and Lactobacillaceae MAGs at day 7, it is likely that the observed ARGs were acquired over time, rather than being encoded as intrinsic resistance mechanisms.

MGEs linked to AMR dissemination. MGEs are an established mechanism for the dissemination of AMR. To determine the function of MGEs in conferring the resolved ARGs under selective pressure, we analyzed the genomic context of the ARGs. The majority of the resistance genes were encoded on the bacterial chromosome (Fig. 3a). To assess the role of bacteriophages and plasmids in AMR transmission, we compared two strategies, where in the first approach, PathoFact was used to predict AMR on phages and plasmids obtained via metaviralSPAdes and metaviralspAdes, respectively. In the second approach, PathoFact was used to identify both MGEs and ARGs from the metagenome assemblies obtained via IMP (“Methods”). We found that the standalone PathoFact detected more MGEs including those associated with AMR, compared to the SPAdes-based approach (Supplementary Fig. 3c). Furthermore, our analyses revealed that ARGs were encoded on both phages and plasmids (Fig. 3a). Interestingly, a reduction in the general abundance of ARGs mediated via plasmids was observed at day 7 in the treated mice compared to the controls, but the overall levels recovered by day 21. In contrast, phages linked to AMR were significantly enriched in the treated mice at day 7, compared to both pre-treatment at day 0 and the control mice at day 7 (Fig. 3a, b). Moreover, when analyzing the specific categories of the resistome, we found an increase in the abundance of phage sequences linked to aminoglycoside, aminoglycoside:aminocoumarin, and beta-lactam resistance, in conjunction with the administered antibiotic cocktail (Fig. 3c). Significant differences in these phage-associated AMR categories were observed in the treated group compared to the controls at day 7 and also within the group when comparing phage-associated ARG levels between days 0 and 7 (Fig. 3c).

To further investigate the effect of HGT on the evolution of AMR we identified and characterized all HGT events within the samples pre- (day 0) and post-treatment (days 7 and 21). Using MetaCHIP and subsequent manual analyses (“Methods”) verifying the full-length matches for horizontally-transferred genes, we assessed whether ARGs were transferred across MAGs within the timepoints. We did not observe differences in the overall number of HGT events between the control and treatment groups at all timepoints (Fisher’s exact test). However, to further assess whether HGT contributes to ARG spread, HGT events were specifically linked to AMR genes. We did not find a significant correlation between AMR and HGT in the control and treated mice across all timepoints, although Akkermansia spp. was involved in AMR-associated HGT events in 6/8 (i.e., 75%) of the mice treated with antibiotics (Supplementary Fig. 4a). Albeit not statistically significant (Fisher’s Exact test, p > 0.1), Akkermansia was only involved in AMR-associated HGT in 3/8 (i.e., 37.5%) of the control mice (Supplementary Fig. 4b).

Integrons mediate AMR in antibiotic-treated mice. To further investigate the effect of antibiotic treatment on the evolution of AMR within the microbiota, we assessed the panogenomes of the significantly enriched and recalcitrant taxa in the treated mice, i.e., Akkermansia muciniphila and Lilgulactobacillus spp. Interestingly, panogenome analyses of Akkermansia muciniphila revealed the acquisition of several genes at day 21 compared to day 0, including those mediated by integrases (Fig. 4a).

Horizontal gene transfer is typically attributed to phages and plasmids in metagenomes. However, integrons, often overlooked, play a key role in AMR dissemination and prevalence. To evaluate the role of integrons in AMR, we assessed the abundance of attC sites and intI genes, both of which are required for efficient integron-mediated activity. We estimated the abundance of these genes on the same contig, including those that were associated with AMR categories. Overall, we found that ARGs abundant in antibiotic-treated mice were transferred via integrons (Fig. 4b and Supplementary Fig. 5a). Of these, there was a significant enrichment (adjusted p-value < 0.05, Wilcoxon rank-sum tests) in ARGs associated with complete (presence of attC and intI genes) integrons in mice at day 21 compared to day 0 and also when compared to the controls (Fig. 4b).

Additionally, these integron-mediated ARGs (complete, gene cassettes, and incomplete) were further analyzed to identify their putative genomic locations on phages or plasmids, since they are known to be carriers of integrons, thus elaborating on the method of integron-mediated AMR transmission. Interestingly, we identified several integron-mediated ARG cassettes encoded on plasmids at day 21 in the antibiotic-treated mice (Fig. 4c and Supplementary Fig. 5b).

As we identified antibiotic-induced changes of the microbial composition, we further investigated the association of AMR-encoding integrons with the microbial community. For this, we linked the AMR-associated ‘complete’ integrons with the reconstructed genomes and found that a substantial number was associated with genomes from families including Akkermansiae, Lachnospiraceae, and Enterobacteriaceae (Fig. 4c and Supplementary Fig. 5c). This finding reinforces our earlier findings with respect to enriched taxa and potential ARG-mediated mechanisms of resistance through integrases.

Discussion
The classes and uses of antibiotics have been extensively developed since the fateful discovery of “mold juice” by Alexander Fleming. Their use, and their overuse, has led to unrecoverable and irreversible states of resistance, resulting in an “arms race” where newer and more potent molecules are becoming a necessity to ward off otherwise-susceptible bacteria. Even though antibiotics may result in the emergence of multi-resistant pathogens, their expanding use in medicine, especially
as a means of modulating the gut microbiome, remains unquestionable. For example, they have also been proposed as prophylactics for treating cancers and modulating the gut microbiota. Antibiotics not targeting Clostridioides difficile infection are commonly used within the first weeks of a fecal microbiota transplantation (FMT) as a standard therapy. Similarly, preoperative antibiotic prophylaxis in humans is a common practice and typically involves three antibiotics (cefazolin, vancomycin, and gentamicin). These are usually administered individually while vancomycin in combination with other antibiotics (e.g., cefazolin) has been proposed for treatment of methicillin-resistant Staphylococcus aureus. Here, we hypothesized that antibiotic treatment would lead to an evolution of AMR in the commensal gut microbiome.

**Fig. 3 Abundance levels of resistome categories.**

- **a** Relative abundance of AMR encoded on the bacterial chromosome and those mediated via MGEs, such as bacteriophages (phages) and plasmids. n = 8 biological replicates per group. The center line denotes the median value (50th percentile), while the outer lines of the box represent the 25th–75th percentiles. The black whiskers mark the 5th and 95th percentiles. Significance was assessed using a two-sided Wilcoxon rank sum test, where, p-values are indicated by *, i.e., * < 0.05, ** < 0.01, *** < 0.001.

- **b** Abundance levels of AMR categories disseminated via phages and plasmids. Categories pre- and post-treatment (day 0 and day 21 respectively).

- **c** Abundance levels of aminoglycoside, aminoglycoside:aminocoumarin, and beta-lactam resistance genes mediated via phages in the control and treated mice (n = 8 biological replicates).

The center line denotes the median value (50th percentile), while the outer lines of the box represent the 25th–75th percentiles. The black whiskers mark the 5th and 95th percentiles. Source data are provided as a Source Data file.
population within a single animal lifespan and tested our hypothesis in a wild-type mouse cohort.

The antibiotics (ampicillin, vancomycin, metronidazole, and neomycin) were chosen given their utility in several mouse models and in varying combinations in line with some clinical procedures. We observed that Akkermansia muciniphila was significantly enriched whilst most taxa were depleted in mice post-treatment, which is in line with other reports where vancomycin treatment alone led to propagation of A. muciniphila or its dominance in the resistant commensal population. The resistance of this taxon can specifically be attributed to the presence of β-lactamase and nitroimidazole resistance genes reported by van Passel et al. These findings also agree with the report by Palleja et al. in which the authors found that species harboring...
β-lactam resistance genes were positively selected during antibiotic exposure\(^{45}\), which is likely the case in our study since we observed higher ARG abundances at day 21 and not prior to antibiotic treatment. Alternatively, the observed ARGs could also be due to the acquisition of resistance genes possibly via lateral or horizontal gene transfer as reported previously by Guo et al.\(^{45}\).

In addition to an overall enrichment in *A. muciniphila*, we also observed an enrichment in the functional complement of *A. muciniphila* with respect to signaling molecules, specifically quorum sensing and cyclic dinucleotide signaling. Microbial communities are characterized by emergent properties that themselves are primarily shaped by microbial interactions\(^{46}\). These interactions include intra- and extracellular signaling, such as quorum-sensing (QS) and cyclic dinucleotide sensing, as a means of adapting to internal and external stimuli\(^{47}\). Due to the paucity of external stimuli from other bacteria following antibiotic treatment, it is plausible that QS functions were selected for as a means for recalcitrant community members to ramp up signaling functions to induce antibiotic tolerance\(^{48}\), contributing towards AMR, which may be mediated via QS molecules leading to bacterial resistance gene expression in a density-dependent manner\(^{48}\). Furthermore, QS molecules have also been reported to regulate secondary metabolite synthesis\(^{49}\). However, we observed a depletion in genes involved in secondary metabolite synthesis following antibiotic treatment which is expected since a majority of the endogenous population is depleted. Alternatively, this phenomenon also suggests that selection for genes involved in signaling and secondary metabolite synthesis are somewhat uncoupled in our experimental murine model and, thus, are subject to different selective sweeps. Overall, our results highlight the role of key functions conferred by specific bacterial taxa in antibiotic-exposed communities and shed light on the shorter-term evolutionary processes shaping community assembly and composition.

Given the nature of the antibiotic cocktail treatment, we found several related ARGs in the metagenomes of the treated mice. More importantly, we observed significantly increased resistance at day 21, against three out of the four antibiotics used in our study protocol: *aminoglycoside* (neomycin), *beta-lactam* (ampicillin), and *glycopeptide* (vancomycin). We, however, did not recover any resistance genes against nitroimidazoles (metronidazole). Additionally, we found that several taxa in mice treated with antibiotics were directly linked to the resistance categories of the antibiotics that they were treated with. This suggests that the selective pressure of the administered antibiotics may lead to real-time evolution of AMR within the gut microbiome. This is in line with a recent report by Xu et al.\(^{50}\) albeit in a different mouse time evolution of AMR within the gut microbiome. This is in line with a recent report by Xu et al.\(^{50}\) albeit in a different mouse standard habitat conditions (humidity: 40–70%, temperature: 22 °C) with a 12:12 light cycle. Two groups of mice were established (control and treatment), and each group contained 8 animals (4 males + 4 females). Antibiotics, ampicillin (1 g/L), vancomycin (500 mg/L), metronidazole (1 g/L), and neomycin (1 g/L) were chosen for their utility in several mouse models\(^{37,58}\) and in line with some preoperative procedures\(^{58}\). They were administered as a cocktail within the drinking water to the treatment group starting at 8 weeks of age. Antibiotics were administered during a period of one week, after which the change was made to regular drinking water for the duration of the recovery period. Fecal samples were collected prior to treatment and subsequently daily for a duration of 19 days (both treatment and recovery phase) starting prior to the antibiotic treatment till take down (Fig. 1a).

**Methods**

### Ethical considerations

The animal experiment was performed according to all applicable laws and regulations, after receiving approval by the institution’s Animal Experimentation Ethics Committee at UL (AEEC) and the Ministry of Agriculture, Verniculture and Rural Development (LUPA 2019/99). The care and use of animals for research purposes was conducted according to the EU Directive 2010/63/EU, as well as the Grand-Ducal Regulation of 11 January 2013 on the protection of animals used for scientific purposes. These included the justification of the use of animals, their welfare and the incorporation of the principles of the 3Rs (Replacement, Reduction and Refinement).

### Power calculation and sample size estimation

To determine the number of animals required per treatment and control group we performed a multifactorial power analysis based on a 2015 study by Raymond et al.\(^{52}\). For this, we estimated the Jensen–Shannon Divergences (JSD) of the microbial profiles of the antibiotic-treated (cefprozil) and control groups. Based on the observed JSD, the inter-group variability was significantly high, thereby highlighting a minimum sample size of three mice per group to attain a power of 80 and a 5% alpha error rate, reflecting changes in microbial composition (Supplementary Fig. 6).

### Mouse model and antibiotic exposure

Eight-week old C57BL/6J mice were purchased from Charles River Laboratories and experiments were performed according to all applicable laws and the regulations described under Ethical considerations. To limit individual variation of the gut microbiome in experimental groups, mice of the same age (8 weeks) were obtained from the same vendor and the same location in the vendor facility. After a 7-day quarantine and subsequent acclimation period of one week, mice were maintained in single housing conditions. Mice were housed in Allentown NexGen Mouse 500 (194 mm × 130 mm × 381 mm) cages (Allentown, USA) with RIS Relohol Corncob bedding. Mice had access to reverse osmosis water with 2 ppm of chlorine fed ad libitum along with standard AIN-93G chow diet (SAFE, France). The animals were maintained under standard conditions (humidity: 40–70%, temperature: 22 °C) with a 12:12 light cycle. Two groups of mice were established (control and treatment), and each group contained 8 animals (4 males + 4 females). Antibiotics, ampicillin (1 g/L), vancomycin (500 mg/L), metronidazole (1 g/L), and neomycin (1 g/L) were chosen for their utility in several mouse models\(^{37,58}\) and in line with some preoperative procedures\(^{58}\). They were administered as a cocktail within the drinking water to the treatment group starting at 8 weeks of age. Antibiotics were administered during a period of one week, after which the change was made to regular drinking water for the duration of the recovery period. Fecal samples were collected prior to treatment and subsequently daily for a duration of 19 days (both treatment and recovery phase) starting prior to the antibiotic treatment till take down (Fig. 1a).

### Fecal processing and nucleic acid extraction

A total of 48 fecal samples were obtained across three timepoints, i.e., prior to treatment: day 0, immediately after treatment: day 7, and after recovery: day 21, from each of the mice. 50 mg of frozen stool samples were aseptically weighed into sterile vials. Genomic DNA was isolated with the DNeasy PowerSoil Kit (Qiagen, USA) including an additional incubation step at 65 °C and milling, as described previously\(^{59}\). A minimum of 200 ng of total DNA was recovered from all the samples, yielding sufficient DNA for metagenomic sequencing including high-resolution, artefact-curated metagenomic data for subsequent analyses\(^{35}\). DNA extracted from all timepoints was thereafter stored at −80 °C until further use.

### DNA sequencing

All DNA samples were subjected to random shotgun sequencing. Briefly, 200 ng of DNA was used for metagenomic library preparation using the Westburg NGS DNA Library Prep Kit (cat. no. WB 9096, Westburg Life...
Integrated Meta-omic Pipeline (IMP; v3 - commitID #6f1badf7)55 was used to process paired forward and reverse reads using the built-in metagenomic workflow as previously described36. The workflow includes pre-processing, assembly, genome reconstruction, and functional annotation of genes based on custom databases in a reproducible manner. After trimming the adapters, the preprocessing step included the removal of Mus musculus (GRCm38.p6 (GCA_000001635.8), retina) from 16-Mus_musculus ftp://www.ensembl.org/Mus_musculus/Info/Index) reads. Thereafter the de novo assembly was performed using the MEGAHIT (version 2.0) assembler37. Default IMP parameters were retained for all samples. Metagenomic operational taxonomic unit (mOTU) profiles were generated from the trimmed and preprocessed reads to generate microbiome profiles for the control and treatment groups using mOTUs v2.5.136. Concurrently, we used MetaBAT59 and MaxBin260 for binning in addition to an in-house binning methodology previously described36 for genome reconstructions, i.e., MAGs. Subsequently, we obtained a non-redundant set of MAGs using DASTool v1.1.461 with a score threshold of 0.7 for downstream analyses, and those with a minimum completeness of 90% and lower was as assessed by CheckM v1.1.3.2. Taxonomy was assigned to the MAGs using the extensive database packaged with gtdbtk v1.7.0.41. To generate pangenomes, we collected all the bins taxonomically identified as Akkermansia muciniphila and used the anvio-based pangenome workflow described by Eren et al. (http://merenlab.org/2016/11/08/pangenome-workflow/)61. On these treatment groups (#1) was excluded from the pangenome analyses due to the unavailability of MAGs.

Identification of antimicrobial resistance genes and association with mobile genetic elements. We used PathoFact v1.0, a pipeline for the prediction of virulence factors and AMR genes, to predict and identify ARGs within our metagenomes. The assembly files from individual samples were used as input for the AMR analyses. To assess the relevance of metaplasmidSAPes and metaviruses identified from the metagenome analysis, we used the anvi’o-based pangenome workflow described by Eren et al. (http://merenlab.org/2016/11/08/pangenome-workflow/)61. On these treatment groups (#1) was excluded from the pangenome analyses due to the unavailability of MAGs.

Data processing for metagenomics, including genome reconstruction.

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Data analysis. Figures for the study including visualizations derived from the taxonomic and functional, were created using version 3.6 of the R statistical software package. GraphPad62 was used to generate the figures for describing the longitudinal weight measurements of the mice. DESeq263 and Wilcoxon rank-sum tests with FDR-adjustments for multiple testing were used to assess significant differences for the AMR and taxonomic analyses whereas a paired two-way ANOVA (Analysis of Variance) within the nime package was used for identifying statistically significant differences in the integron distributions for the HGT events were generated using scripts found within the MetaCHIP package25 while the pangenome visualizations were obtained using anvio65.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Code availability. The open-source tools and algorithms used for the data analyses are reported in the Methods section, including relevant flag used for various tools. The scripts and analysis codes are provided at https://git-r3lab.uni.lu/susheel.busi/intonate.

Data availability. The sequencing data generated for this study are available via NCBI’s SRA under the accession number: PRJNA691897. The metadata file indicating group and timepoint information can be obtained via the same accession ID. Raw counts for taxon abundances and genes generated in this study have been provided in the Source Data file. Source data are provided with this paper.

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S.B.B., E.L., and P.W. conceptualized and designed the study. M.T. performed the animal experiments including housing, antibiotic treatment, and fecal sample collection. L.d.N. and S.B.B. did the DNA extractions, analyzed the data, and created the figures. R.H. and the Sequencing Platform sequenced the samples. All authors contributed to the preparation of the manuscript.

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