

Study links specific gut bacteria to increased risk of severe malaria

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Shotgun metagenomics revealed distinct gut microbiota composition and genetic potential within and between the hyperparasitemia resistant and susceptible mice. a C57BL/6 mice were acquired from four different vendors (N =15/group): Charles River Laboratories (CR), Envigo (Env), Taconic Biosciences (Tac), and Jackson Laboratory (Jax). Mice from Taconic Biosciences were obtained from two different facilities with differential susceptibility to Py hyperparasitemia¹². Five mice from each group were infected with Py while the remaining ten mice were sacrificed to collect ceca content along with mucosa scrapes for shotgun metagenomics. b Parasitemia curve of mice infected with Py. c Number of fastq reads after shotgun sequencing and quality control. d Alpha diversity measured using observed taxonomic units (OTUs) defined at species level. e Alpha diversity measure by Shannon entropy. f Beta diversity shown by Principal coordinate analysis (PCoA) plot using Clark output at species level with Bray-Curtis distance. g Beta diversity shown by PCoA plot using Metaphlan2 output at species level with Bray-Curtis distance. h Beta diversity shown by PCoA plot using MaxBin output at species level with Bray-Curtis distance. i Beta diversity shown by PCoA plot using CZID output at bin level with Bray-Curtis distance. j Number of unique genes detected at 95% sequence homology. All data are mean \pm SE (standard error) unless explicitly stated. Bacterial diversity was performed on normalized data. Mice resistant to hyperparasitemia are encircled (f-i). Alpha diversity significance were calculated with Kruskal Wallis test and beta diversity significance by pairwise Permutational multivariate analysis of variance (PERMANOVA). *p

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