AIEC infection triggers modification of gut microbiota composition in genetically predisposed mice, contributing to intestinal inflammation

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Running head: AIEC induce gut dysbiosis, contributing to inflammation



Figure S1. Transient colonization of $eif2ak4^{+/+}$ and $eif2ak4^{-/-}$ mice with AIEC does not result in modification of the gut microbiota composition at day 1 and 4 post-infection. $eif2ak4^{+/+}$ and $eif2ak4^{-/-}$ mice were challenged by oral gavage for 3 days (once per day) with PBS (N=6 mice per group) or with 10⁹ CFU of the AIEC LF82 strain (N=7 mice per group). Feces were collected at day 1 (**A**) and 4 (**B**) post-infection for the analysis of the bacterial microbiota composition based on Illumina pyrosequencing of the 16S rRNA gene. Mouse fecal bacterial communities were clustered using PCoA of the weighted UniFrac distance matrix. PCoA-1 and PCoA-2 were plotted, and the percentage of the variation explained by the plotted principal coordinates was indicated in the Y-X-axis labels. Comparison between $eif2ak4^{+/+}$ mice + PBS, $eif2ak4^{+/+}$ mice + LF82, $eif2ak4^{-/-}$ mice + PBS and $eif2ak4^{-/-}$ mice + LF82 groups at day 1 (**A**) and 4 (**B**) post-infection. Groups were compared using permanova method. Ns, not significant.



Figure S2: *Eif2ak4* gene deficiency has no impact on the gut microbiota composition of CEABAC10 transgenic (Tg) mice.

 $Tg/eif2ak4^{+/+}$ (N=13) and $Tg/eif2ak4^{-/-}$ (N=13) mice were placed in specific-pathogen free housing conditions. Feces were collected from 12-week old mice for the analysis of the bacterial microbiota composition based on Illumina pyrosequencing of the 16S rRNA gene. Mouse fecal bacterial communities were clustered using PCoA of the weighted UniFrac distance matrix. PCoA-1 and PCoA-2 were plotted, and the percentage of the variation explained by the plotted principal coordinates was indicated in the Y-X-axis labels. Groups were compared using permanova method. Ns, not significant.





Tg/eif2ak4^{+/+} and Tg/eif2ak4^{-/-} mice were challenged by oral gavage for 3 days (once per day) with PBS (N=6 mice per group) or with 10⁹ CFU of the AIEC LF82 strain (N=6 mice per group). Feces were collected at day 1 (**A**) and 4 (**B**) post-infection to analyze the bacterial microbiota composition based on Illumina pyrosequencing of the 16S rRNA gene. Mouse fecal bacterial communities were clustered using PCoA of the weighted UniFrac distance matrix. PCoA-1 and PCoA-2 were plotted, and the percentage of the variation explained by the plotted principal coordinates was indicated in the Y-X-axis labels. Comparison between Tg/eif2ak4^{+/+} mice + PBS, Tg/eif2ak4^{+/+} mice + LF82, Tg/eif2ak4^{-/-} mice + PBS and Tg/eif2ak4^{-/-} mice + LF82 groups at day 1 (**A**) and 4 (**B**) post-infection. Groups were compared using permanova method. Ns, not significant.