

Fra Background

73 In our previous study we have focused on the impact of the various antibiotic treatments on
74 host innate immune response and conjugative R plasmids transfer gene activities by using an
→ 75 experimental zebrafish infection-and-treatment model [5]. The results motivated us to
76 investigate similar processes in a mammalian model. We have therefore assessed the
77 expression of pathogenic ETEC borne pRAS1 transfer genes *in vivo* in piglets, in response to
78 antibiotic, probiotic and Non-steroidal Anti-inflammatory Drug (NSAID) treatments, while
79 simultaneously monitoring selected blood serum proteins.

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81 **Results**

82 **Clinical signs and re-isolation of *E. coli***

83 All the *E. coli* inoculated piglets showed loss or reduction of appetite, acute watery or
84 mucoid-diarrhea, a pale, yellowish colour, increased respiratory activity and reduced
85 movement within 6-8 hours after challenging. The uninfected piglets remained apparently
86 healthy, with normal vigour, appetite and skin colour, and showed no diarrhea.

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88 Macroscopically inflammation of the intestinal mucosa was recorded in both the placebo
89 treated pig and groups treated with ineffective antibiotics, but not with the control group. On
90 the serosa side of the jejunum it was observed severe hyperemia with filled blood vessels and
91 the intestinal wall appeared edematous including firm and thickened to palpate in piglets from
92 all infected groups not given effective treatment. However, also infected piglets from groups
93 that were treated with effective antibiotics had a mild hyperemia visible from the serosa side
94 of the jejunum and also a clear edematous intestinal wall compared to the uninfected control
95 group.

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97 **Histopathology and cultivation from the intestinal content.**