

Supplemental Material to:

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Gut microbial diversity is reduced and is associated with colonic inflammation in a piglet model of short bowel syndrome

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Supplementary table 1: List of primer sequences and Universal ProbeLibrary probe combinations used in this study.

Primer	Sequence 5' to 3'	UPL Probe
RPL32 Forward	aactggccatcagggtcac	#64
RPL32 Reverse	cacaactggaactcctgtctattc	
IL1B Forward	ccaattcagggaccctacc	#19
IL1B Reverse	catggctgcttcagaaacct	
IL8 Forward	ttcttctttatccccaaactgg	#41
IL8 Reverse	ccacatgtcctcaaggtagga	
IL18 Forward	actttactttgtagctgaaaacgatg	#85
IL18 Reverse	tttaggttcaagcttgccaaa	
TNF Forward	ttgtcgctacatcgctgaac	#32
TNF Reverse	ccagtagggcggttacagac	

Figure S1. Rarefaction curve and principle component analysis of samples sequenced by high-throughput DNA sequencing. (A) Rarefaction curves for each group at 97% similarity indicated that the total bacterial diversity present was well represented. Number of operational taxonomic units (OTUs) identified as a function of the number of sequence tags sampled. (B) Principle component analysis revealed that the small bowel resection (SBR) groups at week 2 and week 6 clustered together, and were distinct from the non-operation control (NOC) and sham groups.



