Supplemental Table 1. Final body weights and colon lengths of Wt and *Fat-1* saline control and TNBS-treated mice¹

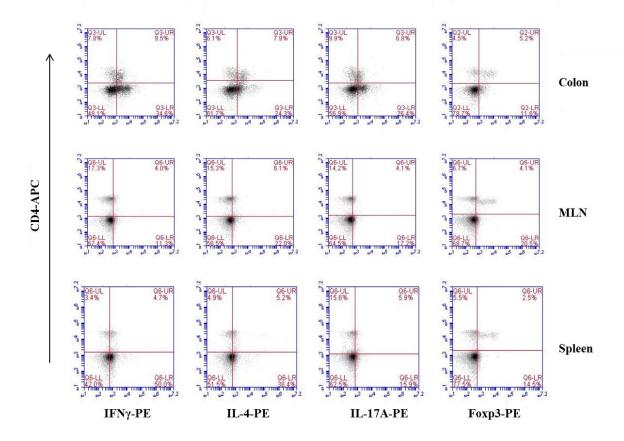
	Saline Control		TNBS		
	Wt	Fat-1	Wt	Fat-1	
Body Weight (g)	$24.09 \pm 0.66^*$	$23.01 \pm 0.82^*$	$18.75 \pm 0.64^{\ddagger}$	$19.89 \pm 0.84^{\ddagger}$	
Colon Length (cm)	$6.11 \pm 0.16^*$	$5.92 \pm 0.20^*$	$4.82 \pm 0.13^{\ddagger}$	$5.14 \pm 0.52^{\ddagger}$	

¹Mean values \pm SEM of saline control (n=4/genotype) and TNBS-treated Wt and *Fat-1* mice (n=9/genotype). Within rows, values not sharing a superscript symbol differ ($P \le 0.05$).

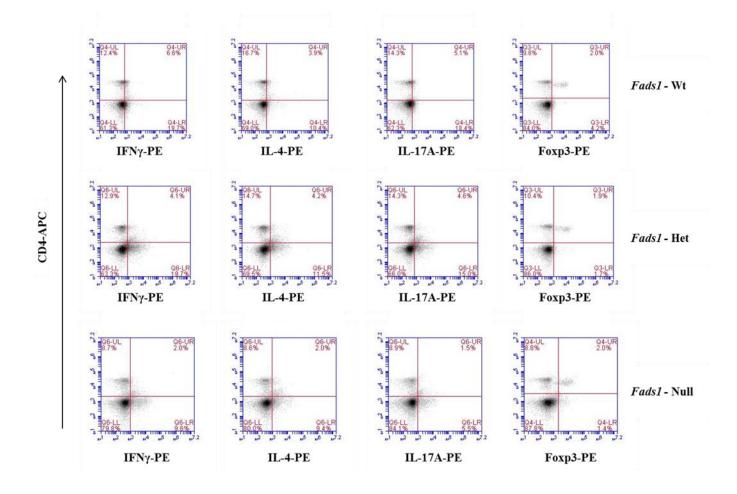
Supplemental Table 2. Final body weights and colon lengths *Fads1* saline control and TNBS-treated mice¹

	Saline Control			TNBS		
_	Wt	Het	Null	Wt	Het	Null
Body Weight (g)	22.53±1.66*	22.13±1.20*	19.23±1.18*	18.86±1.23 [‡]	16.39±0.86 [‡]	16.75±1.14 [‡]
Colon Length (cm)	6.40±0.13*	6.10±0.12*	6.30±0.12*	5.53±0.26 [‡]	5.33±0.25 [‡]	5.48±0.32 [‡]

¹Mean values \pm SEM of Wt, Het and Null *Fads1* saline control (n=4/genotype) and TNBS-treated mice (n=6-9/genotype). Values marked with a different superscript symbol differ (P≤0.05).



Supplemental Figure 1. Representative dot plots deplicting the detection of CD4⁺ T cell subsets in TNBS-treated Wt colon lamina propria (top row), MLN (middle row) and spleen (bottom row). All cells were surface stained with CD4-APC (y axis) followed by intracelluar detection of T cell subset specific markers (x axis): IFNγ-PE (Th1), IL-4-PE (Th2), IL-17A-PE (Th17) and Foxp3-PE (Treg) (see Materials and Methods). Flow cytometric analyses were conducted using an Accuri C6 flow cytometer (Accuri Cytometers).



Supplemental Figure 2. Representative dot plots deplicting the detection of splenic CD4⁺ T cell subsets in TNBS-treated *Fads1* Wt (top row), Het (middle row) and Null mice (bottom row). All cells were surface stained with CD4-APC (y axis) followed by intracelluar detection of T cell subset specfic markers (x axis): IFNγ-PE (Th1), IL-4-PE (Th2), IL-17A-PE (Th17) and Foxp3-PE (Treg) (see Materials and Methods). Flow cytometric analyses were conducted using an Accuri C6 flow cytometer (Accuri Cytometers).