

Combined inhibition of TL1A and integrin $\beta 7$ is superior to either monotherapy in mouse models of colitis and coadministration of SPY001 and SPY002 demonstrates no drug-drug effects on exposure in non-human primates

M. Siegel¹, J. Friedman¹, D. Nguyen¹, J. McNally¹, M. Kennedy¹, O. Ballew¹, E. Lewis¹, D. Giles¹, M. Rose¹, A. Spencer¹

¹Spyre Therapeutics, Waltham MA, United States

Background

- **Combined** use of **targeted biologic agents** could break through the IBD treatment efficacy ceiling while avoiding the risks associated with broad immunosuppression.
- **SPY001** and **SPY002** are **half-life extended antibodies** against validated IBD targets (**$\alpha 4\beta 7$ integrin** and **TL1A**, respectively) which are being evaluated as monotherapies and in combination to treat IBD.

Methods and Results

SPY120 (a combination of SPY001 and SPY002) targets $\alpha 4\beta 7$ and TL1A – orthogonal drivers of IBD

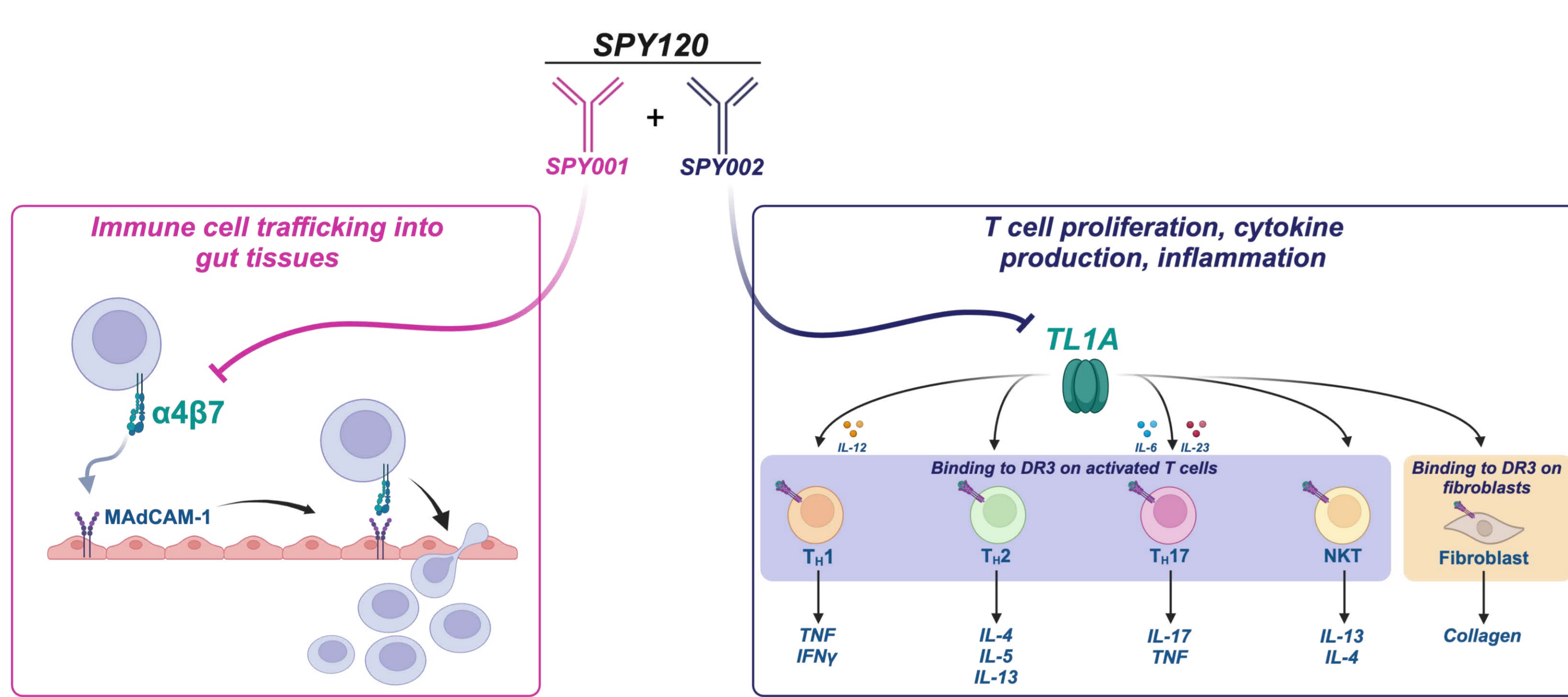


Figure 1: Mechanistic rationale for SPY120. SPY001 is a recombinant humanized anti- $\beta 7$ mAb¹; SPY002 is a fully human anti-TL1A mAb².

Anti-mouse surrogates for SPY120 and its monotherapy components were studied in two IBD mouse models

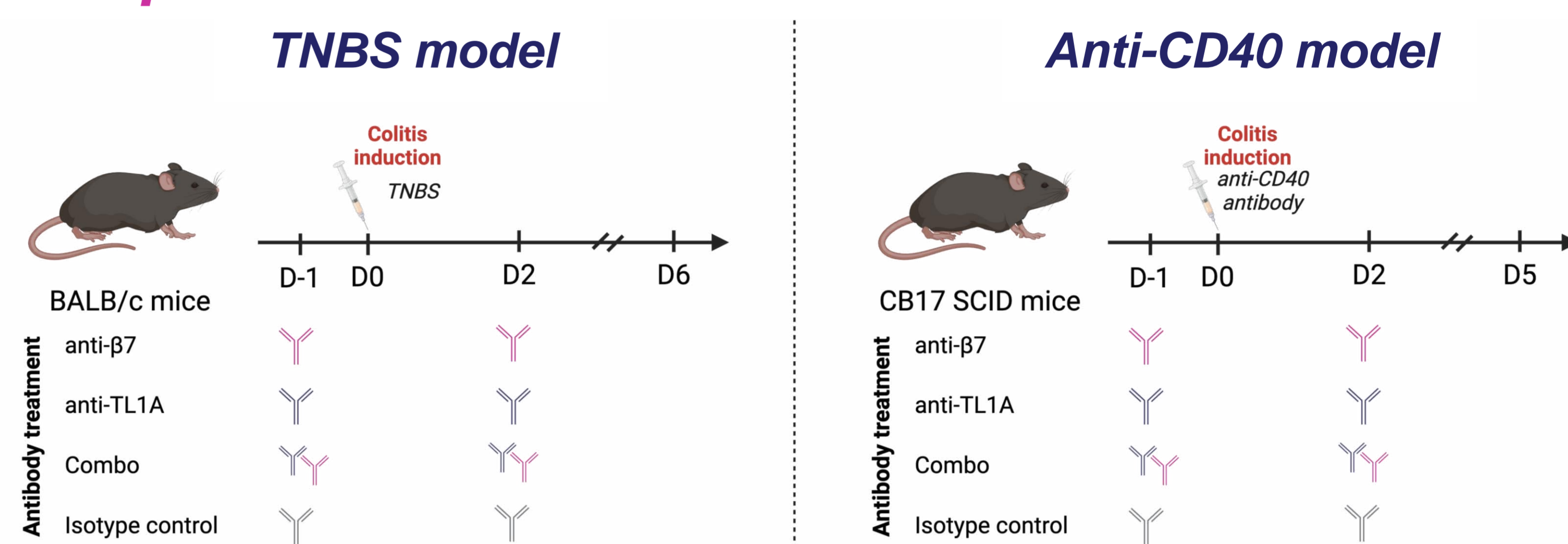


Figure 2: **TNBS model (left)** – BALB/c mice were dosed intravenously with test article (25 mg/kg) on Day -1 and Day 2, with Day 0 representing TNBS administration per prior methods³. **Anti-CD40 model (right)** – CB17 SCID mice were dosed intravenously with test article (25 mg/kg) on Day -1 and Day 2, with Day 0 representing anti-CD40 antibody administration per prior methods⁴.

Anti- $\beta 7$ and anti-TL1A combination improves disease activity score in murine colitis models

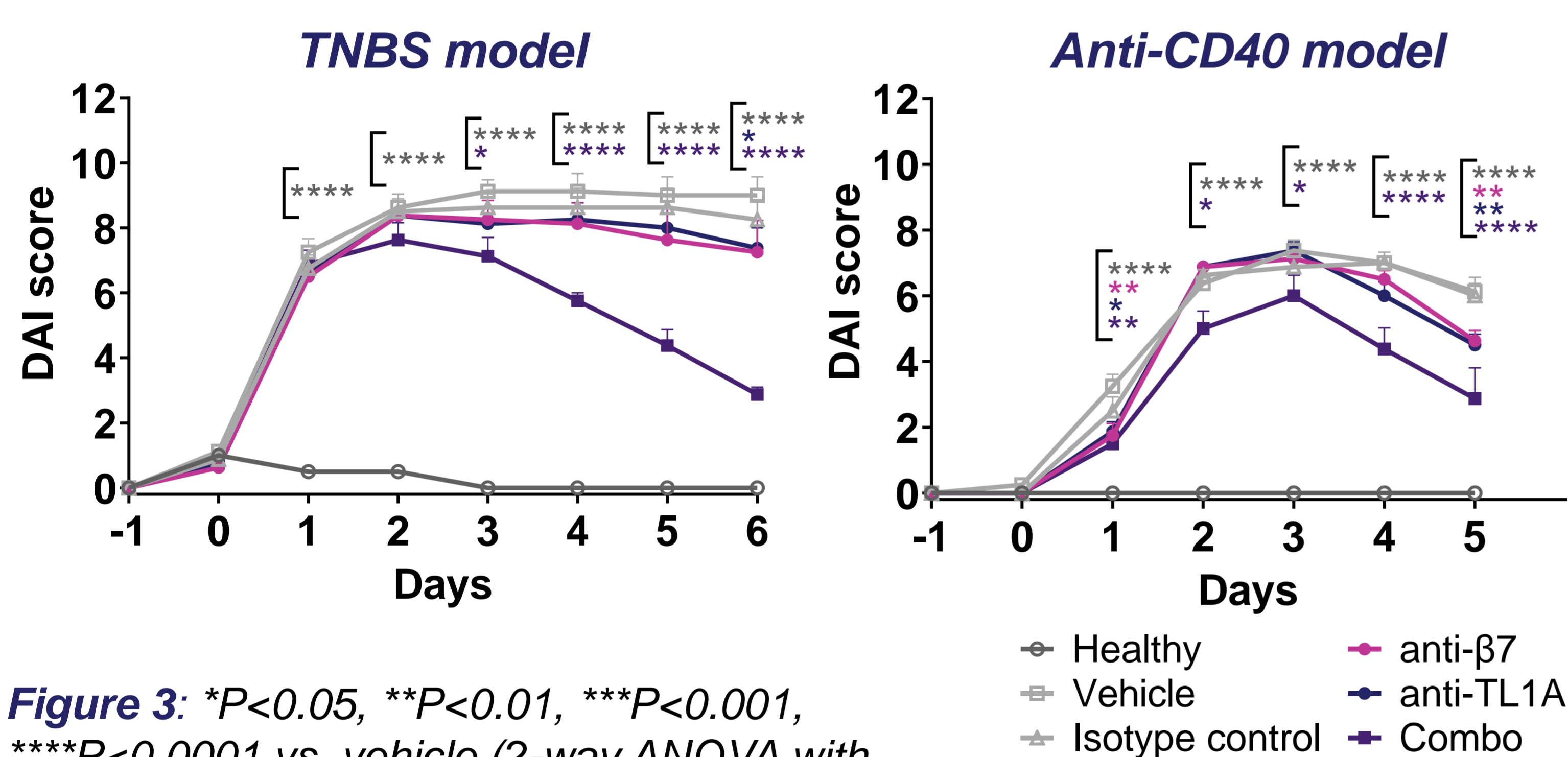


Figure 3: * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$ vs. vehicle (2-way ANOVA with Dunnett's correction). $N = 8$ per group.

Anti- $\beta 7$ and anti-TL1A combination improves histopathology score in murine colitis models

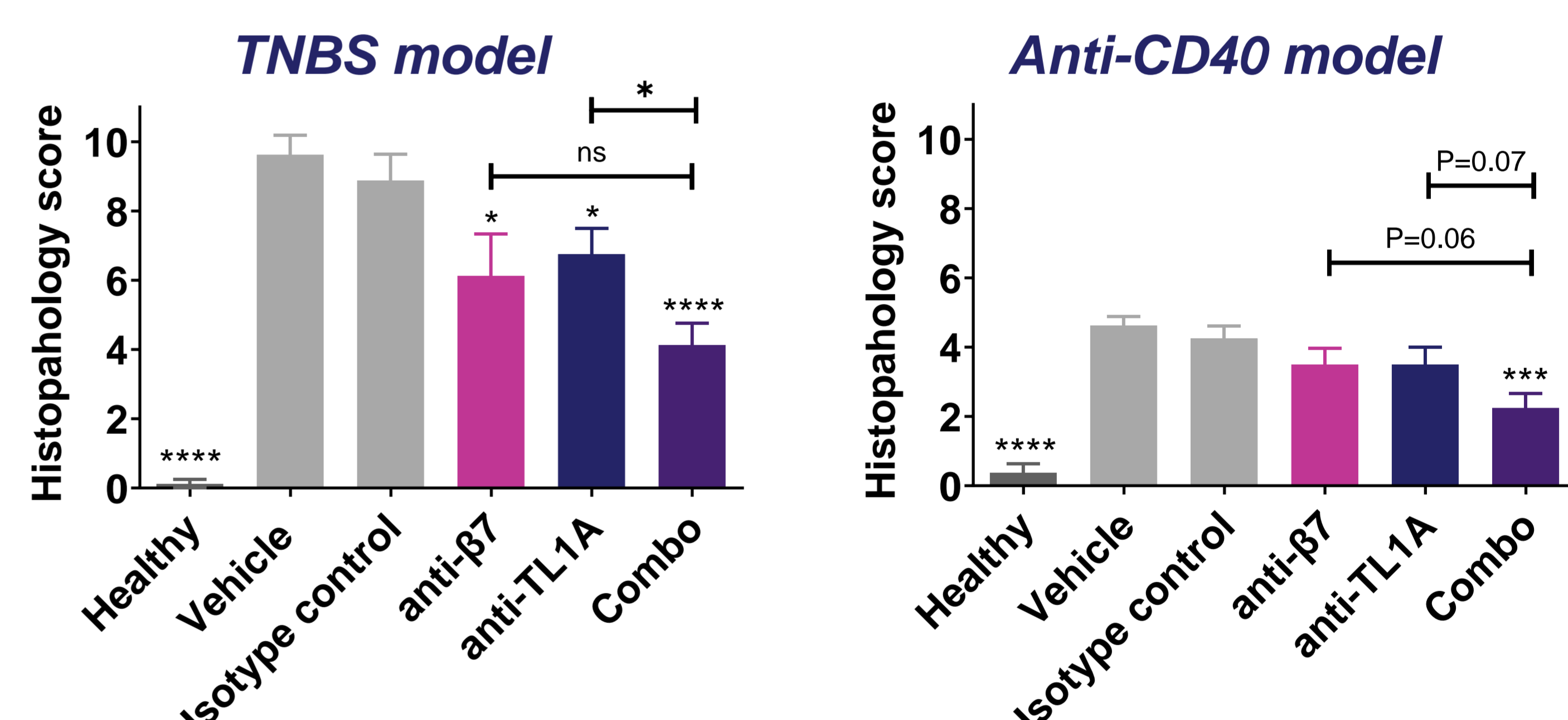


Figure 4: Histopathology scoring of distal colon harvested on D6 (TNBS) or D5 (anti-CD40). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$ vs. vehicle (1-way ANOVA with Dunnett's correction and t-test for individual comparisons). $N = 8$ per group.

All treatments led to a decrease in TNF levels in colons of murine colitis models

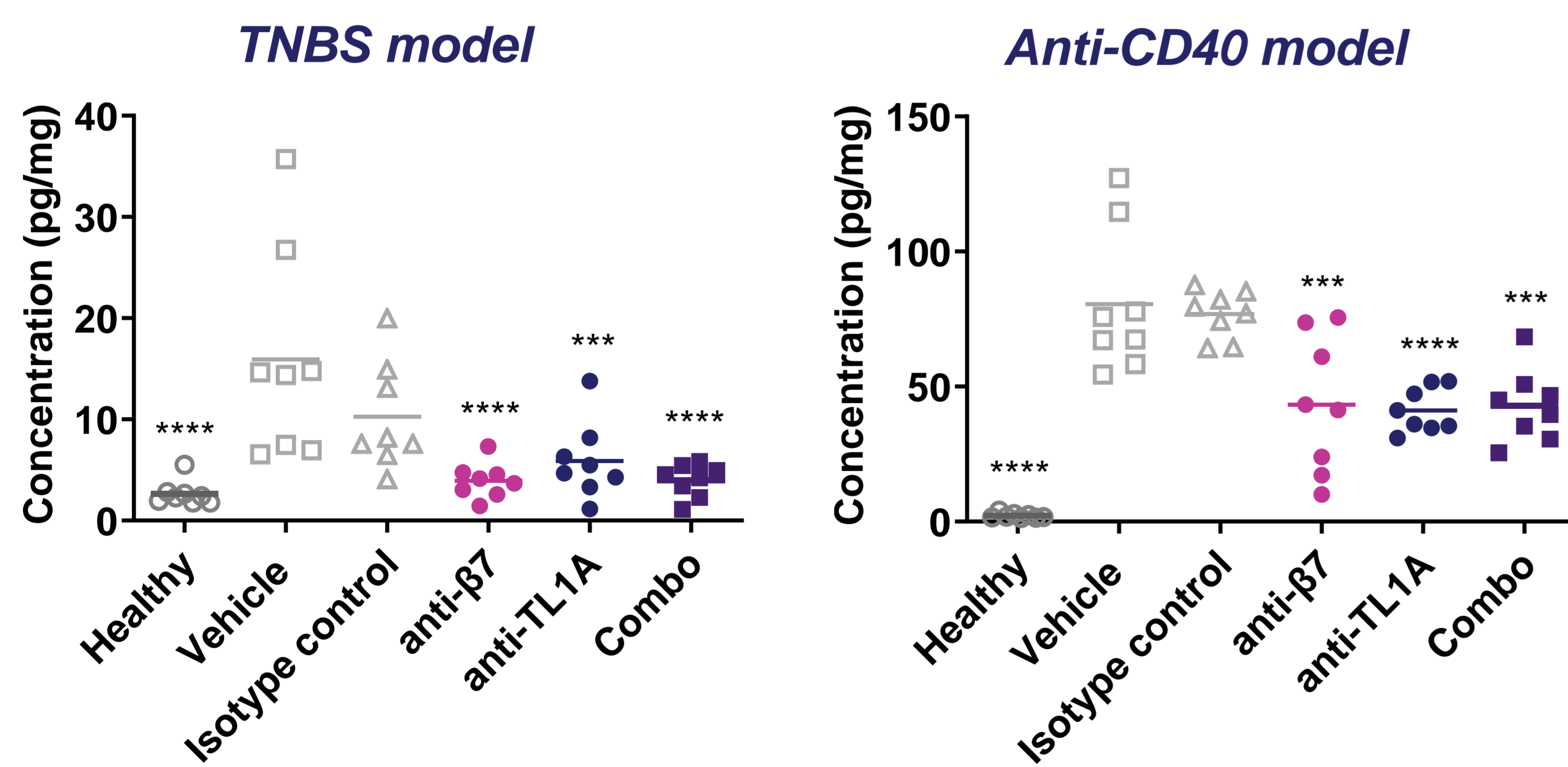


Figure 5: TNF levels in mid-colon harvested on D6 (TNBS) or D5 (anti-CD40). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$, **** $P < 0.0001$ vs. vehicle (1-way ANOVA with Dunnett's correction). $N = 8$ per group.

Serum concentrations of monotherapies and combination therapy were similar in NHPs

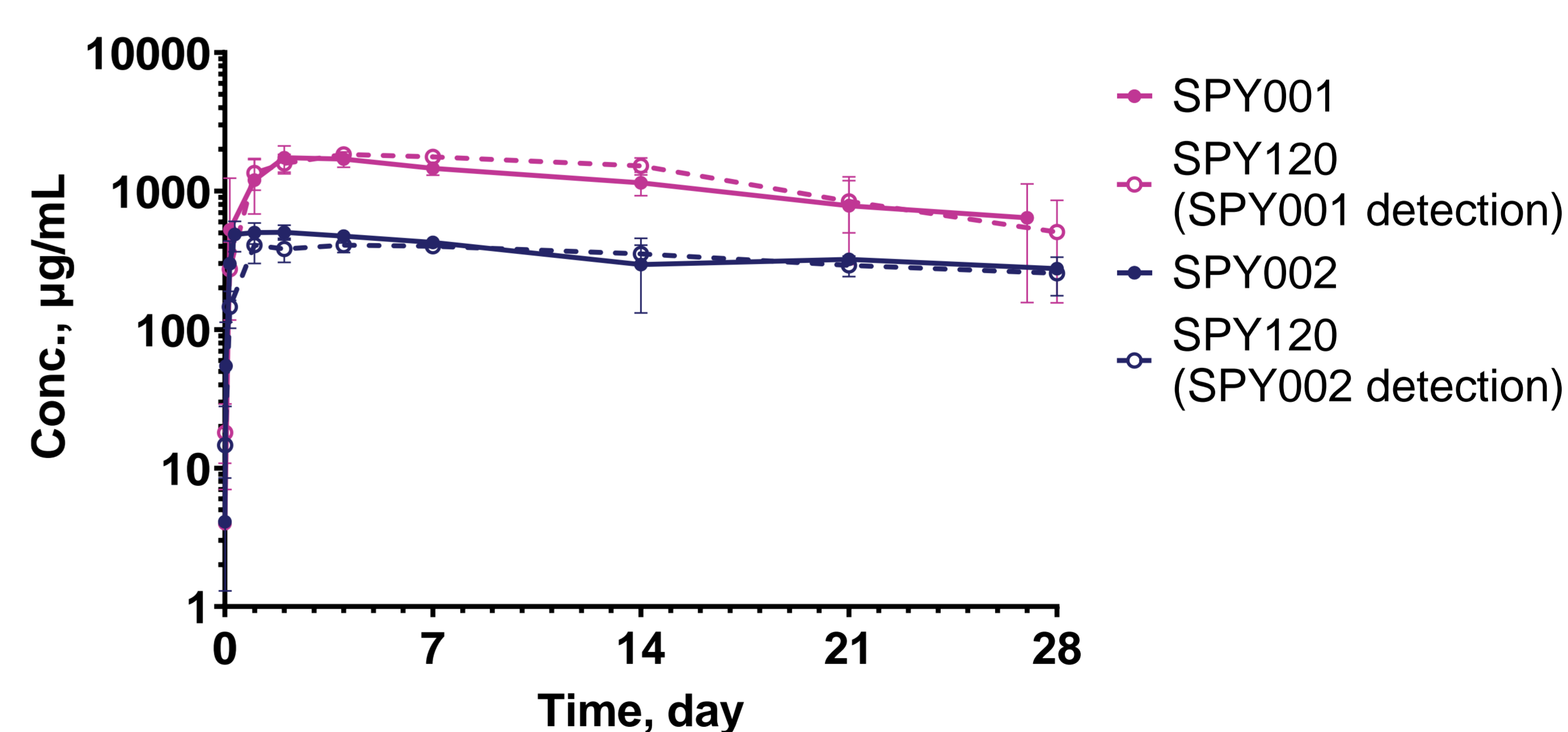


Figure 6: Mean serum concentrations in cynomolgus NHPs following a single SC dose of SPY001 (anti- $\alpha 4\beta 7$, 150 mg/kg), SPY002 (anti-TL1A, 50 mg/kg), or SPY120 (160 mg/kg SPY001 + 50 mg/kg SPY002). $N = 3$ per group.

Conclusions

- Combined **anti- $\beta 7$ integrin** and **anti-TL1A** therapy showed **additive or greater than additive** *in vivo* biological activity relative to either monotherapy in the **TNBS and anti-CD40 mouse colitis models**.
- The **PK profiles** of SPY001 and SPY002 were similar in NHPs whether dosed as monotherapy or in combination, indicating potential for **Q3-6M dosing** for SPY001, SPY002, and SPY120 based on human PK simulations.
- These preclinical results support the **advancement** of the **combination** of SPY001 and SPY002 into **clinical trials**.

References

1. Zhu, E. *et al.* A Novel Monoclonal Antibody Drug Candidate SPY001 Targeting Integrin $\alpha 4\beta 7$ for the Treatment of IBD: In Vitro Properties and Non-Human Primate Pharmacokinetics and Safety. *UEGW*, PP1103 (2024).
2. Zhu, E. *et al.* Characterization of Two Novel Extended Half-life Monoclonal Antibody Drug Candidates Targeting TL1A for the Treatment of IBD. *UEGW*, MP118 (2024).
3. Neurath, M. *et al.* Antibodies to Interleukin 12 Abrogate Established Experimental Colitis in Mice. *JEM* 182, 1281-1290 (1995).
4. Uhlir, H. *et al.* Differential Activity of IL-12 and IL-23 in Mucosal and Systemic Innate Immune Pathology. *Immunity* 25(2), P309-318 (2006).

Disclosures

All authors are employees of Spyre Therapeutics and own equity in Spyre Therapeutics.