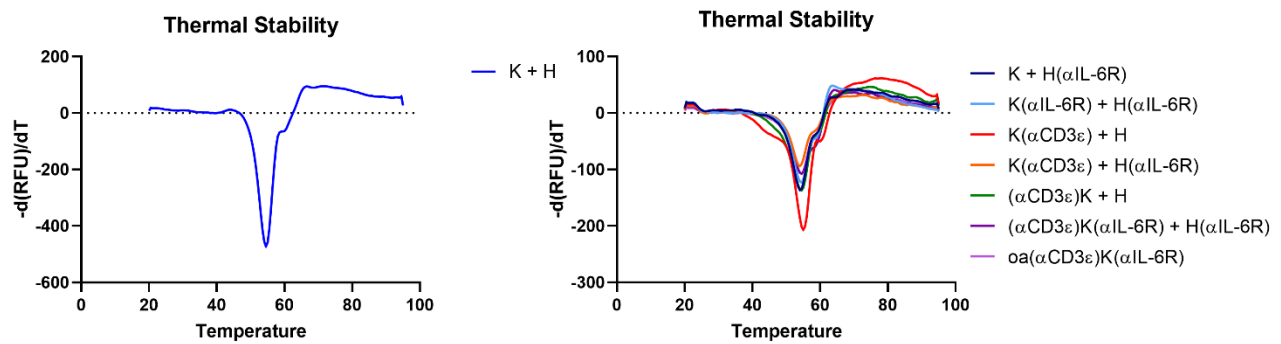


Supplementary Material

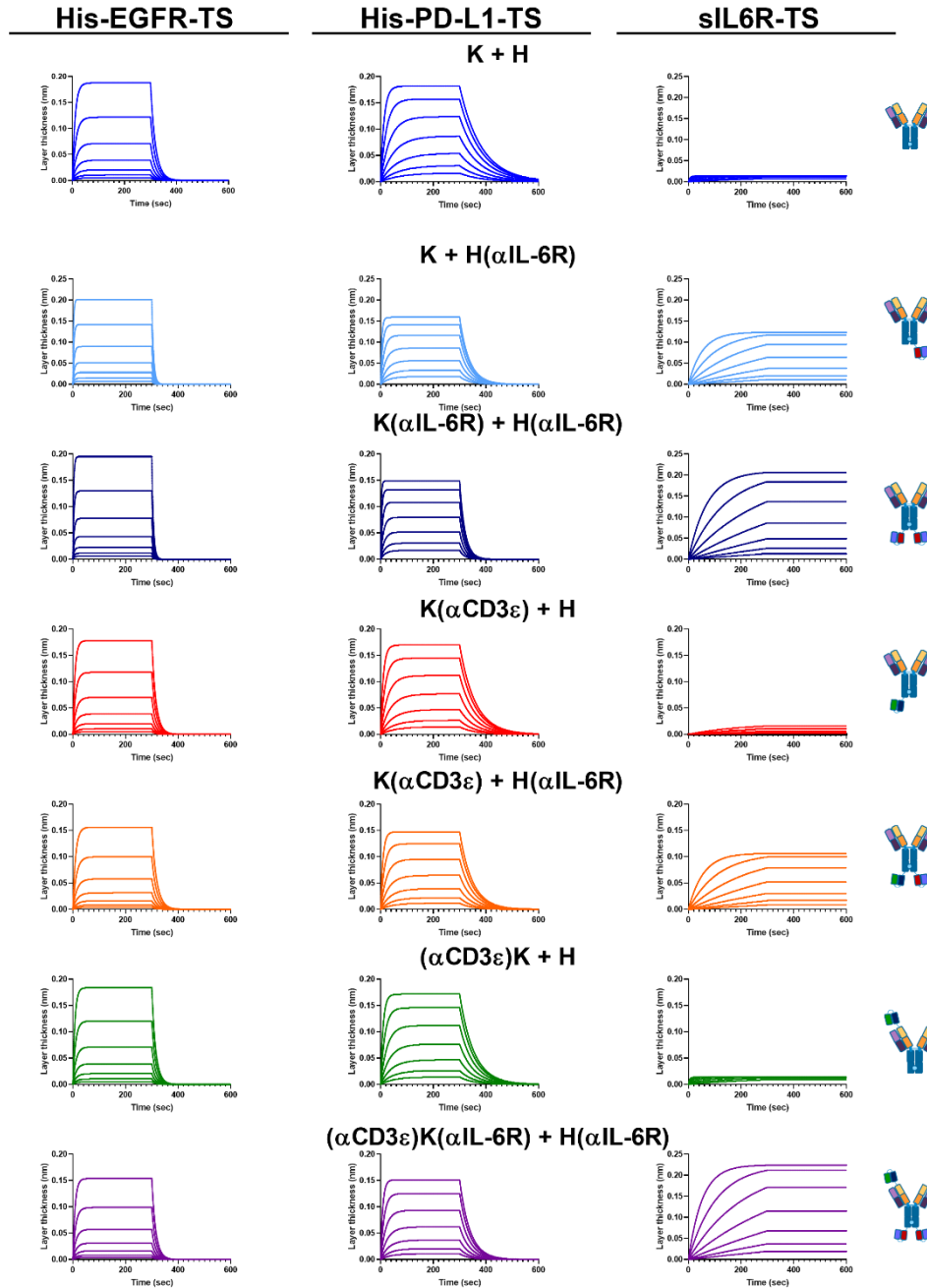
TriTECM: A Tetrafunctional T-Cell Engaging Antibody with Built-In Risk Mitigation of Cytokine Release Syndrome

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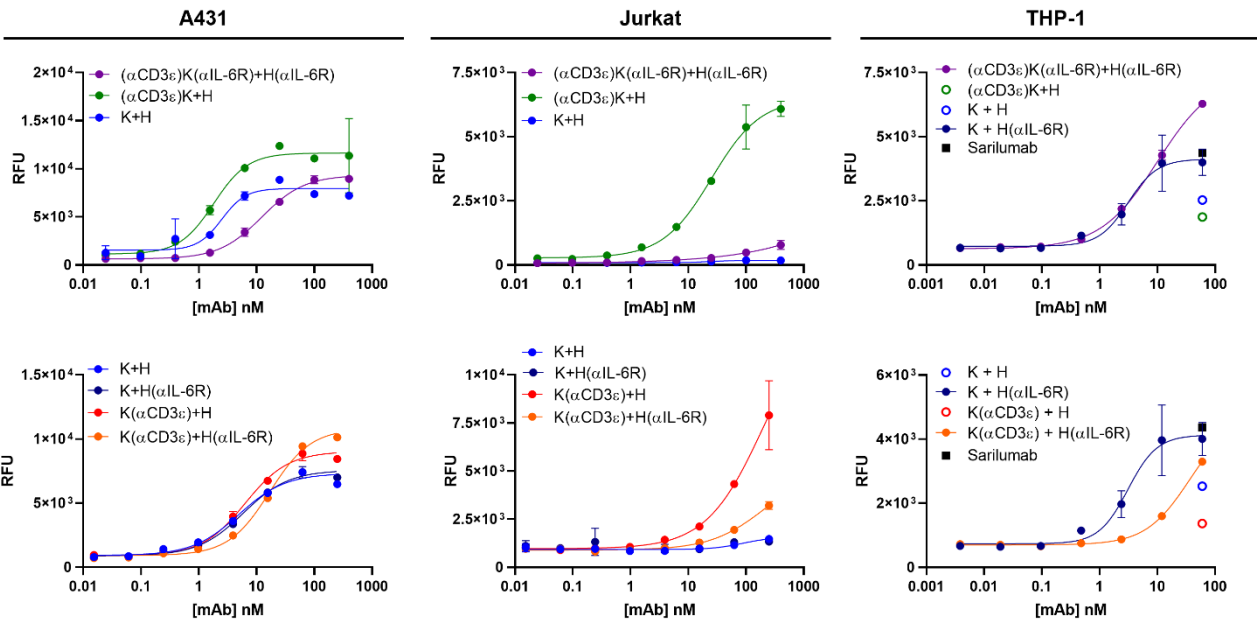
1 Supplementary Figures and Tables



Supplementary Figure 1: Melt peaks after thermal shift assay by SYPRO Orange. RFU – relative fluorescence units. The derivative of RFU is plotted against temperature in a range from 20 – 95 °C. The dotted line represents the threshold set to determine the melt temperature.



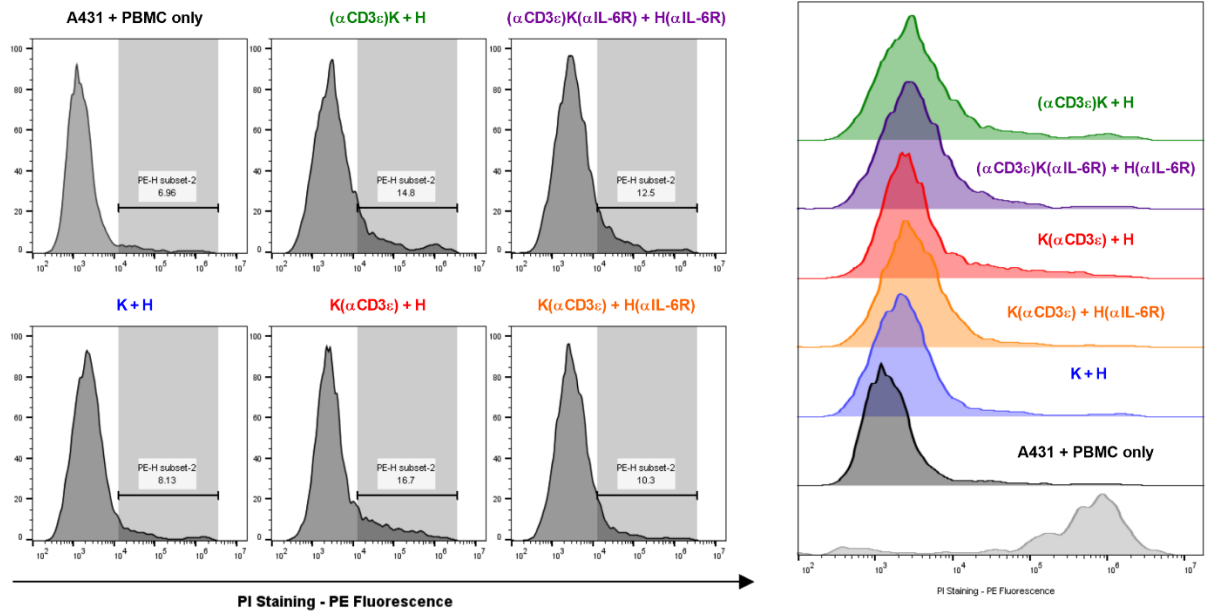
Supplementary Figure 2: Affinity determination by biolayer interferometry. The binding curves for His-EGFR-TS (left), His-PD-L1-TS (middle) or soluble IL6R-TS (right) are displayed for all antibodies. The colour-coding represents the different variants. For His-EGFR-TS and His-PD-L1-TS, a concentration range of 7.8 – 500 nM was measured, while for sIL6R-TS a range from 3.125 – 200 nM was measured.



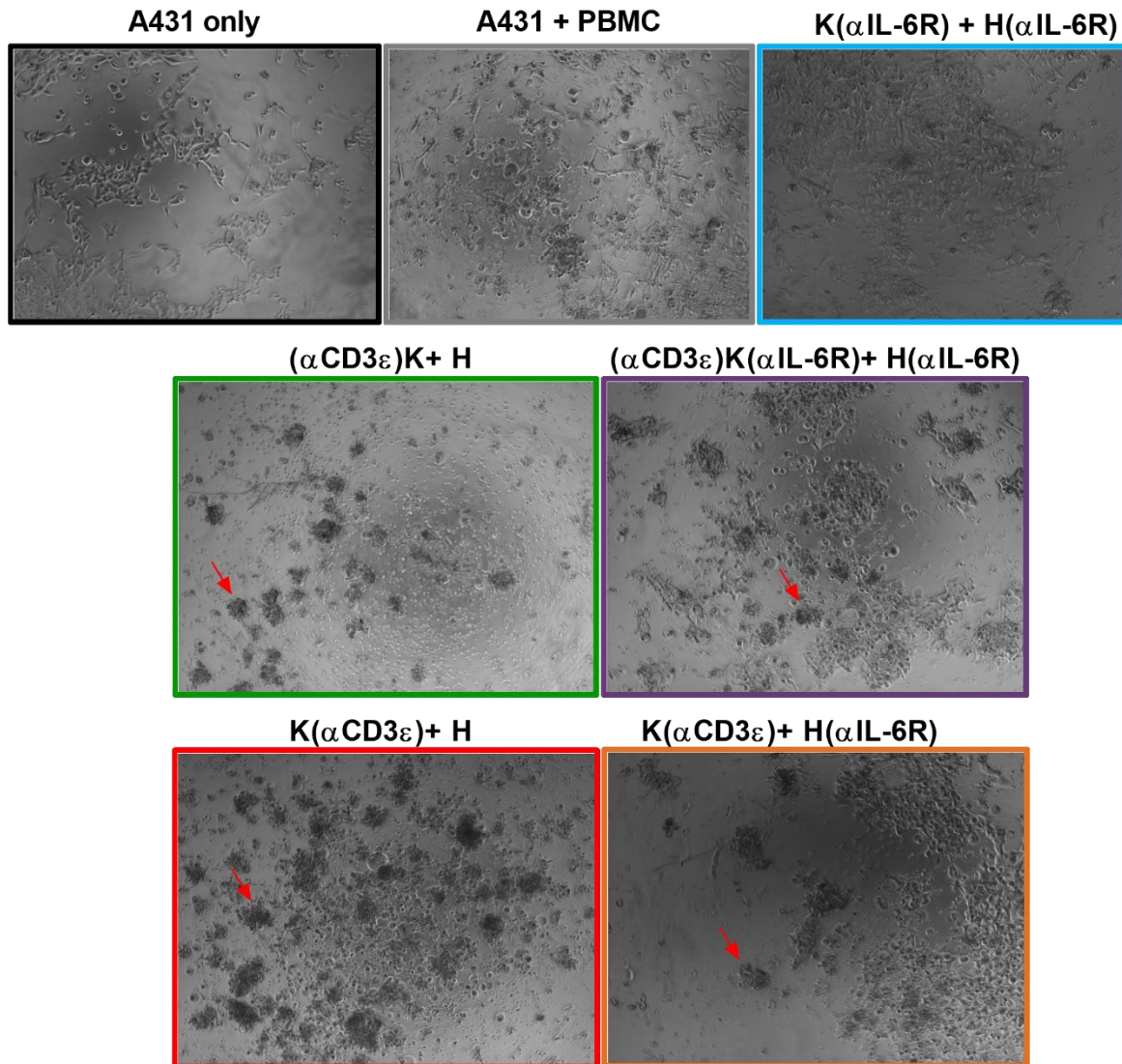
Supplementary Figure 3: On-cell titrations. The antibodies were tested for either CD3-binding (Jurkat), IL6R-binding (THP-1) or EGFR/PD-L1-binding (A431). The top panel show the variants with an N-terminal NI0401 scFv fusion, whereas the bottom panel shows the C-terminal NI0401 scFv variants with the respective controls. The mean fluorescence was determined and plotted using GraphPad prism. A non-linear regression was determined to calculate the on-cell affinities. RFU – relative fluorescence units.

Supplementary Table 1: On-cell affinities on A431 (EGFR⁺⁺⁺/PD-L1⁺), Jurkat (CD3⁺) and THP-1 (IL6R⁺/PD-L1⁺).

| Variant | On-cell affinity (nM) | | |
|---|-----------------------|--------|-------|
| | A431 | Jurkat | THP-1 |
| K + H | 2.50 | - | - |
| K + H(αIL-6R) | 4.30 | - | 3.10 |
| K(αCD3ϵ) + H | 4.50 | 195.90 | - |
| K(αCD3ϵ) + H(αIL-6R) | 21.24 | 213.80 | 36.66 |
| (αCD3ϵ)K + H | 1.82 | 25.38 | - |
| (αCD3ϵ)K(αIL-6R) + H(αIL-6R) | 12.34 | n.d. | 10.08 |



Supplementary Figure 4: Histograms and half-offset overlays for dead cell staining with propidium iodide (PI) after co-culture of A431 and PBMCs for 24 h to measure T-cell-mediated cytotoxicity. Histograms and overlays were generated using FlowJo V10 software.



Supplementary Figure 5: Bright field images of A431 and PBMC co-culture in combination with 20 nM of the indicated antibodies after 48 h incubation. Black dead target cell clusters are exemplified by red arrows in each of the relevant images.