

Supplementary: Relative Contribution of Framework and CDR Regions in Antibody Variable Domains to Multimerisation of Fv- and scFv-Containing Bispecific Antibodies

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Supplementary Data

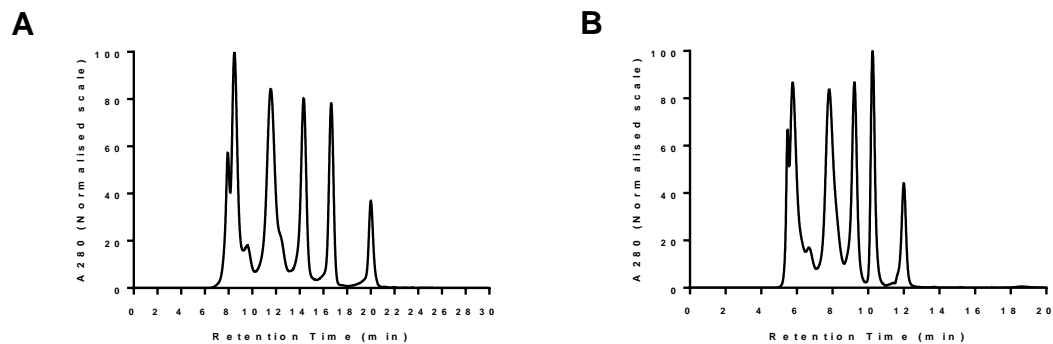


Figure S1. Gel filtration calibration. (A) S200 SEC profile. (B) G3000 SEC profile. Gel filtration standard (BioRad) was loaded for molecular weight estimation of test samples. The mixture contained bovine thyroglobulin (670,000 Da), bovine γ -globulin (158,000 Da), chicken ovalbumin (44,000 Da), equine myoglobin (17,000 Da) and vitamin B12 (1350 Da).

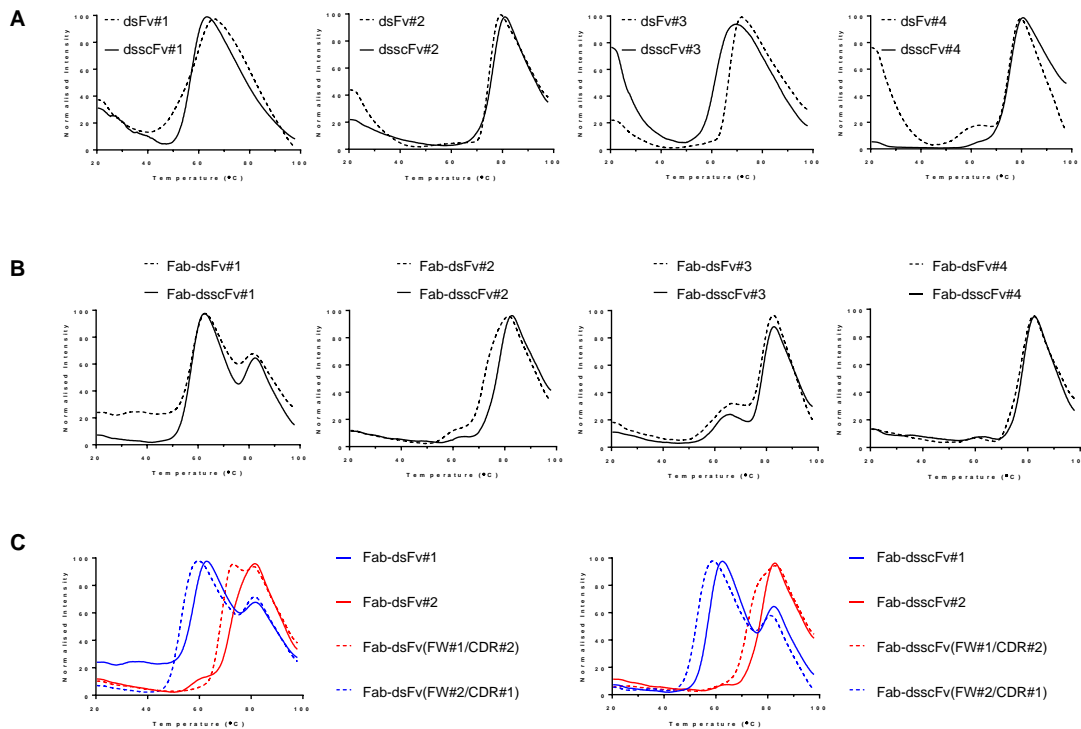


Figure S2. Thermograms of purified proteins. (A) dsFv vs dsscFv. **(B)** Fab-dsFv vs Fab-dsscFv. **(C)** Wild type vs FW/CDR 'swapped' Fab-dsFv and Fab-dsscFv. Fluorescence changes in protein/SYPRO®Orange dye mixtures were analysed on a 7900HT fast real-time PCR System, set at 20 °C to 99 °C with a ramp rate of 1.1 °C/min. The thermograms show unfolding transitions of proteins in PBS pH7.4. The inflection point of the slope(s) was used to generate the T_m .

