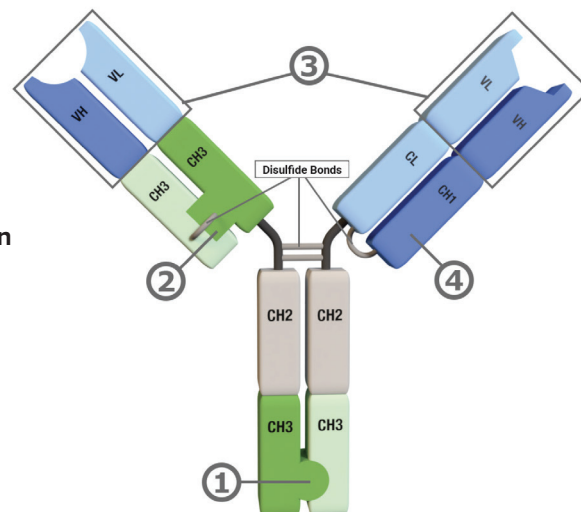


At the heart of multispecific antibody innovation lies the B-Body® Platform—a premier bispecific antibody framework engineered for exceptional yield, purity, and stability. Built entirely on a human IgG-like scaffold, the B-Body Platform addresses critical challenges in antibody discovery and manufacturing, setting new standards in the industry.

## B-Body: innovative by design

### CMC DRIVEN INNOVATIONS

- 1 **Fc Region: Clinically Validated Knobs-into-Holes**
  - Drives heavy chain heterodimerization
- 2 **Fab Arm: Proprietary CH3 Domains to Drive Heterodimerization**
  - Substitutes for CH1/CL in one Fab Arm
  - Solves light chain mispairing issue
  - Novel CH3 mutations, without knobs-into-holes or charge pairs
  - pl asymmetry streamlines polishing steps
- 3 **Heavy & Light Chains: Proprietary Symmetrical Inversions**
  - Enhances expression yield and stability
  - Facilitates one-step purification
  - Compatible with a wide variety of variable domain sources
- 4 **Sole CH1 Domain: Allows a Level of Purification Beyond Protein A/IEX**
  - Facilitates one-step purification with anti-CH1 resins
  - Simplifies in-format discovery
  - Removes homodimers and other impurities in one pass

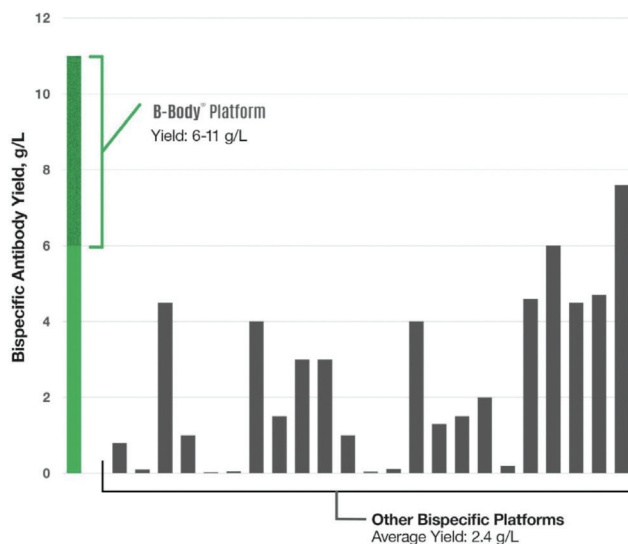


## Unmatched bispecific yield and performance

The B-Body Platform addresses challenges in bispecific antibody discovery and manufacturing, achieving top yields in transient and stable cell lines. Production methods, from engineered CHO pools to fed-batch, yield 6-11 g/L—surpassing other platforms. Perfusion culture can reach 1.8 g/L/day or higher.

### THE B-BODY PLATFORM IS OPTIMIZED FOR:

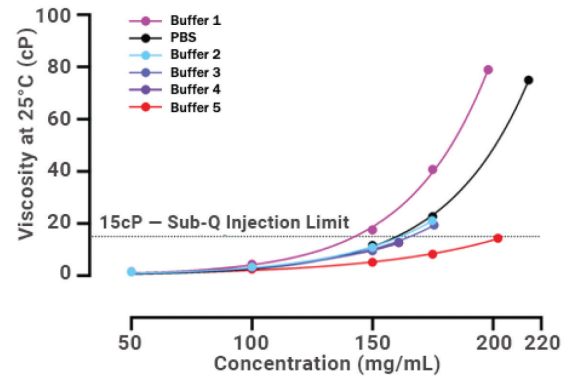
- **Unmatched Bispecific Yield:** Industry-leading production efficiency
- **Validated Production:** Record yields from multiple commercial stable cell lines
- **Proper Assembly:** Optimized heterodimerization for high yield and simple CMC
- **95% Single-Pass Purity:** High purity possible via either CH1 or Protein A in a single pass
- **CMC Compatibility:** Human IgG-like scaffold works with standard purification methods producing high yield and 100% purity with commonly used two-column processes
- **Therapeutics Development:** High solubility, stability and low viscosity for production



# Sub-Q compatibility

B-Body Bispecifics' high solubility, stability, and low viscosity make them ideal for subcutaneous delivery. In tests, both standard and low-fucose variants reached concentrations with viscosity below Sub-Q injection and ultrafiltration/diafiltration limits.

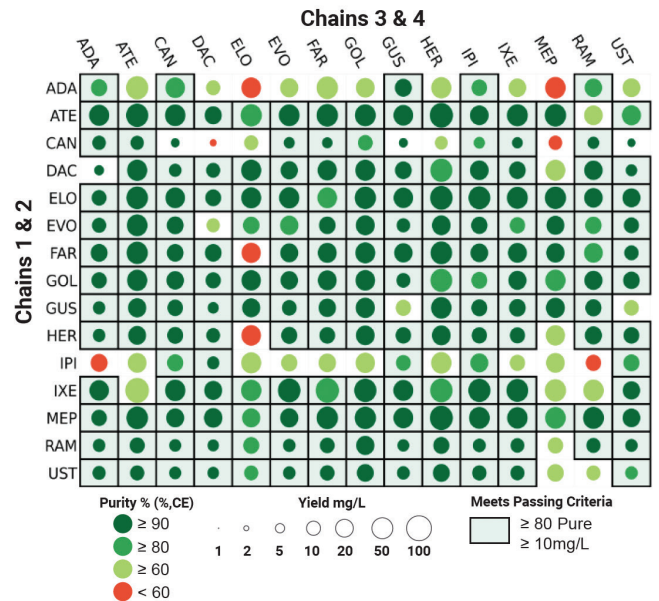
- **Sub-Q Compatible Viscosity:** Up to 200mg/mL or higher without exceeding viscosity limits
- **Within Range of UF/DF Production:** Sub-Q concentrations possible without exceeding UF/DF limits
- **Excellent Solubility:** Higher concentrations (>200mg/mL) can be achieved if required for specific applications



# Validated performance: broad range clinical antibody expression

A 15x15 matrix of clinical-stage antibodies was expressed in 1mL cultures and purified in one step with Anti-CH1 resin. Derived from a variety of heavy and light chain germlines, these bispecifics demonstrate compatibility with a broad range of antibody sources and sequences. Purity was assessed by CE, binding by Octet® BLI, and assembly by mass spec. Over 99% of the combinations tested met purity and yield criteria in at least one orientation, confirming the platform's broad-based performance.

Name	hV Gene	lV Gene	HEK Titer (mg/L)	Fab Tm DSF (°C)	SGAC-SINS AS100 (NH4)2SO4 mM	HIC Retention Time (Min)	SMAC Retention Time (Min)	Poly-Specific. Reag. SMP Score (0-1)	AC-SINS* Δλmax (nm), Average
Adalimumab (ADA)	IGHV3-9*01	IGKV1-27*01	134.9	71.0	900.0	8.8	8.7	0.00	1.1
Atezolizumab (ATE)	IGHV3-23*04	IGKV1-NL1*01	164.1	73.5	300.0	13.4	19.3	0.07	15.0
Canakinumab (CAN)	IGHV3-33*01	IGKV6D-21*02	45.7	72.0	800.0	9.3	8.7	0.00	0.7
Daclizumab (DAC)	IGHV1-46*01	IGKV1-5*01	245.1	74.0	900.0	9.3	8.8	0.00	-0.1
Elotuzumab (ELO)	IGHV3-7*05	IGKV1-27*01	213.2	83.5	700.0	10.3	9.3	0.00	-0.2
Evolocumab (EVO)	IGHV1-18*01	IGLV2-14*01	260.7	65.0	700.0	10.4	9.1	0.20	2.2
Farletuzumab (FAR)	IGHV3-30*03	IGKV1D-33*01	220.8	75.5	800.0	9.5	9.1	0.00	-0.5
Golimumab (GOL)	IGHV3-30*01	IGKV3-11*01	163.2	70.0	0.0	11.4	12.7	0.23	23.0
Guselkumab (GUS)	IGHV5-10-1*04	IGLV1-40*01	167.3	69.5	700.0	11.4	9.2	0.47	3.4
Trastuzumab (HER)	IGHV3-11*05	IGKV1-5*05	159.5	78.5	800.0	9.7	8.8	0.00	2.0
Ipilimumab (IPI)	IGHV3-30*01	IGKV3-20*01	169.6	73.0	400.0	11.6	13.0	0.23	10.4
Ixekizumab (IXE)	IGHV1-46*01	IGKV2D-29*02	97.3	83.0	500.0	10.9	9.1	0.81	20.0
Mepolizumab (MEP)	IGHV2-70*20	IGKV4-1*02	221.5	78.5	900.0	9.2	8.8	0.00	-1.0
Ramucirumab (RAM)	IGHV3-21*01	IGKV1-12*01	90.7	66.0	900.0	9.4	8.7	0.00	0.0
Ustekinumab (UST)	IGHV5-51*01	IGKV1D-16*01	152.7	69.5	1000.0	8.8	8.6	0.15	0.4



Learn more at [twistbioscience.com/twist-biopharma-solutions](https://www.twistbioscience.com/twist-biopharma-solutions) or contact us at [biopharmasales@twistbioscience.com](mailto:biopharmasales@twistbioscience.com)

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