

1500 1st Ave North Birmingham, AL 35203 1-205-453-8242 info@trialtusbioscience.com

Data Sheet

Im7 6B Resin

SKU No.: 10-1060, 10-1062, 10-1065, 10-1071, 10-2060, 10-2065, 10-3060, 10-3065

Description	Im7 ligand (10 kDa) on agarose beads binds the CL7 tag (~16 kDa) fused to
	target proteins. Following cleavage with the appropriate protease, the
	target protein releases from the Im7-bound CL7.
Particle Size	Crosslinked agarose 6B beads (45 - 165 μM)
pH Stability	The Im7 protein is stable on the beads at pH 3-10 (normal working
	conditions). However, CL7/Im7 binding is stable at pH 4.2-10 only.
Salt Stability	≤ 4 M NaCl tested
Binding Capacity	35-40 mg CL7/mL resin
Storage/Shipping	50:50 buffer:resin slurry. 1-, 2-, 5-, and 15-mL of settled resin in 2, 4, 10,
Concentration	and 30 mL of slurry, respectively.
Shipping Conditions	Room temperature / Buffer: 20 mM Tris-Cl pH 8.0, 0.5M NaCl, 5% glycerol,
	0.05% sodium azide
Recommended	4°C or room temperature
Operating Temperature	
Reactivation Details	To remove the CL7 protein and reactivate the resin, wash the column with
	guanidine hydrochloride, exchanging it into physiological buffer.
	Alternatively, wash with Gentle Elution Buffer (3.6 M MgCl₂) to avoid
	refolding.

Additional information: Target protein characteristics (e.g. protein size, conformation, and concentration); flow rate (i.e. lower flow rates may increase the binding capacity); and other parameters (e.g. pH and temperature) can affect the binding capacity.

You can download full protocols from https://trialtusbioscience.com/pages/protein-purification-protocols.

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Licensing Information

TriAltus Bioscience holds the exclusive, worldwide license to the CL7 protein purification technology platform. It was licensed from the University of Alabama at Birmingham (UAB) in Birmingham, Alabama, USA. An international patent filing has been made with protection being sought in the United States, Europe, and other major markets. The CL7 purification technology is available for research use. For commercial use or resale, contact us at sales@trialtusbioscience.com to discuss commercial licensing.

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