

Product datasheet

protag-HiPur TagFP Agarose Beads

Short overview

Cat. No.	89300L
Quantity	2 ml

Product description

Host	Llama/alpaca
Antibody Type	Recombinant, produced in E.coli
Isotype	Single-domain antibody
Clone	1H7
Immunogen	TagFP
Formulation	50% slurry in PBS containing 20% Ethanol
Transfer Vector	> 4 µg mTagBFP per µl of packed beads
Packaging Plasmid	sdAb anti-TagFP clone 1H7
Support	4% cross-linked agarose, bead size 50-150 µm
Buffer compatibility	<ul style="list-style-type: none"> - Common buffer substances at pH 5 to 9 - 2% Triton X-100, 1% Tween-20, 1% NP-40, 1% CHAPS, 1% Deoxycholate, 0.1% SDS - 4 M NaCl, 2 M KCl, 1 M MgCl₂, 100 mM EDTA - 4 M urea - 10 mM DTT, 10 mM 2-Mercaptoethanol - RNase A, DNase I, Benzonase, protease inhibitors
Purification	Affinity chromatography
Storage	2-8°C
Intended use	Research use only
Application	IP
Reactivity	mKate/mKate2, mTagBFP, mTagRFP, mTagRFP657
No reactivity	Dendra2, Dronpa, tdEOS, mEOS3.2, mRuby3, mTFP, GFP, dsRed or their most common derivatives

Applications

Immunoprecipitation (IP)	yes
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Background

protag-HiPur TagFP Agarose Beads are based on a high-affinity single-domain antibodies (sdAb) that are covalently immobilized on 4% cross-linked agarose beads. The sdAbs are attached via a flexible linker which guarantees a high accessibility of the sdAbs and largely eliminates batch-to-batch variations. Due to the single-chain nature of sdAbs and their covalent attachment, no "leakage" of light and heavy chains from IgGs is observed during elution with SDS sample buffer. protag-HiPur TagFP Agarose Beads thus feature high affinity and superior capacity for TagFP fusion proteins while showing negligible non-specific background. protag-HiPur TagFP Agarose Beads immobilize a wide range of TagFP

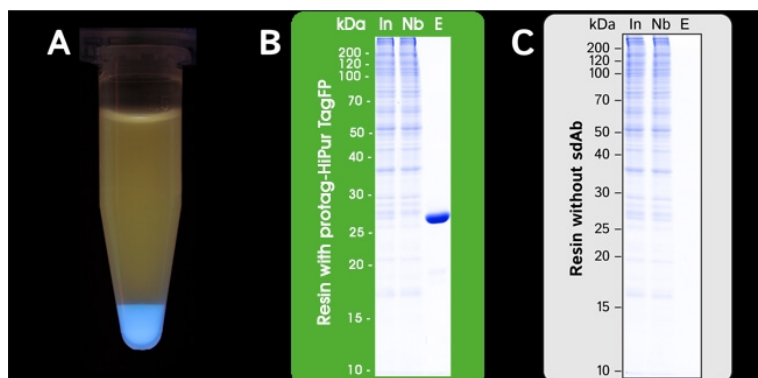
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2024 April 27 / Version: 89300L/DS-040221ibg | Page 1

derivatives, such as mTagBFP, mTagRFP, mTagRFP657 and mKate. Reactivity against other TagRFP derivatives is not analyzed. It does not cross-react with common GFP- or dsRed/RFP derivatives including EBFP. protag-HiPur GFP Agarose Beads are compatible not only with physiological buffers but also with high stringency buffers. With protag-HiPur TagFP Agarose Beads the binding and washing conditions can be adjusted to the experimental needs.

Product images



A) Pull-down of mTagBFP from a mixture of GFP, mCherry and mTagBFP. B) IP of mTagBFP from HeLa lysate. In/Ft: 1/1000 of input and non-bound material. E: Eluate from 1 µl of beads. C) Control experiment using functionalized beads lacking sdAbs.