## For Research Use Only

## HA-Trap Magnetic Agarose, Kit for **Immunoprecipitation**



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Catalog Number: atmak

**Catalog Number: Basic Information** 

**Applications:** IP, Co-IP

Conjugate: Magnetic agarose beads; ~40 um (cross-linked 6% magnetic agarose beads)

**Host:** Alpaca

Type: Nanobody Class:

Recombinant

The HA-Trap Magnetic Agarose Kit is a ready-to-use reagent for the IP of HA-tagged proteins. It consists of an anti-HA-tag Nanobody/VHH coupled to magnetic agarose beads, along with lysis, wash, and elution buffers to use in the IP process. **Description** 

Specificity/Target Binds specifically to the HA-tag (sequence YPYDVPDYA) fused to a protein of interest at N-, C- or internal position. Please note that the affinity is highest for a C-terminal fusion. There is no cross-reactivity to other common peptide tags such as the His6-tag, FLAG-tag, Spot-Tag, V5-tag, Strep-tag, or C-tag (other tags not tested). Background binding to host cell proteins from a range of organisms such as human, mouse and dog cell lines or yeast and plants is low.

**Elution buffer** 2x SDS-sample buffer (Lammli)

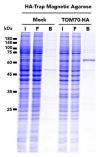
Affinity (K<sub>D</sub>) 6 nM for C-terminal HA-tags and ca. 180 nM for N-terminal fusions.

Storage Storage: Shipped at ambient temperature. Upon receipt store at +4°C. Stable for one year. DO not freeze!

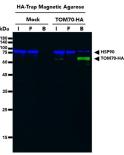
Storage Buffer: 20% ethanol

1(312) 455-8498 (outside USA)

## **Selected Validation Data**



The HA-Trap Magnetic Agarose Kit was used to immunoprecipitate TOM70-HA fusion protein from either untransfected (mock) HEK293T cells or HEK293T cell transfected with full-length TOM70-HA construct. SDS-PAGE analysis was done on samples from the Input (I), Flow-through (F), Bound (B) fractions.



Co-IP using HA-Trap Magnetic Agarose Kit followed by multiplexed WB of TOM70-HA and HSP90 proteins from untransfected (mock) HEK293T cells and HEK293T cells transfected with full-length TOM70-HA construct. WB analysis was done on samples from the Input (I), Flow-through (F) and Bound (B) fractions of the IP. TOM70 Monoclonal Antibody (66593-1-Ig), Multi-rAB CoraLite Plus 488-Goat Anti-Mouse Recombinant Secondary Antibody (RGAM002), HSP90 Polyclonal Antibody (13171-1-AP), and Multi-rAb CoraLite Plus 750-Goat Anti Rabbit Recombinant Secondary Antibody (RGAR006) were used in the WB analysis.