

This product is for research use only (not for diagnostic or therapeutic use)

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Product no AS21 4538

Anti-AGO5c | Argonaute 5c (Zea mays)

Product information

Immunogen KLH-conjugated peptide derived from Zea mays AGO5c, UniProt: A0A1D6GFG2

Host Rabbit

Clonality Polyclonal

Purity Antigen affinity purified serum, in PBS pH 7.4

Format Lyophilized

Quantity 50 μg

Reconstitution For reconstitution, add 50 μl of sterile water.

Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Please, remember to spin tubes briefly prior to opening them to avoid any losses that might occur from lyophilized material adhering to the cap or sides of the tubes.

Application information

Recommended dilution 1:1000 (WB)

Expected | apparent

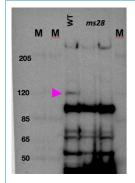
119 kDa

Confirmed reactivity Zea mays

Predicted reactivity | Species of your interest not listed? Contact us

Not reactive in No confirmed exceptions from predicted reactivity are currently known

Selected references To be added when available, antibody available in October 2025.



Samples:

M - Marker

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1 – 68 μg Zea mays WT premeiotic (0.5-1mm) anthers extract (TCA/acetone precipitation)

2 – 30 μg Zea mays ms28 premeiotic anthers (TCA/acetone precipitation)

3-6 8 µg Zea mays ms28 premeiotic anthers (TCA/acetone precipitation)

M – Marker

Extracted amples were separated at RT using Tris-Acetate 3-8% NuPage LDS-PAGE gel, then blotted to PVDF membrane (0.45 μm pore size) using wet transfer O/N in the cold. The day after, blot was stained with Ponceau and dried down. Blot was blocked with 5% milk in TBS-T O/N at 4°C with agitation followed by the incubation with the primary antibody at a dilution of 1:500 with 2% milk in 5mL TBS-T for 1 hr at RT with agitation. Blot was washed with TBS-T 3x at RT with agitation (5 min per wash). Blot was incubated in matching secondary antibody (Goat anti-rabbit IgG (H&L), HRP conjugated, <u>AS09 602</u>, Agrisera) diluted to 1:25 000 in 2% milk in TBS-T for 1 hr at RT with agitation and followed with washes with TBS-T 3x at RT with agitation (5 min per wash), then developed with the Agrisera ECL Bright and SuperBright reagents. Exposure time was 3 minutes.

Target protein band identity was confirmed by IP-MS analysis.