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Datasheet for ABIN7170595 anti-SKA3 antibody (AA 101-195)

3 Images



Overview

Quantity:	100 µg
Target:	SKA3
Binding Specificity:	AA 101-195
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SKA3 antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Spindle and kinetochore-associated protein 3 protein (101-195AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

Target:	SKA3
Alternative Name:	SKA3 (SKA3 Products)
Background:	Background: Component of the SKA1 complex, a microtubule-binding subcomplex of the outer
	kinetochore that is essential for proper chromosome segregation (PubMed:19289083,

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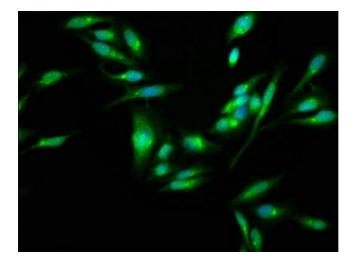
Storage:

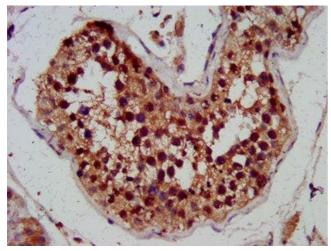
PubMed:19360002, PubMed:23085020). The SKA1 complex is a direct component of the sinetochore-microtubule interface and directly associates with microtubules as oligomeric assemblies (PubMed:19289083, PubMed:19360002). The complex facilitates the processive novement of microspheres along a microtubule in a depolymerization-coupled manner PubMed:19289083). In the complex, it mediates the microtubule-stimulated oligomerization PubMed:19289083). Affinity for microtubules is synergistically enhanced in the presence of the ndc-80 complex and may allow the ndc-80 complex to track depolymerizing microtubules PubMed:23085020). Aliases: C13orf3 antibody, RAMA1 antibody, SKA3 antibody, SKA3_HUMAN antibody, Spindle and kinetochore associated complex subunit 3 antibody, Spindle and kinetochore-associated protein 3 antibody 281X90
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2ecommended dilution: IHC:1:200-1:500 IE:1:50-1:200
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Accommended dilation. In 0.1.200 1.300, II.1.30 1.200,
For Research Use only
iquid
Preservative: 0.03 % Proclin 300
Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
ProClin
his product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be
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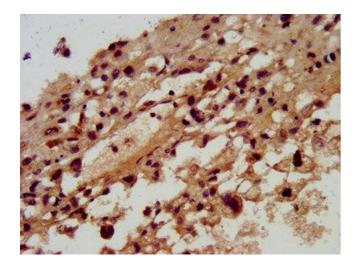
Storage Comment: Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

-20 °C,-80 °C

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Immunofluorescence

Image 1. Immunofluorescence staining of Hela cells with ABIN7170595 at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

Immunohistochemistry

Image 2. IHC image of ABIN7170595 diluted at 1:300 and staining in paraffin-embedded human testis tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Immunohistochemistry

Image 3. IHC image of ABIN7170595 diluted at 1:300 and staining in paraffin-embedded human melanoma performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

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