Anti-LOC4331983 antibody

Product name

Anti-LOC4331983 antibody

Specificity

Rice

Antibody description

Rabbit polyclonal antibody to LOC4331983

Preparation

This antigen of this antibody was synthetic peptide within n-terminal rice d14.

Formulation

Liquid, 1*TBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide.

Storage

Store at +4°C after thawing. Aliquot store at -20°C. Avoid repeated freeze / thaw cycles.

Clonality

Polyclonal

Ig Type

Rabbit IgG

Applications

WB, IHC-P, IF

Dilutions

WB: 1:500-1:1000

IHC-P: 1:50-1:200

IF: 1:50-1:200

Validations

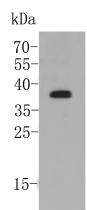


Fig1:; Western blot analysis of D14 on rice tissue lysate. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:5,000 dilution was used for 1 hour at room temperature.

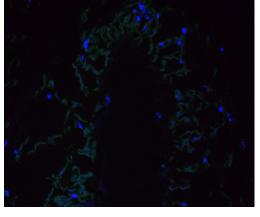


Fig2:; Immunofluorescence staining of paraffinembedded rice tissue using anti-D14 rabbit polyclonal antibody.The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes.(sodium citrate buffer (pH 6.0) for 20 mins.) The tissues were blocked in 10% negative goat serum for 1 hour at room temperature, washed with PBS, and then probed with the antibody () at 1/100 dilution for 10 hours at 4°C and detected using Alexa Fluor[™] 488 conjugate-Goat anti-Rabbit IgG (H+L) Secondary Antibody at a dilution of 1:500 for 1 hour at room temperature.



Anti-LOC4331983 antibody



Catalog Number: 175183

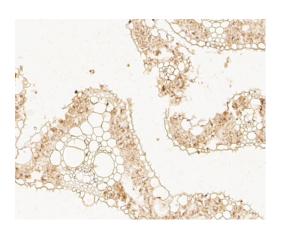


Fig3:; Immunohistochemical analysis of paraffinembedded rice tissue using anti-D14 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes.The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH; 2; O and PBS, and then probed with the antibody () at 1/100 dilution, for 30 minutes at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chrogen. Counter stained with hematoxylin and mounted with DPX.