abcam

Product datasheet

Anti-UMOD antibody [EPR22133-142] - BSA and Azide free (Detector) ab259582

Recombinant RoloWAb

2 Images

Overview

Product name Anti-UMOD antibody [EPR22133-142] - BSA and Azide free (Detector)

Description Rabbit monoclonal [EPR22133-142] to UMOD - BSA and Azide free (Detector)

Host species Rabbit

Tested applications Suitable for: Sandwich ELISA

Species reactivity Reacts with: Mouse

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

General notes ab259582 is a BSA and Azide Free antibody supplied in an unconjugated format and it is suitable for sandwich ELISAs to quantify Mouse Uromodulin. The recommended pair for sandwich

ELISA is:

Capture: ab259581, Mouse Uromodulin Capture Antibody (unconjugated) Detector: ab259582, Mouse Uromodulin Detector Antibody (unconjugated)

The reference range value is 625 - 40000 pg/ml.

Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

The recommended antibody orientation is based on internal optimization for ELISA-based assays. Antibody orientation is assay dependent and needs to be optimized for each assay type. Please note that the range provided for this antibody is only an estimation based on the performance of the product using the recommended antibody pair. Performance of the antibody pair will depend on the specific characteristics of your assay. We guarantee the product works in sandwich ELISA, but we do not guarantee the sensitivity or dynamic range of the antibody in your assay.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C.

Storage buffer Constituent: 100% PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal

Clone number EPR22133-142

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab259582 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Sandwich ELISA		Use at an assay dependent concentration. Can be paired for Sandwich ELISA with Rabbit monoclonal [EPR22133-150] to UMOD - BSA and Azide free (Capture) (ab259581).

Target

Function Not known. May play a role in regulating the circulating activity of cytokines as it binds to IL-1, IL-2

and TNF with high affinity.

Tissue specificity Synthesized by kidney. Most abundant protein in normal human urine.

Involvement in diseaseDefects in UMOD are the cause of familial juvenile hyperuricemic nephropathy type 1 (HNFJ1)

 $\hbox{[MIM:162000]}. \ HNFJ1 \ is \ a \ renal \ disease \ characterized \ by \ juvenil \ onset \ of \ hyperuricemia, \ polyuria,$

progressive renal failure, and gout. The disease is associated with interstitial pathological

changes resulting in fibrosis.

Defects in UMOD are the cause of medullary cystic kidney disease type 2 (MCKD2)

[MIM:603860]. MCKD2 is a form of tubulointerstitial nephropathy characterized by formation of renal cysts at the corticomedullary junction. It is characterized by adult onset of impaired renal

function and salt wasting resulting in end-stage renal failure by the sixth decade.

Defects in UMOD are the cause of glomerulocystic kidney disease with hyperuricemia and isosthenuria (GCKDHI) [MIM:609886]. GCKDHI is a renal disorder characterized by a cystic

dilation of Bowman space, a collapse of glomerular tuft, and hyperuricemia due to low fractional

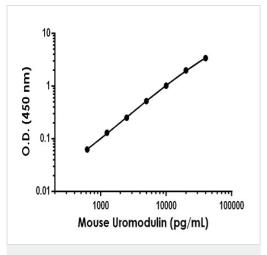
excretion of uric acid and severe impairment of urine concentrating ability.

Sequence similaritiesContains 3 EGF-like domains.

Contains 1 ZP domain.

Cell ular localizationCell membrane. Secreted. Secreted after cleavage in the urine.

Images



Sandwich ELISA - Anti-UMOD antibody [EPR22133-142] - BSA and Azide free (Detector) (ab259582)

Representative standard curve from corresponding SimpleStep ELISA® Kit (ab245726).



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