

Localization – Membrane

Host Species – Mouse

Ig Class – IgG

Intended Use

This antibody is designed for the specific localization of RCC in formalin-fixed, paraffin-embedded (FFPE) tissue sections.

Storage & Handling

Store RTU Vial at 2-8°C. Fresh dilutions for concentrated antibodies, if required, should be prepared prior to use and are stable for up to one day at room temperature (20-26°C).

Working Principle

IHC is a two-step process wherein the primary antibody binds to the antigen of interest and that binding is detected by a chromogen. The primary antibody may be used in IHC using manual techniques or any Automated Staining System. Positive and negative controls should always be run simultaneously with all patient specimens.

Product Description

In normal renal tissue, gp200 is localized along the brush border of both the pars convolute and pars recta segments of the proximal tubule, as well as focally along the luminal surface of Bowman’s capsule adjacent to the emerging proximal tubule. Among other examined normal tissues, gp200 expression is observed along the luminal surfaces of breast lobules and ducts, the luminal surface of the epididymal tubule epithelium, within the cytoplasm of parathyroid chief cells, and focally within the colloid of thyroid follicles. No expression or cross-reactivity is detected in 31 other normal tissue types. Notably, gp200 is expressed in 93% of primary and 84% of metastatic renal cell carcinomas. The monoclonal antibody 66.4.C2 may thus serve as a valuable tool for identifying carcinomas exhibiting proximal nephron differentiation, particularly those with tubular epithelial features.

Material Supplied

RCC antibody is affinity purified and diluted in PBS, pH 7.4, containing 1% BSA and 0.09% sodium azide.

Material required But Not Supplied

- Xylene
- DI Water
- Control Tissues
- Isopropyl alcohol
- Antigen retrieval buffers
- Hematoxylin
- Positive charged slides
- Blocking Reagents
- Mounting media
- Wash Buffer
- Detection System
- Cover glass

Working Reagent Procedure

- Ready-to-Use antibodies have been optimized for use with the recommended protocols and should not require further dilution.
- Concentrated antibodies must be diluted in accordance with the recommended protocol.

Recommended Protocol

Refer the following table for the details on specific recommended protocol for this antibody.

ORDERING INFORMATION	
CATALOG#	DESCRIPTION
DH228-01C	0.1 ML Concentrated Antibody Vial
DH228-05C	0.5 ML Concentrated Antibody Vial
DH228-1C	1 ML Concentrated Antibody Vial
DH228-3R	3 ML Ready-to-Use Antibody Vial
DH228-6R	6 ML Ready-to-Use Antibody Vial
DH228-12R	12 ML Ready-to-Use Antibody Vial

Control Tissue	Normal kidney or renal cell Carcinoma
Dilution factor	1:20-50 (Antibody Diluent: DH144)

Antibody Incubation Time	30-60 Minutes at RT
Retrieval Pre-treatment	Tris-EDTA based HIER (AR9 Buffer: DH020)

Precautions

This product should be used by qualified and trained professional users only.

Avoid microbial contamination of reagents to minimize non-specific staining. Never pipette reagents by mouth. Avoid contact of reagents and specimens with skin. If reagents or specimens come into contact with sensitive area, wash with sufficient amounts of water. Dispose of the unused reagents. This kit contain sodium azide at concentrations of less than 0.1%. Sodium azide is not classified as a hazardous chemical at these concentrations, but proper handling protocols should be observed. For more information on product hazards, precautions and waste disposal, *Material Safety Data Sheets* are available upon request.

Limitations

Improper tissue handling and processing prior to immunostaining can lead to inconsistent results. Variations in embedding and fixation or the nature of the tissue may lead to variations in results. Endogenous peroxidase activity or pseudo peroxidase activity in erythrocytes and tissue biotin may result in non-specific staining based on the detection system employed. Tissues containing Hepatitis B Surface Antigen (HBsAg) may give false positive with horseradish peroxidase systems. Improper counterstaining and mounting may compromise the interpretation of results. Interpretation of the staining result is solely the responsibility of the user. Experimental results should be confirmed by a medically-established diagnostic product or procedure. Evaluation must be performed by a qualified pathologist.

Troubleshooting

For Technical Support contact us at +91 - 7506501122 or info@dygnova.com or your local distributor to report unusual staining.

Doc No: DH/DS/PSA228Rev.00

	Manufacturer Details		Use by Date	LOT	Lot/Batch Number
	Manufacturing Date		Consult Instructions for Use	REF	Catalogue Number
	Temperature Limits		Sufficient for 'n' assays / tests	IVD	In-vitro Diagnostic Medical Device