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### anti-TJP1 antibody (AA 1551-1702)



7

Images



**Publications** 



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Quantity:	100 μL
Target:	TJP1
Binding Specificity:	AA 1551-1702
Reactivity:	Human, Mouse, Rat, Pig, Cow
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This TJP1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Flow Cytometry (FACS), Immunohistochemistry (Paraffinembedded Sections) (IHC (p)), Immunofluorescence (Paraffin-embedded Sections) (IF (p)), Immunofluorescence (Cultured Cells) (IF (cc)), Immunohistochemistry (Frozen Sections) (IHC (fro))

#### **Product Details**

Immunogen:	KLH conjugated synthetic peptide derived from human ZO-1
Isotype:	IgG
Cross-Reactivity:	Cow, Human, Mouse, Pig, Rat
Predicted Reactivity:	Dog,Cow,Chicken,Rabbit,Guinea Pig
Purification:	Purified by Protein A.

#### **Target Details**

Target: TJP1

### **Target Details**

Alternative Name:	ZO-1 (TJP1 Products)
Background:	Synonyms: ZO-1, Tight junction protein ZO-1, Tight junction protein 1, Zona occludens protein
	Zonula occludens protein 1, TJP1, ZO1
	Background: The N-terminal may be involved in transducing a signal required for tight junction
	assembly, while the C-terminal may have specific properties of tight junctions. The alpha
	domain might be involved in stabilizing junctions. Plays a role in the regulation of cell migration
	by targeting CDC42BPB to the leading edge of migrating cells.
Gene ID:	7082
UniProt:	Q07157
Pathways:	Carbohydrate Homeostasis, Cell-Cell Junction Organization
Application Details	
Application Notes:	WB 1:300-5000
	ELISA 1:500-1000
	FCM 1:20-100
	IHC-P 1:200-400
	IHC-F 1:100-500
	IF(IHC-P) 1:50-200
	IF(IHC-F) 1:50-200
	IF(ICC) 1:50-200
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 μg/μL
Buffer:	0.01M TBS( pH 7.4) with 1 % BSA, 0.02 % Proclin300 and 50 % Glycerol.
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be
	handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Expiry Date:

12 months

#### **Publications**

Product cited in:

Zhuang, Zhong, Bian, Fan, Chen, Liu, Liu: "Rhein ameliorates lipopolysaccharide-induced intestinal barrier injury via modulation of Nrf2 and MAPKs." in: **Life sciences**, Vol. 216, pp. 168-175, (2019) (PubMed).

Kulhankova, Kinney, Stach, Gourronc, Grumbach, Klingelhutz, Salgado-Pabón: "The Superantigen Toxic Shock Syndrome Toxin-1 Alters Human Aortic Endothelial Cell Function." in: **Infection and immunity**, (2018) (PubMed).

Zou, Lin, Li, Wu, He, Huang, Wang, Ye, Cheng, Ding, Zheng, Chi: "Huangqin-tang ameliorates dextran sodium sulphate-induced colitis by regulating intestinal epithelial cell homeostasis, inflammation and immune response." in: **Scientific reports**, Vol. 6, pp. 39299, (2018) (PubMed).

Zhao, Qin, Han, Wang, Zhang, Liu: "?-Conglycinin reduces the tight junction occludin and ZO-1 expression in IPEC-J2." in: **International journal of molecular sciences**, Vol. 15, Issue 2, pp. 1915-26, (2014) (PubMed).

Ruan, Liu, Zhou, Mi, Liu, Wu, Yao, Assaad, Deng, Hou, Wu, Yin: "Chlorogenic acid decreases intestinal permeability and increases expression of intestinal tight junction proteins in weaned rats challenged with LPS." in: **PLoS ONE**, Vol. 9, Issue 6, pp. e97815, (2014) (PubMed).

There are more publications referencing this product on: Product page





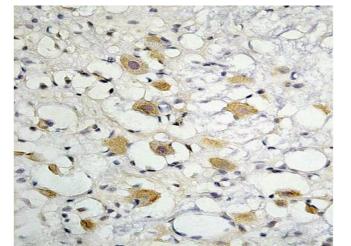






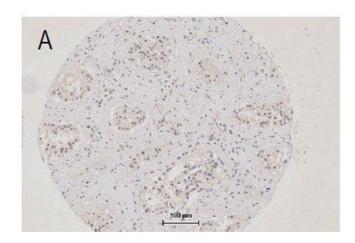
#### **Immunohistochemistry (Paraffin-embedded Sections)**

**Image 1.** HQT regulates epithelial proliferation in the colonic mucosa of mice with DSS-induced acute and chronic colitis.Mice were administered regular water (control) or 3.5 % DSS for 7 days followed by treatment with HQT for 7 days. (a-c) Immunohistochemical analysis of tight junction proteins occluding, ZO-1 and proliferating cells detected based on Ki67 in colon sections in DSS-induced acute colitis mice. (magnification, x200). Graphical representation of the percentage of occluding, ZO-1 and Ki67-positive cells in the mid-colon. Results are expressed as means±SD of three independent experiments, n=8 mice per group. \*p<0.05, \*\*p<0.001 vs. the control group, p<0.05, p<0.001 vs. DSStreated mice. Mice received three cycles of DSS treatment (2.5 %), each cycle consisting of 7 days of water containing DSS followed by 14 days of tap water, followed by treatment with HQT for 7 days. (d) Immunohistochemical analysis of proliferating cells detected based on Ki67 in colon sections in mice with DSS-induced chronic colitis (magnification, x200). Graphical representation of the percentage of Ki67positive cells in the mid-colon. Results are expressed as means±SD of three independent experiments, n=8 mice per group. \*p<0.05, \*\*p<0.001 vs. the control group, p<0.05, p<0.001 vs. DSS-treated mice. - figure provided by CiteAb. Source: PMID27982094



#### **Immunohistochemistry**

**Image 2.** Formalin-fixed and paraffin embedded rat brain labeled with Anti- ZO-1 Polyclonal Antibody, Unconjugated (ABIN675024) followed by conjugation to the secondary antibody and DAB staining



#### **Immunohistochemistry**

**Image 3.** Image kindly submitted by Kamlesh Gupta as part of the free sample program. Caco-2 cell lysate and mouse kidney lysate probed with ZO-1 primary antibody (ABIN675024) at 1:500 overnight at 4 °C. Followed by incubation with anti-rabbit HRP-conjugated secondary antibody at 1:5000 dilution for 1h at 25 °C. Predicted and observed band size: ~200kDa.

Please check the product details page for more images. Overall 7 images are available for ABIN675024.





#### Successfully validated (Immunohistochemistry (IHC))

by Immunohistochemistry Core, NYU Langone

Report Number: 029577

Date: Jan 18 2014

Lot Number:	130913
Method validated:	Immunohistochemistry (IHC)
Positive Control:	Testis
Negative Control:	See Controls section for negative controls
Notes:	Signal was detected in positive control tissue, and not detected in a tissue expected to express very low levels of the target antigen.
Primary Antibody:	- Antibody: human Tight Junction Protein 1 (Zona Occludens 1) (TJP1) - Catalog number: ABIN675024 - Supplier: Bioss - Supplier number: bs-1329r - Lot number: 130913
Secondary Antibody:	- Antibody: Biotinylated goat anti-rabbit/anti-mouse (Kit) - Catalog number: 760-091 - Supplier: Ventana Medical Systems - Lot number: D05923BA
Isotype:	- Antibody: Rabbit IgG isotype control - Catalog number: 790-4795 - Supplier: Ventana Medical Systems - Lot number: C11487
Controls:	<ul> <li>Tissues stained came from a human formalin-fixed paraffin embedded (FFPE) tissue microarray (12-003d):</li> <li>Positive control (specimen known to contain the target protein): human testis, which is expected to express high levels of the antigen.</li> <li>Negative Control (specimen known to not contain the target protein): Indeterminate; protein located on cytoplasmic membrane surface of intercellular tight junctions. The protein may be involved in signal transduction at cell-cell junctions. Expression is expected to be widespread. Thymus, expected to express much lower levels of the antigen as compared to testis, is shown to demonstrate a tissue with no detectable expression.</li> <li>Primary antibody isotype control: Testis (specimen known to contain the target protein) treated with primary antibody isotype control instead of the primary antibody.</li> <li>Secondary antibody only control: Testis (specimen known to contain the target protein) treated with secondary antibody only (no primary antibody).</li> </ul>
Protocol:	<ul> <li>Immunohistochemistry was performed on a Ventana NexES automated platform, instrument manufacturer specific reagents are italicized.</li> <li>1. Slides were preheated in convection oven at 60°C for 30 minutes</li> </ul>

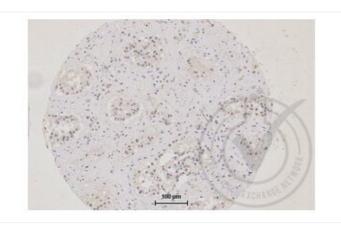
- 2. Deparaffinization procedure: 3 changes of Xylene, 5 minutes each 3 changes of 100% Ethanol, 3 minutes each - 3 changes of 95% Ethanol, 3 minutes each - Rinsed in distilled water, 3 changes
- 1. Heat retrieval procedure Slides retrieved in 10.0 mM Citrate, pH6.0 in a 1000W microwave oven (~100°C) for 15 minutes. - Slides were allowed to cool (in citrate) for 30 minutes. -Slides were washed x 3 in Distilled water
- 1. NexES instrument procedure, iVIEW DAB paraffin protocol (\*abridged\*): Slide chamber warmed to 37°C
- 1. Slides rinsed with \*reaction buffer\* x 3
- 1. \*iVIEW Inhibitor (H2O2)\* applied and incubated for 4 minutes
- 1. Slides rinsed with \*reaction buffer\*
- 1. Antibody Application Primary antibody diluted 1:250 in PBS (100 microliters applied/slide) - Ventana Isotype control applied neat - Slides incubated overnight at room temperature (~12 hours ~25°C)
- 1. Slides rinsed with \*reaction buffer\* x3
- 1. \*iVIEW Biotinylated IgG\* applied and incubated for 8 minutes
- 1. Slides rinsed with \*reaction buffer\*
- 1. \*iVIEW Streptavidin-Horseradish Peroxidase\* applied and incubated for 8 minutes
- · 1. Slides rinsed with \*reaction buffer\*
- 1. \*iVIEW DAB/H2O2\* applied and incubated for 8 minutes
- · 1. Slides rinsed with \*reaction buffer\*
- 1. \*iVIEW Copper\* applied and incubated for 4 minutes
- · 1. Slides rinsed with \*reaction buffer\*
- 1. Slides washed in Dawn Detergent/tap water
- 1. Counterstain Procedure Hematoxylin (Leica 560 MX) 30 seconds Slides washed in tap water, 1 minute - Decolorized (10% Acetic Acid in 70% ethanol), 1 minute - Slides washed in tap water, 1 minute - Bluing (Austin Clear Ammonia), 1 minute - Slides washed in tap water, 1 minute
- 1. Dehydration/coverslipping procedure: 3 changes of 95% Ethanol, 3 minutes each 3 changes of 100% Ethanol, 3 minutes each - 3 changes of Xylene, 5 minutes each - Mounted with Permount
- 1. Imaging: Leica SCN 400F Whole Slide Scanner with Digital Image Hub and Leica Slidepath software

#### **Experimental Notes:**

- Deviations from protocol/procedure supplied by manufacturer (attached).
- Step 1: Heated tissue 60°C for 30 minutes; manufacturer heats for 45 minutes.
- · Step 2: No ethanol wash was performed during deparaffinization; manufacturer includes 1 wash of 80% ethanol for 3 minutes.
- Step 3.1: Slides were heated for 15 minutes; manufacturer provides a range of 15-20 minutes.
- Step 3.2: Slides were cooled for 30 minutes; manufacturer cools for 20 minutes.
- Step 4: Italicized reagents and incubation time are fixed instrument parameters.
- Step 5: Secondary species-specific serum block not used; manufacturer blocks with 5% normal goat serum for 2 hours.

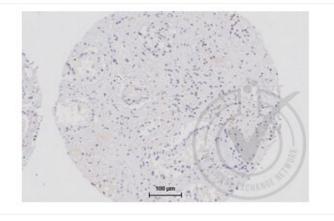
- Step 8.1: Antibody diluted in PBS at 1:250; manufacture did not recommend diluent or dilution.
- Step 8.2.1: Primary antibody incubated at room temperature overnight; manufacturer incubates overnight 4°C with agitation.
- Tissue Interpretation (limited): TJP1: Under the staining parameters described above, testis stained weakly (ducts) positive (Figure 1). Substantial signal detected in limited number of other tissues, including: breast, normal (NOS); pancreatic cancer (NOS), and stomach, normal (NOS). Most tissues showed low level of specific signal. Thymus did not have any detectable signal (Figure 4).
- · I-NC (Isotype negative control): No signal detected
- B-NC (Blank negative control): No signal detected
- · Signal Localization: Signal to noise was adequate with cytoplasmic, nuclear subcellular localization observed. Rare inner-membrane and no distinct membrane signal observed.

#### Images for Validation report #029577



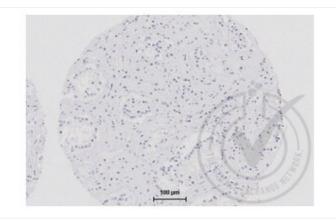
Validation image no. 1 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 1: TJP1 immunostaining of human testis (brown). Counterstain in blue.



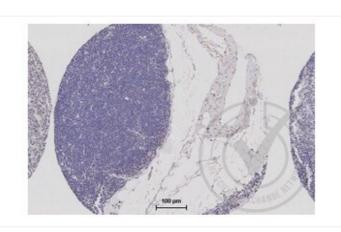
Validation image no. 2 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 2: Isotype control immunostaining of human testis (brown). Counterstain in blue.



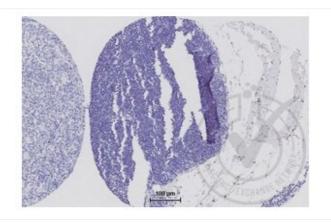
### Validation image no. 3 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 3: Secondary only control immunostaining of human testis (brown). Counterstain in blue.



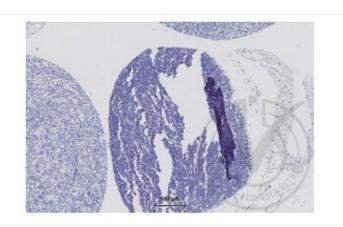
## Validation image no. 4 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 4: TJP1 immunostaining of human thymus (brown). Counterstain in blue.



# Validation image no. 5 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 4: Isotype control immunostaining of human thymus (brown). Counterstain in blue.



### Validation image no. 6 for anti-Tight Junction Protein 1 (TJP1) (AA 1551-1702) antibody (ABIN675024)

Figure 6: Secondary only immunostaining of human thymus (brown). Counterstain in blue.