### bs-0448R

- DATASHEET -

## [ Primary Antibody ]

Isotype: IgG

# **NIS Rabbit pAb**

Host: Rabbit



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Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500)

Reactivity: Mouse, Rat (predicted: Human)

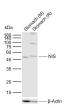
Predicted MW.: <sup>68 kDa</sup>

Subcellular Location: Cytoplasm

Clonality:	Polyclonal		
GenelD:	6528	SWISS: Q92911	
Target:	NIS		
Immunogen:	1: KLH conjugated synthetic peptide derived from human NIS: 525-618/618. < Cytoplasmic >		
Purification: affinity purified by Protein A			
Concentration: 1mg/ml			
Storage:	0.01M TBS (pH7.4) with 1% BSA Glycerol. Shipped at 4°C. Store at -20°C for freeze/thaw cycles.	, ,	
Background:	catalyzes Na+/I- symporter acti and thyroid hormone generatic	vity plays a role in iodide transport on.	

and thyroid hormone generation. Human Sodium lodide Symporter (hNIS) is responsible for iodide concentrating ability within thyroid follicular cells. It is a membrane bound glycoprotein with 13 membrane spanning domains and 14 extramembranous domains. It may represent an autoantigen in thyroid.

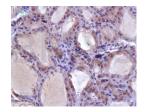
### — VALIDATION IMAGES



Sample: Lane 1: Mouse Stomach tissue lysates Lane 2: Rat Stomach tissue lysates Primary: Anti-NIS (bs-0448R) at 1/200 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 68 kDa Observed band size: 48 kDa



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0 ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-NIS Polyclonal Antibody, Unconjugated(bs-0448R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Tissue/cell: rat thyroid gland; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer ( 0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-NIS Polyclonal Antibody, Unconjugated(bs-0448R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining

### - SELECTED CITATIONS -------

- [IF=6.291] Siying Tang. et al. GLIS3 mediated by the Rap1/PI3K/AKT signal pathway facilitates real-ambient PM2.5 exposure disturbed thyroid hormone homeostasis regulation. Ecotox Environ Safe. 2022 Mar;232:113248 WB ;MOUSE. 35093813
- [IF=5.467] Jiang B.. et al. Primary human thyrocytes maintained the function of thyroid hormone production and secretion in vitro. J ENDOCRINOL INVEST. 2023 May;:1-12 ICC ;Human. 37133653
- [IF=4.872] Dong X et al. PM2.5 disrupts thyroid hormone homeostasis through activation of the hypothalamicpituitary-thyroid (HPT) axis and induction of hepatic transthyretin in female rats 2.5Ecotoxicol Environ Saf.2021 Jan 15;208:111720. IHC,WB ;Rat. 33396051

- [IF=4.848] Shasha Hou. et al. Downregulation of miR-146b-3p Inhibits Proliferation and Migration and Modulates the Expression and Location of Sodium/Iodide Symporter in Dedifferentiated Thyroid Cancer by Potentially Targeting MUC20. Front Oncol. 2020; 10: 566365 WB,IF ;Human. 33489878
- [IF=4.488] Niu Mengda. et al. Evaluation of [18F]tetrafluoroborate as a Potential PET Imaging Agent in a Sodium Iodide Symporter-Transfected Cell Line A549 and Endogenous NIS-Expressing Cell Lines MKN45 and K1. Mol Imaging. 2022;2022:2679260 WB ;Human. 10.1155/2022/2679260