

# Product Datasheet

## TetR Antibody - BSA Free NB600-234

Unit Size: 0.1 ml

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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**NB600-234**

TetR Antibody - BSA Free

**Product Information**

<b>Unit Size</b>	0.1 ml
<b>Concentration</b>	1.0 mg/ml
<b>Storage</b>	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
<b>Clonality</b>	Polyclonal
<b>Preservative</b>	0.02% Sodium Azide
<b>Isotype</b>	IgG
<b>Purity</b>	Immunogen affinity purified
<b>Buffer</b>	PBS and 30% Glycerol

**Product Description**

<b>Host</b>	Rabbit
<b>Gene ID</b>	4924774
<b>Species</b>	Non-species specific
<b>Immunogen</b>	A synthetic peptide made to an internal portion of Escherichia coli TetR protein (between residues 25-75) [UniProt P04483]

**Product Application Details**

<b>Applications</b>	Western Blot, ELISA
<b>Recommended Dilutions</b>	Western Blot 1:100, ELISA 1:100-1:2000
<b>Application Notes</b>	This TetR antibody is useful for Western Blot and ELISA.

**Images**

Western Blot: TetR Antibody [NB600-234] - Analysis of TetR using 23 kDa recombinant protein. Note the presence of a dimer.

250>  
150>  
100>  
75>  
50>  
37>  
25>  
20>  
15>  
10>

**Publications**

Kobylarz MJ, Goodwin JM, Kang ZB et al. An iron-dependent metabolic vulnerability underlies VPS34-dependence in RKO cancer cells PLoS ONE 2020-08-24 [PMID: 32833964]

Vercauteren K, Pasko RA, Gleyzer N et al. PGC-1-related coactivator: immediate early expression and characterization of a CREB/NRF-1 binding domain associated with cytochrome c promoter occupancy and respiratory growth. Mol Cell Biol. 2006-10-01 [PMID: 16908542] (WB)

Details:

WB (TR-expressing and wild-type control BALB/3T3 cells): Fig 8. Disease-specific gene repositioning in breast cancer.FISH, Table 1 and various figures throughout the publication.

## Procedures

### Serum protocol for TetR Antibody (NB600-234)

TetR Antibody:

Western Blot Protocol

1. Perform SDS-PAGE on samples to be analyzed, loading 40 ug of total protein per lane.
2. Transfer proteins to membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
3. Stain according to standard Ponceau S procedure (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
4. Rinse the blot.
5. Block the membrane using standard blocking buffer for at least 1 hour.
6. Wash the membrane in wash buffer three times for 10 minutes each.
7. Dilute primary antibody in blocking buffer and incubate 1 hour at room temperature.
8. Wash the membrane in wash buffer three times for 10 minutes each.
9. Apply the diluted HRP conjugated secondary antibody in blocking buffer (as per manufacturers instructions) and incubate 1 hour at room temperature.
10. Wash the blot in wash buffer three times for 10 minutes each (this step can be repeated as required to reduce background).
11. Apply the detection reagent of choice in accordance with the manufacturers instructions.

Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%.

\*The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.





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### **Products Related to NB600-234**

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NB600-234PEP	TetR Antibody Blocking Peptide
HAF008	Goat anti-Rabbit IgG Secondary Antibody [HRP]
NB7160	Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]
NBP2-24891	Rabbit IgG Isotype Control

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### **Limitations**

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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