

Anti-JMY [HMY117]

Catalogue number: 152747

Sub-type: Primary antibody

Images:

Contributor

Inventor: Helen Turley

Institute: University of Oxford

Images:

Tool details

***FOR RESEARCH USE ONLY**

Name: Anti-JMY [HMY117]

Alternate name:

Class: Monoclonal

Conjugate: Unconjugated

Description: The anti-JMY [HMY117] antibody recognises uman JMY. JMY is a p300-binding protein with dual action: by enhancing P53 transcription in the nucleus, it plays an important role in the cellular response to DNA damage, while by promoting actin filament assembly in the cytoplasm; it induces cell motility in vitro. It might act either as tumor suppressor or as oncogene. This antibody may be used in research but also as a diagnostic tool.

Purpose:

Parental cell:

Organism:

Tissue:

Model:

Gender:

Isotype: IgG1

Reactivity: Human

Selectivity:

Host: Mouse

Immunogen: Synthetic peptide from the c terminus of the peptide sequence, identical for both isoform 1 and 2

Immunogen UNIPROT ID:

Sequence:

Growth properties:

Production details:

Formulation:

Recommended controls: MCF7, HeLa

Bacterial resistance:

Selectable markers:

Additional notes:

Target details

Target: JMY

Target alternate names:

Target background: The anti-JMY [HMY117] antibody recognises uman JMY. JMY is a p300-binding protein with dual action: by enhancing P53 transcription in the nucleus, it plays an important role in the cellular response to DNA damage, while by promoting actin filament assembly in the cytoplasm; it induces cell motility in vitro. It might act either as tumor suppressor or as oncogene. This antibody may be used in research but also as a diagnostic tool.

Molecular weight:

Ic50:

Applications

Application: IHC ; WB

Application notes:

Handling

Format: Liquid

Concentration: 1 mg/ml

Passage number:

Growth medium:

Temperature:

Atmosphere:

Volume:

Storage medium:

Storage buffer: PBS with 0.02% azide

Storage conditions: -15° C to -25° C

Shipping conditions: Shipping at 4° C

Related tools

Related tools:

References

References: Ward et al. 1996. Proc Natl Acad Sci U S A. 93(4):1524-8. PMID: 8643665.

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