Anti-Human Ataxin-3 [4B8]

Catalogue number: 156506 Sub-type: Primary antibody

Images:

Contributor

Inventor: Patrick Loll

Institute: Drexel University

Images:

Tool details

*FOR RESEARCH USE ONLY

Name: Anti-Human Ataxin-3 [4B8]

Alternate name: Machado-Joseph Disease (MJD) Protein 1; Spinocerebellar Ataxia Type 3

Class: Monoclonal

Conjugate: Unconjugated

Description: Ataxin-3 is a deubiquitinating enzyme involved in the clearance of misfolded proteins. The enzyme functions in proteasome degradation and autophagy. Mutated forms of Ataxin-3 can lead to misfolded protein aggregation which cause the neurodegenerative disorder spinocerebellar ataxia type-3 (SCA3), also known as Machado-Joseph disease.

ols.org

Purpose: Marker Parental cell: Organism: Tissue: Model:

Gender: Isotype:

Reactivity: Human

Selectivity: Host: Mouse

Immunogen: Recombinant Human Ataxin-3
Immunogen UNIPROT ID: ATX3_HUMAN

Sequence:

Growth properties: Production details:

Formulation:

Recommended controls: IgG1 kappa

Bacterial resistance:

Selectable markers: Additional notes:

Target details

Target: Ataxin-3

Target alternate names:

Target background: Ataxin-3 is a deubiquitinating enzyme involved in the clearance of misfolded proteins. The enzyme functions in proteasome degradation and autophagy. Mutated forms of Ataxin-3 can lead to misfolded protein aggregation which cause the neurodegenerative disorder spinocerebellar ataxia type-3 (SCA3), also known as Machado-Joseph disease.

Cancer Tools.org

Molecular weight: 42 kDa

Ic50:

Applications

Application: ELISA; IP **Application notes:**

Handling

Format: Liquid

Concentration: 0.9-1.1 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:

Cancer Tools.org