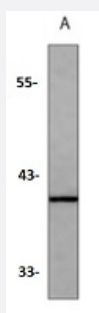


RecomAb™

Doublecortin recombinant monoclonal antibody

Catalog # RAB02406 Size 100 uL

Applications



Western Blot (Tissue lysate)

Western blot analysis of rat brain (A) whole cell lysates with Doublecortin recombinant monoclonal antibody (Cat # RAB02406).

Specification

Product Description	Rabbit recombinant monoclonal antibody raised against recombinant Doublecortin.
Antibody Species	Rabbit
Immunogen	Original antibody is raised against recombinant protein of human Doublecortin.
Theoretical MW (kDa)	41
Reactivity	Rat
Specificity	Recognizes endogenous levels of Doublecortin protein.
Form	Liquid
Purification	Immunogen affinity chromatography
Isotype	IgG
Recommend Usage	Western Blot (1:500-1:1000)
Storage Buffer	In 50mM Tris-Glycine, pH 7.4 (0.15M NaCl, 50% Glycerol, 0.01% Sodium azide and 0.05% BSA)

Storage Instruction

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

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

- Western Blot (Tissue lysate)

Western blot analysis of rat brain (A) whole cell lysates with Doublecortin recombinant monoclonal antibody (Cat # RAB02406).

Gene Info — DCX

Entrez GeneID[1641](#)**Protein Accession#**[Q9ESI7](#)**Gene Name**

DCX

Gene Alias

DBCN, DC, LISX, SCLH, XLIS

Gene Description

doublecortin

Omim ID[300067](#) [300121](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

In the developing cortex, cortical neurons must migrate over long distances to reach the site of their final differentiation. The protein encoded by this gene is a cytoplasmic protein which appears to direct neuronal migration by regulating the organization and stability of microtubules. The encoded protein contains two doublecortin domains, which bind microtubules. In addition, the encoded protein interacts with LIS1, the regulatory gamma subunit of platelet activating factor acetylhydrolase, and this interaction is important to proper microtubule function in the developing cortex. Mutations in this gene are a cause of X-linked lissencephaly. Multiple transcript variants encoding at least three different isoforms have been found for this gene. [provided by RefSeq]

Other Designations

OTTHUMP00000062892|doublecortex|doublin|lissencephalin-X

Disease

- [Autistic Disorder](#)
- [Epilepsy](#)

- [Genetic Predisposition to Disease](#)