

# DPYSL2 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02242

## Basic Information

**Catalog No.**

RM02242

**Category**

Cell Line

**Parental Cell line**

HeLa

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

DPYSL2

**Species**

Human

**Gene ID**

1808

**Swiss Prot**

Q16555

**Synonyms**CRMP-2; CRMP2; DHPRP2; DRP-2; DRP2;  
N2A3; ULIP-2; ULIP2

## Contact

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## Background

This gene encodes a member of the collapsin response mediator protein family. Collapsin response mediator proteins form homo- and hetero-tetramers and facilitate neuron guidance, growth and polarity. The encoded protein promotes microtubule assembly and is required for Sema3A-mediated growth cone collapse, and also plays a role in synaptic signaling through interactions with calcium channels. This gene has been implicated in multiple neurological disorders, and hyperphosphorylation of the encoded protein may play a key role in the development of Alzheimer's disease. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. [provided by RefSeq, Sep 2011]

## Product Information

**Description**

DPYSL2 Knockout cell line is engineered from HeLa cell line with Gene-Editing Technology.  
Allele-1:53bp deletion in exon2  
Allele-2:53bp deletion in exon2  
Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

**Packaging**

1 vial parental cell line and 1 vial knockout cell line

**Shipping Conditions**

Dry ice

**Amount**1~5x10<sup>6</sup> cells/vial**Storage**

Stored in liquid nitrogen for a long time less than -130°C. Minimizing freeze-thaw cycles.

**Protocol**

Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at 37°C with 5% CO<sub>2</sub> condition.

1. Thaw the vial in 37°C water bath, and shake it to melt as soon as possible.
2. Transfer the cell suspension to a 15mL conical tube with pre-warmed 5mL complete medium and centrifuge 1000rpm for approximately 5 minutes at room temperature.
3. Remove and discard the supernatant.
4. Resuspend the cell pellet with 1mL pre-warmed complete medium and seed in 10cm dish.
5. Add 8-10mL of complete medium.
6. Incubate the culture at 37°C incubator with 5% CO<sub>2</sub>.
7. A subcultivation ratio of 1:2-1:4 is recommended.

## Sequencing data

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WT CGAGGCCACTCCC\*\*\*\*\*CAGGGAATGACGTC  
Mut CGAGGCCACTCCC\*\*\*Deletion\*\*\*CAGGGAATGACGTC  
Allele-1: 53bp deletion in exon2  
WT CGAGGCCACTCCC\*\*\*\*\*CAGGGAATGACGTC  
Mut CGAGGCCACTCCC\*\*\*Deletion\*\*\*CAGGGAATGACGTC  
Allele-2: 53bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and DPYSL2 knockout (KO) HeLa cells, using sanger sequencing.