# MEF ULK1 KO (SIM) Cell Line

Catalogue number: 151715

Sub-type: Images:

#### Contributor

**Inventor:** Sharon Tooze

Institute: Cancer Research UK, London Research Institute: Lincoln's Inn Fields

Images:

#### **Tool details**

#### \*FOR RESEARCH USE ONLY

Name: MEF ULK1 KO (SIM) Cell Line

Alternate name:

Class:

Conjugate:

Cancer Tools.org **Description:** The MEF ULK1 KO (SIM) cell line can be used to study ULK1-dependent processes, including autophagy. A more complete phenotype requires depletion of ULK2 by RNAi . Mouse embryonic fibroblasts derived from a ULK1 homozygous knock out mouse embryo and immortalized by serial passaging (spontaneous transformation).

**Purpose:** Parental cell: Organism: Mouse Tissue: Embryo Model: Knock-Out

Gender: Isotype: Reactivity: Selectivity: Host:

Immunogen:

**Immunogen UNIPROT ID:** 

Sequence:

**Growth properties:** Autophagy, fibroblast

Production details: Primary embryonic fibroblasts were isolated from the embryos of a pregnant female Ulk1-/- mouse at day 13p.c. The MEFs were immortalised by SIM using a standard serial passaging protocol.

Formulation:

**Recommended controls: Bacterial resistance:** Selectable markers: Additional notes:

### **Target details**

Target: ULK1

**Target alternate names:** 

**Target background:** 

Molecular weight:

Ic50:

# **Applications**

**Application:** 

**Application notes:** 

## Handling

Format: Frozen **Concentration:** Passage number:

Growth medium: DMEM + 20% FCS + 2mM Glutamine + pen/strep

**Temperature: Atmosphere:** Volume:

Storage medium: Storage buffer: Storage conditions:

Shipping conditions: Dry ice

### Related tools

Related tools: MEF ULK1/2 WT (SIM) Cell Line; MEF ULK2 KO (SIM) Cell Line; MEF ULK1 ULK2 DKO (SIM) Cell Line; MEF ULK1 ULK2 DKO (SV40) Cell Line; MEF ULK1/2 WT(SV40) Cell Line; MEF ULK1 KO (SV40) Cell Line; MEF ULK2 KO (SV40) Cell Line

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#### References

**References:** McAlpine et al. 2013. Autophagy. 9(3):361-73. PMID: 23291478.; Regulation of nutrient-sensitive autophagy by uncoordinated 51-like kinases 1 and 2.; Chan et al. 2009. Mol Cell Biol. 29(1):157-71. PMID: 18936157.; Kinase-inactivated ULK proteins inhibit autophagy via their conserved C-terminal domains using an Atg13-independent mechanism.

