U2OS-CRISPR-SNAPf-Nup358/RanBP2

300663

GENERAL INFORMATION

Description

U2OS-CRISPR-SNAPf-Nup358/RanBP2 □□ osteosarcoma □□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ U2OS □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ RANBP2 (□□□□□□□□□□□□□□□□ □□□□□ NUP358) □□ CRISPR/Cas9 □□□□□□□□□□□□□□□□□□□□□□□□ SNAPf □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ Nup358/RanBP2 □□□□□□□□□□□□□□□□□□□□

GMO status details

GMO-S1: This human osteosarcoma cell line (U2OS-CRISPR-SNAPf-Nup358/RanBP2) contains a CRISPR-engineered SNAPf-Nup358/RanBP2 fusion enabling precise labeling of nuclear pore cytoplasmic fibrils. The modification is stably integrated. This classification applies only within Germany and may differ elsewhere.

Biosafety level	1

CHARACTERISTICS

Morphology	□□□□□□□□□□□□□□□□□□□□□□
Cell type	□□□□□□□□□□□□□□□□ (□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□)
Doubling time	□□□□□□ 24 □□□ 36 □□□□□□□

IDENTIFIERS

Citation

U2OS-CRISPR-SNAPf-Nup358/RanBP2 (□□□□□□□□□□□□□□□□□ Cytion 300663)

NCBI TaxID	9606
Cellosaurus	Not assigned (CRISPR-modified U2OS derivative; parental U2OS CVCL_0042)
Depositor	□□□□□□□□□□□□□□□□□□□□□□□□□□□□ (EMBL)

EXPRESSION AND SUSCEPTIBILITY

Protein expression

Nup358/RanBP2, SNAPf-□□□□

HANDLING AND CULTURE

Culture medium

McCoys 5a, w: 3.0 □□□□/□□□□ □□□□□□, w: □□□□□□ □□□□□□□□, w: 2.0 □□□□□□□□□□□ □□□□□□□□□□□□□□□, w: 2.2 □□□□/□□□□ NaHCO3 (□□□□□□□□□□□□□□ Cytion 820200a)

Incubation atmosphere	37°C, 5%CO2, □□□□□□□□□□□□□

Supplements

□□□ FBS 10%, □□□□□□ 3.0 □□□□/□□□□, □□□□□□□□□□□□□□□□□□, □□□□□□□□□□□□□□□ 2.0 □□□□□□□□□□□□, □□□□□□□□□□□□□□□□□□□□□□□□□□ 2.2 □□□□/□□□□, NEAA 1%

Dissociation reagent	□□□□□□□□

Subculturing

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Thawing and culturing

Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
Upon receipt, either store the cryovial immediately at temperatures below -150°C to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required.
For immediate culturing, swiftly thaw the vial by immersing it in a 37°C water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains.
Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently.
Centrifuge the mixture at 300 x g for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium.
Gently resuspend the cell pellet in 10 ml of fresh culture medium. For adherent cells, divide the suspension between two T25 culture flasks; for suspension cultures, transfer all the medium into one T25 flask to promote effective cell interaction and growth.
Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.

QUALITY CONTROL AND CRYOPRESERVATION

Sterility

□□ PCR □□□□□□□□□□□□□□□□□□□□

Warranty

□□□Cytion □□□□