

Product sheet

MDA-MB-175-VII | 305825

NCBI_TaxID 9606

CellosaurusAccession CVCL_1400

XXXXXXXXXX XXXX-XXXXXXXXXXXXXXXXXX

Isoenzymes AK-1, 1 ES-D, 1 G6PD, B GLO-I, 1-2 PGM1, 2 PGM3, 1-2

Tumorigenic X, X, XXXXXXXX XXXXXXXX XXXX 21 XXX XXXXXXXX XX 100% (5/5) XXXXXXXX XXXXXXXX XXXXXXXX XXX XX-XXXXXX 10(7) XXXX

Mutational profile XXXXXXXX: XXXXXXX XXXXX, NRG1 + HGNC, TENM4, XX/XXXXX=TENM4-NRG1, DOC4-NRG1, XXXX=XXXXXX

Karyotype XXXX XXX = 84; XXXX = 82 XX 89

XXXXXXXX

Culture Medium DMEM:Ham's F12 (1:1), w: 3.1 g/L XXXXXXX, w: 2.5 mM L-XXXXXXX, w: 15 mM HEPES, w: 0.5 mM XXXX XXXXXXX, w: 1.2 g/L NaHCO3 820400a)

Supplements XXXX XXXXX 10% FBS + XXXXXXXX (5 XXXXXXXXX/XX' X)

Dissociation Reagent XXXXXXX

Doubling time 112 XXXX

Fluid renewal 2 XX 3 XXXXXXX XXXXXXX

Freeze medium XXXXXXX XXXXXXX XXXXXXX, XXX XXXXXXX XXXXXXX XXXXXXX XXX (XXXX FBS) + 10% DMSO XXX XXXXXXX XXXXXXX XXXXXXX XXXXXXX, XX C

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Thawing and Culturing Cells

1. Thaw the vial rapidly in a water bath at 37°C. Do not allow the cells to reach room temperature. Transfer the cells to a pre-warmed medium.
2. Seed the cells into a pre-warmed medium in a 150 cm² flask. The cell density should be approximately 1.5 x 10⁶ cells per flask.
3. Incubate the cells in a humidified atmosphere of 5% CO₂ at 37°C. The medium should be changed every 3-4 days.
4. Harvest the cells when they reach 70-80% confluency.
5. Seed the cells into a 15 cm² flask at a density of 1.5 x 10⁶ cells per flask.
6. Harvest the cells when they reach 70-80% confluency.
7. Seed the cells into a 10 cm² flask at a density of 1.5 x 10⁶ cells per flask.
8. Harvest the cells when they reach 70-80% confluency.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Harvest cells at 70-80% confluency and freeze in a freezing medium.

Shipping Conditions Store at -78°C.

Storage Conditions Store at -150°C for up to 196 days.

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Sterility The cells are free of mycoplasmas and PCR detectable agents.