[illegible] ME-180

300196

[illegible] ME-180 [illegible] 66 [illegible] ME-180 [illegible]

[illegible] ME-180 [illegible] (TNF alpha) [illegible] ME-180 [illegible] (HPV) [illegible] HPV-68 [illegible] HPV-18 [illegible] HPV

[illegible] ME-180 [illegible] ME-180 [illegible] ME-180 [illegible]

GENERAL INFORMATION

Description

[illegible] ME-180 [illegible] 66 [illegible] ME-180 [illegible]

[illegible] ME-180 [illegible] (TNF alpha) [illegible] ME-180 [illegible] (HPV) [illegible] HPV-68 [illegible] HPV-18 [illegible] HPV

[illegible] ME-180 [illegible] ME-180 [illegible] ME-180 [illegible]

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Banking information
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IBAN: DE35 6725 0020 0009 2076 86
BIC: SOLADES1HDB

Tissue	มดลูก , ปากมดลูก
Disease	มะเร็งผิวหนังชนิดเอพิเดอร์โมอิด
Metastatic site	เยื่อไขมันในช่องท้อง

Synonyms

- Me-180
- ME 180
- ME180

CLASSIFICATION

Organism	Human
Sex	Female
Growth properties	Adherent
Biosafety level	2

CHARACTERISTICS

Morphology	ลักษณะคล้ายเยื่อบุผิว
Cell type	เยื่อบุผิว

IDENTIFIERS

Citation

ME-180 (หมายเลขแคตตาล็อก Cytion 300196)

NCBI TaxID	9606
Cellosaurus	[CVCL_1401]

EXPRESSION AND SUSCEPTIBILITY

Viruses

HPV68 เป็นบวก

HANDLING AND CULTURE

Culture medium

McCoys 5a, w: 3.0 กรัม /ลิตร กลูโค □, w: เ □ถียร กลูตามีน , w: 2.0 มิลลิโมลาร์ โซเดียมไพรูเวต , w: 2.2 กรัม / ลิตร NaHCO3 (หมายเลขบทความ Cytion 820200a)

Incubation atmosphere	37°C, 5%CO2, บรรยากาศชื้น

Supplements

เติมอาหารเลี้ยงเชื้อด้วย FBS 10%

Dissociation reagent	แอคคูเท □

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Subculturing

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Seeding density	1 x 10^4 □□□□□/□□.^²
Fluid renewal	2 □□□ 3 □□□□□□□□□□□□□□□□
Post-thaw recovery	□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□ 5 x 10⁴ □□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□ □□ □□□□□□□□□ 24 □□□□□□□

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

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Freeze medium	□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ (□□□□□□ FBS) + 10% DMSO □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□ CM-1 (□□□□□□□ □□□□□□□□□ Cytion 800100) □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□ □□□ □□□□

Storage conditions

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Shipping conditions

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DONOR

Donor age	66 □□
Ethnicity	□□□□□□□□□□

LEGAL AND SAFETY

Safety precautions

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Warranty

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Material Transfer Agreement required	Yes

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