[illegible] HL-60

300209

[illegible] HL-60 [illegible] 36 [illegible] promyelocytic [illegible]

[illegible] HL-60 [illegible] (DMSO) [illegible]

[illegible] HL-60 [illegible] c-myc [illegible] (TNF-alpha)[illegible] HL-60 [illegible]

[illegible] HL-60 [illegible] MAPK [illegible]

[illegible] HL-60 [illegible]

GENERAL INFORMATION

Description

[illegible] HL-60 [illegible] 36 [illegible] promyelocytic [illegible]

[illegible] HL-60 [illegible] (DMSO) [illegible]

[illegible] HL-60 [illegible] c-myc [illegible] (TNF-alpha) [illegible] HL-60 [illegible]

[illegible] HL-60 [illegible] MAPK [illegible]

[illegible] HL-60 [illegible]

Tissue	[illegible]
Disease	[illegible]

Applications

[illegible]

Synonyms

- HL 60
- HL.60
- HL60

CLASSIFICATION

Organism	Human
Sex	Female
Growth properties	Suspension
Biosafety level	1

CHARACTERISTICS

Morphology	[illegible]
Cell type	[illegible]

IDENTIFIERS

Citation

HL-60 (หมาย □ลบแคตตาล็อก Cytion 300209)

NCBI TaxID	9606
Cellosaurus	CVCL_0002

EXPRESSION AND SUSCEPTIBILITY

Receptors expressed

คอม □พล □ก □ ์ , □อฟ □ ี

Isoenzymes

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

Oncogenes

ไมค์พลัส

Reverse transcriptase	□ชิงลบ

Products

ตัวทำลาย□ □นื้องอก (TNF) หรือที่รู้จักในชื่อตัวทำลาย □ □นื้องอกอัลฟา (TNF-alpha, TNF alpha) หลังจากการกระตุ้นด้วยกรดฟอโบลไมริสติก

HANDLING AND CULTURE

Culture medium

RPMI 1640, w: 2.0 mM กลูตามีน □สถียร , w: 2.0 กรัม /ลิตร NaHCO3 (หมาย □ลบบทความ Cytion 820700a)

Incubation atmosphere	37°C, 5%CO2, บรรยากาศชื้น

Supplements

□ติมอาหาร □ลี้ยง □ชื้อด้วย FBS ที่ทำให้ไม่□ทำงานทางชีวภาพด้□วยความร้อน 10%

Subculturing

รักษาวัฒนธรรมโดยการ □ติมหรือ □ปลี่ยนอาหาร □ลี้ยง □ชื้อ □ป็นระยะๆ □ริ่มต้นวัฒนธรรมด้วยความหนาแน่น 5 x 10^5 □ซลล์ /มล . และรักษาความ □ข้มข้นของ □ซลล์ให้อยู่ในช่วง 3 x 10^5 ถึง 1 x 10^6 □ซลล์ /มล . □พื่อการ □จริญ □ติบโตที่ดีที่สุด

Seeding density	2 x 10^5 □ซลล์ /มิลลิลิตร
Fluid renewal	2 ถึง 3 ครั้งต่อสัปดาห์

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.

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Banking information
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BIC: SOLADES1HDB

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QUALITY CONTROL AND CRYOPRESERVATION

Sterility

[illegible] PCR [illegible]

[illegible]

Freeze medium	[illegible] (FBS [illegible]) + 10% DMSO [illegible] CM-1 ([illegible] Cytion 800100) [illegible]

Storage conditions

[illegible] −150 [illegible] −196 °C [illegible] −80 °C [illegible]

Shipping conditions

[illegible] −78 °C [illegible]

DONOR

Donor age	36 [illegible]
Ethnicity	[illegible]

LEGAL AND SAFETY

Safety precautions

[illegible]

Warranty

[illegible]Cytion [illegible]
[illegible]
60 [illegible]
[illegible]
[illegible] [illegible]/[illegible]
[illegible]
<a href=""/Who-we-are/Warranty/"" hreflang=""x-default"">[illegible].

Material Transfer Agreement required	Yes

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