

CellosaurusAccession CVCL_0342

XXXXXXXXXX XXXX-XXXXXXXXXXXXXXXXXX

XXXXXXXXXX

Culture Medium	DMEM, w: 4.5 g/L D-glucose, w: 4 mM L-glutamine, w: 3.7 g/L NaHCO ₃ , w: 1.0 mM β-mercaptoethanol (Gibco Cytion 820300a)
Supplements	FBS 10%
Dissociation Reagent	Trypsin
Subculturing	Cells are grown until reaching confluence. Cells are trypsinized using Trypsin-EDTA solution, washed with PBS, and resuspended in DMEM supplemented with 10% FBS. Cells are seeded into new flasks at a density of 2 × 10 ⁴ cells per flask.
Seeding density	2 x 10 ⁴ cells/flask
Fluid renewal	Medium is changed every 3 days.
Freeze medium	For freezing, cells are harvested by trypsinization, washed with PBS, and resuspended in DMEM supplemented with 10% FBS + 10% DMSO. Cells are then seeded into cryovials and frozen.

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Thawing and
Culturing Cells

1. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
2. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
3. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
4. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
5. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
6. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
7. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。
8. 将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。

Incubation
Atmosphere

37°C, 5% CO₂, 饱和湿度

Flask Coating

无涂层

Freezing
Procedure

将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。

Shipping
Conditions

将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。

Storage
Conditions

将细胞悬液缓慢加入含有培养基的细胞培养瓶中，轻轻混匀，然后将培养瓶置于37°C CO₂培养箱中培养。

细胞培养与转染

Sterility

细胞培养与转染过程中，应严格遵守无菌操作规范，避免污染。PCR扩增过程中，应使用无菌试剂和耗材，避免污染。