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Product sheet

Hep-74.3A | 400208

Protein expression	8, 18, 18
Tumorigenic	, C3H/HE
Mutational profile	P53 wt
Culture Medium	Ham's F12, w: 1.0 mM , w: 1.0 mM , w: 1.1 g/L NaHCO3 (Cytion 820600a)
Supplements	10% FBS
Dissociation Reagent	
Subculturing	cells are seeded into 25 cm ² flasks in DMEM/F12 supplemented with 10% FBS. When cells reach confluence, they are trypsinized and seeded into new flasks. The medium is changed every 3-5 days. Cells are passaged every 2-3 weeks.
Seeding density	1 x 10 ⁴ cells/cm ²
Fluid renewal	3-5 times
Post-Thaw Recovery	cells are thawed and seeded into 25 cm ² flasks in DMEM/F12 supplemented with 10% FBS. The medium is changed every 3-5 days. Cells are passaged every 2-3 weeks.
Freeze medium	DMEM/F12 supplemented with 10% FBS + 10% DMSO

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

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

**Shipping
Conditions**

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

**Storage
Conditions**

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

细胞培养

Sterility

细胞培养液经过严格的灭菌处理，确保无菌。PCR 试剂和模板 DNA 均经过严格的灭菌处理，确保无菌。