

## HepG2 Cell Line

#Cat: NB-19-0060

Size:  $1 \times 10^6$  cells/vial

HepG2 is a cell line exhibiting epithelial-like morphology that was isolated from hepatocellular carcinoma of a 15-year-old, white, male youth with liver cancer. The cells express 3-hydroxy-3-methylglutaryl-CoA reductase and hepatic triglyceride lipase activities. The cells demonstrate decreased expression of apoA-I mRNA and increased expression of catalase mRNA in response to gramoxone (oxidative stress). There is no evidence of a Hepatitis B virus genome in this cell line.

### Characteristics

<b>Cell Name</b>	Hep G2 Cell Line
<b>Organism</b>	<i>Homo sapiens</i> , Human
<b>Morphology</b>	Epithelial-like
<b>Tissue</b>	Liver
<b>Cell type</b>	Hepatocellular
<b>Disease</b>	Carcinoma
<b>Growth Properties</b>	Adherent
<b>Derivation</b>	HepG2 was derived from a liver hepatocellular carcinoma of a 15 year old Caucasian male
<b>Age</b>	15 years adolescent
<b>Ethnicity</b>	White
<b>Gender</b>	Male
<b>Karyotype</b>	modal number = 55 (range = 50 to 60); has a rearranged chromosome 1
<b>Tumorigenic</b>	No
<b>STR profile</b>	<b>AMEL:</b> X, Y <b>CSF1PO:</b> 10,11 <b>D13S317:</b> 9,13 <b>D16S539:</b> 12,13 <b>D18S51</b> : 13,14 <b>D21S11</b> : 29,31 <b>D3S1358:</b> 15,16 <b>D5S818</b> : 11,12 <b>D7S820</b> : 10 <b>D8S1179:</b> 15,16 <b>FGA:</b> 22,25 <b>PENTA D:</b> 9,13 <b>PENTA E:</b> 15,20 <b>THO1:</b> 9 <b>TPOX:</b> 8,9 <b>vWA:</b> 17
<b>Biosafety Level</b>	1
<b>Product format</b>	Frozen
<b>Storage conditions</b>	Vapor phase of liquid nitrogen

## Culture conditions

<b>Complete medium</b>	The base medium for this cell line is Minimum Essential Medium (MEM) with stable glutamine (Catalog No. NB-58-0091). To make the complete growth medium, add fetal bovine serum to a final concentration of 10%
<b>Temperature</b>	37°C
<b>Atmosphere</b>	Air, 95%; CO <sub>2</sub> , 5%
<b>Subculturing</b>	Volumes are given for a 75 cm <sup>2</sup> flask, increase or decrease the amount of medium needed proportionally for culture vessels of other sizes <ol style="list-style-type: none"><li>1. Remove and discard culture medium</li><li>2. Briefly rinse the cell layer with DPBS solution to remove all traces of serum that contains trypsin inhibitor</li><li>3. Add 2 mL of Trypsin-EDTA solution to the flask and observe cells under an inverted microscope until cell layer is dispersed (usually within 5 to 10 minutes). Cells that are difficult to detach may be placed at 37°C to facilitate dispersal</li><li>4. Add 8 mL of complete growth medium and aspirate cells by gently pipetting</li><li>5. Add appropriate aliquots of the cell suspension to new culture vessels</li></ol>
<b>Seeding density</b>	2 x 10 <sup>4</sup> to 6 x 10 <sup>4</sup> viable cells/cm <sup>2</sup>
<b>Subculture ratio</b>	1:4-1:6
<b>Medium renewal</b>	Twice per week
<b>Cryopreservation</b>	Basal medium with 30% FBS and 10% (v/v) DMSO

## Handling information

<b>Unpacking</b>	<ol style="list-style-type: none"><li>1. Check all containers for leakage or breakage</li><li>2. Remove the frozen cells from the dry ice packaging and immediately place the cells at a temperature below -130°C, preferably in liquid nitrogen vapor, until ready for use</li></ol>
<b>Procedure</b>	To ensure the highest level of viability, thaw the vial and initiate the culture as soon as possible upon receipt. If upon arrival, continued storage of the frozen culture is necessary, it should be stored in liquid nitrogen vapor phase and not at -70°C. Storage at -70°C will result in loss of viability <ol style="list-style-type: none"><li>1. Quickly thaw the vial by gentle agitation in a 37°C water bath within 1-2 minutes</li><li>2. Remove the vial from the water bath as soon as the contents are thawed and decontaminate by dipping in or spraying with 70% ethanol. All operations from this point on should be carried out under strict aseptic conditions</li><li>3. Carefully open the vial and transfer the cell suspension to a centrifuge tube containing 8 mL complete culture medium (room temperature). Centrifuge at 300 x g for 5-7 minutes and carefully discard the supernatant containing residual freezing medium</li><li>4. Gently resuspend the cell pellet in 10 ml fresh complete medium and dispense into a 75 cm<sup>2</sup> culture flask</li><li>5. Incubate the culture at 37°C in a suitable incubator for 24-48 hours and check under the microscope. All further steps are described in the subculture section</li></ol>

## **Required Products**

These products are vital for the proper use of this item and have been confirmed as effective in supporting functionality. If you use alternative products, the quality and effectiveness of the item may be affected.

**Neo MEM with Earle's Salts, with Stable Glutamine** (NB-58-0091)

**Neo FBS, Collected in South America** (NB-58-0001A, NB-58-0001B)

**Neo Trypsin-EDTA (0.05 %) in DPBS (1x)** (NB-58-0108)

**Neo Freeze 1, Cryopreservation Medium with Fetal Bovine Serum** (NB-58-0065)

**Neo Dulbecco's PBS (1x), w/o Ca & Mg, w/o Phenol Red** (NB-58-0022)

## **Related Products**

**Human genomic DNA (Hep G2)** (NB-19-0060-DNA)