# DATA SHEET <br> Human Embryonic Hepatocyte Primary Cell Culture - T150 Plated Cells 

Catalog number: 33003-07-T150
Description:Human Embryonic Hepatocyte Primary Cell Culture T150 Plated Cells. Also available in T25, T75, and T225 tissue culture flask with plated cells, shipped at room temperature or in Frozen Vial, shipped in dry ice. The Clonal Human Embryonic Hepatocyte Primary Cell Culture was derived from Human Hepatocyte. They were maintained in Celprogen's Human Embryonic Hepatocyte Primary Cell Culture Complete Growth Medium and sub-cultured every 24 to 48 hours on Human Embryonic Hepatocyte Primary Cell Culture Extra-cellular Matrix.

Source: Human Hepatocyte
Donors: All donors from which the cells were derived were pre-screened; donors tested negative for the usual blood donation infectious disease panel ABO/RH, Hepatitis B Surface Antigen, HIV1 and 2, Syphilis, hepatitis B core, Human T Lymphocyte Virus 1and 2, Hepatitis C Virus, Antibody Screen, Nucleic Amplification Test for HIV 1, HCV, West Nile Virus and Antibodies to Trypanosoma cruzi (the agent of Chagas disease).

Mycoplasma test: Negative-PCR and mycoplasma agar methods
Sterility: $\quad$ Negative for bacteria, yeast, and mold
Storage Conditions: Liquid nitrogen vapor phase for frozen Ampule of Human Embryonic Hepatocyte Primary Cell Culture. For plated cells in tissue culture flask, upon receipt of the cells wipe the flask with $70 \%$ ethanol and transfer to sterile tissue culture hood. In the tissue culture hood remove the media from the cells and wash the cells with 1 X PBS sterile solution, for 2-3 minutes, remove the 1 X PBS solution and then Trypsinize. After Trypsinization of the Cells neutralize the Trypsin with equal volume of Human Embryonic Hepatocyte Primary Cell Culture Complete Growth Media with Serum and collect the Cell suspension in sterile conical centrifuge tube in the tissue culture hood. Centrifuge the cell suspension at 100 g for 7 minutes in centrifuge. Plate cells $5 \times 10^{5}$ cells per pre-coated flasks with Human Embryonic Hepatocyte Primary Cell Culture Extra-cellular Matrix for Expansion in Human Embryonic Hepatocyte Primary Cell Culture Complete Growth Medium.

## Positive Markers: CD133, Albumin, CK14, CK18, Alpha actin, alpha Fetal Protein, CYT450

Morphology \& Proliferation: Mixed population of cells with approximately $95 \%$ attached cells and the other $5.0 \%$ in suspension, need to change cell culture media every day after 48 hours of initial cell culture or when the media starts changing color to slight yellow from pink. Fast growing cell culture, change media with Celprogen's Human Embryonic Hepatocyte Primary Cell Culture Complete Growth Medium with the appropriate Human Embryonic Hepatocyte Primary Cell Culture Extra-cellular Matrix. Temperature $37^{\circ} \mathrm{C}$ in $5 \% \mathrm{CO}_{2}$ humidified incubator.

## Sub-culturing:

1. Remove flask and wipe with $70 \%$ ethanol. Then transfer to the tissue culture hood. Refer to protocols, flow diagrams and videos for more detail.
2. Transfer the supernatant contents to a sterile centrifuge tubes, this will contain detached cells.Centrifuge at 100 g for 7 minutes to obtain cell pellet. Plate cells in a T150 flask pre-coated with ECM.
3. Add pre-warmed Human Embryonic Hepatocyte Primary Cell Culture Growth Media to flask that has attached cells and incubate for two hours in the incubator. After 2 hours wash the cells with 1X PBS, remove $100 \%$ 1X PBS solution.
4. Trypsinize the attached cells with 3 ml 1 X Trypsin EDTA for 2-3 minutes. At the end of Trypsinization add equal volume of complete growth media with serum. Centrifuge the cells at 100 g for 7 minutes to obtain cell pellet.
5. Add the 500 ul of cells to T150 flask pre-coated with Human Embryonic Hepatocyte Primary Cell Culture Extracellular Matrix with 7 ml of Human Embryonic Hepatocyte Primary Cell Culture Complete Growth Medium. Depending on the size of the pellet you may utilize a new flask or you may combine the cells to the flask that had the supernatant cells.
6. Incubate the cells in the T150 flask at $37^{\circ} \mathrm{C}$ in a $5 \% \mathrm{CO}_{2}$ humidified incubator. Perform $100 \%$ Media Change every 24 to 48 hours.
7. Medium renewal every other day or 2-3 days, sub-culturing ratio: 1:2 or 1:3 depending on the cell density.
8. Refer to protocols, flow diagrams and videos for more detail. http://celprogen.com/tech.htm

Freezing Medium: Available for purchase Cat\# M33003-07FM
Trypsin: Available for purchase Cat\# T1509-014
IX PBS: Available for Purchase Cat\# P1408-013
Storage temperature: Liquid nitrogen vapor phase
Product Orders: Before submitting an order you will be asked to read and accept the terms and conditions of Celprogen's Material Transfer Agreement (MTA).

Permits/Forms: In addition to the MTA mentioned above, other CELPROGEN and/or regulatory permits may be required for the transfer of this CELPROGEN material. Anyone purchasing CELPROGEN material is ultimately responsible for obtaining the permits.

## Biosafety Level: 1

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