

Datasheet for HCT116/EGFR L858R cancer marker mutation cell lines (Homozygote)

Catalog number: SL703

Product: HCT116/EGFR L858R cancer marker mutation cell lines (Homozygote) with spike-in

signature at 849Q and 850H locus

Description: This product is a sgRNA and donor vectors (puromycin selection) for the EGFR L858R

genome editing cell line. This cell line can be used in vitro as a reference for gene

mutation detection or other related applications.

Spike-in*: with spike-in signature at 849Q and 850H locus

Quantity: 1 vial of 2 x 10⁶ cells; frozen

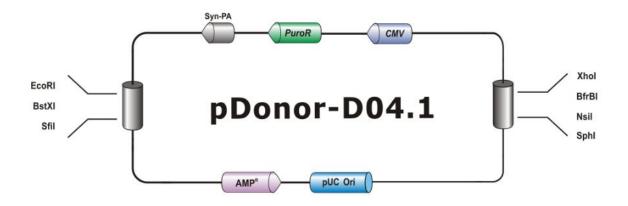
Shipping conditions: Dry ice

Storage conditions: Liquid nitrogen vapor phase. Remove the item from the dry ice packaging and

check all items for damage and leakage. Place immediately into storage at or

below -140 °C, preferably into the liquid nitrogen vapor phase, until use.

Transgene integration:



Source of parental line: HCT116

Organism: Homo sapiens, human

Tissue: colon

Cell Type: epithelial

^{*} IP protected by 2019111467778 which was owned by GeneCopoeia, Inc.



Quality control: >95% viability before freezing. All cells were tested and found to be free of

mycoplasma, bacteria, viruses, and other toxins.

Safety instructions: To ensure safety, protective gloves, clothing, and a face mask should be worn

when handling frozen vials. Some leakage may occur into the vial during storage. The liquid nitrogen will be converted to gas upon thawing. Due to the nature of nitrogen gas, pressure may build within the vial and possibly result in the vial

exploding or losing its cap. This may cause flying debris.

Thawing procedure: The vial of cells should be thawed in a 37 °C water bath with gentle agitation. For

optimal performance, the vial should be thawed in under two minutes. Ensure that the cap of the vial did not loosen upon thawing, and re-tighten as needed. Spray the vial with 70% EtOH and wipe off. Repeat. Using aseptic technique, add the contents of the vial to 9 ml of complete growth medium (without selection). Centrifuge for 5 min. at 250 x g. Aspirate the medium, being careful not to disturb the pellet. Resuspend in 10 mL of complete growth medium, and place into a culture vessel of your choice. Only add selection to the medium after 24

hours in culture.

Culture conditions:

Complete Growth Medium

The base medium for this cell line is RPMI1640. For optimal growth and maintenance of selection, add the following components to the base medium: fetal bovine serum to a final concentration of 10%.

Selection

puromycin to a final concentration of 0.6 μg/mL

Culture temperature:

37 °C with 5% CO₂

Subculture:

Replace culture medium with selection-free medium and incubate for up to 6 hours. Rinse the cells and split at 1 x 10^5 viable cells/mL to 1 x 10^6 cells/mL.



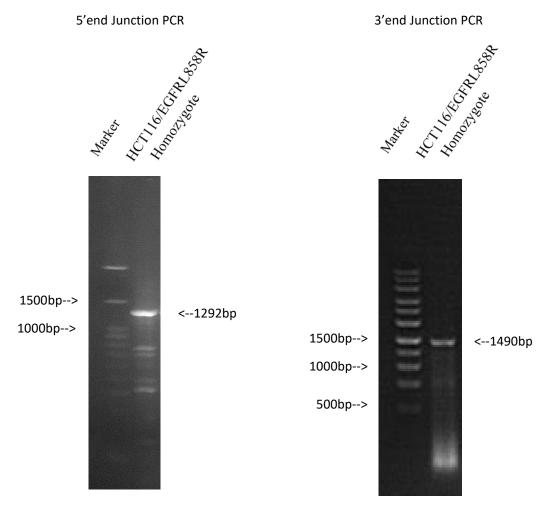
Cryopreservation: Freeze slowly in complete growth medium supplemented with 5% (v/v) DMSO.

Mycoplasma: Negative

(MycoAllert Mycoplasma Detection Kit from Lonza)

Product QC:

(a) Junction PCR result:



(b)Junction PCR positive statistics:

5'end Junction PCR positive amount (rate)	3'end Junction PCR positive amount (rate)	5'end and 3'end Junciton PCR double positive amount (rate)	Total sample amount
135(70.3%)	101(52.6%)	87(45.3%)	192



(c)FISH of EGFR probes on HCT cells:

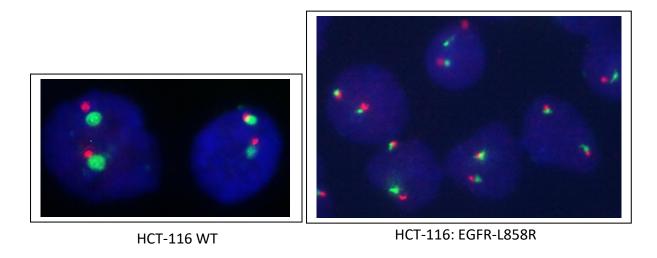
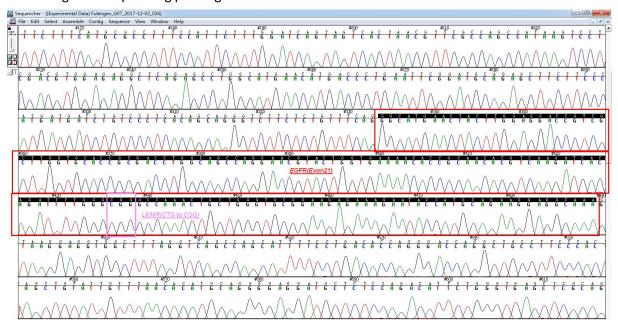


Fig.1: FISH of Human HCT-116 cells: GCI VividFISHTM FISH probes CEP7/EGFR were hybridized on HCT-116 wild type(WT) and HCT-116 EGFR-L858R mutated cells. The two green signals are from Chromosome 7 centromeres, while the two orange signals are representing the EGFR gene on chromosome 7. Both of them have two alleles of EGFR. The cell nuclear are counter stained with DAPI which is shown in blue.

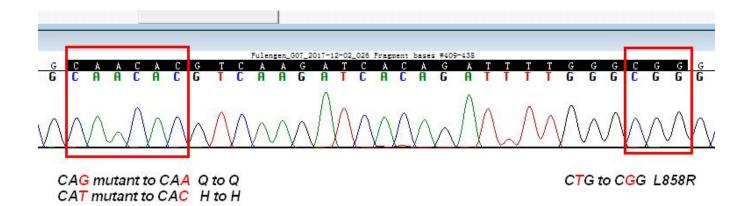
(d)DNA Sequencing peak figure:

EGFR L858R fragment sequencing peak figure:









Citation of product: If use of this item results in a publication, please use this information:

HCT116/EGFR L858R cancer marker mutation cell lines (SL703, GeneCopoeia,

Inc., Rockville, MD).

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