

14-538ACL: TLR9/HEK293 Stable Cell Line

Application : Functional Assay

Description

TLR9/HEK293 Stable Cell Line is a stably transfected HEK293 cell line which expresses human Toll-like receptor 9 (TLR9, also designated as CD289). TLR9 mediates immune responses by recognition of unmethylated CpGs of bacterial or viral origin, of which activation involves in various human diseases including systemic lupus erythematosus, erythema nodosum leprosum, inflammatory bowel disease, rheumatoid arthritis, sarcoidosis and Alzheimer's disease.

Sequence data: Human TLR9 (accession number NP_059138)

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MGFCRSALHPLSLLVQAIMLAMTLALGTLPAFLPCELQPHGLVN
CNWLFKSVPHFSMAAPRGVNTSLSLSSNRIHHLHDSDFAHLPRLHNLKWNCPVVG
LSPMHFPCMTIEPSTFLAVPTLEELNLSYNNIMTVPALPKSLISLSHTNILMLDS
ASLAGLHALRFLFMDGNCYKNPCRQALEVAPGALLGLGNLTHLSLYNNLTVVPRNL
PSSLEYLLLSYNRIVKLAPEDLANLTALRVLDVGGNCRRCDHAPNPCMECPRHFPQLH
PDTFSHLRLEGLVLKDSLSWLNASWFRGLGNLRVLDLSENFYKCYTKTKAFQGLT
QLRKLNLNFNYQKRVSAHLSLAPSGSLVALKELDMHGIFRSLDETTLRPLARLPM
LQTLRLQMFNINQAQLGIFRAFPGLRYVDLSNRISGASELTATMGEADGGEKVWLQP
GDLAPAPVDTPSSEDFRPNCTLNFTLDLSRNNLTVQPEMFAQLSHLQCLRLSHNCI
SQAVNGSQFLPLTGLQVLDLSHNKLDLYHEHSFTELPRLEALDSYNSQPFGMQGVGH
NFSFVAHLRTRLRHLSLAHNNIHSQVSQQLCSTSLRALDFSGNALGHMWAEGDLYLHFF
QGLSGLIWLDSLQNRHLTLPLQTLRNLPSKSLQVLRDLNYLAFFKWWSLHFLPKLEVL
DLAGNQLKALTNGSLPAGTRLRRLDVSCNSISFVAPGFFSKAKELRELNLSANALKTV
DHSWFGPLASALQILDVSANPLHCACGAAMDFLLEVQAAVPGPLSRVKCGSPGQLQG
LSIFAQDLRLCLDEALSWDCAFSLLAVALGLGVPMHLHLCGWDLWYCFHLCLAWLPW
RGRQSGRDEDALPYDAFVVDKTSQSAVADWVYNELRGQLEECRGRWALRLCLEERDWL
PGKTLFENLWASVYGSRKTLFVLAHTDRVSGLLRASFLAQRLLEDKDVVVLVILS
PDGRRSRYVRLRQRLCRQSVLLWPHQPSGQRSFWAQLGMALTRDNHHFYNRNFCQGPT
AE
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Product Info

Amount : 1 Vial
Content : Each vial contains $2 \sim 3 \times 10^6$ cells in 1 ml of 90% FBS + 10% DMSO
Storage condition : Immediately upon receipt, store in liquid nitrogen.

Application Note

Application:

- Functional assay.

Culture conditions:

Cells should be grown at 37°C with 5% CO₂ using DMEM medium (w/ L-Glutamine, 4.5g/L Glucose and Sodium Pyruvate) supplemented with 10% heat-inactivated FBS and 1% Pen/Strep, plus 10 µg/ml of Blasticidin.

It is recommended to quickly thaw the frozen cells upon receipt or from liquid nitrogen in a 37°C water-bath, transfer to a tube containing 10 ml of growth medium without Blasticidin, spin down cells, resuspend cells in pre-warmed growth medium without Blasticidin, transfer resuspended cells to T25 flask and culture in 37°C-CO₂ incubator.

Leave the T25 flask in the incubator for 1~2 days without disturbing or changing the medium until cells completely recover viability and become adherent. Once cells are over 90% adherent, remove growth medium and passage the cells through trypsinization and centrifugation. At first passage, switch to growth medium containing Blasticidin. Cells should be split before they reach complete confluence.

To passage the cells, detach cells from culture vessel with Trypsin/EDTA, add complete growth medium and transfer to a tube, spin down cells, resuspend cells and seed appropriate aliquots of cells suspension into new culture vessels. Subcultivation ratio = 1:10 to 1:20 weekly. To achieve satisfactory results, cells should not be passaged over 16 times.

LIMITED USE RESTRICTIONS:

THIS PRODUCT IS SOLELY FOR IN VITRO RESEARCH USE ONLY. NOT FOR DIAGNOSTIC OR THERAPEUTIC USE.

By use of this product, user agrees to be bound by the terms of this limited use statement.

This product is solely for Internal Research Purposes and not for Commercial Purposes. Commercial Purposes include, but are not limited to (1) use of the cell line in manufacturing; (2) use of the cell line to provide a service, information or data; (3) use of the cell line for therapeutic, diagnostic or prophylactic purposes; or (4) resale of the cell line whether or not such cell lines are resold for use in research. The buyer cannot sell, give or otherwise transfer this product to a third party.

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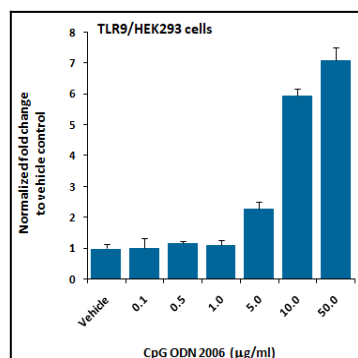


Fig-1: Functional analysis of the TLR9/HEK293 cell line. TLR9/HEK293 cells as well as parental HEK293 cells were transfected with NF-κB/Renilla luciferase reporter plasmid for 16 h. Cells were stimulated with different doses of CpG ODN-2006 (Abeomics, Cat. #15-1017) for 16 h followed by luciferase assay using Leeporterä Renilla Luciferase Assay Reagent (Abeomics, Cat. #17-1101).