
Cell Line Data Sheet for TC-32

Disease: Primitive neuroectodermal tumor (PNET)
Phase of Therapy: Diagnosis
Treatment: None
Disease Stage:
Gender: Female
Age at diagnosis: 31 months
Race: N/A
Age at sample collection: N/A
Source of Culture: Solid Tumor
Primary Tumor Site:
Date Established:

EWS/FLI1 Status: ERG
p53 functionality: Non-Functional
Karyotype: +5,+10,i(1q),t(11;22)(q24;q12)
Modal No: 48 (46-49)

IC90 (DIMSCAN*):

<u>VNC (ng/ml)</u>	<u>L-PAM (µg/ml)</u>	<u>ETOP (ng/ml)</u>	<u>RAP (ng/ml)</u>
N/A	N/A	N/A	N/A

VNC, vincristine; L-PAM, melphalan; ETOP, etoposide; RAP, rapamycin

Growth Conditions:
 Please see Protocols section at <https://www.cccells.org/protocols.php>
 5% CO₂, 20% O₂, 37.0°C

Media Formulation:
 Please see Protocols section at <https://www.cccells.org/protocols.php>
 Cells are grown in a base medium of Iscove's Modified Dulbecco's Medium plus the following supplements (to a final concentration): 20% Fetal Bovine Serum, 4mM L-Glutamine, 1X ITS (5 µg/mL insulin, 5 µg/mL transferrin, 5 ng/mL selenous acid)

Doubling Time:
Growth Properties: 24 hours
STR Profile: Teardrop-shaped cells with processes Adherent, grow mostly in clumps
 May be obtained at <https://strdb.cccells.org/>

Notes:

All COG Repository cell lines are antibiotic-free, mycoplasma-free, and cryopreserved in 50% FBS / 7.5% DMSO. Each vial label contains the cell line name, passage number, total viable cell count (usually 5-10e6), the overall cell viability, and date frozen. All cell lines are validated with original patient sample by STR analysis.



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References:

1. Reynolds CP, Brodeur GM, Tomayko MM, Donner L, Helson L, Seeger RC, Triche TJ: Biological classification of cell lines derived from human extra-cranial neural tumors. *Prog Clin Biol Res.*271:291-306, 1988. PubMed ID: 3406003
2. Wang Y, Einhorn P, Triche TJ, Seeger RC, Reynolds CP. Expression of Protein Gene Product 9.5 and Tyrosine Hydroxylase in Childhood Small Round Cell Tumors. *Clin Cancer Res.* 6, 551-558, 2000. PubMed ID: 10690538
<https://clincancerres.aacrjournals.org/content/6/2/551.long>
3. Batra S, Reynolds CP, Maurer BJ. Fenretinide cytotoxicity for Ewing's sarcoma (ES) and primitive neuroectodermal Tumor (PNET) cell lines is decreased by hypoxia and synergistically enhanced by ceramide modulators. *Cancer Research* 64: 5415-5424, 2004. PubMed ID: 15289350
<https://cancerres.aacrjournals.org/content/64/15/5415.long>
4. Thiele, Carol J., McKeon, Catherine, Triche, Timothy J., Ross, Robert A., Reynolds, C. Patrick, and Israel, Mark A. (1987). Differential Protooncogene Expression Characterizes Histopathologically Indistinguishable Tumors of the Peripheral Nervous System. *J Clin Invest* 80: 804-811. PubMed ID: 2887586
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC442306/>
5. Whang-Peng, J., Triche, T.J., Knutsen, T., Miser, J., Kao-Shan, S., Tsai, S., and Israel, M. A. (1986). Cytogenetic Characterization of Selected Small Round Cell Tumors of Childhood. *Cancer Genet Cytogene* 21: 185-208. PubMed ID: 3004699
<https://www.sciencedirect.com/science/article/pii/0165460886900014?via%3Dihub>



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Cell Line Name: TC-32

Low confluency (10x magnification)

High confluency (10x magnification)

Low confluency (20x magnification)

High confluency (20x magnification)

