

## Cell Line Data Sheet for CHLA-10

**Disease:** Primitive neuroectodermal tumor (PNET)  
**Phase of Therapy:** Post-Chemotherapy (Progressive Disease)  
**Treatment:** 4 cycles of cisplatin, doxorubicin, cyclophosphamide, etoposide  
**Disease Stage:**  
**Gender:** Female  
**Age at diagnosis:** 168 years  
**Race:** N/A  
**Age at sample collection:** N/A  
**Source of Culture:** Solid tumor (thoracic lymph node)  
**Primary Tumor Site:** Thorax  
**Date Established:** 1987

**EWS/FLI1 Status:** FLI1  
**p53 functionality:** Non-Functional  
**Karyotype:**  
**Modal No:**

<b>R-IC50 (DIMSCAN*):</b>	<u>Vincristine (ng/ml)</u>	<u>Melphalan (µg/ml)</u>	<u>Etoposide (ng/ml)</u>	<u>Rapamycin (ng/ml)</u>
*see reference 3	0.39 ± 0.05	7.27 ± 1.20	0.12 ± 0.01	0.62 ± 0.08

**Growth Conditions:** Please see Protocols section at <https://www.cccells.org/protocols.php>  
5% CO<sub>2</sub>, 20% O<sub>2</sub>, 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:** 32 hours  
**Growth Properties:** Teardrop-shaped cells with processes, adherent, grow mostly in clumps

**STR Profile:** May be obtained at <https://strdb.cccells.org/>

**Notes:** The Childhood Cancer Repository has a matching direct-to-culture diagnosis cell line available from this same patient – CHLA-9

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.



CHILDREN'S  
ONCOLOGY  
GROUP



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### References:

1. Thompson PM, Maris JM, Hogarty MD, Seeger RC, Reynolds CP, Brodeur GM, White PS. Homozygous deletion of CDKN2A (p16INK4a/p14ARF) but not within 1p36 or at Other Tumor Suppressor Loci in Neuroblastoma. *Cancer Res.* 61, 679-686, 2001. PubMed ID: 11212268  
<https://cancerres.aacrjournals.org/content/61/2/679.long>
2. 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>
3. Kang MH, Smith MA, Morton CL, Keshelava N, Houghton PJ, Reynolds CP. National Cancer Institute Pediatric Preclinical Testing Program: Model Description for In Vitro Cytotoxicity Testing. *Pediatr Blood Cancer* 56: 239-249, 2011. PubMed ID: 20922763  
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3005554/>

SEE NCI Pediatric Preclinical Testing Program references.



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**Cell Line Name: CHLA-10**

**Low confluency (10x magnification)**

**High confluency (10x magnification)**

**Low confluency (20x magnification)**

**High confluency (20x magnification)**