EPD-210FHTC

Clinical annotation:

Age: 10 Gender: Male

Location: Posterior fossa, left cerebellar hemisphere, extending to brain-

stem and through foramen magnum

Diagnosis: Anaplastic Ependymoma
Pre-treatment: Radiation and chemotherapy

Source: Recurrent, autopsy

Stage: M0 at diagnosis, metastatic recurrence

EFS (months): 31.5 from diagnosis OS (months): 85.4 from diagnosis



Histology of PDO

Pathology of human tumor: From initial diagnosis: The sections reveal sheets of cells with small round to oval nuclei with finely speckled chromatin and eosinophilic cytoplasm. The nuclei exhibit palisading and frequently are arranged in pseudorosettes around thin-walled vessels and occasional true rosettes. No significant pleomorphism or mitotic activity is noted, and no definite necrosis is identified. From autopsy: Increased mitotic activity (10/10 high power fields), larger and more hyperchromatic nuclei, and higher cellularity than the prior specimen.

Model information:

Mouse strain: Cell line

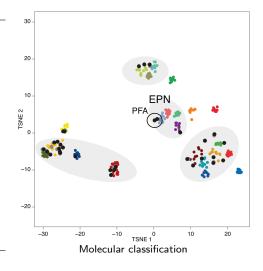
Site of transplantation: -

Protocol: Olson lab tissue culture protocol

Days to P0/P1/P2:

PI: James M. Olson

Contact: Request model at www.btrl.org

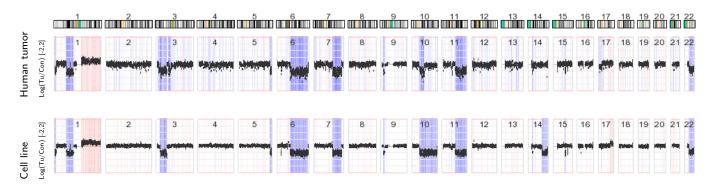


Molecular information:

Entity: Ependymoma
Subgroup: PFA
Curated lesions: chr1q gain

Detailed information: Explore molecular data in PDX explorer

Explore genomic data of pediatric PDOX cohort



Copy-number ratio (tumor vs. pseudo-control)

Last updated: 08/07/18