# The Open Pediatric Cancer Project

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## **Authors**

#### • Eric Wafula

**(D)** 0000-0001-8073-3797

Department of Bioinformatics and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Sangeeta Shukla

**(D)** 0000-0002-3727-9602

Department of Bioinformatics and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Krutika S. Gaonkar

**D** 0000-0003-0838-2405

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Department of Bioinformatics and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Run Jin

**D** 0000-0002-8958-9266

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Komal S. Rathi

© 0000-0001-5534-6904

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Department of Bioinformatics and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Yuankun Zhu

**1** 0000-0002-2455-9525

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Bailey K. Farrow

**(D)** 0000-0001-6727-6333

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Daniel P. Miller

**(D)** 0000-0002-2032-4358

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Mariarita Santi

**(iii)** 0000-0002-6728-3450

Department of Pathology and Laboratory Medicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Department of Pathology and Laboratory Medicine, University of Pennsylvania Perelman School of Medicine, Philadelphia, PA, 19104, USA

### · Adam A. Kraya

**(D)** 0000-0002-8526-5694

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Xiaoyan Huang

© 0000-0001-7267-4512

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Bo Zhang

**D** 0000-0002-0743-5379

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Brian M. Ennis

© 0000-0002-2653-5009

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Ryan J. Corbett

© 0000-0002-3478-0784

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Sharon J. Diskin

**(D)** 0000-0002-7200-8939

Division of Oncology, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Department of Pediatrics, University of Pennsylvania, Philadelphia, PA, 19104, USA

### • Nicholas Van Kuren

**(D)** 0000-0002-7414-9516

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Noel Coleman

© 0000-0001-6454-1285

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### · Christopher Blackden

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Jennifer L. Mason

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Saksham Phul

© 0000-0002-2771-2572

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### Miguel A. Brown

**(D)** 0000-0001-6782-1442

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Alex Sickler

**(D)** 0000-0001-7830-7537

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

#### Adam C. Resnick

**(D)** 0000-0003-0436-4189

Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA · Funded by Children's Brain Tumor Network; NIH 3P30 CA016520-44S5, U2C HL138346-03, U24 CA220457-03; NCI/NIH Contract No. 75N91019D00024, Task Order No. 75N91020F00003; Children's Hospital of Philadelphia Division of Neurosurgery

### • Jo Lynne Rokita^ ■

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Center for Data-Driven Discovery in Biomedicine, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Division of Neurosurgery, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA; Department of Bioinformatics and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA · Funded by NCI/NIH Contract No. 75N91019D00024, Task Order No. 75N91020F00003

### Kelsey Keith

**(D)** 0000-0002-7451-5117

Department of Biomedical and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

### • Asif T Chinwalla

**1** 0000-0001-7831-3996

Department of Biomedical and Health Informatics, Children's Hospital of Philadelphia, Philadelphia, PA, 19104, USA

# **Contact information**

^Lead Contact: Jo Lynne Rokita <u>rokita@chop.edu</u>

⊠Correspondence: Jo Lynne Rokita <u>rokita@chop.edu</u>

### In Brief

# **Highlights**

# **Summary**

# **Keywords**

# Introduction

# **Results**

# **Discussion**

# Acknowledgments

# **Author Contributions**

Author	Contributions
Eric Wafula	Formal analysis, Software
Sangeeta Shukla	Data curation, Formal analysis, Investigation, Methodology, Software, Writing – Original draft, Writing - Review and editing
Krutika S. Gaonkar	Data curation, Formal analysis, Investigation, Methodology, Software, Writing – Original draft, Writing - Review and editing
Run Jin	Data curation, Formal analysis, Visualization, Writing – Original draft, Writing - Review and editing
Komal S. Rathi	Formal analysis, Investigation, Methodology, Writing – Original draft
Yuankun Zhu	Data curation, Formal analysis, Investigation, Methodology, Supervision
Bailey K. Farrow	Data curation, Software
Daniel P. Miller	Formal analysis
Mariarita Santi	Investigation, Validation, Writing - Review and editing
Adam A. Kraya	Methodology
Xiaoyan Huang	Formal analysis
Bo Zhang	Data curation, Formal analysis
Brian M. Ennis	Data curation, Formal analysis
Ryan J. Corbett	Formal analysis
Sharon J. Diskin	Investigation, Supervision, Validation, Funding acquisition, Writing - Review and editing
Nicholas Van Kuren	Data curation, Software
Noel Coleman	Data curation
Christopher Blackden	Resources
Jennifer L. Mason	Supervision
Saksham Phul	Data curation, Methodology, Formal analysis
Miguel A. Brown	Data curation, Methodology, Formal analysis
Alex Sickler	Methodology, Formal analysis

Author	Contributions
Adam C. Resnick	Conceptualization, Funding acquisition, Resources, Supervision
Jo Lynne Rokita^	Conceptualization, Data curation, Formal analysis, Funding acquisition, Investigation, Methodology, Software, Supervision, Writing – Original draft, Writing - Review and editing
Kelsey Keith	Software, Writing - original draft, API, Formal Analysis, Data Curation, Visualization
Asif T Chinwalla	Project Administration, Supervision, Methodology, Investigation, Validation, API

# **Declarations of Interest**

# **Figure Titles and Legends**

# **Table Titles and Legends**

### **OPENPEDCAN METHODS**

### **RESOURCE AVAILABILITY**

### Lead contact

Requests for access to OpenPedCan raw data and/or specimens may be directed to, and will be fulfilled by Jo Lynne Rokita (rokita@chop.edu).

# **Materials availability**

This study did not create new, unique reagents.

# Data and code availability

Within OpenPedCan (OPC), we harmonized, aggregated, and analyzed data from multiple sources. We harmonized data from the Therapeutically Applicable Research to Generate Effective Treatments (TARGET cohort) Initiative, an NCI-funded collection of disease-specific projects that seeks to identify the genomic changes of pediatric cancers [1]. We included already harmonized neuroblastoma samples from the Gabriella Miller Kids First (GMKF cohort) Pediatric Research Program, a large-scale effort to accelerate research and gene discovery in pediatric cancers and structural birth defects [2]. Additionally, we re-harmonized all samples from the Open Pediatric Brain Tumor Atlas (OpenPBTA, PBTA cohort), an open science initiative led by Alex's Lemonade Stand Foundation Childhood Cancer Data Lab and the Center for Data-Driven Discovery (D3B) at the Children's Hospital of Philadelphia (CHOP), which genomically characterized pediatric brain tumor data from the Children's Brain Tumor Network (CBTN), and the Pacific Pediatric Neuro-oncology Consortium (PNOC) [3,4]. Building on the work of OpenPBTA, OPC added the PBTA X01 data [5], the Chordoma Foundation data [6/], and the MI-ONCOSEQ Study [7], donated to CBTN by the University of Michigan, to the PBTA cohort. Finally, OPC includes the Children's Hospital of Philadelphia (CHOP) P30 Panel data generated by CHOP's Division of Genomic Diagnostics (DGD cohort) which includes fusion panel data [3]. In addition to pediatric cancer data, OpenPedCan contains adult data from large science consortiums as references.

For normal gene expression, GTEx [8] was used, and for comparison with adult cancers, The Cancer Genome Atlas (TCGA) [9] was included.

Merged summary files for OpenPedCan v12 are openly accessible in <u>CAVATICA</u> or via download-data.sh script in the <a href="https://github.com/PediatricOpenTargets/OpenPedCan-analysis">https://github.com/PediatricOpenTargets/OpenPedCan-analysis</a> repository. Cancer group summary data are visible within the NCI's pediatric <a href="Molecular Targets Platform">Molecular Targets Platform</a> and cohort, cancer group, and individual data are visible within <a href="PedcBioPortal">PedcBioPortal</a>

OpenPedCan analysis modules were developed within OpenPBTA [4], modified based on OpenPBTA, or newly created and can be found within the following publicly available repositories. OpenPBTA module analyses can be found at <a href="https://github.com/AlexsLemonade/OpenPBTA-analysis">https://github.com/AlexsLemonade/OpenPBTA-analysis</a>. OpenPedCan module analyses can be found at

https://github.com/PediatricOpenTargets/OpenPedCan-analysis. OpenPedCan api code can be found at https://github.com/PediatricOpenTargets/OpenPedCan-api.

All original code was developed within the following modules in the OpenPedCan analyses repository as listed below. Links to the modules are available here, and within each module is a detailed README that describes the purpose and intended usage of the scripts, along with pointers to the results from the data those scripts process.

<u>chromosomal-instability</u> <u>cnv-frequencies</u>
cnv-frequencies
<u>collapse-rnaseq</u>
<u>compare-gistic</u>
copy number consensus call
<u>create-subset-files</u>
data-pre-release-qc
<u>efo-mondo-mapping</u>
<u>filter-mtp-tables</u>
focal-cn-file-preparation
<u>fusion-frequencies</u>
<u>fusion-summary</u>
fusion filtering
gene-set-enrichment-analysis
gene match
<u>immune-deconv</u>
<u>independent-samples</u>
long-format-table-utils
methylation-preprocessing
methylation-summary
molecular-subtyping-ATRT
molecular-subtyping-CRANIO

List of OpenPedCan Analyses Modules	
molecular-subtyping-EPN	
molecular-subtyping-EWS	
molecular-subtyping-HGG	
molecular-subtyping-LGAT	
molecular-subtyping-MB	
molecular-subtyping-NBL	
molecular-subtyping-chordoma	
molecular-subtyping-embryonal	
molecular-subtyping-integrate	
molecular-subtyping-neurocytoma	
molecular-subtyping-pathology	
<u>mtp-annotations</u>	
<u>mtp-tables-qc-checks</u>	
<u>mutational-signatures</u>	
pedcbio-cnv-prepare	
pedcbio-sample-name	
pedot-table-column-display-order-name	
<u>rna-seq-expression-summary-stats</u>	
<u>rnaseq-batch-correct</u>	
<u>run-gistic</u>	
<u>snv-frequencies</u>	
<u>tmb-calculation</u>	
<u>tp53 nf1 score</u>	
<u>tumor-gtex-plots</u>	
<u>tumor-normal-differential-expression</u>	

Software versions are documented in **Table XX**.

### **Data releases**

We maintained a data release folder on Amazon S3, downloadable directly from S3 or our openaccess CAVATICA project, with merged files for each analysis (See data and code availability section). As we produced new results that we expected to be used across multiple analyses, or identified data issues, we created new data releases in a versioned manner.

### **EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS**

Please refer to the OpenPBTA manuscript [4] for additional details for the CBTN and PNOC cohorts and project-specific study details for GMKF, TARGET, and TCGA cohorts. For the PBTA X01 cohort, libraries were prepped using the Illumina TruSeq Strand-Specific Protocol to pull out poly-adenylated transcripts.

### cDNA Library Construction

Total RNA was quantified using the Quant-iT™ RiboGreen® RNA Assay Kit and normalized to 5ng/ul. Following plating, 2 uL of ERCC controls (using a 1:1000 dilution) were spiked into each sample. An aliquot of 325 ng for each sample was transferred into library preparation. The resultant 400bp cDNA went through dual-indexed library preparation: 'A' base addition, adapter ligation using P7 adapters, and PCR enrichment using P5 adapters. After enrichment, the libraries were quantified using Quant-iT PicoGreen (1:200 dilution). Samples were normalized to 5 ng/uL. The sample set was pooled and quantified using the KAPA Library Quantification Kit for Illumina Sequencing Platforms.

Illumina Sequencing

Pooled libraries were normalized to 2nM and denatured using 0.1 N NaOH prior to sequencing. Flowcell cluster amplification and sequencing were performed according to the manufacturer's protocols using the NovaSeq 6000. Each run was a 151bp paired-end with an eight-base index barcode read. Data was analyzed using the Broad Picard Pipeline which includes de-multiplexing and data aggregation.

### **METHOD DETAILS**

## Nucleic acids extraction and library preparation

### **Data generation**

### **DNA WGS Alignment**

Please refer to the OpenPBTA manuscript for details [4].

# **Quality Control of Sequencing Data**

Please refer to the OpenPBTA manuscript for details [4].

# SNP calling for B-allele Frequency (BAF) generation

Please refer to the OpenPBTA manuscript for details [4].

# **Somatic Mutation Calling**

## SNV and indel calling

Please refer to the OpenPBTA manuscript for details [4].

#### VCF annotation and MAF creation

**Gather SNV and INDEL Hotspots** 

**Consensus SNV Calling** 

**Somatic Copy Number Variant Calling (WGS samples only)** 

**Consensus CNV Calling** 

We adopted the consensus CNV calling described in OpenPBTA manuscript [doi:10.1016/j.xgen.2023.100340] with minor adjustments. For each caller and sample with WGS performed, we called CNVs based on consensus among Control-FREEC ([10]; [11]), CNVkit ([doi? 10.1371/journal.pcbi.1004873]), and GATK ([doi? 10.1101/gr.107524.110]). Sample and consensus caller files with more than 2,500 CNVs were removed to de-noise and increase data quality, based on cutoffs used in GISTIC ([12]). For each sample, we included the following regions in the final consensus set: 1) regions with reciprocal overlap of 50% or more between at least two of the callers; 2) smaller CNV regions in which more than 90% of regions were covered by another caller. For GATK, if a panel of normal was not able to be created (required 30 male and 30 female with the same sequencing platform), consensus was not run for tumors with WGS performed on that sequencing platform. We defined copy number as NA for any regions that had a neutral call for the samples included in the consensus file. We merged CNV regions within 10,000 bp of each other with the same direction of gain or loss into single region. Any CNVs that overlapped 50% or more with immunoglobulin, telomeric, centromeric, segment duplicated regions, or that were shorter than 3000 bp were filtered out. The CNVKit calls for WXS samples were appended to the consensus CNV file.

### **Somatic Structural Variant Calling (WGS samples only)**

Please refer to the OpenPBTA manuscript for details [4].

## **Methylation Analysis**

## Methylation array preprocessing

We preprocessed raw Illumina 450K and EPIC 850K Infinium Human Methylation Bead Array intensities using the array preprocessing methods implemented in the minfi Bioconductor package [13]. We utilized either preprocessFunnorm when an array dataset had both tumor and normal samples or multiple OpenPedcan-defined cancer\_groups and preprocessQuantile when an array dataset had only tumor samples from a single OpenPedcan-defined cancer\_group to estimate usable methylation measurements (beta-values and m-values) and copy number (cn-values). Some Illumina Infinium array probes targeting CpG loci contain single-nucleotide polymorphisms (SNPs) near or within the probe [14], which could affect DNA methylation measurements [15]. As the minfi preprocessing workflow recommends, we dropped probes containing common SNPs in dbSNP (minor allele frequency > 1%) at the CpG interrogation or the single nucleotide extensions.

Details of methylation array preprocessing are available in the <u>OpenPedCan methylation-preprocessing module</u>.

## Methylation beta-values summaries

We comprehensively summarized gene-level and isoform-level metrics for the methylation betavalues estimated by array preprocessing to provide insight into the variations in overall genomic DNA methylation levels observed across different pediatric tumors by computing CpG probe-level summary metrics in each cancer group within a cohort, including 1) beta-values quantiles, 2) gene expression (TPM) and methylation (beta-values) correlation, 3) TPM median expression, and 4) transcript representation - a proxy for percent isoform expression in a gene. In addition, each CpG probe was annotated with a gene feature to identify the genomic regions likely involved in regulating gene expression.

Details of the analysis are available in the OpenPedCan methylation-summary module.

### Methylation sample classification

We ran the <u>dkfz's brain classifier version 12.5</u>, a comprehensive DNA methylation-based classification of CNS tumors across all entities and age groups [16]. Unprocessed IDAT-files from the <u>Children's Brain Tumor Network (CBTN)</u> Infinium Human Methylation EPIC (850k) BeadChip arrays were used as input and the following information was compiled into the histologies.tsv file: dkfz\_v12\_methylation\_subclass (predicted methylation subtype), dkfz\_v12\_methylation\_subclass\_score (classification score), dkfz\_v12\_methylation\_mgmt\_status (*MGMT* methylation status), and dkfz\_v12\_methylation\_mgmt\_estimated (estimated *MGMT* methylation fraction).

### **Gene Expression**

The <a href="tmor-normal-differential-expression">tumor-normal-differential-expression</a> module perfoms differential expression analyses for all sets of Disease (cancer\_group) and Dataset (cohort) across all genes found in the gene-expression-rsem-tpm-collapsed.rds table. The purpose of this analysis is to highlight the correlation and understand the variability in gene expression in different cancer conditions across different histological tissues. For OpenPedCan v12 data release, this module performs expression analysis over 102 cancer groups across 52 histological tissues for all 54,346 genes found in the dataset. This analysis was performed on the Children's Hospital of Philadelphia HPC and was configured to use 96G of RAM per CPU, with one task (one iteration of expression analysis for each set of tissue and cancer group) per CPU (total 102\*52=5304 CPUs) using the <a href="R/DESeq2">R/DESeq2</a> package. Please refer to script run-tumor-normal-differential-expression.sh in the module for additional details on Slurm processing configuration. The same analysis can also be performed on CAVATICA, but requires further optimization. The module describes the steps for CAVATICA set up, and scripts to publish an application on the portal. The required data files are also available publicly on CAVATICA under the <a href="Open Pediatric Cancer (OpenPedCan) Open Access">Open Access</a>. Refer to the module for detailed description and scripts.

#### **Abundance Estimation**

Among the data sources used for OpenPedCan, GTEx and TCGA used GENCODE versions v26 and v36, respectively. Moreover, the gene symbols used in these different GENCODE versions also varied. Therefore, the gene symbols had to be harmonized for compatibility to map unique gene identifiers to their gene symbols. ENSG IDs from each data source were pulled and mapped to the GTF/GFF3 file from GENCODE v39 to extract unique gene symbols and remove duplicates. Additionally, the gene expression matrices had some instances where multiple Ensembl gene identifiers mapped to the same gene symbol. This was dealt with by filtering the expression matrix to only genes with [FPKM/TPM] > 0 and then selecting the instance of the gene symbol with the maximum mean [FPKM/TPM/Expected\_count] value across samples. This enabled many downstream modules that require RNA-seq data have gene symbols as unique gene identifiers. Refer to collapse-rnaseq module for scripts and details.

### **Gene Expression Summary Statistics**

We generated RNA-Seq gene expression (TPM) summary statistics for independent tumor samples from the combined OpenPedCan gene expression matrices, including cancers from pediatric cohorts (PBTA, GMKF, and TARGET) and adult cancers from the TCGA cohort. We grouped selected samples into two groups containing samples from a cancer group in either each cohort or all cohorts, and calculated TPM means, standard deviations, gene-wise z-scores, group-wise z-scores, and ranks for each group as described in the <a href="OpenPedCan rna-seq-expression-summary-">OpenPedCan rna-seq-expression-summary-</a>

stats module in detail. The resulting gene-wise and group-wise summary statistics tables were annotated with EFO and MONDO disease codes associated with the cancer groups.

### **Gene fusion detection**

# **QUANTIFICATION AND STATISTICAL ANALYSIS**

### Focal Copy Number Calling (focal-cn-file-preparation analysis module)

Please refer to the OpenPBTA manuscript for details on assignment of copy number status values to CNV segments, cytobands, and genes [4]. We applied criteria to resolve instances of multiple conflicting status calls for the same gene and sample, which are described in detail in the <u>focal-cn-file-preparation</u> module. Briefly, we prioritized 1) non-neutral status calls, 2) calls made from dominant segments with respect to gene overlap, and 3) amplification and deep deletion status calls over gain and loss calls, respectively, when selecting a dominant status call per gene and sample. These methods resolved >99% of duplicated gene-level status calls.

Gene Set Variation Analysis (gene-set-enrichment-analysis analysis module)

Please refer to the OpenPBTA manuscript for details [4].

Fusion prioritization (fusion\_filtering analysis module)

Mutational Signatures (mutational-signatures analysis module)

Tumor Mutation Burden (snv-callers analysis module)

**Clinical Data Harmonization** 

**WHO Classification of Disease Types** 

**Molecular Subtyping** 

Here, we build upon the molecular subtyping performed in OpenPBTA [4].

### High-grade gliomas.

A new high-grade glioma entity called infant-type hemispheric gliomas (IHGs), characterized by distinct gene fusions enriched in receptor tyrosine kinase (RTK) genes including *ALK*, *NTRK1/2/3*, *ROS1* or *MET*, was identified in 2021 [doi? 10.1038/s41467-019-12187-5]. To identify IHG tumors, first, tumors which were classified as "IHG" by the DKFZ methylation classifier or diagnosed as "infant type hemispheric glioma" from pathology\_free\_text\_diagnosis were selected [16]. Then, the corresponding tumor RNA-seq data were utilized to seek the evidence for RTK gene fusion. Based on the specific RTK gene fusion present in the samples, IHGs were further classified as "IHG, ALK-altered", "IHG, NTRK-altered", "IHG, ROS1-altered", or "IHG, MET-altered". If no fusion was observed, the samples were identified as "IHG, To be classified".

### Atypical teratoid rhabdoid tumors.

Atypical teratoid rhabdoid tumors (ATRT) tumors were categorized into three subtypes: "ATRT, MYC", "ATRT, SHH", and "ATRT, TYR" [17]. In OpenPedCan, the molecular subtyping of ATRT was based solely on the DNA methylation data. Briefly, ATRT samples with a high confidence DKFZ methylation subclass

score (>= 0.8) were selected and subtypes were assigned based on the DKFZ methylation subclass [doi10.1038/nature26000?]. Samples with low confidence DKFZ methylation subclass scores (< 0.8) were identified as "ATRT, To be classified".

#### Neuroblastoma tumors.

Neuroblastoma (NBL) tumors with a pathology diagnosis of neuroblastoma, ganglioneuroblastoma, or ganglioneuroma were subtyped based on their MYCN copy number status as either "NBL, MYCN amplified" or "NBL, MYCN non-amplified". If pathology\_free\_text\_diagnosis was "NBL, MYCN non-amplified" and the genetic data suggested MYCN amplification, the samples were subtyped as "NBL, MYCN amplified". On the other hand, if pathology\_free\_text\_diagnosis was "NBL, MYCN amplified" and the genetic data suggested MYCN non-amplification, the RNA-Seq gene expression level of MYCN was used as a prediction indicator. In those cases, samples with MYCN gene expression above or below the cutoff (TPM >= 140.83 based on visual inspection of MYCN CNV status) were subtyped as "NBL, MYCN amplified" and "NBL, MYCN non-amplified", respectively. MYCN gene expression was also used to subtype samples without DNA sequencing data. If a sample did not fit none of these situations, it was denoted as "NBL, To be classified".

### Integration of brain tumor methylation classifications

# TP53 Alteration Annotation (tp53\_nf1\_score analysis module)

Please refer to the OpenPBTA manuscript for details [4].

### Prediction of participants' genetic sex

Please refer to the OpenPBTA manuscript for details [4].

# Selection of independent samples (independent-samples analysis module)

For analyses that require all input biospecimens to be independent, we use the OpenPedCan-analysis <u>independent-samples</u> module to select only one biospecimen from each input participant. For each input participant of an analysis, the independent biospecimen is selected based on the analysis-specific filters and preferences for the biospecimen metadata, such as experimental strategy, cancer group, and tumor descriptor.

# **Supplemental Information Titles and Legends**

# Consortia

# References

- 1. **dbGaP Study** <u>https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?study\_id=phs000218.v23.p8</u>
- 2. **dbGaP Study** <a href="https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?">https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?</a> study id=phs001436.v1.p1
- 3. **dbGaP Study** <a href="https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?study\_id=phs002517.v1.p1">https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?study\_id=phs002517.v1.p1</a>
- 4. OpenPBTA: The Open Pediatric Brain Tumor Atlas

Joshua A Shapiro, Krutika S Gaonkar, Stephanie J Spielman, Candace L Savonen, Chante J Bethell, Run Jin, Komal S Rathi, Yuankun Zhu, Laura E Egolf, Bailey K Farrow, ... Jaclyn N Taroni *Cell Genomics* (2023-05) <a href="https://doi.org/gr92p6">https://doi.org/gr92p6</a>
DOI: <a href="https://doi.org/gr92p6">10.1016/j.xgen.2023.100340</a>

- 5. **FY21 X01 Projects for the Gabriella Miller Kids First Program** (2021-09-02) <a href="https://commonfund.nih.gov/kidsfirst/2021x01projects">https://commonfund.nih.gov/kidsfirst/2021x01projects</a>
- 6. Home
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   https://www.chordomafoundation.org/
- 7. Michigan Center for Translational Pathology <a href="https://mctp.med.umich.edu">https://mctp.med.umich.edu</a>
- 8. **dbGaP Study** <u>https://www.ncbi.nlm.nih.gov/projects/gap/cgi-bin/study.cgi?study\_id=phs000424.v9.p2</u>
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