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Osteosarcoma

PRINCIPAL INVESTIGATOR: Karen Pollok

CONTRACTING ORGANIZATION: Trustees of Indiana University

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14. ABSTRACT- Osteosarcoma (OS) is extremely challenging to treat for 25-50% of the patients already have metastatic disease at the time of diagnosis. The purpose of this proposal is to improve therapy for OS by expanding upon our initial data to target replication stress (RS), a hallmark of OS. We previously used a systems biology approach to select OS patient-derived xenograft models (PDX) that harbor RS signatures such as copy number gains in MYC and RAD21 and demonstrated the feasibility of targeting replication stress via CHK1 and BET inhibitors (Pandya et al., Cancers, 2020). The scope of the work now uses a panel of modeling approaches including metastatic OS to analyze mechanisms-of-action, efficacy, and safety. To address biological variables of race, gender, and age, a diverse panel of pediatric and AYA OS models were developed and characterized by our team. In terms of major findings in year one, we published an integrated bioinformatics and in-vivo validation study showing the fidelity of the molecular signatures between original tumors and PDX as well as promising efficacy in response to BET inhibition as a single agent (Pandya et al., Cancers, 2022). We extended those studies to demonstrate that BET inhibition in combination with salvage therapy further increases efficacy and survival in a model of aggressive OS. The significance of the work to date is that studying clinically relevant in-vivo models will advance OS research and patient care by identifying how these drugs in combination promote OS cell death. These studies will lead to new mechanistic insights and potentially identify new biomarkers of response and additional therapeutic vulnerabilities as well as provide rationale for clinical trial development.					
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1. INTRODUCTION:

Osteosarcoma (OS) is the subject for these studies as it is the most common bone cancer in children as well as in adolescents and young adults (AYA). In OS, treatment has not improved in the past 3 decades, underscoring the urgency for developing efficacious and safe therapies. This is due, in part, to copy number variations (CNVs) which can lead to dysregulation of multiple signaling pathways and uncontrolled cancer cell growth. Analysis of pediatric and AYA OS CNVs from public and Riley Hospital for Children databases identified a large number of CNVs associated with poor overall survival in OS patients many of which were associated with replication stress (RS). In OS patients harboring increased copies of genes associated with RS such as MYC and RAD21, this correlated with poor overall survival (n=85 patients analyzed). The dysregulated process of RS can be exploited by using drugs that will uncouple RS in the cancer cells from strategies that the cancer cells typically use to survive. In this proposal, our scope and strategy is to focus on two types of drugs that inhibit critical regulators of replication stress called bromodomain and extraterminal (BET) proteins and CHK1. The purpose of this project is to improve therapeutic outcomes in OS by designing safe combination therapies based on the mutational profile of the OS tumors. Information gained here will provide strong translational rationale for targeting RS and moving this approach forward for clinical testing in aggressive and metastatic OS. This research will benefit children and grandchildren of military members, veterans, or children and AYA who are the beneficiary of a military member. Successful completion of this work will maximize the time for active Service member to remain on duty by either allowing them to return to duty earlier or not be called home in the event that their child/grandchild/beneficiary have OS because it is well-managed and not an emotional and economic burden to the family.

2. KEY WORDS:

Osteosarcoma, pediatric, adolescents and young adults (AYA), replication stress, BETs, CHK1, patient-derived xenografts (PDXs), copy number variations (CNVs), orthotopic, intratibial, lung colonization, cell line derived xenograft (CDXs), xenoline, DNA damage response (DDR).

3. ACCOMPLISHMENTS:

What were the major goals of the project?

For year one, analysis of sequencing data of OS pediatric and adolescent OS PDX models at the DNA and RNA levels was completed (100%). We evaluated in vitro OS response to CHK1+BET inhibitors (80%), assessed DNA damage kinetics (90%), conducted dose-finding studies in vivo (100%), and PK-PD studies on PDX models using identified doses combined with evaluation of tumor response and kinome reprogramming via MIB/MS following single-agent and combination therapy (75%). We reprioritized a couple of year two goals for year one to evaluate BET dependency in CHK1i-or chemotherapy-mediated OS cell death (100%) and began downstream target modulation studies (25%).

What was accomplished under these goals?

Major activities included in vitro mechanism studies and first phase of in vivo studies which included dose-finding studies and combination studies. **Specific objectives** were to develop efficacious and safe therapies with a focus on underlying mechanisms of response to therapy that will exacerbate RS and increase cell death. **Key outcomes** were that the OS PDX models recapitulate the molecular signatures of the original tumors overall with retainment of oncogenic signatures and a portion of these data have now been published (Pandya et al., Cancers 2022) as well as workflow strategy for identification of potential therapeutic response biomarkers, of which RS signatures are evident in the OS models. In vitro assays indicate that OS PDX are very sensitive to the BETi AZD5153 as a single agent (Fig. 1) and use of topotecan can sensitize OS cells and PDX to BETi (Fig. 2). Sensitivity correlates with increased RS and cell death. **One goal not completely** met for year one is the analysis of all the kinome samples which were delayed due to equipment repairs. The machine is now repaired, and samples are in the queue. Methodologies in year one include whole genome sequencing, RNAseq, Incucyte ZOOM to track cell growth, Western blot, flow cytometry, alkaline comet assay, MIB/MS Kinome Profiling, RNA-seq, and efficacy and safety studies in mouse models. We have found additive to synergistic inhibition of growth in a panel of OS cell lines using BETi+CHK1i or BETi+salvage therapy which results in increased RS, DNA damage, and apoptosis. In a short-term efficacy study using PDX96, we found decreased tumor growth and increased RS as indicated by the DNA damage marker, γ H2AX. In a long-term survival study in an OS PDX model of refractory disease we found increased efficacy with single agent BETi/AZD5153 and further improvement of survival in combination BETi + AZD5153+ topotecan (Fig. 2).

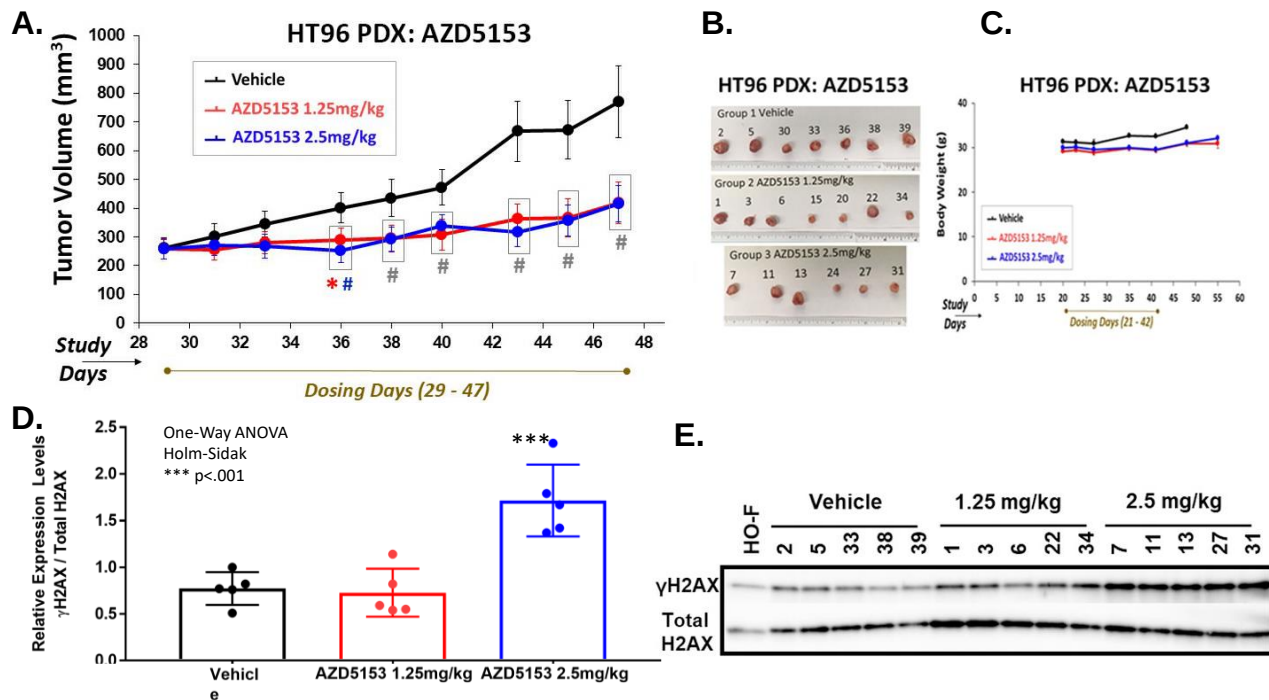
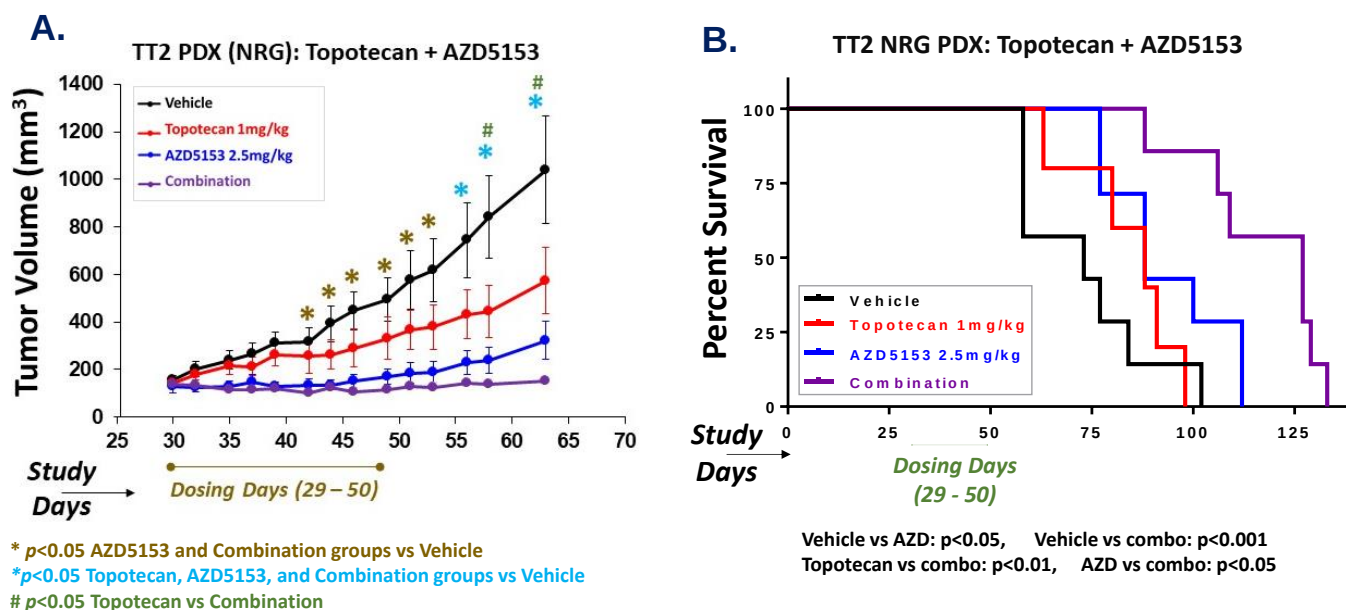


Figure 1. BET inhibition via AZD5153 suppresses OS PDX growth is well tolerated and enhances replication stress in vivo. A. tumor volumes over time. **B.** tumor weights. **C.** body weights over time. **D-E.** quantification and western blot of γ H2AX as a marker of RS-mediated DNA damage following AZD5153 treatment. Mouse IDs are above each lane. HO-F – human osteoblasts from femur. PDX96 was derived from a naïve OS patient with aggressive disease.



What opportunities for training and professional development has the project provided?

- Pankita H. Pandya (Co-I; Assistant Research Professor) has gained training on how to analyze multiplexed inhibitor bead kinome enrichment assays by working with Dr. Angus (co-I) and she has continued her training with Dr. Kun Huang in bioinformatic analysis (RNAseq). She has had many professional development opportunities to present on this project which include presenting at Sarcoma Translation Research Working group meeting at our institution, as well as presenting as a poster at the work at American Association of Cancer Research (AACR) international meeting. Pankita has also participated in Global Cancer Consortium Omics workshop to continue her bioinformatics training. She also attended the annual CTOS meeting.
- Nikan Riyahi is a 4th year graduate student who is working on this project for his PhD. He has had many training opportunities this year. He was trained by the Preclinical Modeling Therapeutics Core (PMTTC) to conduct in-vivo studies associated with the proposal. He has also been trained to conduct the cellular response and mechanisms-of-action studies for the in vitro aspects of the grant. Nikan has been able to present the work completed in this grant through his departmental seminar, 2023 AACR with a IU Simon Comprehensive Cancer Center (IUSCCC) Travel Award, and a Paradise travel award through Pharm/Tox. He placed 3rd place in the October 2022 IUSCCC Research Day and was an invited speaker for IUSCCC Cancer Research Day May 2023.

How were the results disseminated to communities of interest?

We participated in a number of venues to disseminate information to the community that enhances understanding and provides educational opportunities to young scientists from diverse backgrounds to science. Our team participated in Tyler Trent Foundation (Tee off for Tyler) golf outing to raise awareness for osteosarcoma pediatric cancer research. We attended the Riley Children's Foundation Cancer Research Touch Point and the Mighty Mason fundraiser to share information about research program to the community. We attended the Purdue Riley Dance Marathon stewardship brunch to provide an update on our OS research and the Pediatric Precision Genomics Program at the Riley Hospital for Children. We mentored fifty high-school students selected state-wide to participate in a 2-day interactive science program called Molecular Medicine In Action (MMIA); our module focused on sarcoma research and the precision genomics program at the Riley Hospital for Children. We participated in a video on OS cancer research updates at the IU School of Medicine YouTube channel: <https://youtu.be/JcnWFKrYnts>

What do you plan to do during the next reporting period to accomplish the goals?

We will continue to analyze kinetics of DNA damage and downstream target modulation both in vitro and in vivo. We plan to use a new assay validated in the lab called the growth delay assay where one can expose cultures to multiple doses of drugs to better simulate what would happen in vivo. We believe this approach may have improved predictive value in contrast to finite cultures with one drug exposure. With doses determined of the CHK1i and BETi, we will conduct in-vivo studies in PDX models complemented by RNAseq, kinome analysis, and validation to evaluate tumor responses. We will begin to evaluate dosing in the metastatic modeling approaches.

4. IMPACT:

What was the impact on the development of the principal discipline(s) of the project?

Our findings will be impactful by increasing our knowledge base on how to best treat aggressive OS. We were able to molecularly characterize OS cell lines, xenolines (derived from pediatric and adolescent OS PDXs), and PDXs. Such molecular characterization will help the scientific community that focus on pediatric and AYA osteosarcoma research or treatment to know the genomic landscape of the models they are working with and also give us insight on molecular biomarkers of interest within our own models. Use of high-risk molecularly characterized OS PDXs and xenolines that are linked to clinical history will provide additional insight into patient stratification. Collectively, these studies will provide critical feasibility data that has high translational value and great potential to improve survival and quality of life for patients with high-risk OS. We have successfully demonstrated the promise of using combination therapies that increase RS-mediated DNA damage and OS cell death. Additionally, kinome profiling will help increase our understanding on how BET+/-CHK1 inhibition therapy reprograms the OS tumors and may provide new opportunities for therapeutic intervention. Lastly, development of orthotopic, syngeneic immunocompetent, and lung colonization models of osteosarcoma will help provide more knowledge on the role of the tumor microenvironment and therapeutic responses in this disease.

What was the impact on other disciplines?

While the primary discipline impacted this research is biology/medicine/health (particularly pediatric and AYA osteosarcoma), other disciplines such as the military can benefit from these studies. For instance, this research will benefit not only young adult military Service members who may develop OS, but also the children and grandchildren of military Service members, Veterans, or patients who are the beneficiary of a military Service member. Improving therapies for aggressive OS will maximize the time for active service member to remain on duty by either allowing them to return to duty earlier or not be called home in the event that their relative's OS is well-managed. Such studies can also positively impact and give more insight into adult cancers and other diseases that high moderate levels of replication stress.

What was the impact on technology transfer?

Nothing to report.

What was the impact on society beyond science and technology?

Nothing to report

5. CHANGES/PROBLEMS:

Changes in approach and reasons for change

Nothing to report

Actual or anticipated problems or delays and actions or plans to resolve them

Nothing to report

Changes that had a significant impact on expenditures

Nothing to report

Significant changes in use or care of human subjects, vertebrate animals, biohazards, and/or select agents

Nothing to report

Significant changes in use or care of human subjects

Nothing to report

Significant changes in use or care of vertebrate animals

Nothing to report

Significant changes in use of biohazards and/or select agents

Nothing to report

6. PRODUCTS:

• **Publications, conference papers, and presentations**

Journal publications.

P.H. Pandya, A.J. Jannu, K. Bijangi-Vishehsaraei, E. Dobrota, B.J. Bailey, †#F. Barghi, H.E. Shannon, †#N. Riyahi, †#N.P. Damayanti, C. Young, †#R. Malko, †#R. Justice, E. Albright, G.E. Sandusky, L. D. Wurtz, C.D. Collier, M.S. Marshall, R.I. Gallagher, J.D. Wulfschle, E.F. Petricoin, K. Coy, M. Trowbridge, A.L. Sinn, J.L. Renbarger, M.J. Ferguson, K. Huang, J. Zhang, M.R. Saadatzaheh, K.E. Pollok. 2022. Integrative multi-OMICS identifies therapeutic response biomarkers and confirms fidelity of clinically-annotated, serially-passaged patient-derived xenografts established from primary and metastatic pediatric and AYA solid tumors. *Cancers* 2022. Dec 30;15(1):259. doi: 10.3390/cancers15010259. PMID: 36612255, PMCID: PMC9818438.
Acknowledgement of federal support: yes

Books or other non-periodical, one-time publications.

Nothing to report

Other publications, conference papers and presentations.

1. Riyahi N, Pandya PH, Bailey BJ, Dobrota EA, Young C, Shannon HE, Barghi F, Malko R, Bijangi-Vishehsaraei K, Trowbridge M, Coy K, Kennedy FM, Sinn AL, Angus S, Ferguson MJ, Saadatzaheh MR, Pollok KE. "Therapeutic Targeting of BET Bromodomain Proteins in Osteosarcoma." Wells Center Retreat. Indianapolis, Indiana., USA Aug 2022 also presented at IU Simon Comprehensive Cancer Center Cancer Research Day (Oct 2022) Indianapolis, Indiana., USA; *3rd place Award- translational research category for graduate students.*
2. *Dobrota E, Pandya PH, Jannu AJJ, Bijangi-Vishehsaraei K, Bailey BJ, Barghi F, Shannon HE, Riyahi N, Damayanti NP, Young C, Malko R, Justice R, Albright E, Sandusky Gem Wurtz LD, Collier CD, Petricoin E, Coy K, Trowbridge M, Sinn AL, Ferguson MJ, Huang K, Zhang J, Saadatzaheh MR, Pollok KE. "Multi-omics analytic pipeline of pediatric and AYA solid tumor patient-derived xenografts reveals biomarkers of therapeutic sensitivity and resistance." IU Simon Comprehensive Cancer Center Cancer Research Day (Oct 2022) Indianapolis, Indiana., USA
3. N. Riyahi, P.H. Pandya, B.J. Bailey, E.A. Dobrota, C. Young, H.E. Shannon, F. Barghi, R. Malko, K. Bijangi-Vishehsaraei, M.A. Trowbridge, K. Coy, F.M. Kennedy, A.L. Sinn, S. Angus, M.J. Ferguson, M. R. Saadatzaheh, K.E. Pollok. Therapeutic targeting of BET bromodomain proteins increases DNA damage and potentiates salvage therapy in osteosarcoma xenografts derived from patients with replication stress signatures. Annual AACR Meeting 2023, Orlando, Fla., April 2023.
4. N. Riyahi, P.H. Pandya, B.J. Bailey, E.A. Dobrota, C. Young, H.E. Shannon, F. Barghi, R. Malko, K. Bijangi-Vishehsaraei, M.A. Trowbridge, K. Coy, F.M. Kennedy, A.L. Sinn, S. Angus, M.J. Ferguson, M. R. Saadatzaheh, K.E. Pollok. Therapeutic targeting of BET bromodomain proteins increases DNA damage and potentiates salvage therapy in osteosarcoma xenografts derived from patients with replication stress signatures. Annual AACR Meeting 2023, Orlando, Fla., April 2023. *Selected for Oral Presentation at the IU Simon Comprehensive Cancer Center Cancer Research Day (May 2023) Indianapolis, Indiana., USA*
5. Dobrota E, Pandya PH, Jannu AJJ, Bijangi-Vishehsaraei K, Bailey BJ, Barghi F, Shannon HE, Riyahi N, Damayanti NP, Young C, Malko R, Justice R, Albright E, Sandusky G.E. Wurtz LD, Collier CD, Petricoin E, Coy K, Trowbridge M, Sinn AL, Ferguson MJ, Huang K, Zhang J, Saadatzaheh MR, Pollok KE. "Integrative Analysis of Multi-omics Data on Pediatric and AYA solid tumor Patient-Derived Xenografts Unveils Biomarkers of Therapeutic Response." IU Simon Comprehensive Cancer Center Cancer Research Day (May 2023) Indianapolis, Indiana., USA; *2nd Place Award – Research Associate category*

- **Website(s) or other Internet site(s)**

We participated in a video on OS cancer research updates at the IU School of Medicine YouTube channel: <https://youtu.be/JcnWFKrYnts>

- **Technologies or techniques**

Nothing to report

- **Inventions, patent applications, and/or licenses**

Nothing to report

- **Other Products**

Nothing to report

7. PARTICIPANTS & OTHER COLLABORATING ORGANIZATIONS

What individuals have worked on the project?

Name:	Karen Pollok
Project Role:	Co-PI (contact)
Researcher Identifier (ORCID ID):	0000-0001-6031-5707
Nearest person month worked:	2.7
Contribution to project:	Dr. Pollok supervised all experimental design, execution, and analysis. She was involved in all aspects of manuscript construction.
Funding Support:	No change

Name:	D. Wade Clapp
Project Role:	Co-PI
Researcher Identifier (ORCID ID):	0000-0001-9465-0917
Nearest person month worked:	1.0
Contribution to project:	Dr. Clapp advised on experimental design to increase clinical relevance for in vivo studies and data analysis.
Funding Support:	No change

Name:	Pankita Pandya
Project Role:	Co-I
Researcher Identifier (ORCID ID):	0000-0002-2021-5821
Nearest person month worked:	3.6
Contribution to project:	Dr. Pandya is in charge of all bioinformatics analyses. She works with Dr. Pollok on experimental design, execution, and analysis. She was involved in all aspects of manuscript construction.
Funding Support:	No change

Name:	Steve Angus
Project Role:	Co-I
Researcher Identifier (ORCID ID):	0000-0001-9465-0917
Nearest person month worked:	1.0
Contribution to project:	Dr. Angus oversees global kinome profiling data acquisition and analysis. He worked with Dr, Pandya on preparation of data for publication.
Funding Support:	No change

Name:	Nikan Riyahi
Project Role:	Graduate student
Researcher Identifier (ORCID ID):	0000-0002-9488-477X
Nearest person month worked:	12.0
Contribution to project:	Mr. Riyahi works with Ms. Dobrota on bench experiments including analysis of tumor tissue from in-vivo studies. He coordinates all aspects of in vivo experiments and has been trained this past year to conduct drug dosing and track tumor growth. He helps with manuscript construction.
Funding Support:	No change

Name:	Erika Dobrota
Project Role:	Research Analyst
Researcher Identifier (ORCID ID):	Not available
Nearest person month worked:	3.6
Contribution to project:	Ms. Dobrota is in charge of conducting bench research for downstream target validation studies and processing samples from in vivo experiments for RNAseq and kinome analysis.
Funding Support:	No change

Name:	Barb Bailey
Project Role:	Research Analyst
Researcher Identifier (ORCID ID):	Not available
Nearest person month worked:	2.0
Contribution to project:	Ms. Bailey prepares cells, conducts in vitro drug screens, and works with Mr. Riyahi on execution of in vivo studies and on DNA damage analysis assay.
Funding Support:	No change

Has there been a change in the active other support of the PD/PI(s) or senior/key personnel since the last reporting period?

Nothing to report

What other organizations were involved as partners?

Nothing to report

8. SPECIAL REPORTING REQUIREMENTS

COLLABORATIVE AWARDS:

QUAD CHARTS:

9. APPENDICES: Meeting abstracts and Pandya et al., Cancers 2022.

Article

Integrative Multi-OMICs Identifies Therapeutic Response Biomarkers and Confirms Fidelity of Clinically Annotated, Serially Passaged Patient-Derived Xenografts Established from Primary and Metastatic Pediatric and AYA Solid Tumors

Pankita H. Pandya ^{1,2,†} , Asha Jacob Jannu ^{3,†}, Khadijeh Bijangi-Vishehsaraei ^{1,2,‡}, Erika Dobrota ², Barbara J. Bailey ², Farinaz Barghi ^{2,4}, Harlan E. Shannon ², Niknam Riyahi ^{2,5}, Nur P. Damayanti ⁶, Courtney Young ², Rada Malko ², Ryli Justice ², Eric Albright ⁷, George E. Sandusky ⁷ , L. Daniel Wurtz ⁸, Christopher D. Collier ⁸, Mark S. Marshall ¹, Rosa I. Gallagher ⁹, Julia D. Wulfkühle ⁹, Emanuel F. Petricoin ⁹, Kathy Coy ¹⁰, Melissa Trowbridge ¹⁰, Anthony L. Sinn ¹⁰, Jamie L. Renbarger ¹, Michael J. Ferguson ¹, Kun Huang ³, Jie Zhang ^{4,*}, M. Reza Saadatzaheh ^{1,2,‡}  and Karen E. Pollok ^{1,2,*}

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† These authors contributed equally to this work.

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Simple Summary: Solid tumors account for ~60% of pediatric, as well as adolescent and young adult (AYA), cancers, and outcomes for patients with these progressive diseases remain poor. This highlights the critical need to develop tumor models from patients with aggressive cancers so that oncogenic signatures can be identified for therapeutic testing. Thus, patient-derived xenografts (PDXs) were established from sarcoma and Wilms tumor patients at diagnosis or following treatment. Overall, the molecular landscape of serially passaged PDXs recapitulated the original tumor based on an integrated multi-OMICS pipeline that cross-validated cancer-associated pathways. Actionable mechanisms of tumor progression were identified. CDK4/6 and BETs were prioritized as biomarkers of therapeutic response for in vivo validation. In osteosarcoma PDXs harboring pertinent molecular signatures, inhibition of CDK4/6 or BETs decreased growth. This systematic approach that links patient disease history to data generated from its corresponding PDX provides a foundation to discover improved therapies for patients with high-risk cancers.

Abstract: Establishment of clinically annotated, molecularly characterized, patient-derived xenografts (PDXs) from treatment-naïve and pretreated patients provides a platform to test precision genomics-guided therapies. An integrated multi-OMICS pipeline was developed to identify cancer-associated pathways and evaluate stability of molecular signatures in a panel of pediatric and AYA PDXs following serial passaging in mice. Original solid tumor samples and their corresponding PDXs were evaluated by whole-genome sequencing, RNA-seq, immunoblotting, pathway enrichment

analyses, and the drug–gene interaction database to identify as well as cross-validate actionable targets in patients with sarcomas or Wilms tumors. While some divergence between original tumor and the respective PDX was evident, majority of alterations were not functionally impactful, and oncogenic pathway activation was maintained following serial passaging. CDK4/6 and BETs were prioritized as biomarkers of therapeutic response in osteosarcoma PDXs with pertinent molecular signatures. Inhibition of CDK4/6 or BETs decreased osteosarcoma PDX growth (two-way ANOVA, $p < 0.05$) confirming mechanistic involvement in growth. Linking patient treatment history with molecular and efficacy data in PDX will provide a strong rationale for targeted therapy and improve our understanding of which therapy is most beneficial in patients at diagnosis and in those already exposed to therapy.

Keywords: pediatric; adolescents and young adults (AYA); patient-derived xenografts (PDXs); osteosarcoma (OS); rhabdomyosarcoma (RMS); Wilms tumor; multi-OMICS; precision genomics; CDK4/6; BETs

1. Introduction

Pediatric cancers, which account for approximately 1% of new cancer diagnoses each year, remain the leading cause of disease-related mortality in children [1,2]. While solid tumors are quite rare in this patient population, solid tumors represent approximately 60% of all malignant neoplasms observed in children (0–14 years), as well as adolescents and young adults (AYA; 15–39 years) in the United States [3,4]. Pediatric and AYA solid tumors highlighted in our study include osteosarcoma (OS), the most common malignant bone cancer in the pediatric and AYA populations [5]; rhabdomyosarcoma (RMS), the most common soft-tissue sarcoma in children [6]; and Wilms tumor, the most common renal cancer in children [7].

There is a critical need to identify efficacious and safe therapies to benefit pediatric and AYA patients with aggressive solid tumors (metastatic, relapsed, or recurrent cancers), wherein the 5-year survival rate is still only 0–30% [8]. In addition, there is a great desire in the oncology community to ultimately replace chemotherapies that can be overly toxic, mutagenic, and lead to long-term side effects that diminish quality of life in cancer survivors [1,9]. Despite this urgency, development of anti-cancer therapy drugs specifically for children significantly lags behind progress obtained in adults [8]. This is evident from the fact that, in 1980–2017, the US Food and Drug Administration approved only 11 anti-cancer therapy drugs for pediatric purposes [8]. The slow progress towards achieving improved treatment options for high-risk pediatric solid tumors can be, in part, attributed to factors such as the rarity of these diseases, which makes advancing potential therapeutic interventions into clinical trials rather difficult due to low patient numbers for clinical trials [8]. Furthermore, the vast amount of mechanistic and molecular genetic heterogeneity present among pediatric and AYA cancers further complicates development of targeted therapies [8].

Another major hurdle in improving pediatric cancer patients' care is attributed to the limited availability of reliable biologically relevant preclinical models [10]. While cell lines are accepted as essential preclinical tools in research, they do not always faithfully recapitulate the molecular landscape observed in the primary human tumors [11,12]. There are some cases wherein the PDX model differentiates from the original human tumor. This can be attributed to sampling bias of genetically heterogeneous tumors that may inadvertently occur [13,14]. The dissected piece of the tumor for PDX development may not fully represent the molecular landscape of the entire tumor [15], which, ultimately, may cause further clonal evolution and genetic drift compared to the original tumor [15–19]. It has also been reported that PDXs developed from advanced tumors have higher take-rate than those developed from less aggressive tumors (at early stages of disease progression), which may not fully mimic the full range of molecular landscapes present in the original tumor [13,20–23]. Additionally, since PDXs lack the immune system as well as the vasculature and stromal

components from the host, tumors that predominately rely on stromal components for growth may not provide good fidelity between the PDX and the original tumor [15].

Despite some PDX models not truly recapitulating the molecular characteristics of the original human tumor, overall concordance between key oncogenes in PDXs and their respective original human tumor are evident in the majority of PDXs shown here, as well as in other comprehensive, large-scale genomic characterization studies conducted in PDXs encompassing a variety of solid tumors, such as carcinomas, sarcomas, and melanomas [24]. In many instances, these models can be established from limited tumor tissue through serial passaging in immunodeficient mice, providing an opportunity to interrogate tumor-specific molecular signatures and test new therapies precisely designed for that tumor [11,12,25]. Therefore, molecular characterization of the original human tumor specimens (P0) with their derived PDXs is an essential step in establishing model validity [12]. In the context of pediatric and AYA cancer PDX models, Rokita et al. [26] identified mutational signatures within the PDXs that recapitulate key alterations found in pediatric CNS tumors, extracranial solid tumors, and hematologic malignancies [26]. Furthermore, Nanni and colleagues proved that serial passaging of pediatric bone sarcomas, such as OS, recapitulated the morphology and the RNA expression profile of the original patient tumor [27].

While advancements in next-generation sequencing (NGS) have shed light on identification of targeted therapies based on genomic alterations of the DNA and/or RNA, not all observed genomic alterations will result in consequential outcomes in terms of protein function [28] relevant to cancer pathogenesis. Indeed, in recent clinical studies with molecularly targeted therapies against important targets, such as AKT, pre-treatment levels of the activation/phosphorylation levels of AKT itself predicted the response; however, genomic markers that infer AKT pathway activation such as mutations in *PI3K*, *PTEN*, and *AKT1,2* did not [29].

Targeted therapies identified by single-level -OMICS analyses at the DNA, RNA, or protein/phosphoprotein level have the potential to miss therapeutic options that could be effective in heterogeneous cancers like pediatric and AYA solid tumors. Therefore, to fully understand the molecular stability of PDXs following serial passaging and to exploit the power of using novel clinically annotated PDX models to uncover therapeutic options, we developed a multi-OMICS pipeline encompassing DNA, RNA, protein, and pathway-enrichment analysis. Therapeutic response signatures/biomarkers CDK4/6 and bromodomain extraterminal domain proteins (BETs) were selected as proof-of-concept targets to explore mechanisms of tumor growth and validate anti-tumor response to small-molecule-inhibitor monotherapy in OS PDXs. This will set the foundation for obtaining further insights into the molecular mechanisms that contribute to the pathogenesis of these diseases.

2. Materials and Methods

2.1. *NOD.Cg-Prkdc Scid Il2rgtm1Wjl/SzJ* (NSG) Mice

NSG mice were obtained from the on-site breeding colony maintained by the Preclinical Modeling and Therapeutics Core (PMTC) at Indiana University Simon Comprehensive Cancer Center (IUSCCC). All procedures were approved by the Institutional Animal Care (IACUC) and Use Committee at the Indiana University School of Medicine (IUSM, IACUC studies #19052 and 22028). Animals were maintained under pathogen-free conditions and maintained on a Teklad Lab Animal Diet (TD 2014, Harlan Laboratories, Indianapolis, IN, USA) with ad libitum access to sterile tap water under a 12 h light-dark cycle at 22–24 °C.

2.2. Development of PDXs from OS, RMS, or Wilms Tumor Specimens

Tumor samples for development of the PDXs were obtained from patients following consent under IUSM IRB protocol #1501467439. The PDXs were developed as previously described by Mattar et al. [30] with some modifications. The original tumor specimens from a patient (P0) were cut into fragments that were either flash-frozen ($5 \times 5 \text{ mm}^2$), cryopreserved ($3 \times 3 \text{ mm}^2$), or implanted ($2 \times 2 \text{ mm}^2$) into the flanks of NSG mice gender-matched to that of the patient to develop PDX passage 1 (P1). Following implantation, growth kinet-

ics were tracked by electronic caliper connected to StudyLog software (South San Francisco, CA, USA). Briefly, the mice were monitored at least twice weekly for overall health and more frequently if clinical signs were noted. During expansion phase, tumor volumes were measured twice weekly by electronic caliper for up to 300 days or until tumor size was 1000–1500 mm³. Tumor volumes in this range were preferred to minimize necrotic tissue in the tumors. Immediately after P1 PDX extraction, the tumors were cut to 2 × 2 mm² fragments under aseptic conditions, and then 1–2 PDX tumor tissue fragments were implanted into the flank of 6–8-week-old gender-matched NSG mice to generate PDX passage 2 (P2) tumors. The same process was conducted to obtain PDX passage 3 (P3) tissue. All P0 and corresponding PDXs were authenticated for their identity/species by DNA fingerprinting analysis using a 16- marker short-tandem repeat (STR) analysis (IDEXX BioResearch, Columbia, MO, USA). The first priority for tumor samples at our institution is for diagnosis and specimens required for ongoing clinical trials. Generation of PDX is the next priority. Tissue quantity can vary greatly from patient to patient. The PDXs characterized in this study are a subset of our larger pediatric and AYA PDX panel of rare tumors. Over the past several years, our success rate was defined as expansion of PDX at least through passage 2. In OS, 51.4% cases resulted in successful PDX generation. For RMS cases, 53.8% led to successful PDX generation, and 80% of Wilms tumor cases gave rise to successful Wilms tumor PDXs. For the biopsy samples used for PDX generation, the surgeon's expertise and tumor scans were used to identify viable tumor tissue. For resection samples used for PDX generation, the pathologists release viable tumor based on their expert analysis of the tissue sent by the surgical team. To transfer the tissue sample to the lab for further processing, MACS tissue storage medium was used (Miltenyi Biotec, cat # 130–1100-008).

2.3. DNA Extraction

Total genomic DNA was isolated from flash-frozen P0 and PDX samples (*n* = 30 samples) using the QIAamp DNA Mini Kit (Qiagen, Germantown, MD, USA) or DNeasy Blood and Tissue Kit (Qiagen, Germantown, MD, USA), as indicated by the manufacturer's instructions. Subsequently, DNA quantity and was assessed by nanodrop, QUBIT Fluorometer, and Agilent 2100 Bioanalyzer.

2.4. Bioinformatic Data Analysis

2.4.1. Somatic Whole-Genome Sequencing (WGS) Analysis

WGS was performed at New York Genome Center (NYGC, New York, NY, USA) on the following PDXs: HT72, HT77, HT87, HT96, HT74, and HT98. For these samples, TruSeq PCR-free preparation was done. WGS on HT120 and HT139 was performed at IUSM Center for Medical Genomics core using the Illumina Nextera DNA Flex Library Prep Kit. For all P0 specimens and PDX samples, the sequencing depth/coverage was 90×. All SNV and CNV data will be uploaded to the dbGaP database (<https://www.ncbi.nlm.nih.gov/gap/>). OS, RMS, and Wilms tumor cohorts were sequenced using the NovaSeq 6000 Illumina platform to generate raw reads in FASTQ format. The raw sequencing data for the control sample, NA12878, were obtained from the International Genome Sample Resource and used as the normal reference human genome. Adapter sequences, primers, and other unwanted sequences were first identified and trimmed using Cutadapt [31] and later filtered against mouse genome using BBsplit [32] from BBtools suite to remove mouse-originated reads. The raw sequencing reads were aligned to the human reference genome hg38 with Burrows–Wheeler aligner (BWA) [33].

The objective of the bioinformatics analyses was to compare the copy number variations (CNVs) and single-nucleotide variations (SNVs) of each P0 versus respective PDX passages as well as to compare within each set of PDX passages.

2.4.2. Copy Number Variation (CNV) Analysis

CNV analysis was performed using CNVkit [34]. The copy number profile of NA12878 was used as the reference to correct the biases in regional coverage and GC content in P0 and

its respective PDX passages of HT72, HT77, HT87, HT96, HT74, HT98, HT120, and HT139. Using the circular binary segmentation, genome-wide segmented copy number estimates for each sample were generated based on fixed thresholds. Genes with significant deletions and duplications in each call set were determined using the “genometrics” command with default thresholds on log2 ratios and minimum number probes per segment. Within each P0–PDX cohort with NA12878 as the reference, the genes with differential copy number estimates in at least one of the PDX passages were called. To identify cancer-associated genes, 723 genes from the Catalogue of Somatic Mutations in Cancer (COSMIC) database [35] were used to further filter the copy number call set of each PDX cohort. R packages such as “ape” was used to plot the dendrogram to understand the genomes of PDXs, “pheatmap” to generate heatmaps of differential copy number profiles for each cohort and “ggplot” to visualize the overview of genome-wide copy number alteration distribution on the chromosomal level. Correlation bar plots were created to visualize the P0–PDX cohorts. A copy number of 3–5 was designated as copy number gain, whereas a copy number ≥ 6 was referred to as copy number amplification [36]. Anything fewer than two copies was designated as copy number loss or deletion.

2.4.3. Single-Nucleotide Variation (SNV) Analysis

Sentieon Suite was used for data analysis to generate the SNV/Indels calls in each P0–PDX cohort for HT72, HT77, HT87, HT96, HT74, HT98, HT120, and HT139 [37]. The SNVs identified by the variant calling process, and their variant consequences were annotated using Ensembl Variant Effect Predictor (VEP) with loaded genomic prediction algorithms and databases, such as Ensembl’s Variant Consequences, SIFT, PolyPhen, dbSNP, ClinVar, and COSMIC. For SIFT, if the probability score is <0.05 , then the variant is predicted to alter protein function and score of >0.05 is predicted to be tolerated and not impact protein function [38,39]. Polyphen scores >0.908 for variants are characterized as “probably damaging” [39,40]. Variants with scores between 0.556–0.908 are designated as “possibly damaging” [40], and scores ≤ 0.446 suggests that those variants are “benign”, indicating no impact on protein function [39]. Variant haplotypes were called using NA12878 as the reference, which included any variants in P0 or the corresponding PDX passages in each tumor. They were verified by their presence of nucleotide mutations and/or significant shifts in variant allele frequencies, such as losses of heterozygosity (LOH), when compared to NA12878. We further filtered the variants according to the calls’ quality and the impact on function. Therefore, only the variants of high quality (depth ≥ 10 , genotype quality ≥ 20 , allelic balance between 0.2–0.8) that are predicted to be of high impact and to have a deleterious effect on functions (polyphen score >0.15 and SIFT ≤ 0.05) were retained [41]. The mutations in genes involved in important molecular processes, as well as oncogenic pathways, were determined. An in-house R package was developed to generate oncoprint plots to visualize a differential mutation profile in tumor samples compared to NA12878. MAFtools and ggplot2 were used to summarize the SNV statistics, i.e., count of variants per variant class, consequences, and transitions/transversions.

2.4.4. RNA-Seq Data Analysis

RNA extraction and paired-end RNA-seq sequencing were conducted at Quick Biology, Inc (Monrovia, CA USA; Illumina sequencing of 30M, PE150). Human Adipose Tissue Poly A+ RNA (Takara Cat# 636162) was used as the filtering control. The sequencing reads were first mapped to the latest UCSC transcript set using Bowtie2 version 2.1.0 [42], and the gene expression level was estimated using RSEM v1.2.15 [43]. The BBsplit function in the BBmap tool was utilized to remove the mouse reads. Differentially expressed genes were identified using the edgeR program [44]. Genes showing altered expression with $p < 0.05$ and more than 1.5-fold changes were considered differentially expressed. Pearson correlation analysis was performed to study the similarity between the somatic sample, P0, and the PDXs, for each dataset. Goseq [45] was used to perform the gene ontology (GO) enrichment analysis,

and KOBAS was used to perform the pathway analysis [46]. Additionally, two other pathway databases (KEGG and Reactome) were also utilized for enrichment analysis.

2.5. Western Blot Analysis

Protein extraction of PDX tumor samples was conducted using either UREA or radioimmunoprecipitation assay (RIPA), based on reports suggesting that protein extraction from solid tumor samples require different methods (i.e., UREA and/or RIPA) due to differences in the cytoplasmic location, solubility, and molecular weights of the protein of interest [47]. Due to the limited availability of P0 samples, they were not used for Western blot analysis. Flash-frozen PDX tumor samples were pulverized by smashing twice with a cryo-PREP® Automated Dry Pulverizer (Covaris, Woburn, MA, USA). Approximately 25 mg PDX powder was lysed in 300 µl 8 M Urea Buffer (8 M urea, 100 mM Tris pH 8, 5 mM DTT) or 300 µl RIPA buffer containing protease and phosphatase inhibitors and processed with a Bioruptor® Plus sonication device on high setting for 5 cycles [30 s on; 30 s off] at 4 °C (Diagenode, Denville, NJ, USA). Debris was spun out, and the protein concentration of the cleared lysate was determined by RCDC Protein Assay (Bio-Rad, Hercules, CA, USA) and quantified using BioTek Synergy H4 (BioTek, Winooski, VT, USA). For Western blot analysis, proteins were separated on TGX Stain-Free™ gels (Bio-Rad, Hercules, CA, USA) along with Precision Plus Protein™ All-Blue Standards (Bio-Rad, Hercules, CA, USA) and transferred to LF PVDF membrane using the Trans-Blot Turbo Transfer System (Bio-Rad, Hercules, CA, USA). Membranes were blocked for 1 h at room temperature in 5% non-fat dry milk in TBS-T (137 mM NaCl, 20 mM Tris, and 0.1% Tween 20). Membranes were washed following antibody incubation with TBS-T for a total of three 10 min washes. The correct molecular weight for each protein was confirmed by the Precision Plus All Blue Standard (Bio-Rad, Hercules, CA, USA). The following antibodies were diluted in either 5% non-fat dry milk or 5% BSA in TBS-T per manufacturer's instruction and used for detection: rabbit anti-RAD21 (130 kDa, cat# 4321, Cell Signaling Technology, Boston, MA, USA); mouse anti-c-MYC [9E10] (67 kDa, cat# sc-40, Santa Cruz Biotechnology, Inc., Dallas, TX, USA); mouse anti-p53 [D0-1] (53 kDa, cat# sc-126, Santa Cruz Biotechnology Inc.); mouse anti-Cyclin D3 (31 kDa, cat# 2936, Cell Signaling Technology, Boston, MA, USA); mouse anti-Cyclin E1 (48 kDa, cat# 4129, Cell Signaling Technology, Boston, MA, USA); rabbit anti-CDKN2A/p16^{INK4A} (17 kDa, cat# ab108349, Abcam, Waltham, MA, USA); rabbit anti-PTEN (54 kDa, cat# 9559, Cell Signaling Technology, Boston, MA, USA); rabbit anti-RAC1 (21 kDa, cat# 4651, Cell Signaling Technology, Boston, MA, USA); rabbit anti-HSP90Aβ1 (96 kDa, cat# NBP2-68937, Novus Biologicals, Centennial, CO, USA); anti-phospho-H2AX serine 139 [γH2AX-Ser139] (15 kDa, cat# 2577, Cell Signaling Technology, Boston, MA, USA); rabbit anti-total H2A.X [D17A3] (15 kDa, cat# 7631, Cell Signaling Technology, Boston, MA, USA); rabbit anti-RB1 [D20] (110 kDa, cat# 9313, Cell Signaling Technology, Boston, MA, USA); rabbit anti-phospho-RB1 Ser795 (110 kDa, cat# 9301, Cell Signaling Technology, Boston, MA, USA); rabbit anti-BRD4 [E2A7X] (200 kDa, cat# 13440, Cell Signaling Technology, Boston, MA, USA); rabbit anti-CDK4 [D9G3E] (30 kDa, cat# 12790, Cell Signaling Technology, Boston, MA, USA); rabbit anti-CDK6 [D4S8S] (36 kDa, cat# 13331, Cell Signaling Technology, Boston, MA, USA); rabbit anti-vinculin [E1E9V] (124 kDa, cat# 13901, Cell Signaling Technology, Boston, MA, USA), and rabbit anti-GAPDH [14C10] (37 kDa, cat# 2118, Cell Signaling Technology, Boston, MA, USA). Blots were then incubated with the appropriate horseradish peroxidase-conjugated secondary antibody, diluted 1:5000 in 5% non-fat dry milk-TBST for 1 h at room temperature (Anti-mouse IgG HRP-conjugate, cat# W4021, Promega, Madison, WI, USA; Anti-rabbit IgG HRP-conjugate, cat# W4011, Promega). Membranes were again washed following secondary antibody incubation with TBS-T for a total of three 10 min washes. Proteins were detected using SuperSignal Western Chemiluminescent Substrate (Thermo Fisher Scientific; Waltham, MA, USA) and imaged using the Bio-Rad ChemiDoc Imaging System (Bio-Rad, Hercules, CA, USA). Human vertebral mesenchymal stem cells (HVMSCs; CP7550 ScienCell Research

Laboratories, Carlsbad, CA, USA), as well as human osteoblasts-femoral (HOB-F; CP4610 ScienCell Research Laboratories, Carlsbad, CA, USA), were utilized as our controls.

2.6. Histologic Characterization of Original Tumor Specimen and PDX

For analysis of tissue integrity, tumor tissues from P0 and their corresponding passaged PDXs were fixed in 10% neutral buffered formalin at 4 °C for 72 h and then embedded in paraffin. As previously described by our group, 5 µm sections were cut and flattened on a heated water bath, floated onto microscope slides, and dried as previously described [5]. For routine assessment, slides were stained with hematoxylin and eosin [5]. Microscopic analysis of hematoxylin and eosin (H&E) stains was conducted in a blind fashion by a board-certified pathologist. Slides were scanned using Aperio whole-slide imaging software (Sausalito, CA, USA).

2.7. Protein Pathway Activation Mapping via Reverse Phase Protein Array (RPPA) Analysis of FFPE Xenograft Samples

2.7.1. Xenograft Tissue Processing and Generation of Whole-Tissue Lysates

FFPE-embedded xenograft tissue blocks were sectioned at 8 µm onto glass slides. RPPA analysis was conducted on original P0 tumors when available and their respective PDXs for HT72, HT77, HT87, HT96, HT74, HT98, HT120, and HT139. Between 2 and 6 sections per sample were de-paraffinized in xylene, rehydrated in 100% and 95% ethanol, briefly fixed in 70% ethanol containing protease inhibitors (Complete Mini EDTA-free, Roche, Indianapolis, IN, USA), then dehydrated in 95% ethanol, 100% ethanol and then lysed directly from slides in an appropriate volume of extraction buffer containing 50% Tissue Protein Extraction Reagent (T-PER, Thermo Fisher Scientific; Waltham, MA, USA), 47.5% 2× Tris-Glycine SDS sample buffer (Invitrogen, Waltham, MA, USA), and 2.5% β-mercaptoethanol (Thermo Fisher Scientific, Waltham, MA, USA). The resulting whole-tissue lysates were heated for 8 min at 100 °C. Samples were stored at −80 °C and reheated at 100 °C for 2 min, vortexed, and briefly centrifuged just prior to printing.

2.7.2. Array Printing and Analysis

The total protein concentration in each sample was estimated by printing onto a nitrocellulose slide (Avid, Grace Biolabs, Bend, OR, USA) along with a BSA standard concentration curve using a Quanterix 2470 Arrayer equipped with 180 µm pins at a deposition volume of ~11 nL per sample. Total protein levels were assessed by staining with Sypro Ruby Protein Blot Stain (Invitrogen) according to the manufacturer's instructions. Tissue lysate concentrations were adjusted to a maximum of 250 µg/mL with extraction buffer. RPPA printing for antibody staining was performed as described above, and antibody staining and image analysis of xenograft samples was conducted as previously described [48–51]. Briefly, signaling pathway activation was evaluated by staining the arrays with 123 antibodies against signaling endpoints, mainly phosphorylated and cleaved protein products. Before use for RPPA analysis, antibody specificity was confirmed by Western blot and analysis as previously described [51]. Antibody staining intensities were generated using the MicroVigene v5 Software Package (Vigenetech; Carlisle, MA, USA).

2.7.3. RPPA Statistical Analysis

Each data value represents the average of 3 replicate printed spots. All final data values were normalized to total protein following the subtraction of negative control. Additionally, all data values within an endpoint, such as a specific kinase, can be compared to each other, but data values between different endpoints such as different kinases or signaling proteins cannot be directly compared. If a zero value is obtained, then those are true zero values where spot intensities were either not above local background or not above negative-control spot intensities. To ensure data reliability, we prioritized RPPA data wherein coefficients of variation (CV) among the replicate spots were ≤10% [52].

2.8. Compounds

Palbociclib (CDK4/6 inhibitor) and OTX015 (BET inhibitor) were both purchased from Chemietek (Indianapolis, IN, USA). For in vivo studies, palbociclib was formulated in 50 mmol/L sodium lactate, pH 4, and OTX015 was formulated in 10% DMSO, 20% PEG400, 5% Tween 80, and 65% water.

2.9. Screening of OS PDX with Small-Molecule Inhibitors Based on Therapeutic Response Biomarkers

Through the multi-OMICS analytical pipeline described in this study, high-risk molecular signatures in OS PDXs derived from patients with aggressive disease were prioritized based on our interrogation of the Drug Gene Interaction Database (DGIdb; <https://www.dgldb.org/> (accessed on 7 July 2022)), as well as evidence from the literature for in vivo screening. For these in vivo efficacy studies, HT77 and HT96 OS PDX tumor fragments were implanted into gender-matched NSG mice. Once tumor volumes reached 100–200 mm³, mice were randomized and treated with either 120 mg/kg palbociclib or 25 mg/kg OTX015. HT77 PDX mice were treated with vehicle ($n = 8$) or palbociclib ($n = 7$) at 120 mg/kg for 5 consecutive days PO, SID, daily for 3 weeks. HT96 PDX mice were treated for ~6 weeks with 25 mg/kg OTX015 ($n = 5$) or vehicle control ($n = 6$), BID, PO, daily for 5 consecutive days. For all in vivo studies, tumor volumes and body weights were evaluated 2–3 times per week.

2.10. Statistical Analysis

All data are reported as mean $\pm$ SEM. Data were considered significant at $p < 0.05$. For in vivo studies, tumor growth data was evaluated by two-way ANOVA with a Holm–Sidak multiple comparison test where * $p < 0.05$, # $p < 0.001$, ^ $p < 0.0001$. GraphPad Prism Software (GraphPad Inc., San Diego, CA, USA) was used for data analysis and visualization.

3. Results

3.1. Comprehensive Clinical Annotations of PDX Models Established from OS, RMS, and Wilms Tumor Patients at Different Phases of Therapy

The workflow of the PDX pipeline consists of the direct implant of the original tumor samples into NSG mice, followed by serial passaging of the samples to expand PDX tissue for study (graphical abstract). This is followed by STR authentication and multi-OMICS analyses of P0 versus the corresponding PDX passages (P1, P2, and/or P3 when available). Cross-validation of -OMICS data such as CNVs at the protein level provides further confirmation of the molecular signatures. Importantly, this pipeline links the patient disease journey prior to and after sample collection to -OMICS data, integrated pathway analysis, and actionable signatures with target-specific drugs through the Drug Gene Interaction Database. This provides a solid platform to prioritize targeted therapies and evaluate in vivo efficacy and safety. In partnership with the Pediatric Precision Genomics Program at Riley Children’s Hospital, we established a panel of pediatric and AYA solid-tumor PDXs from tumor specimens of naïve and pre-treated patients diagnosed with OS, RMS, or Wilms tumor. Comprehensive clinical annotations for each patient including diagnoses, demographics, and disease status are shown in Table 1. Details of treatment plans and therapeutic responses prior to and following the acquisition of tumor samples for PDX development are provided in Table S1.

HT72 and HT77 PDXs were derived from metastatic sites in a heavily pre-treated male OS patient at different time points during salvage therapy. HT87 PDX was established from a treatment-naïve female OS patient that was presented with localized disease classified histologically as high-grade OS, received MAP therapy (methotrexate, doxorubicin, and cisplatin), and at the time of publication was off therapy. HT96 PDX was derived from a localized tumor site from a treatment-naïve male OS patient that presented with metastatic disease at the time of diagnosis. After the sample was acquired for PDX development, the patient underwent frontline and multiple salvage therapies but eventually succumbed to

disease and is now deceased. HT74 PDX was generated prior to therapy from a female patient with an unclassified high-grade RMS with widely metastatic and advanced disease at diagnosis that did not respond to therapy and is deceased. Wilms tumor models were selected from patients with disease classifications ranging from Stages II–IV. Both Wilms tumors (HT98 and HT139) exhibited localized disease; however, complete response to chemotherapy and nephrectomy was achieved in both cases. Wilms tumor HT120 was derived from a metastatic lesion in the liver of a pre-treated female patient that has presented with multiple recurrences of metastatic disease since the PDX was established.

Table 1. Overview of patient characteristics for the PDX panel.

PDXs	Diagnosis	Primary or Progressive Sample	Biopsy or Resection	Gender	Race	Age	Disease Status
HT72	OS	Progressive	Biopsy	Male	Caucasian	18	Deceased
HT77	OS	Progressive	Resection	Male	Caucasian	18	Deceased
HT87	OS	Primary	Biopsy	Female	Caucasian	17	High-grade
HT96	OS	Primary	Biopsy	Male	Caucasian	9	Deceased
HT74	RMS	Primary	Biopsy	Female	Caucasian	14	Deceased
HT98	Wilms tumor	Primary	Biopsy	Male	Caucasian	3	C.R.
HT120	Wilms tumor	Progressive	Resection	Female	Caucasian	9	Deceased
HT139	Wilms tumor	Primary	Resection	Male	Caucasian	8	C.R.

C.R. = complete response.

3.2. Maintenance of Histological Fidelity in PDX Models

Sections of the serially passaged PDXs were analyzed by H&E and fidelity evaluated by a blinded pathologist. Overall, the PDX-passaged tumors maintained the histological fidelity of their original tumors (Figure S1). Though, as has been reported by others, human stromal components are lost following engraftment and serial passaging [53]. Once solid tumors engraft and propagate in the murine host, the PDX tumors are comprised of human tumor cells and mouse stromal tissues [54–56]. WGS analysis indicated that the percentage of mouse reads was similar in the majority of the PDX passages (8–12%) with one exception. HT87 mouse reads were 21–27% mouse across the HT87 P1–P3 passages (Figure S2). It has been reported that murine stromal content can either be conserved or vary among tumors irrespective of whether they originate from similar or different cancers [57].

3.3. Genome-Wide Analysis of P0 Tumor Specimens versus Their Respective PDX Serial Passages

Genome-wide variant analysis was done to identify and compare the molecular signatures at the DNA level within each P0–PDX cohort. Multiple levels of comparative DNA analysis were conducted using WGS data of each P0–PDX cohort. In the first level of analysis, an overview of the genomic landscape where similarities and differences in DNA sequences of the P0 versus the respective passages and among the PDX passages for each individual model are described irrespective of potential impact on protein expression and function. This is complemented with a focused analysis on to what extent oncogenic-associated genes are differentially impacted by serial PDX passaging.

3.4. Comparative Analysis of CNV Profiles across the Genome

Circular binary algorithm was used to segment the genomic regions with similar average log₂ ratios across all genes for P0 tumor specimens and the corresponding PDXs for HT72, HT77, HT87, HT96, HT74, HT98, HT120, and HT139. NA12878 served as the normal human genome filtering control, as it is a well-recognized benchmark dataset [58]. To evaluate overall genomic stability of serially passaged PDXs, the chromosomal distribution of all CNVs across the genome were visualized by chromplots (Figures S3–S10) [59]. Compared to OS PDXs (HT72, HT77, HT87, HT96) and RMS PDX (HT74), the Wilms tumor PDXs (HT98, HT120, and HT139) had fewer CNV events (losses/deletions or gains/amplifications), consistent with the fact that Wilms tumors tend to be more genomically stable tumors, as illustrated by chromplots (Figures S3–S10) and in the detailed analysis below.

CNV similarities versus differences between the P0 sample and its subsequent passages (P1, P2, and/or P3), as well as among the passages for each model, were quantified by Pearson's correlation coefficient (PCC) score and plotted on correlation heatmap plots (Figure 1) [60,61]. A Pearson's positive correlation coefficient (PPCC) score of 1 signifies complete concordance between the datasets and score of 0 indicates complete discordance [17,61]. PPCC scores are classified as weak (0–0.3), moderately positive (0.3–0.5), strongly positive (0.5–0.7), or very strongly positive (0.7–1) [61]. OS PDXs HT72 and HT77 (from same patient at different phases of therapy), as well as HT96 cohorts, exhibited a very strong positive correlation between P0 and passaged PDXs for total CNVs (PPCC scores, 0.9–0.98 for HT72 and HT77; PPCC scores of 0.72–0.94 for HT96 Figure 1A), and COSMIC-associated CNVs (PPCC scores 0.90–0.98 for HT72 and HT77; PPCC scores of 0.75–0.95 for HT96 Figure 2A). While OS PDX HT87 exhibited relatively more discordance than the other OS PDX, a strong positive correlation between P0 and its respective PDX passages was still evident for total CNVs (PPCC scores, 0.57–0.97; Figure 1A) and HT87 COSMIC-associated CNVs (PPCC scores 0.56–0.97; Figure 2A). Whether this discordance is attributable to the lower number of human reads that could be called due to high murine stromal levels (21–27% mouse reads across the passages, Figure S2) and/or clonal divergence is not clear. The highest degree of divergence between the P0 and the PDX passages was found in the RMS HT74 PDX. PPCC scores for the RMS HT74 PDX were moderately positive for the total CNVs (P0 versus P1, PPCC = 0.32; P0 versus P2, PPCC = 0.34; Figure 1B). Moreover, the COSMIC-associated CNVs were even more discordant than the total CNVs presenting weak PPCC scores (P0 versus P1, PPCC = 0.25; P0 versus P2, PPCC = 0.21; Figure 2B), consistent with genetic drift of the serially passaged PDX at least at this level of DNA analysis. Of note, once the sarcoma PDXs were established, CNVs across the passages of all PDXs stabilized. For Wilms tumor PDXs (HT98, HT120, and HT139), very strongly positive correlation was observed between each P0 and respective PDX passages for total CNVs (PPCC scores = 0.94–0.96 for HT98, PPCC scores of 0.91–0.98 for HT120, and PPCC scores of 0.93–0.98 for HT139; Figure 1C). Similarly, COSMIC-associated CNVs also showed strong positive correlation in Wilms tumor samples (PPCC scores = 0.96–0.99 for HT98, PPCC scores of 0.89–0.97 for HT120, and PPCC scores of 0.92–1 HT139; Figure 2C), thus highlighting the relatively low levels of inherent genome instability in these tumors [52].

To provide additional insight into the degree of changes found in the P0–PDX sets, the absolute number of losses and gains/amplifications of the CNVs in each COSMIC-associated gene are presented in Table S2A–H. At this level of analysis, it becomes evident that most differences in copy number losses, as well as gains/amplifications in the COSMIC-associated genes, are modest. As an example, all amplifications (>6 copies) and copy number gains (<6 copies) exhibited only minor differences of 1–2 copies across the P0 and the respective PDX passages (Table S2A–H). Allelic losses were overall maintained with some exceptions highlighted below.

CNV profiles of 25 genes from the COSMIC database were prioritized based on their expression and the role they have in that particular cancer type. For example, across pediatric and AYA OS PDXs, *ATRX* gene deletions (0–1 copies present) were evident (Figure 3A–D; Table S2A–D). Previous studies have shown that deletions in *ATRX* resulting in loss of protein expression can contribute to pediatric and AYA OS tumorigenesis [62,63]. OS PDXs also had increased *MYC* and *RAD21* copy number gains (3–7 copies; Figure 3A–D; Table S2A–D), both of which are known cancer markers with multi-functions leading to increased DNA replication and oncogenic-induced replication stress, a hallmark of aggressive OS [5]. Notably, HT72, HT77, and HT87 acquired a deletion of *RB1* in one allele during the PDX serial passaging (Figure 3A–C; Table S2A–C), which was not seen in the respective P0 sample, whereas HT96 maintained the *RB1* deletion found in the original tumor (Figure 3D; Table S2D). A bi-allelic *PTEN* deletion was evident in the OS HT72 PDX that was not found in the P0 sample or the OS PDX HT77 derived from the same patient at a later resection (Table S2A,B). It should be noted that not every gene will appear in each PDX cohort. As an example, *ATM* has a copy number gain present in P0, P1, P2, and P3 of HT96 that was not observed

in NA12878 normal control sample that was used as the filter. However, ATM copy number gain, amplification, or deletion in Figure 3A–C (HT72, HT77, and HT87) is not listed, since the copy number change was not different between NA12878 vs. P0, P1, P2, and/or P3. In the unclassified RMS HT74, *CCND1* gain (3–5 copies) and a *CDKN2A* deletion (0 copies) were found in both the P0 and the respective passages, which is a common alteration found in RMS [64,65] (Figure 3E; Table S2E). *CCND1* had copy numbers of 3 and 5 in HT74-P1 and HT74-P2, respectively, and both alleles of *CDKN2A* were deleted in HT74-P1 and HT74-P2 (Figure 3E; Table S2E). In primary encapsulated Wilms tumors (HT98 and HT139), deletion of both alleles of *AMER1* was detected in both the P0 and respective PDXs (Figure 3F,H; Table S2F,H). *WTX* has been reported to be mutated, truncated, or deleted in 30% of Wilms tumor cases [66,67]. However, clinical significance of *WTX* alterations are still unclear. When assessing the CNV profile of HT120, which was derived from a metastatic Wilms tumor of the liver (Figure 3G; Table S2G), no deletions for *WTX* were evident. Though CNVs in genes involved with cell cycle regulation were evident, such as *CDK4* and *CCND2* copy number gains (three copies of each gene) compared to primary Wilms tumor samples HT98 (Table S2F) and HT139 (Table S2H). Of note, copy number gains of *KRAS* (3 copies) were found in the metastatic Wilms tumor HT120-P0 and corresponding passaged PDXs (Figure 3C; Table S2G). Other groups have reported activating mutations of *KRAS* in human Wilms tumor samples that correlate with hyperactivation of the WNT pathway [66,68].

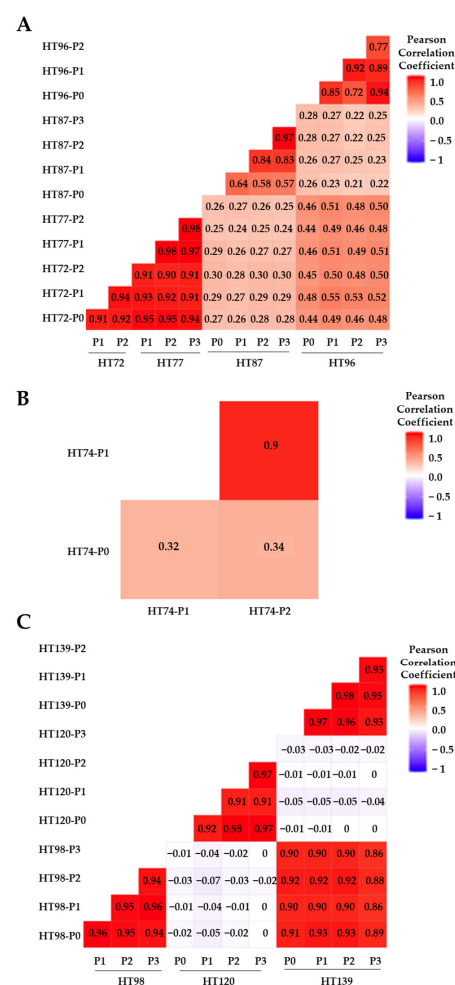


Figure 1. Analysis of congruence in total genes containing CNVs in the original P0 tumor and corresponding PDX passages. Heatmap shows PPCC scores for CNVs in P0 tumors and serial passages of (A) OS PDXs HT72, HT77, HT87, HT96, (B) RMS PDX HT74, and (C) PDXs for Wilms tumors HT98, HT120, and HT139.

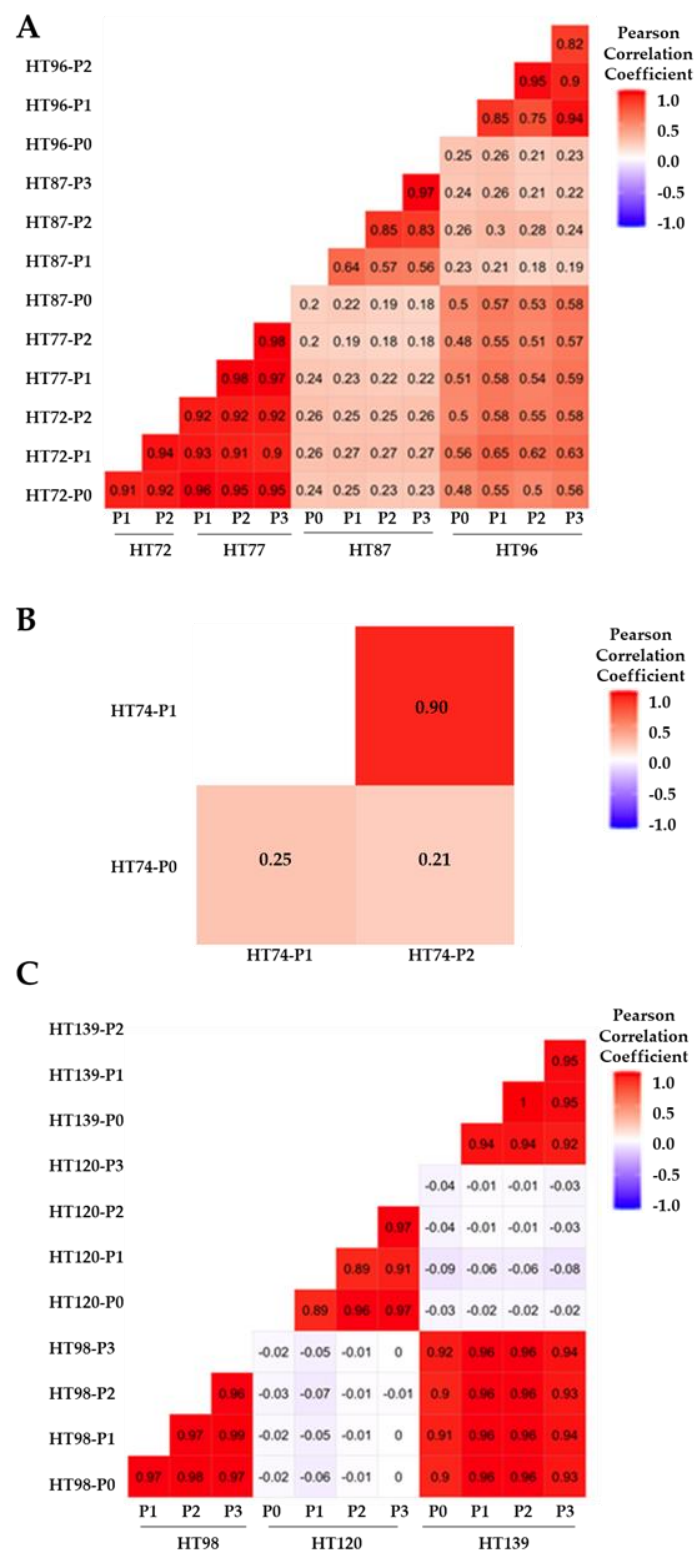


Figure 2. Analysis of similarity in COSMIC genes containing CNVs in the original P0 tumor and corresponding PDX passages. Heatmap shows PPCC scores for CNVs in P0 tumors and serial passages of (A) OS PDXs HT72, HT77, HT87, HT96, (B) RMS PDX HT74, and (C) PDXs for Wilms tumors HT98, HT120, and HT139.

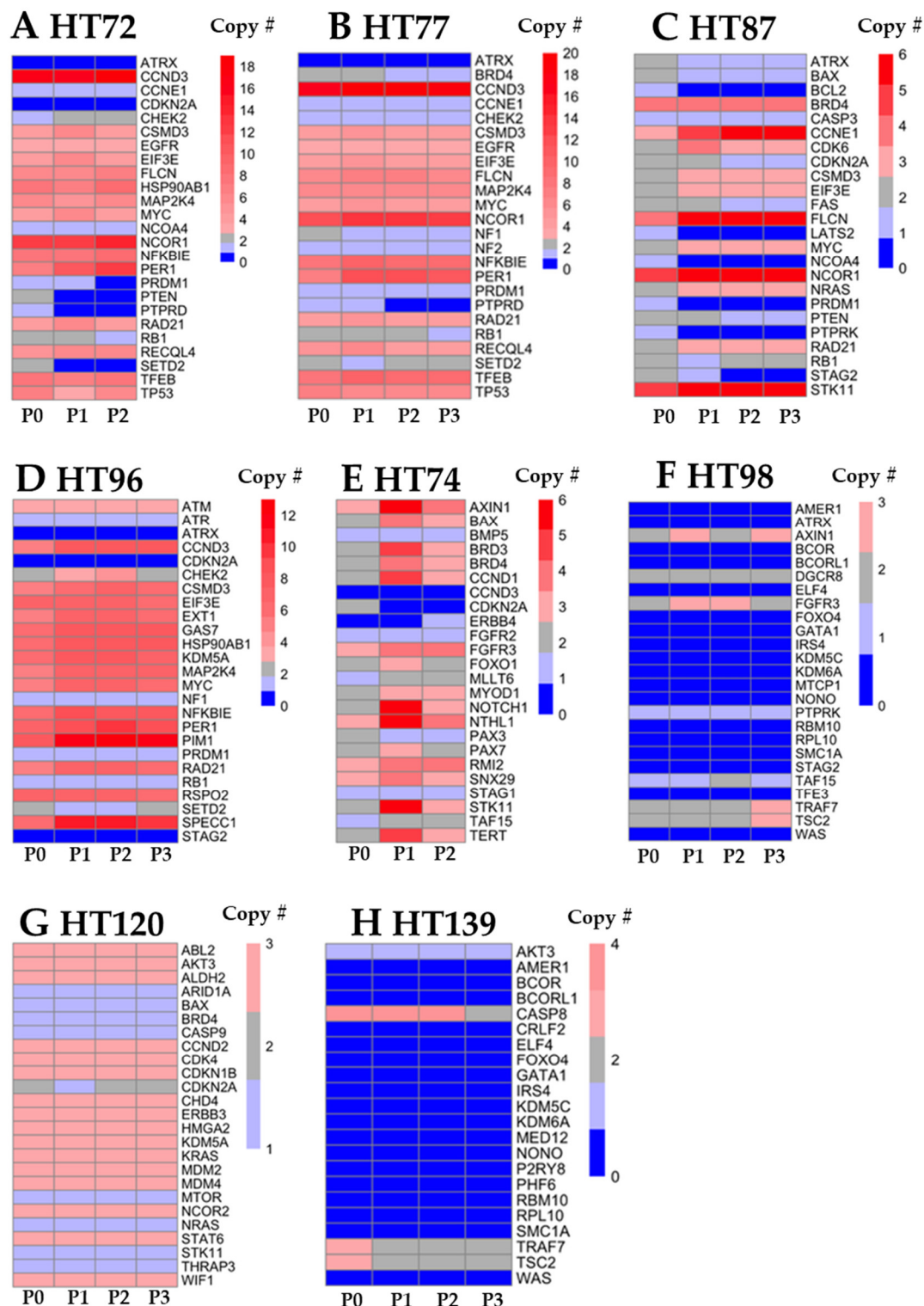


Figure 3. Distribution of copy number events involving 25 COSMIC-associated genes in P0 tumors and corresponding PDX passages. (A) HT72 (OS), (B) HT77 (OS), (C) HT87 (OS), (D) HT96 (OS), (E) HT74 (RMS), (F) HT98 (Wilms tumor), (G) HT120 (Wilms tumor), and (H) HT139 (Wilms tumor). All data were filtered against normal control (NA12878). Copy # = copy numbers; Neutral = 2 copies, Deletion < 2 copies, Gains = 3–5 copies, Amplification > 6 copies. Note: HT72 and HT77 PDX were developed from the same patient at different time points. Due to low quantities of P0 HT77 tissue, P0 HT72 was used for both HT72 and HT77 analyses.

3.5. Integration of Prioritized Cancer-Associated CNVs in OS with Corresponding Protein Expression

To validate the impact of copy number gains or amplifications and losses on gene expression, protein levels were analyzed for cancer-associated pathways in the OS PDXs (Figure 4). Due to limited availability of P0 samples, they were prioritized for WGS and RNAseq. Specific CNV-protein associations (Figure 4) were selected due to their importance in OS pathogenesis. RAD21, MYC, γ H2AX, total H2AX, and P53 all have been reported to play roles in the DNA replication, replication stress, and genome instability in OS [5,69]. Additionally, amplification of cell cycle proteins cyclin D3 (CCND3) and cyclin E1 (CCNE1), as well as CDKN2A deletion, are common signatures found in high-risk OS [70–72], all of which result in uncontrolled cell cycle progression in OS [72]. Moreover, roles for RAC1 [73] and HSP90A β 1 [74] have also been reported in OS pathogenesis.

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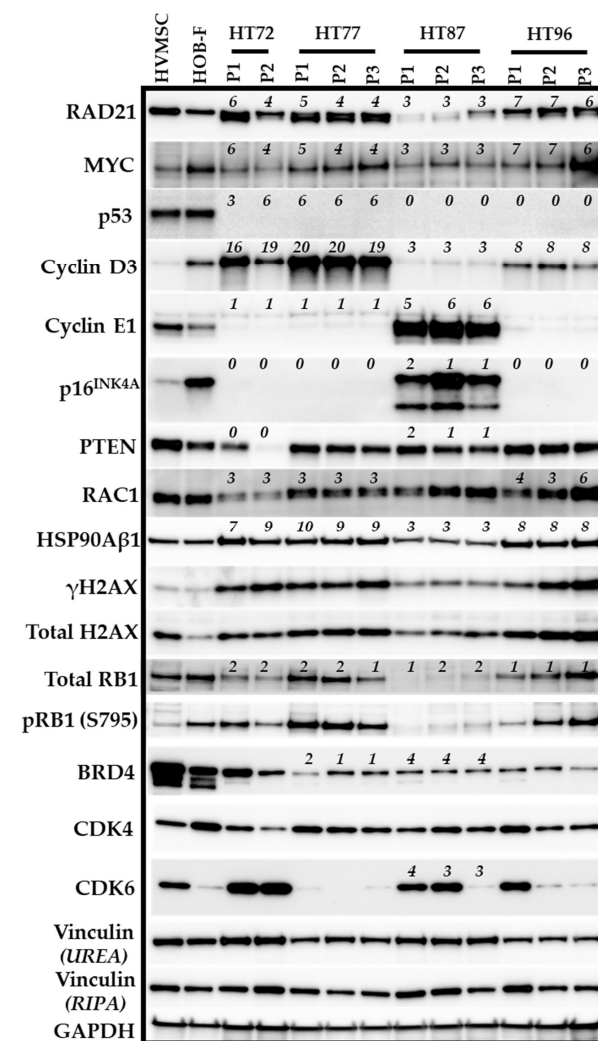


Figure 4. Impact of CNVs in oncogenic-associated genes on corresponding protein levels. CNV-protein links were evaluated by Western blots on tumor lysates from passages (P1, P2, and P3) of the OS PDXs (HT72, HT77, HT87, and HT96). Baseline levels of selected proteins of interest were: RAD21, MYC, p53, Cyclin D3, Cyclin E1, p16^{INK4A}, PTEN, RAC1, HSP90A β 1, γ H2AX, total H2AX, p-RB1 (S795), total RB1, BRD4, CDK4, and CDK6. Vinculin served as the loading control and GAPDH as the housekeeping gene. Copy numbers for genes containing CNVs are the top of each immunoblot. This experiment was conducted one time. Copy numbers for genes containing CNVs are the top of each immunoblot.

Cross validation of CNVs (Table S2) and protein expression (Figure 4) demonstrated congruence of copy numbers and protein expression for *CCND3* in the HT72 and HT77 OS PDXs (Figure 4, copy number of 16–20), as well as for *CCNE1* in the HT87 PDXs (Figure 4, copy number of 5–6). In some cases, copy number gains/amplifications or losses/deletions (Table S3A,B) do not correlate with the corresponding protein expression (Figure 4). For example, as mentioned above, *PTEN* in HT72 is reported to have zero copies in both passages (Table S2A). However, *PTEN* protein was still present in HT72-P1 but absent in HT72-P2, which correlates with the lack of both *PTEN* alleles (Figure 4). While WGS data for *TP53* indicated copy numbers of 3–6 (Table S3A,B), the P53 protein was not detectable (Figure 4). Detailed analysis of the *TP53* gene locus revealed that multiple SNVs in the upstream and downstream regions of *TP53* gene were present, which likely disrupt the expression of P53. Additionally, the *RB1* gene was found to have two wild-type copies in OS HT87-P2 and HT87-P3, but protein expression levels were low relative to other PDX models, highlighting the fact that intact *RB1* alleles do not necessarily equate to higher *RB1* protein levels. In OS HT96-P1-P3, one allele of *RB1* was deleted based on CNV data; however, protein levels were still maintained (Figure 4). BET BRD4 protein was relatively stable across the OS PDX passages. Moreover, while *CDK4* protein levels were maintained across the OS PDX passages, *CDK6* levels varied. In the paired HT72 and HT77 OS PDXs, *CDK6* was quite high in HT72 but absent in the HT77 PDXs. Furthermore, *CDK6* levels in OS PDX HT87 and OS PDX HT96 decreased during the serial passaging, suggesting that *CDK6* is not necessary for growth and survival of the PDX. Differential expression of *CDK4* and *CDK6* was evident in the OS PDXs. Dai et al. previously demonstrated that *CDK4* and *CDK6* can have distinct biological functions that regulate cancer growth and metastasis [75]. Functional diversity of *CDK4* and *CDK6* have not been investigated in sarcomas and warrants further investigation. Notably, these disparities between linking CNVs and corresponding protein expression indicate the need to validate protein expression since the efficiency of sequencing reads, coupled with factors regulating transcription, translation, and post-translation, could lead to inconsistencies between CNV and protein levels [76,77].

3.6. Comparative Analysis of Single-Nucleotide Variations (SNVs), Somatic Variants, and Single-Base-Pair Substitution Variants in P0 Tumor Specimens versus Their Respective PDXs

Quantitative data with descriptive analysis [78] on the characterization of common genomic features (SNVs, somatic variants, and transitions/transversions) was conducted. In this analysis, genomic deviations are depicted in the P0 and the corresponding PDX passages. While some alterations were predicted to be pathogenic, many deviations were found to either not occur in cancer-associated genes or to occur in genes with unknown or not fully characterized functions.

Stability of small (SNVs, Indels) and large (insertions, deletions) structural variants, as well as somatic base substitutions for COSMIC genes, were overall preserved between the P0 tumor and the corresponding passaged PDX cohorts (Figure 5).

Scheme 10 base pairs of DNA [79]. At the total SNV level, all sarcoma PDXs exhibited a very strongly positive correlation between P0 and respective passaged PDXs and among the PDX passages for each individual OS model (PPCC scores of, 0.8–1 obtained HT72, HT77, and HT87 cohorts; PPCC score of 0.8–0.9 for HT96; Figure 6A). PPCC scores for the RMS HT74 PDX were strongly positive for SNVs (P1 versus P2, PPCC = 0.94; Figure 6B). However, P0 versus P1 or P2 had weak PPCC scores of 0.10 and 0.17, respectively; Figure 6B). For Wilms tumor PDXs (HT98, HT120, and HT139), very strongly positive correlation was observed for SNVs between each P0 and respective PDX passages for total CNVs (PPCC scores = 0.96–0.97 for HT98, PPCC scores of 0.92–0.94 for HT120, and PPCC scores of 0.94–0.95 for HT139; Figure 6C).

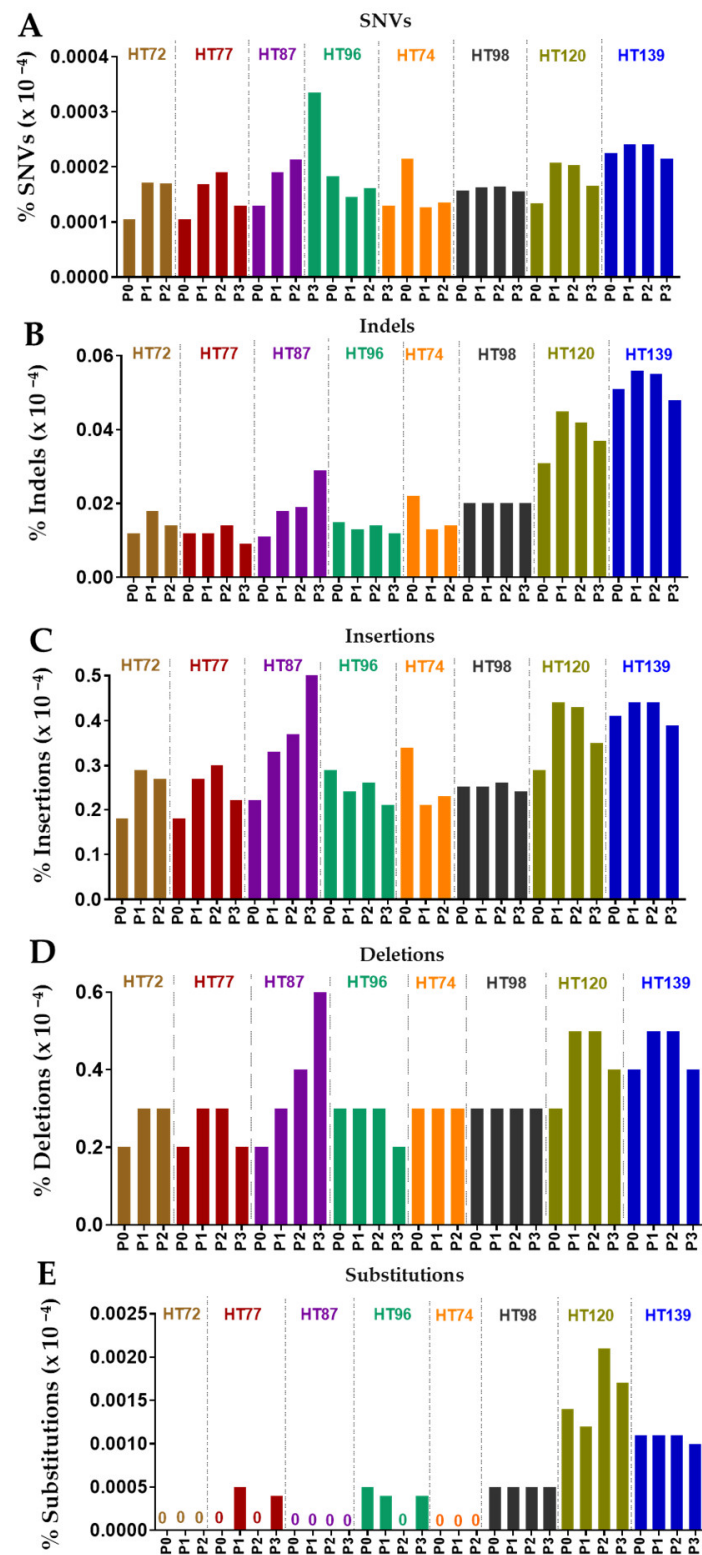


Figure 5. Total number of events identified in COSMIC genes as common variant classes such as (A) SNVs, (B) indels, (C) insertions, (D) deletions, and (E) substitutions. The data is normalized to total mappable variant counts specific to that particular sample. All data are normalized proportions to 10^{-4} , as indicated by y-axis. The variant frequency for each PDX was normalized to sequencing depth.

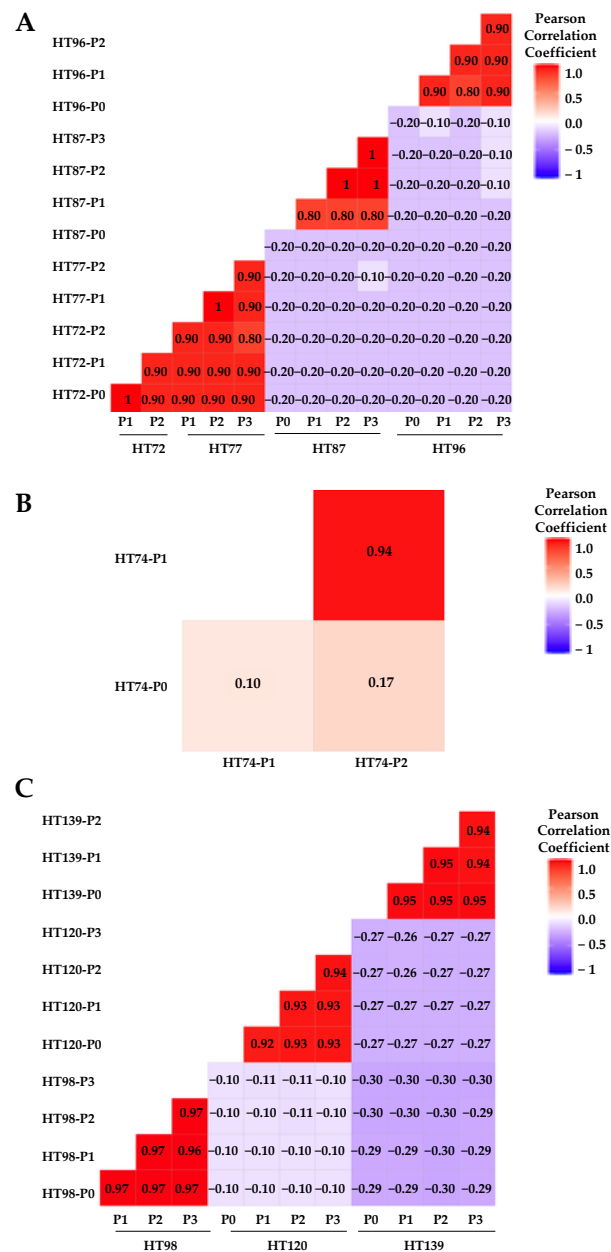


Figure 6. SNVs were preserved between P0 and their respective passaged PDXs and among the PDX of each tumor cohort. PPCC score was calculated for SNV profiles in all genes between original tumors and their corresponding PDX passages for (A) OS PDXs HT72, HT77, HT87, HT96, (B) RMS PDX HT74, and (C) PDXs for Wilms Tumors HT98, HT120, and HT139.

In P0 tumors and their respective passaged PDXs, most of the SNVs in P0 (Figure 7) were preserved in successive PDX passages. UpSet plots demonstrate intersections of SNVs that are the same between the P0 and its subsequent PDX passages. For example, in HT72, 87.4% of SNVs are common between P0, P1, and P2 (Figure 7A), whereas HT77 illustrates 88.9% common SNVs among P0, P1, P2, and P3 (Figure 7B). In HT87 and HT96, 78.4% (Figure 7C) and 76.8% (Figure 7D) SNVs, respectively, are commonly found between P0 and their subsequent passages. Similarly, RMS PDX HT74 also has about 76.9% SNVs occurring between the P0, P1, and P2 (Figure 7E). For Wilms tumors, SNVs that are commonly occurring between P0 and their respective passages are all > 90% (HT98 = 94.7%, HT120 = 91.9%, HT139 = 96.3%; Figure 7F,H). In addition, there were small numbers of SNVs that were only found in subsets of a particular cohort (Figure 7).

illustrates 88.9% common SNVs among P0, P1, P2, and P3 (Figure 7B). In HT187 and HT196, 78.4% (Figure 7C) and 76.8% (Figure 7D) SNVs, respectively, are commonly found between P0 and their subsequent passages. Similarly, RMS PDX HT74 also has about 76.9% SNVs occurring between the P0, P1, and P2 (Figure 7E). For Wilms tumors, SNVs that are commonly occurring between P0 and their respective passages are all > 90% (HT98 = 94.7%, HT120 = 91.9%, HT139 = 96.3%; Figure 7F,H). In addition, there were small numbers of SNVs that were only found in subsets of a particular cohort (Figure 7).

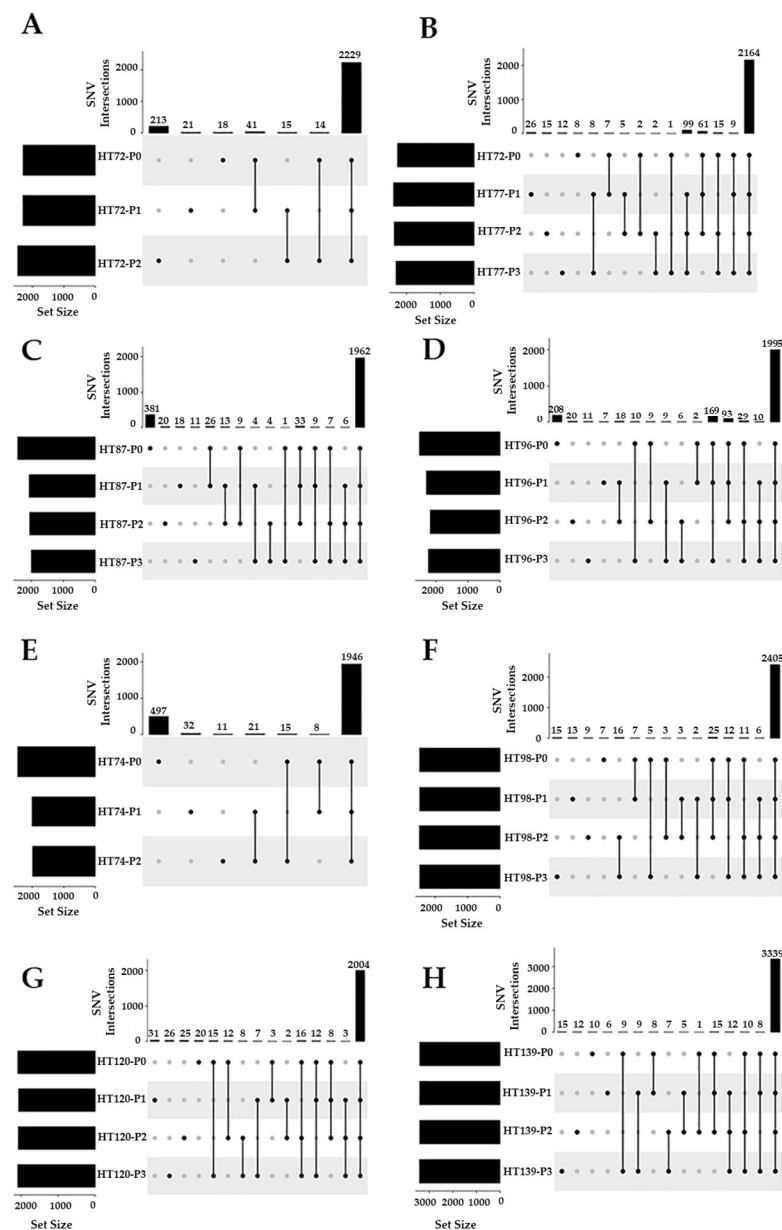


Figure 7: UpSet plots of SNVs indicating the intersection size of SNVs between P0 and their corresponding passages. The majority of the SNVs were retained following serial passaging. (A) HT72 (OS), (B) HT77 (OS), (C) HT87 (OS), (D) HT96 (OS), (E) HT74 (RMS), (F) HT98 (Wilms tumor), (G) HT120 (Wilms tumor), and (H) HT139 (Wilms tumor). The intersection size refers to the frequency of the combination depicted below the bar [80]. The set size corresponds to the frequency of associations [80] with P0 and its subsequent PDX passages.

3.7. Single-Base-Pair Substitutions Patterns across PDX Passages

This variant type includes nucleotide transitions and transversions [81]. Transitions involve the exchange of a two-ring purine with another two-ring purine [i.e., adenine (A) with guanine (G) and vice versa] or one-ring pyrimidine with another one-ring pyrimidine [thymine (T) with cytosine (C) and the reverse] [81]. Transversions occur when a two-ring purine is interchanged with a one-ring pyrimidine or vice-versa [81]. Generally, transitions are less likely to alter the amino acid sequences, but they occur at higher frequency than transversions [81,82]. Our data was consistent with previous studies in colorectal cancers [83].

Across the P0 tumors and their corresponding PDX cohorts, the number of transversion events (Figure 8A–H) was lower than the transition events (Figure 8I–L). Overall frequency of transversions and transitions in all genes remained stable in the serial PDX passages compared to its respective P0 tumor (Figure 8) [78]. In terms of maintaining the same transversions or transitions within the same genes, 52–88% of the transversion and 55–88% of the transitions were maintained in OS P0 tumor and its corresponding OS PDX passages (Table S3A,B). Up to 49% of transversions and 51% of transitions from P0 were preserved between RMS P0 and its respective passaged PDXs (Table S3A,B). The Wilms tumor had a maximum of 94% and 97% of transversions and transitions respectively that were conserved in P0 tumor and its subsequent PDX passages (Figure 8; Table S3A,B).

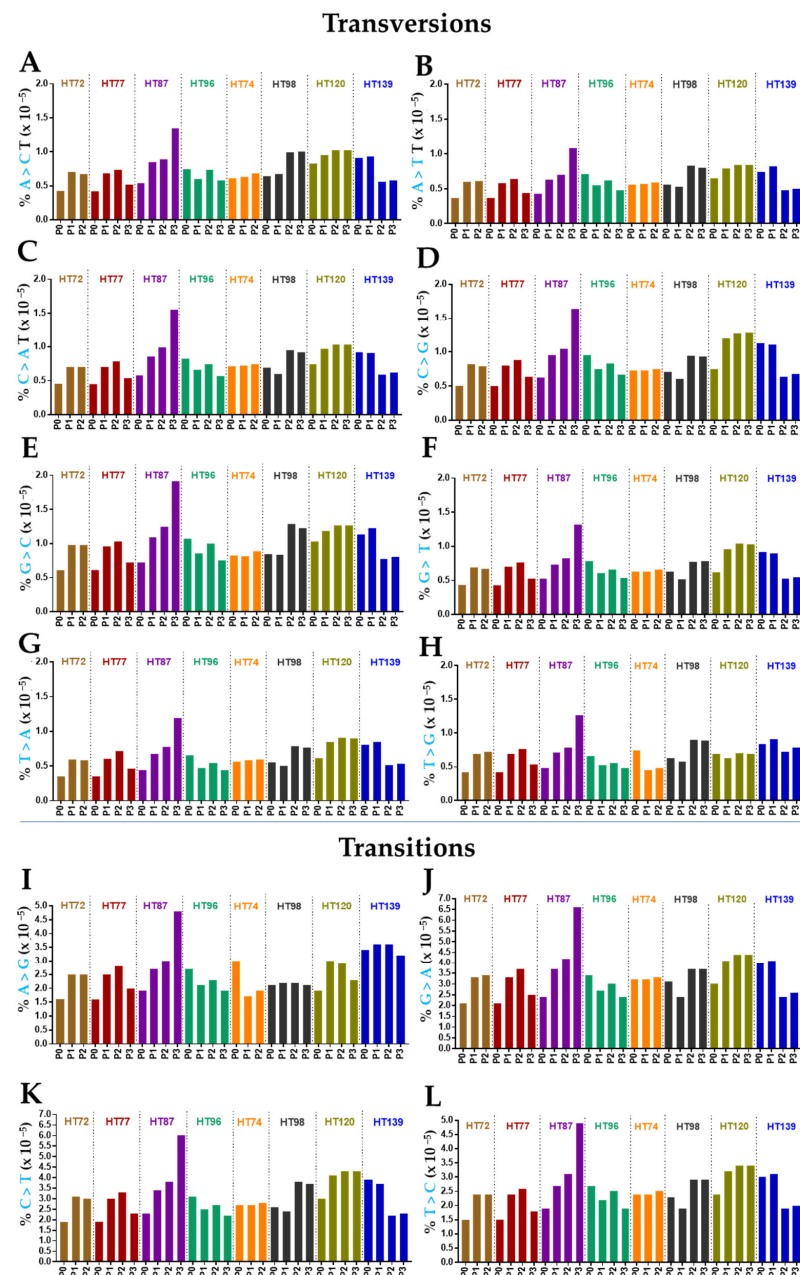


Figure 8. Percentage of transversion and transition events in all genes among P0 and their corresponding PDXs. Transition events: (A) A > C, (B) A > T, (C) C > A, (D) C > G, (E) G > C, (F) G > T, (G) T > A, (H) T > C. Transversion events: (A) A > C, (B) A > T, (C) C > A, (D) C > G, (E) G > C, (F) G > T, (G) T > A, and (H) T > C. The data is normalized against sequencing depth.

3.8. Potential Functional Impact of Somatic Variants in the P0 Tumor and Respective Passaged PDXs

Potential functional effects of the somatic variants were assessed between the P0 tumor and the corresponding PDX passages (Figure 9). Consequences of these variants were predicted based on the ensemble plus sequence ontology predictions and organized in order of severity in terms of predicted effect on transcription (Figure 9) [84]. Splice acceptor variants (Figure 9A), stop-gains (Figure 9B), start losses (Figure 9C), and inframe deletions (Figure 9D) rarely occurred in the P0–PDX cohorts. In contrast, more frequent occurrences of in-frame insertions (Figure 9E), frameshift variants (Figure 9F), and missense variants (Figure 9G) were found. Notably, upstream and downstream gene variants were by far the most common types of variant consequences observed in the PDX models (Figure 9H,I). In the sarcomas, 50–82% of the variants in the P0 were maintained in the corresponding PDXs (Table S4). In Wilms tumors, high fidelity of variants was observed in the P0 and corresponding PDX passages with 91–96% of the variants retained (Table S4). Moreover, approximately 50% of the frameshift variants in OS, 60% in RMS, and 53% in Wilms tumor were passed from P0 to their corresponding PDX passages (Figure S11). As for missense variants, a high degree of variability was observed among the P0–PDX cohorts with 21–84% retention frequency of these variants in the P0 and all the corresponding PDXs (Figure S12), with the lowest retention frequency in the OS HT96 PDX and the highest in the Wilms tumor HT98 PDX.

3.9. Functional Predictions of Somatic Variants from the P0 Tumor and the Respective Serially Passaged PDXs

Somatic variants from the P0 tumor and the respective passaged PDXs were filtered against the reference NA12878 dataset. Thus, only variants found within the original tumor and/or the respective passaged PDXs were retained and evaluated by sorting tolerant from intolerant (SIFT) algorithm and [polymorphism phenotyping (PolyPhen) algorithm to predict the functional outcome of these variants [38,85].

For the PDX datasets, genes with variants considered to be damaging or deleterious based on SIFT and/or Polyphen scores are presented (Figure 10; Table S5). Variant allele frequencies (VAFs) and oncoplots are presented for each variant when the frequency was 0.20 across in at least one of the samples (P0 tumor and/or the respective passages). While oncoplots of each P0 tumor and the respective passaged PDXs depict deleterious missense mutations in many genes, it is noteworthy that deleterious missense mutations in *MUC16* are present among all of the P0 tumors and their respective passaged PDXs (Figure 10A–H). A repeating peptide epitope of *MUC16*, a mucosal protein, known as cancer antigen 125 (CA125) is overexpressed in adult cancers, such as ovarian, lung, and pancreatic cancer, where it contributes to immunoprotection, metastasis, oncogenic signaling, and disease [86,87]. Missense variants in *MUC16* are one of the most common mutations observed in this gene [88,89]. However, two deleterious single nucleotide polymorphisms observed in all our PDXs cohorts include: rs76810971 and rs769311065 (Table S5), and further investigation needs to be conducted to elucidate the roles of these polymorphisms, especially in the context of pediatric and AYA OS, RMS, and Wilms tumors. Additionally, in the Wilms tumor PDX HT139 oncoplot (Figure 10H), there is a “start-loss” variant for *MYCL* where the normal “A” of start codon “AUG” is switched with a “G” so the proper translation is not initiated [90]. Thus, this predicts that no *MYCL* protein is generated in this Wilms tumor PDX due to translational cessation.

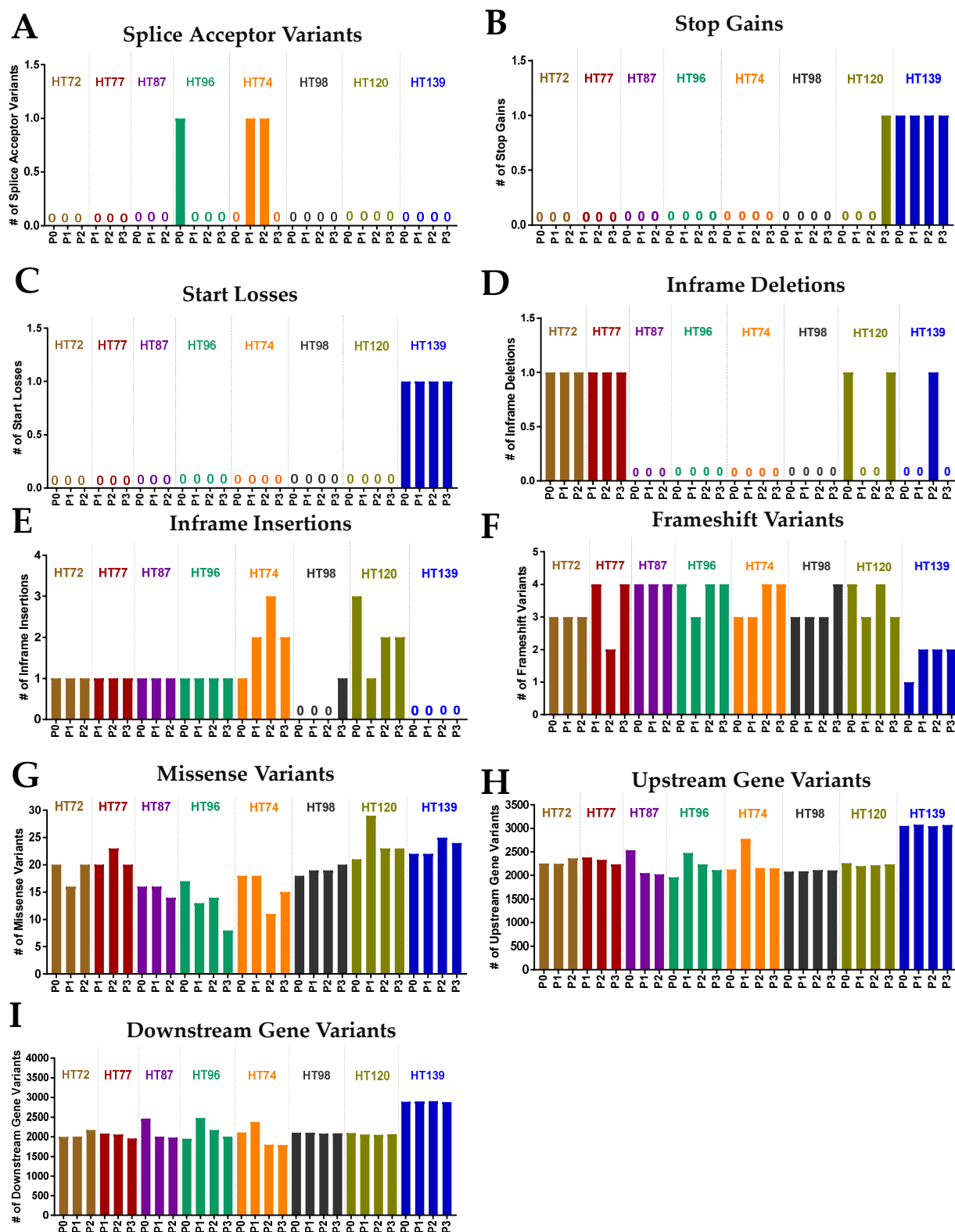


Figure 9. Number of variant consequences in COSMIC genes of P0 and their corresponding PDXs. (A) Splice acceptor variants, (B) stop gains, (C) start losses, (D) in-frame deletions, (E) in-frame insertions, (F) frameshift variants, (G) missense variants, (H) upstream gene variants, (I) downstream gene variants. The data were normalized against sequencing depth.

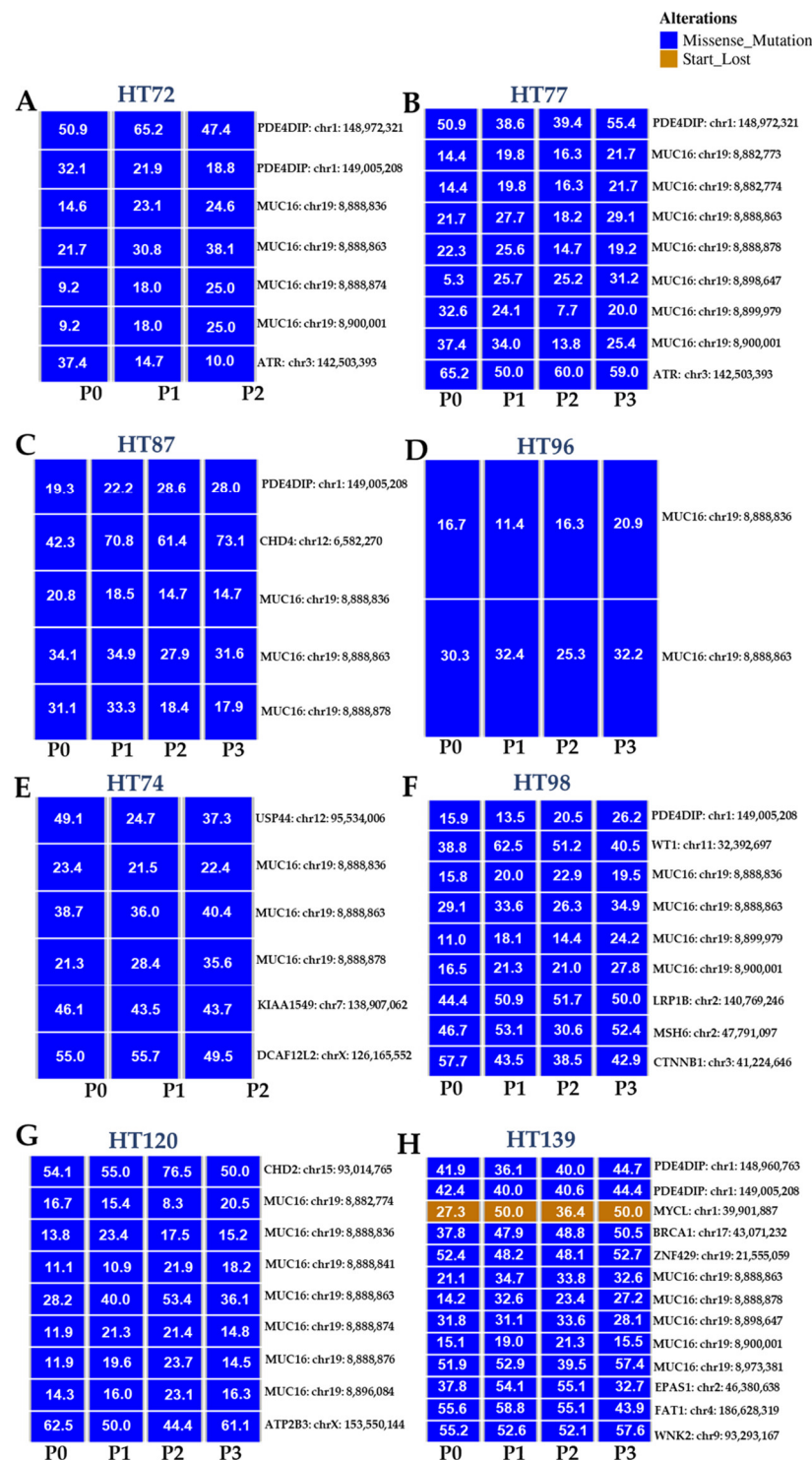


Figure 10. Identification of deleterious predictions of variant consequences in specific genes. SIFT and Polyphen algorithms were used to identify potential deleterious/damaging consequence of the variant on gene function. VAFs and oncoplots are presented for each variant when the frequency was ≥ 0.20 for at least one of the samples (P0 and/or its respective passaged PDXs). As previously mentioned, variants of high qualities (depth ≥ 10 , genotype quality ≥ 20 , allelic balance between 0.2–0.8) that are predicted to be of high impact and deleterious effect on functions (polyphen score > 0.15 and SIFT ≤ 0.05) were retained [41]. (A) HT72 (OS), (B) HT77 (OS), (C) HT87 (OS), (D) HT96 (OS), (E) HT74 (RMS), (F) HT98 (Wilms tumor), (G) HT120 (Wilms tumor), and (H) HT139 (Wilms tumor).

3.10. Transcriptome-Based Enrichment Pathway Analyses in Sarcoma PDXs

Comparison of the sarcoma P0 and the respective PDX-P2 RNA-seq datasets indicated positive correlation of global gene expression with HT72, HT77, and HT96 exhibiting high transcriptome correlation (Figure S13, $r = 0.87\text{--}0.96$). P0 and PDX-P2 RNA-seq datasets derived from HT87 and HT74 were positively correlated but to a lesser extent (Figure S13, $r = 0.7\text{--}0.78$).

Prioritized gene targets of interest from the RNA-seq in P0 and PDX-P2 sarcomas were interrogated using pathway databases (GO, KEGG, and Reactome). The top 15 pathways exhibiting high statistical significance are illustrated with Reactome enrichment bubble plots for HT72 (Figure 11A,B), HT77 (Figure 11C), HT87 (Figure 12), and HT96 (Figure 13). Herein, gene ratio (number of genes in that particular pathway vs. total number DEG genes) and p -values are presented for the P0 tumor, as well as PDX-P2. Notably, genes for all passaged PDXs and original tumor pairs demonstrated high degree of enrichment in the cell cycle and mitotic cell cycle pathways that are commonly dysregulated in these tumor types (Figures 11–13, p -values of $\leq 2 \times 10^{-5}$) [91–94]. Furthermore, irrespective of the type of pediatric and AYA solid tumor present, all original P0 tumors and passage 2 (P2) of the PDXs indicated that cell cycle pathways were highly statistically significant and also had the highest numbers of differentially expressed genes.

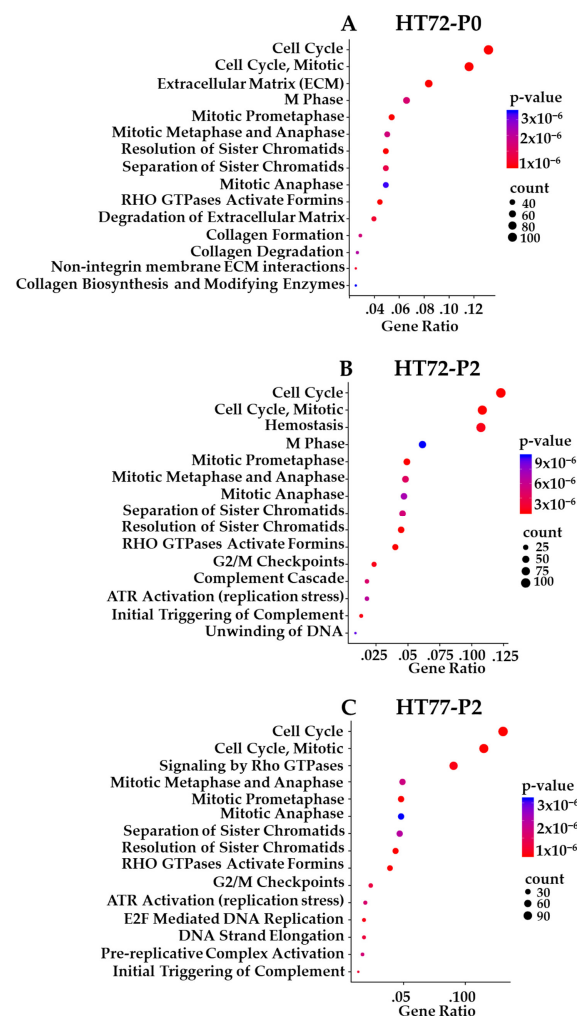


Figure 11. In (A) HT72-P0, (B) HT72-P2, and (C) HT77-P2, cell cycle pathway is identified as key player following pathway enrichment analyses of transcriptome. Genes that were statistically significant ($p < 0.05$) and >1.5 -fold change were selected for pathway enrichment analysis using GO, KEGG, and Reactome.

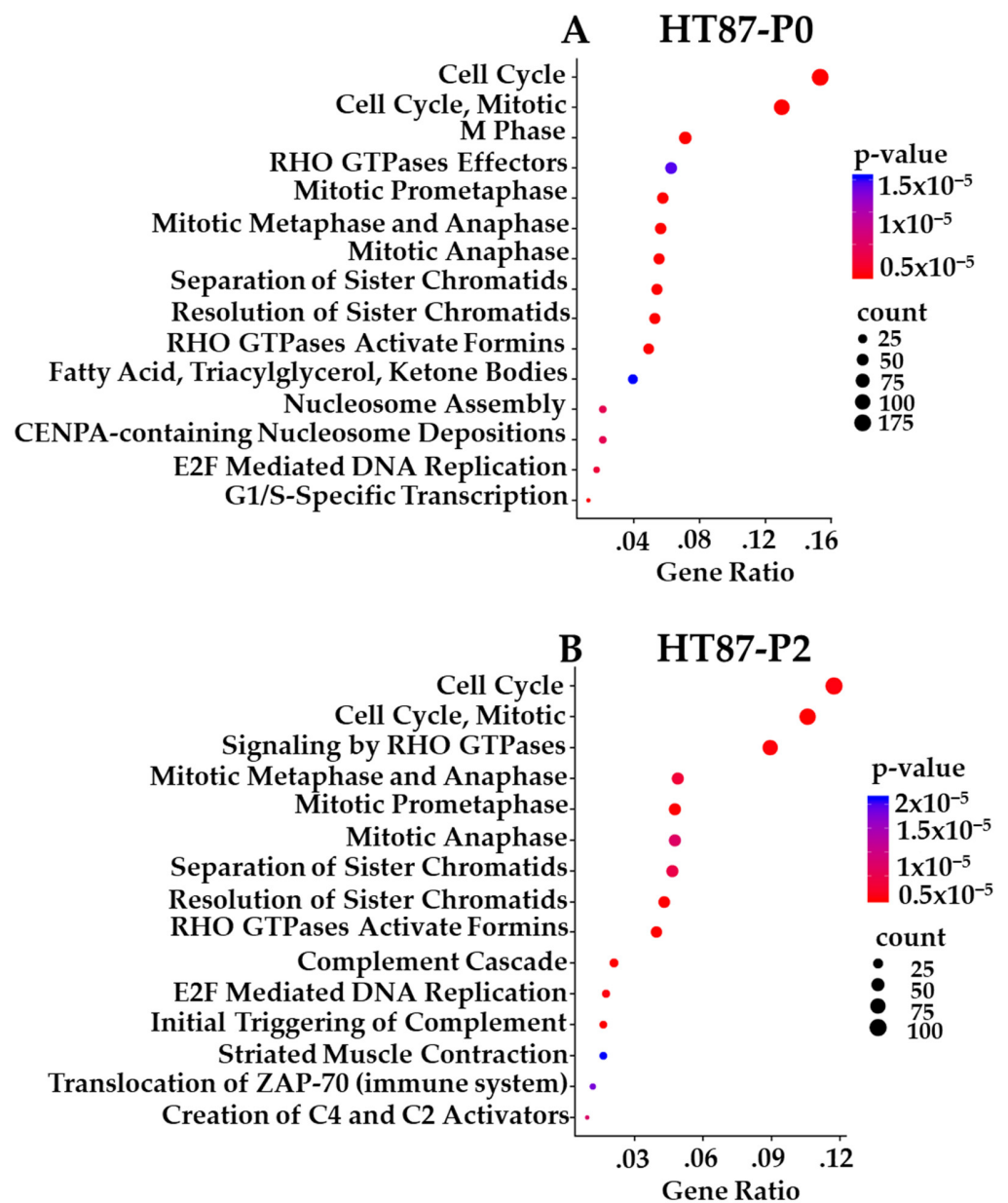


Figure 12. Pathway enrichment analyses of transcriptome for (A) HT87-P0 and (B) HT87-P2 illustrate cell cycle pathway as one of the most significant pathways of interest. Genes that were statistically significant ($p < 0.05$) and >1.5 -fold change were selected for pathway enrichment analysis using GO, KEGG, and Reactome.

Whether the transcript levels correlated with copy number of genes and/or protein expression from Figure 4's Western blot really depended on the gene interest and which PDX was being investigated (Table S6). For example, in HT72, HT77, and HT96, transcript levels and copy numbers for CCND3 were increased. Additionally, an increase in Cyclin D3 protein expression for HT72-P2, HT77-P2, and HT96-P2 was also evident (Figure 4). For CDK6, increased transcript levels correlated within increased protein expression in HT72-P2 and HT87-P2 (Figure 4, Table S6). However, for other genes, such as TP53, transcript levels did not necessarily correlate with copy numbers and/or protein expression. Such discrepancies may be attributed to post-transcriptional and/or post-translational modifications.

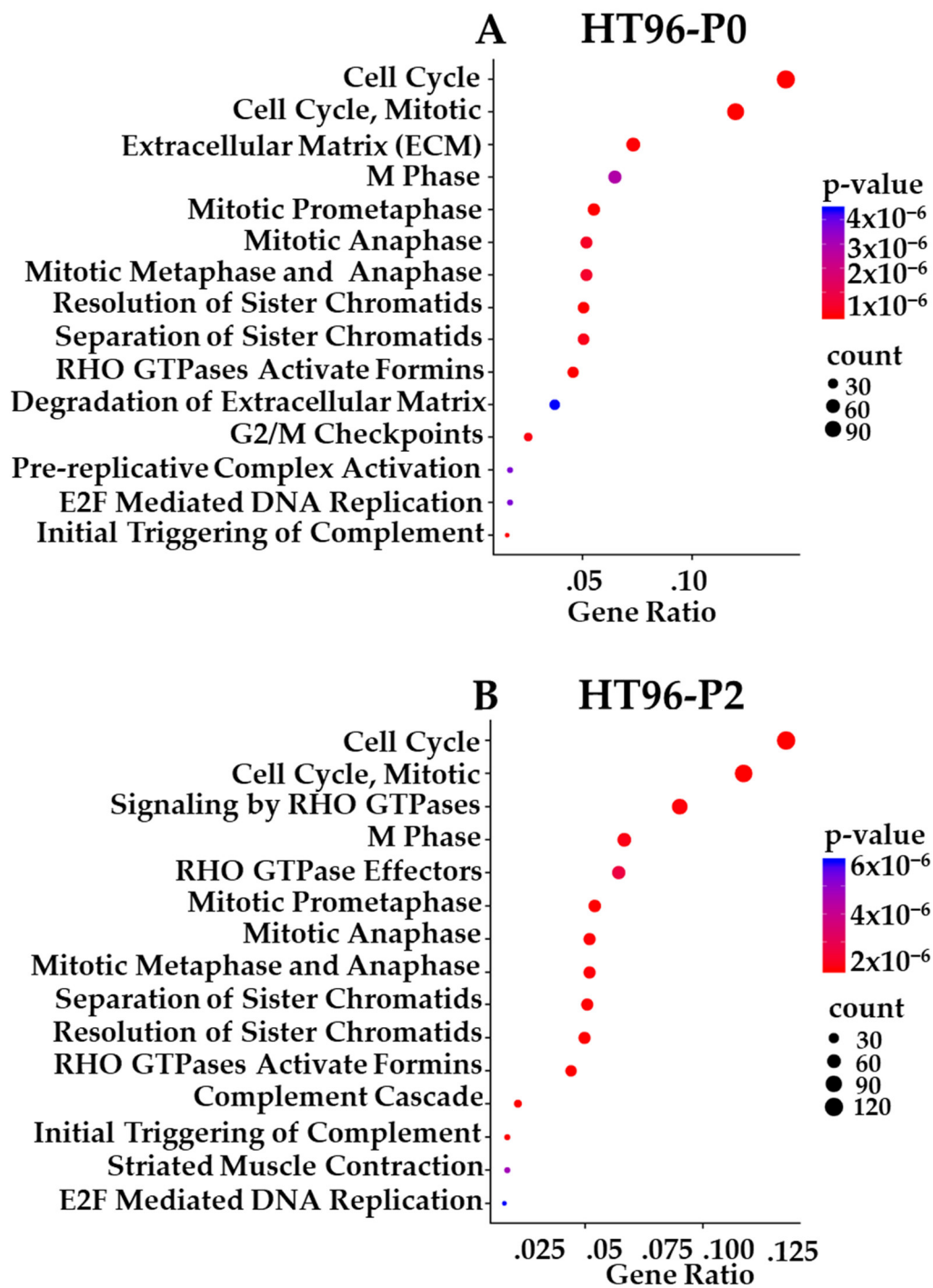


Figure 13. (A) HT96-P0 and (B) HT96-P2 pathway enrichment analysis highlights cell cycle pathway as a critical player for this PDX and the original tumor (P0). Genes that were statistically significant ($p < 0.05$) and >1.5 -fold change were selected for pathway enrichment analysis using GO, KEGG, and Reactome.

Reactome pathway enrichment data is presented as bubble plots for each primary tumor and subsequent PDX passage 2 of HT72-P0 (Figure 11A; OS), HT72-P2 (Figure 11B; OS), HT77-P2 (Figure 11C; OS), HT87-P0 (Figure 12A; OS), HT87-P2 (Figure 12B; OS), HT96-P0 (Figure 13A; OS), and HT96-P2 (Figure 13B; OS). RNA-seq was conducted on the original tumor specimen ($n = 1$) and PDX-P2 ($n = 3$ biological replicates) of each PDX. Human Adipose Tissue Poly A+ RNA was used as the filtering control for the RNA-seq

data. Differentially expressed genes (DEGs) with a >1 counts per million, fold change of >1.5, and p value of <0.05 were selected for subsequent pathway enrichment analysis via (GO, KEGG, and Reactome). Since all three databases showed overlapping pathway enrichment results, Reactome was selected as the representative database. The bigger the bubble plot the greater the number of genes that are affected in that specific pathway.

3.11. Comparative Analysis of Functional Protein/Phosphoprotein-Based Cell Signaling Activation Architecture and Pathway Enrichment in P0 Tumor Specimens vs. Their Respective PDX Passages

Different-OMIC datasets can be linked and used to cross-validate oncogenic pathway activation in cancer. The original P0 (when sufficient high-quality tissue was available) and subsequent PDX passages (P1–P3) were used for $\geq$ RPPA-based quantitative protein and phosphoprotein analysis and pathway activation mapping. RPPA was performed on 123 key proteins and phosphoproteins known to be “actionable” targets and/or associated with oncogenic pathways, with CV of <10% as the cut-off for positivity. Under these parameters, $n = 13$ total proteins and $n = 58$ phosphoproteins yielded positive signals of significance (Figure 14 and Figure S14) [95]. It should be noted that, in Figure 14 and Figure S14, while relative levels of proteins can be compared for each individual protein across a row, total/phosphoprotein levels cannot be compared between different proteins down each column. Profiling of select kinase activities via RPPA indicated significant activation of major pathways known to contribute to cancer survival, growth, and metastasis in the P0 and their passaged PDX. Of interest, Notch 1, a transmembrane receptor involved in growth and invasion was detected in all the P0 and respective PDX cohorts (OS and RMS Figure 14 and Wilms tumor, Figure S14) [96–98]. In contrast to the P0 tumor, Wilms tumor HT139 PDX passages display relatively decreased levels of the downstream effector for mTORC1, p70S6K, which is phosphorylated at serine 371 (S371) [99], as well as relatively increased cofilin phosphorylated at serine 3, which was previously demonstrated to contribute to cancer metastasis [100–103] and be associated with human glomerular disease [103]. OS HT77 PDX exhibited increased protein kinase C alpha subunit (PKC α) phosphorylated at T638 and protein kinase C beta II subunit (PKC β II) phosphorylated at T648 compared to P0. This signature is consistent with the invasive nature of the HT77 passaged PDXs; for PKC, it has been reported to promote metastasis in human and canine OS [104,105]. Using a stringent cut-off of CV ≤ 10 [52] as a positive signal for the RPPA data, multiple pathways contributing to autophagy, apoptosis, cell cycle, DNA damage, proliferation, survival, angiogenesis, invasion, migration, and cytoskeletal remodeling are activated, such as CDK4/6, DNA damage/oncogenic stress, MEK-ERK, and PI3K/AKT/mTOR.

We next integrated these data to develop a diagram depicting how pathway activation from the RPPA-based signaling analysis operative in both sarcoma and Wilms tumors can be used to construct biological networks that likely contribute to sarcoma pathogenesis (Figure 15). Many of the proteins are involved in multiple pathways, such as autophagy, apoptosis, cell cycle control, DNA damage, proliferation, survival, angiogenesis, invasion, migration, and cytoskeletal remodeling (Figure 15) [106–132]. Furthermore, Figure 15 integrates oncogenic-associated pathways identified by COSMIC-associated CNVs (Figure 3 and Table S2A–H) and/or proteome analyses. For example, in OS tumors COSMIC-associated CNV data indicated copy number gains/amplifications in *MYC*, *RAD21*, *Cyclin D3*, *Cyclin E1*, as well as loss of *CDKN2A* and *TP53* genes (Figures 3 and 4, Table S2A–H) that were then validated for corresponding proteins by Western blot (Figure 4) and/or RPPA (Figure 14).

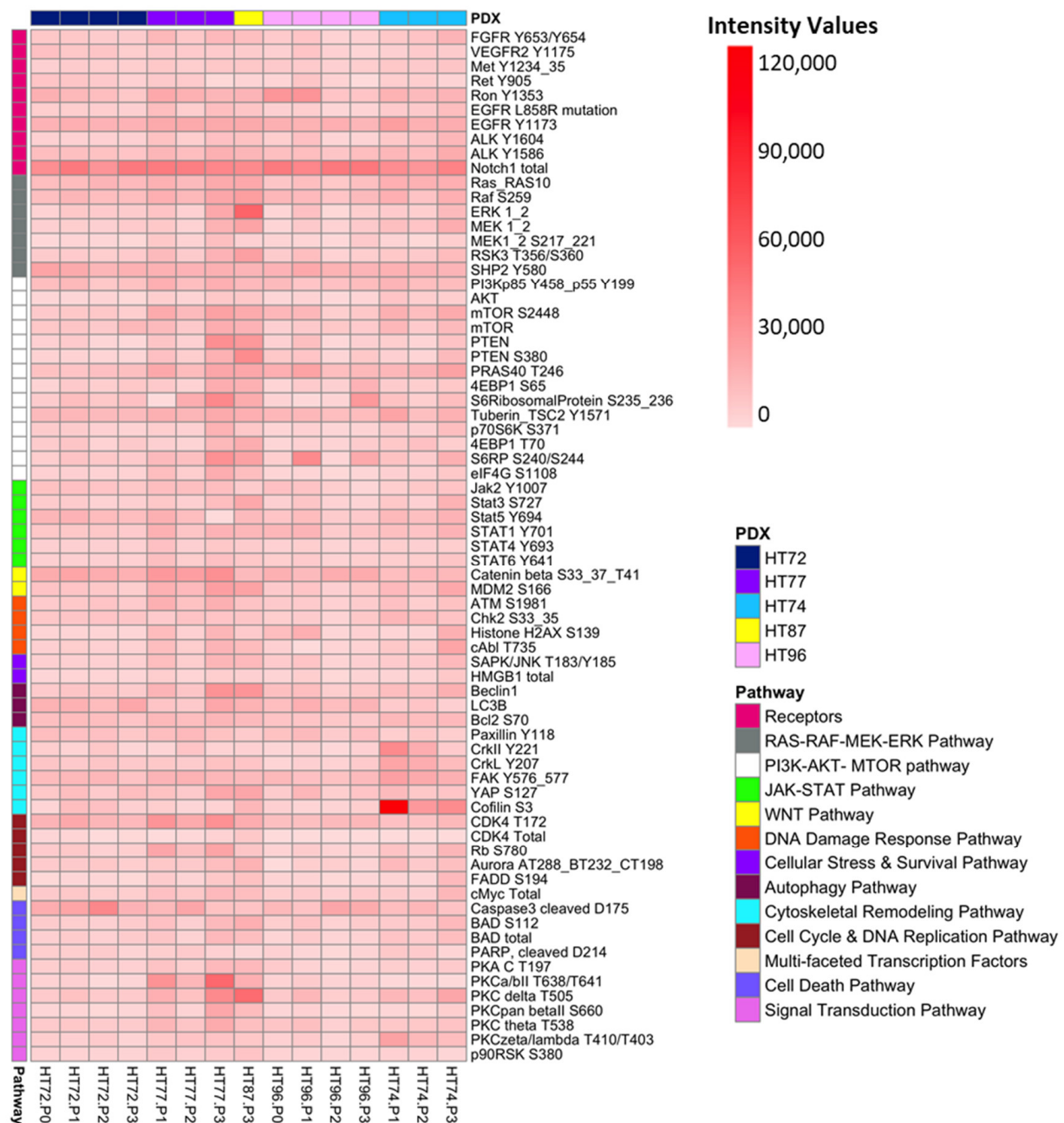
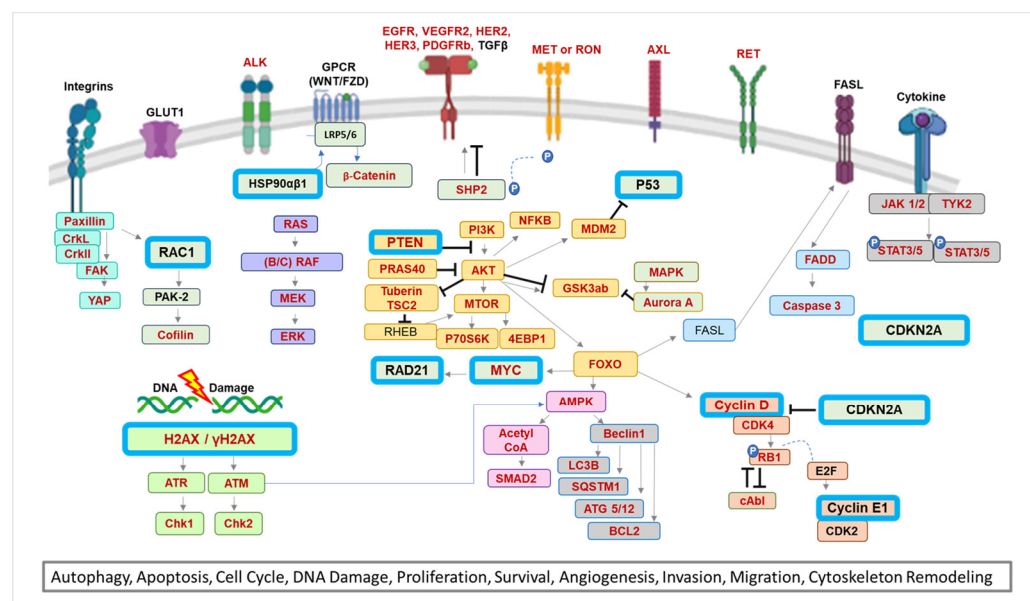


Figure 14. Integration of Western blot and RPPA data for OS. Expression of total and phosphoproteins, as evident by either RPPA or Western blot analysis (refer to figure key), were used to map out a network of biological pathways that may contribute to disease pathogenesis. Some of the selected proteins, such as MYC, PTEN, Cyclin D, and H2AX, were cross-validated by Western blot. For example, COSMIC-associated CNV data indicated copy number gains/amplifications in *MYC*, *RAD21*, *Cyclin D3*, *Cyclin E1*, as well as loss of *CDKN2A* and *P53* genes that were then validated for corresponding proteins by Western blot and/or RPPA.



KEY

Proteins in red text and without a blue box outline are validated only in RPPA data

Proteins in red text and outlined by blue box are validated by RPPA and Western blots

Proteins in black text and outlined by blue box are validated only by Western blot

Figure 15. Activated pathways are preserved among P0 and their respective PDX passages. Heatmap of total and phosphoproteins via RPPA. Sarcomas (HT72, HT77, HT87, and HT74) and Wilms tumor (HT98, HT120, and HT139). Each data point represents the average of 3 replicate printed spots. All final data values were normalized to total protein following the subtraction of negative control. Coefficient of variation (CV) among the replicate spots of <10% [23] was used as the cut-off for positive signals. Arrows indicate targets that are downstream in the pathway.

Based on the RRPA data, there is evidence that a number of actionable cancer-associated pathways are indeed activated in all of the PDX samples. However, there are relative differences in the activation level of specific pathways across the PDX panel (Table S7). Whether these subtle differences in pathway activation equate to differential therapeutic responses in each PDX warrants further investigation.

3.12. Proof-of-Concept In Vivo Studies in OS PDX: Monotherapy Screens to Explore Mechanisms of Tumor Growth Based on Therapeutic Response Biomarkers

By utilizing the multi-OMICS analytical pipeline described in this study, we compiled high-risk molecular signatures in the PDX derived from patients with aggressive disease based on our interrogation of the Drug Gene Interaction Database (DGIdb; <https://www.dgiddb.org/> (accessed on 7 July 2022)), as well as evidence from the literature. Attractive candidates for therapeutic targeting are summarized for HT72 (Table S8A), HT77 (Table S8B), HT87 (Table S8C), HT96 (Table S8D), HT74 (Table S8E), and HT120 (Table S8F) [5,65,133–154]. Wilms tumor patients from which primary localized tumors were used to develop HT98 and HT139 PDXs responded well to frontline treatment (Table S1). Therefore, linking molecular signatures to therapeutic response biomarkers and drug candidates in these tumors was not prioritized. Based on our multi-OMICS pipeline, CDK4/6 hyperactivation and BETs were prioritized as targets of interest for in vivo screening since both signatures are present in OS PDX derived from patients with aggressive and metastatic disease. CDK4/6 plays an integral role in OS growth, survival, and invasion [5,65]. Targeting CDK4/6 was one priority, since these proteins function as major regulators of the cell cycle pathway, which is activated based on CNVs analysis (Figure 2). This includes CCND3 amplification, lack of CDKN2A/p16^{INK4A}, and presence of RB1+ where 1–2 wild-type alleles

are present and the expression of RB1 (Figure 4). This is further supported by the fact that activation of cell cycle pathways is evident in the transcriptome-based enrichment pathway analyses (Figures 11–13), as well as the protein pathway enrichment analysis (Figure 14). For example, in PDX HT77 as well as other OS PDXs, the RPPA data show increased total plus phosphorylated CDK4 (T172), as well as increased expression of phosphorylated RB (S780), which serve as therapeutic response biomarkers and predict sensitivity to CDK4/6 inhibitors such as palbociclib [65]. Additionally, lack of *CDKN2A/p16^{INK4A}* and overexpression of other components in this pathway, such as *CCND3* (Cyclin D3) and RB1 (Figure 4), also predict sensitivity to CDK4/6 inhibitors as a means to block cell cycle entry [65]. PDX HT77 was derived from a metastatic lesion in a heavily pretreated 18-year-old OS patient (Figure 16A, Table S1) that was established prior to treatment with palbociclib. While the patient was treated late in the disease course with palbociclib, he eventually succumbed to the disease. In PDX HT77, palbociclib was well tolerated based on body weights (Figure S15A) and significantly blocked tumor growth compared to vehicle as long as the mice remained in treatment (Figure 16B). In this study, we were not only able to validate growth dependency on CDK4/6 but also provided evidence that the PDX HT77 was indeed responsive to palbociclib at a point in the disease history when the PDX was established.

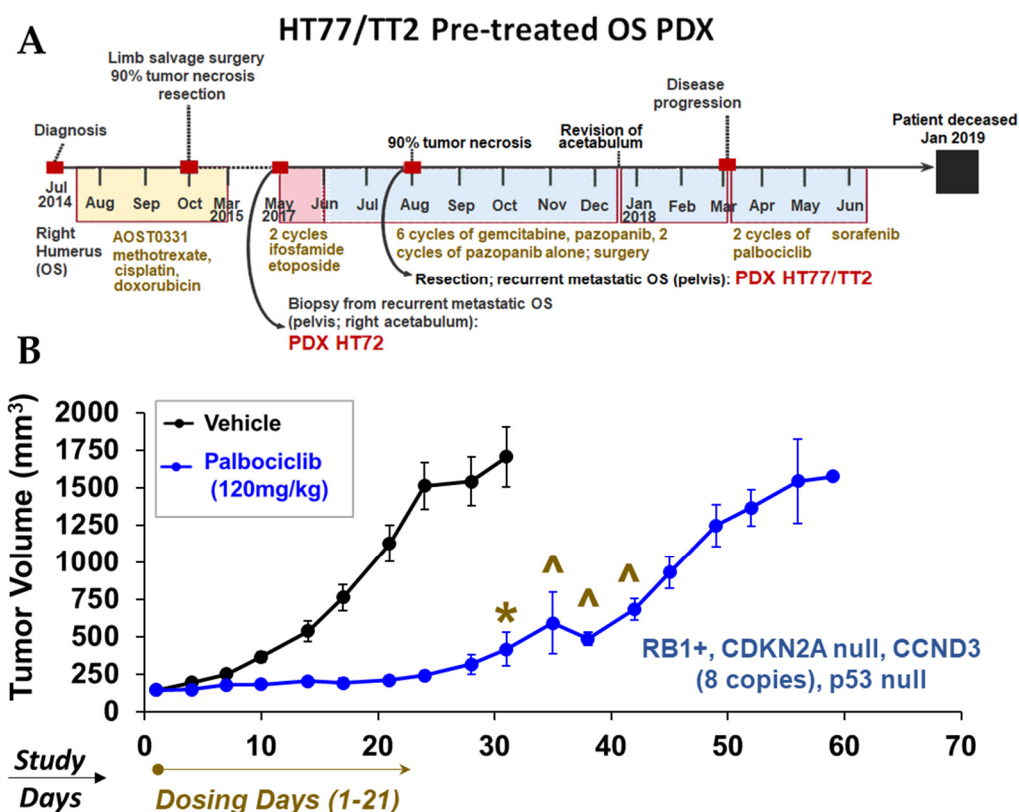


Figure 16. CDK4/6 inhibitor, palbociclib, significantly blocks growth of PDX HT77. (A) HT77 PDX was derived from a metastatic lesion of a pretreated 18-year-old male OS patient harboring molecular signatures indicative of CDK4/6 activation. (B) Mice were treated with vehicle ($n = 8$) of palbociclib ($n = 7$) at 120 mg/kg for 5 consecutive days PO, SID, daily for 3 weeks. All tumor volumes are represented as mean $\pm$ standard error mean (SEM). Two-way ANOVA with Holm–Sidak multiple comparison test; * $p < 0.05$, ^ $p < 0.0001$. Note: For HT77, statistical analyses were conducted up to the point that all mice were still in the study (day 41).

Targeting BET proteins was also prioritized, since BET-BRD4 is a therapeutic target of emerging interest in sarcomas. BETs contribute not only to transcriptional regulation by binding to acetylated lysines, but also, in the appropriate molecular contexts, can lead

to an imbalance in transcription–replication kinetics and induce non-tolerated levels of replication stress in cancer [155]. Studies by us demonstrated that BET inhibition can lead to increased replication stress and increased death in models of OS treated with CHK1 inhibitors [5]. Moreover, MYC, a common downstream target of BETs, is sensitive to BET inhibitors in some cellular contexts [5,156,157]. As previously mentioned, OS PDX HT96 is derived from a treatment-naïve 9-year-old OS patient that ultimately succumbed to metastatic disease (Figure 17A; Table S1). This model exhibits replication stress (i.e., BRD4 and γ H2AX) and expresses MYC (Figure 4). Treatment with BET inhibitor, OTX015, lead to a transient decrease in OS PDX96 growth compared to vehicle control (Figure 17) and was well tolerated, as indicated by stable body weights over time (Figure S15). As mentioned above, molecular signature tables of the PDXs derived from patients with aggressive disease are provided in Table S8A–F. Data obtained from our multi-OMICS pipeline and pathway integration provide numerous biomarkers of therapeutic response and drug candidates of actionable pathways that could be used for development and testing of other targeted therapies, which will be the focus of future investigations.

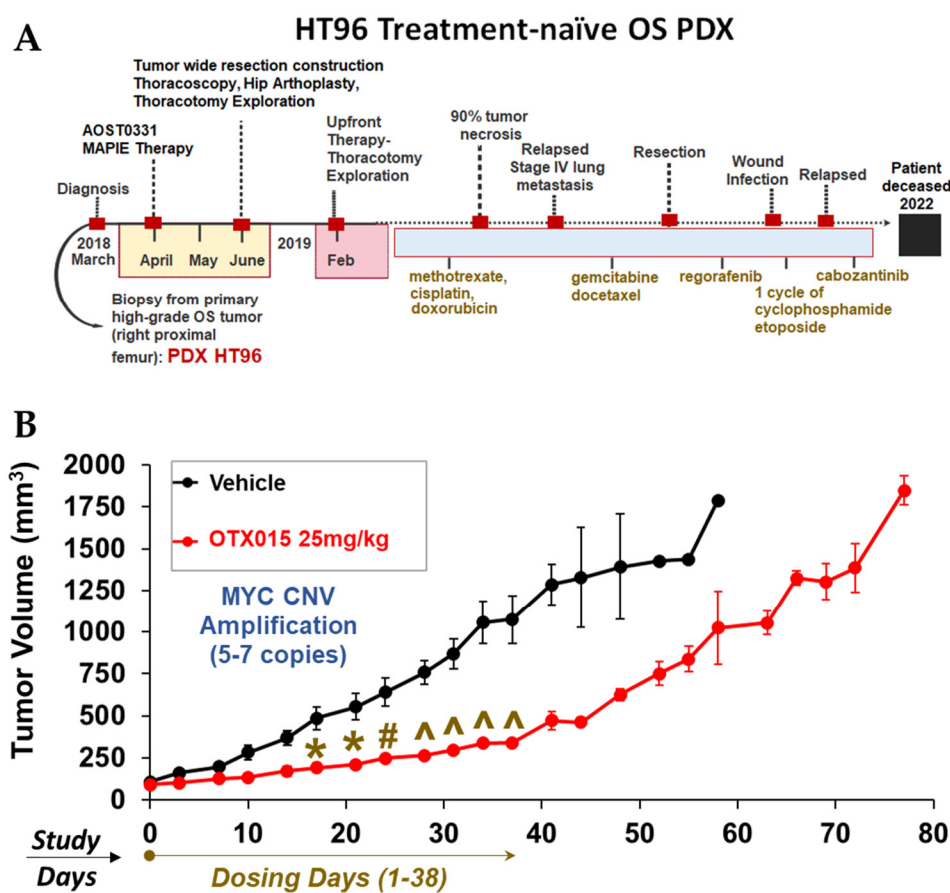


Figure 17. BET inhibitor, OTX015, decreases growth of PDX HT96. **(A)** HT96 PDX was derived from a treatment-naïve biopsy sample from a 9-year-old male OS patient. **(B)** HT96 PDX mice were treated for ~6 weeks with 25 mg/kg OTX015 ($n = 5$) or vehicle control ($n = 6$), BID, PO, daily for 5 consecutive days with a 2-day dosing holiday. All tumor volumes are represented as mean $\pm$ standard error mean (SEM). Two-way ANOVA with Holm–Sidak multiple comparison test; * $p < 0.05$, # $p < 0.001$, ^ $p < 0.0001$. Note: For HT96 study, statistical analyses were conducted up to the point that all mice were still in the study (day 37).

4. Discussion

Precision genomics has emerged as a critical component of clinical decision making in treatment of patients with aggressive disease. Identifying the optimal targeted therapy

for each patient based on the molecular signature of the tumor has the potential to change long-term treatment paradigms in hard-to-treat cancers. The field now needs to use relevant preclinical oncology models that are linked to the patient's disease journey in a de-identified manner to validate targeted therapy such that earlier intervention with efficacious and safe molecularly guided therapy can be realized.

Therefore, there is great impetus to design feasible, cost-effective, and reliable approaches to uncover and cross-validate therapeutic targets as well as develop preclinical models that recapitulate the original tumor. Our objective for this study was two-fold. First, we sought to develop an integrated multi-OMICS approach for identifying molecular signatures (DNA, RNA, protein, and/or phosphoprotein) in PDX models developed at our institution in which disease history prior to and after tumor collection is linked in a de-identified manner to PDX datasets. In this study, PDX models derived from treatment naïve and pre-treated solid-tumor patients were evaluated. It is important to note that one limitation in our study is that, by chance, all of our PDXs were developed from Caucasian patients (Table 1) during the initial stage of program development at our institution. It has been reported that racial and ethnic diversities can have an impact on tumor mutational burden, as well as on response to immune-based therapies [158,159]. Furthermore, tumor mutational burdens for races, such as African American patients, have been reported to be inflated when publicly available germline variant databases are used for filtering non-somatic polymorphisms, demonstrating the need to use patient-specific germlines [159]. Second, to elucidate if genome instability impacted the molecular stability of original tumors following engraftment and serial propagation, PDXs derived from tumors with inherent genome instability (OS, RMS) and relatively more stable genomes (Wilms tumors) were selected for deep molecular characterization.

In depth surveillance and cross-validation of -OMICS data provides insights in gaining a holistic view of the molecular landscape of the cancer. Furthermore, analysis of pathway activation at the protein expression and protein activation/phosphorylation levels provides insights into prioritization of therapeutic targets [160]. RPPA-based pathway activation mapping is one such approach by which an unbiased multiplexed and comprehensive array of targeted total and phosphoproteins can be evaluated in a limited amount of sample [28,161]. This multiplex analytical method can allow more samples to be tested at one time, which has proven to be more cost-effective compared to other molecular proteome platforms. RPPA-based analysis of individual tumor samples could be a major advantage in the clinical setting, for it has the potential to provide a quantitative read-out of the expression and functional activation state of hundreds of protein drug targets that verify target and pathway activation using slides obtained from paraffin-embedded tissue blocks [28,161]. In our study, total and phosphoprotein data from RPPA analysis was used to highlight activation of a number of key signaling pathways involved in these diseases (Figure 14).

Generation of novel clinically-annotated PDXs provided the opportunity to have sufficient tissue to validate CNV effects on protein expression (Figure 4). Notably, based on the RPPA data, expression of total PTEN and MYC (c-MYC) was the highest in HT77-P3 and HT87-P3 (Figure 14), although Figure 4 illustrates that PTEN and MYC levels were detected in most OS PDXs by Western blot. As previously mentioned, *PTEN* in HT72 is reported to have zero copies in both passages (Table S2A). However, PTEN protein was still present in HT72-P1 but absent in HT72-P2, which correlates with the lack of both *PTEN* alleles (Figure 4). It is important to note that, in our RPPA data (Figure 14), while the presence of total PTEN in its inactive form based on its phosphorylation at S380 [162] is evident for HT72 P0-P3, these levels are relatively low (Figure 14). These variations may lie in the differences between the Western blot and RPPA detection techniques, such as the sample processing procedures, amount of input sample, number of freeze/thaw cycles, or antibodies for RPPA vs. Western blots recognizing different epitopes [163–165], rather than inter-slide variability, as the criteria positivity was stringent with a cut-off of $CV \leq 10$ [52].

Our study highlights the value of integrating proteomic/phosphoprotein analysis as a multi-OMICS approach in conjunction with assessment of DNA and RNA alterations through PDX passages. While DNA and RNA can provide insight into impactful mutations and/or gene products involved in oncogenic pathway dysregulation, they do not always correlate with protein expression or function due to post-transcriptional and/or post-translational modifications, such as phosphorylation [166]. Our results show that, in general, serial propagation of these pediatric and AYA solid tumor PDXs, a congruence with the original tumors in terms of their histological, molecular, and proteomic characteristics was preserved, particularly in the detailed analyses of functionally impactful DNA sequences.

Our data indicate that eight out ten of our PDX models maintain the biological fidelity similar to that of the original patient tumors. In the two models [1 OS (HT87) and 1 RMS (HT74)] wherein the PDX passages show more discordance when compared to their respective original tumor specimens, it is important to note that many molecular signatures are still maintained, as is evident from CNV, SNV, RPPA, and Western blot data. The discordance at the DNA level observed in some of the P0–PDX cohorts may or may not be of significance in terms of protein function and pathway activation. However, further investigations are needed to determine if genetic alterations in cancer-associated and non-cancer-associated genes have deleterious/damaging impact on their respective protein function and if that contributes to disease pathogenesis in ways not understood at this point in time. There are many caveats to be considered when characterizing a PDX and determining its suitability for efficacy and safety testing from the standpoint of similarities and dissimilarities to the original P0 tumor. As previously mentioned in the introduction, a number of factors can contribute to the differences observed between the pairs of P0 tumor and passaged PDXs, such as clonal evolution, genetic drift, intra- and inter-tumoral heterogeneity, presence of human stroma and immune system in original tumors versus increased murine stromal content in passaged PDXs, and insufficient amount of the original tumor sequenced, as well as lineage splitting due to tumors' spatial heterogeneity during PDX propagation [16–19]. Additionally, it should also be noted that it is impossible to know if clonal drift or evolution that appears in some PDXs over time might also naturally occur in that same patient at some point in time over the course of their disease.

Other challenges also exist that still hinder the use of PDXs in precision medicine [167], such as their lack of functional immune system and absence of a human tumor microenvironment [168]. While serial passaging of PDXs results in acceptable preservation of the human patient's tumor cells, the primary tumor-associated stroma is replaced by murine stroma [57]. The ratio of tumor cells to stroma cells varies between individual tumors regardless of the cancer type [57]; thus, if a particular tumor, such as HT87, has more murine stromal components, this could indicate that the stromal compartments are critical for growth of this tumor. This may also provide insight into possible therapeutic interventions, as it may suggest that targeting the tumor microenvironment may be more beneficial in terms of decreasing growth for this particular tumor. Despite these challenges, PDXs provide a highly valuable preclinical tool for drug screens in pediatric cancers. These models can also be linked to clinically relevant information regarding the original tumor, as we have done here, which provides additional insights into the disease course and clinical outcomes [18] and helps us prioritize targets for therapeutic interventions.

As such, we utilized the integrated multi-OMICS approach to select targets of interest, such as CDK4/6 or BETs, to validate *in vivo* mechanisms of disease progression in OS PDX models (Figures 16 and 17). Notably, clinical history for patient HT77 indicated that the CDK4/6 inhibitor palbociclib was given to the patient, albeit at a much later phase in the disease's journey of progression (Figure 16A), such that the therapy may not have been able to compete against the increased tumor burden. However, when palbociclib was given at an earlier stage of the disease, as indicated by the HT77 PDX *in vivo* study (Figure 16B), tumor growth decreased. Studies are in progress to identify additional therapeutic vulnerabilities and explore combination targeted therapies in these models.

5. Conclusions

In this study, we established an integrated clinical- and research-based platform where molecular data can be obtained on pediatric and AYA patients with pediatric solid tumors to facilitate identification, prioritization, and validation of genomically guided treatments that can benefit patients. Oncogenic heterogeneity, as well as non-standardized methodologies to conduct -OMICS analyses on pediatric and AYA solid tumor PDXs, contribute to the significant lag in improving therapeutic and prognostic outcomes, especially in the relapsed setting [26,169]. Thus, novel clinically annotated, serially passaged PDX models that recapitulate the molecular landscape of the original tumor sample and can be cross-validated will help improve decision making of molecularly guided therapy. Linking patient treatment history with PDX data offers an exciting and underexplored opportunity to not only test new combination therapies based on -OMICS data but to also gain a better understanding of what therapy may be most beneficial to a patient who has already had therapy. Comprehensive identification and prioritization of actionable targets and signaling pathways using integrated multi-OMICS helps identify and cross-validate relevant therapeutic targets in aggressive pediatric and AYA solid tumors that could have been missed from evaluation of only a single layer of the tumor molecular landscape. Efficacy data obtained using these types of well-curated PDX models can provide rationale for moving targeted therapies to future clinical trials earlier in the disease course and at a time before development of chronic adverse events and disease progression, a stage that is nearly impossible to treat [170]. A major benefit of such study is to expand pediatric and AYA PDX panels for future patients and not necessarily for the patients from which the PDX was derived. The field in pediatric and AYA solid tumors is at the point where scientists are generating efficacy and safety data to provide rationale for earlier intervention in future patients. Depending on the clinical situation, tumors are sequenced either upfront or following resection. In the future, we envision the molecular signature and the clinical history of a new patient will be compared and aligned to similar datasets of previous patient(s) from which PDX(s) were also generated. While efficacy data from a PDX would not be the defining component of a clinical decision, it could provide supportive data to help guide decision making for the newly diagnosed or relapsed patient. As such, studies like this continue to be critical to identifying high-yield targets and focusing clinical trial efforts on innovative therapeutics most likely to benefit these underserved pediatric populations. With that said, however, as more PDX models are developed and data is linked to the corresponding patient's clinical history outcomes, these models may eventually become a more prominent and accepted component of clinical decision making. As such, studies like this continue to be critical to identifying high-yield targets and focusing clinical trial efforts on innovative therapeutics most likely to benefit these underserved pediatric populations.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/cancers15010259/s1>. Table S1: Comprehensive clinical information on pediatric and AYA patients donating solid tumors for PDXs, Figure S1: Histological characteristics are preserved in original tumors and their respective PDX passages. Tumor tissues from OS PDXs (HT72, HT77, HT87, and HT96), RMS PDXs (HT74), and Wilms tumor PDXs (HT98, HT120, and HT139) were harvested, formalin fixed, pro-cessed and stained with H&E. The H&E stains are representative of 3 mice per group. All images are at 20× magnification, Figure S2: Percentage of mouse reads filtered out from the established PDX passages. Highly conserved regions between mouse and human genome may result in false positive variants due to presence of mouse stromal tissue. Thus, to overcome this challenge any se-quencing reads that have been mapped to the mouse genome have been removed during WGS of each PDX passage for accurate depiction of the molecular profile. Percentage of mouse reads that were present and removed from each PDX passage are illustrated here. P0 represents the original human patient sample and will have 0% of mouse reads since it has not been introduced in the murine host, Figure S3: HT72 chromplots compare CNV distribution across genome of A P0 tumor and B-C corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0; gray = background noise, Figure S4: HT77 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. HT72 and

HT77 are from the same patient, thus P0 tumor of HT72 was used for comparison. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S5: HT87 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S6: HT96 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S7: HT74 chromplots compare CNV distribution across genome of A parental tumor and B-C corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S8: HT98 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S9: HT120 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Figure S10: HT139 chromplots compare CNV distribution across genome of A P0 tumor and B-D corresponding PDXs. Orange = statistically significant amplification if >0 or deletion if <0 ; gray = background noise, Table S2A: Absolute copy numbers identified in COSMIC genes in HT72 cohort, Table S2B: Absolute copy numbers identified in COSMIC genes in HT77 cohort, Table S2C: Absolute copy numbers identified in COSMIC genes in HT87 cohort, Table S2D: Absolute copy numbers identified in COSMIC genes in HT96 cohort, Table S2E: Absolute copy numbers identified in COSMIC genes in HT74 cohort, Table S2F: Absolute copy numbers identified in COSMIC genes in HT98 cohort, Table S2G: Absolute copy numbers identified in COSMIC genes in HT120 cohort, Table S2H: Absolute copy numbers identified in COSMIC genes in HT139 cohort, Table S3A-B: Descriptive analysis of transversions and transitions in each P0-PDX cohort, Table S4: Descriptive analysis of total variants in P0-PDX subset, Figure S11: UpSet plots of frameshift mutations occurring on the same genomic location between paired PDX passages and their P0. A HT72 (OS), B HT77 (OS), C HT87 (OS), D HT96 (OS), E HT74 (RMS), F HT98 (Wilms tumor), G HT120 (Wilms tumor), and H HT139 (Wilms tumor), Figure S12: UpSet plots of missense variants occurring on the same genomic location between paired PDX passages and their P0. A HT72 (OS), B HT77 (OS), C HT87 (OS), D HT96 (OS), E HT74 (RMS), F HT98 (Wilms tumor), G HT120 (Wilms tumor), and H HT139 (Wilms tumor), Table S5A: HT72 oncoplot annotation part 1, Table S5B: HT72 oncoplot annotation part 2, Table S5C: HT77 oncoplot annotation part 1, Table S5D: HT77 oncoplot annotation part 2, Table S5E: HT87 oncoplot annotation part 1, Table S5F: HT87 oncoplot annotation part 2, Table S5G: HT96 oncoplot annotation part 1, Table S5H: HT96 oncoplot annotation part 2, Table S5I: HT74 oncoplot annotation part 1, Table S5J: HT74 oncoplot annotation part 2, Table S5K: HT98 oncoplot annotation part 1, Table S5L: HT98 oncoplot annotation part 2, Table S5M: HT120 oncoplot annotation part 1, Table S5N: HT120 oncoplot annotation part 2, Table S5O: HT139 oncoplot annotation part 1, Table S5P: HT139 oncoplot annotation part 2, Figure S13: Gene expression profiles are maintained between P0 and its respective PDXs. This is evident by RNA-seq correlation plots for A HT72 (OS), B HT77 (OS), C HT87 (OS), D HT96 (OS), and E HT74 (RMS). The closer the correlation coefficient (R) is to 1 the more positive linear relationship exists between groups being compared [165], Table S6: Integration of transcript levels to copy numbers for prioritized targets, Figure S14: Activated pathways in Wilms tumors are preserved among P0 and their respective PDX passages. Heatmap of total and phosphoproteins via RPPA in Wilms tumor (HT98, HT120, and HT139) is presented. Each data point represents the average of 3 replicate printed spots. All final data values were normalized to total protein following the subtraction of negative control. Coefficient of variation (CV) among the replicate spots of $<10\%$ [23] was used as the cut-off for positive signals, Figure S15: Body weights are comparable between A vehicle vs palbociclib in HT77 and B between vehicle vs OTX015 treated groups and for HT96 PDXs. All body weights are represented as mean $\pm$ SEM. Table S7: Intensity values of protein expression obtained from RPPA data following pathway groupings of proteins based on their function, Table S8A: Prioritized therapies for HT72 based on therapeutic response biomarkers and actionable targets, Table S8B: Prioritized therapies for HT77 based on therapeutic response biomarkers and actionable targets, Table S8C: Prioritized therapies for HT87 based on therapeutic response biomarkers and actionable targets, Table S8D: Prioritized therapies for HT96 based on therapeutic response biomarkers and actionable targets, Table S8E: Prioritized therapies for HT74 based on therapeutic response biomarkers and actionable targets, Table S8F: Prioritized therapies for HT120 based on therapeutic response biomarkers and actionable targets.

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Conflicts of Interest: H.E.S. is a retiree and shareholder of Eli Lilly and Co. J.D.W. is a consultant to and shareholder of Theralink Technologies, Inc. E.F.P. is a consultant to and shareholder of Theralink Technologies, Inc. and Perthera, Inc. All other authors declare that they have no conflict of interest.

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Riyahi N, Pandya PH, Bailey BJ, Dobrota EA, Young C, Shannon HE, Barghi F, Malko R, Bijangi-Vishehsaraei K, Trowbridge M, Coy K, Kennedy FM, Sinn AL, Angus S, Ferguson MJ, Saadatzaheh MR, Pollok KE. "Therapeutic Targeting of BET Bromodomain Proteins in Osteosarcoma." Wells Center Retreat. Indianapolis, Indiana., USA Aug 2022 also presented at IU Simon Comprehensive Cancer Center Cancer Research Day (Oct 2022) Indianapolis, Indiana., USA; *3rd place Award- translational research category for graduate students.*

Osteosarcoma (OS) is an aggressive pediatric cancer with ~35% of patients developing metastasis over time. The survival rate for metastatic and relapsed OS patients is <30% and there is currently no standardized salvage therapy. Lack of efficacy is attributed to extensive genetic complexity present in OS that is partly due to moderate levels of replication stress (RS). While high levels of RS can induce cell death, moderate RS levels may cause genomic instability that contributes to progression of OS. Therefore, induction of RS to high levels, especially in genetically complex cancers like OS, could be a promising therapeutic strategy. Bromodomain and extra-terminal domain (BET) proteins (BRD2,3,4) are a family of epigenetic readers that not only regulate gene expression networks, but also regulate DNA replication and hence, RS. BRD4 directly regulates major factors involved in DNA replication and checkpoint signaling. Thus, disruption of BRD4 function should exacerbate RS to levels that cause cell death. The objective of this study is to test the hypothesis that BET inhibition potentiates the efficacy of current salvage therapy through RS induction in aggressive OS. The effects of BET inhibitors (BETi), AZD5153 and OTX-015, as single agents and in combination with drugs used in salvage therapy such as Topotecan were evaluated for effects on OS cell growth, PARP cleavage, and the DNA damage repair network. BET knockdown experiments were performed to evaluate target selectivity and dependency. In-vivo efficacy and safety studies focused on patient-derived xenografts (PDXs) of relapsed OS. TT2-77 xenoline, Saos2, G292, and U2OS cell lines were selected for *in vitro* experiments. Combination index and Bliss independence analyses demonstrated additive to synergistic cell growth inhibition upon treatment with clinically relevant concentrations of BETi+topotecan. Significant increase in PARP cleavage was observed in the combo compared to single agent, indicating enhancement of apoptosis. Moreover, Western analyses demonstrated that BETi induces its effect, at least partly, via decreased CHK1 activation and increased DNA damage. Selective siRNA treatments illustrated that transient knockdown of individual BET proteins was not sufficient for potentiation of topotecan-induced cell death in OS cells, indicating that simultaneous knockdown of BETs may be required. Dose-finding studies of AZD5153 in relapsed OS PDXs that harbor replication stress signatures (TT2-77 and PDX96) indicated that daily doses of 1.25 or 2.5 mg/kg AZD5153 were well tolerated and effective in partially suppressing tumor growth compared to vehicle ($p < 0.05$, Multiple t-test; Holm-Sidak). *In vivo* combination treatments of BET+topotecan are in progress. These data collectively suggest that BET inhibition alongside salvage therapy holds promise as a novel treatment strategy for inducing RS-mediated cell death in aggressive OS.

N. Riyahi, P.H. Pandya, B.J. Bailey, E.A. Dobrota, C. Young, H.E. Shannon, F. Barghi, R. Malko, K. Bijangi-Vishehsaraei, M.A. Trowbridge, K. Coy, F.M. Kennedy, A.L. Sinn, S. Angus, M.J. Ferguson, M. R. Saadatzaheh, K.E. Pollok. Therapeutic targeting of BET bromodomain proteins increases DNA damage and potentiates salvage therapy in osteosarcoma xenografts derived from patients with replication stress signatures. Annual AACR Meeting 2023, Orlando, Fla., April 2023. *Selected for Oral Presentation at the IU Simon Comprehensive Cancer Center Cancer Research Day (May 2023) Indianapolis, Indiana., USA*

Osteosarcoma (OS) is an aggressive cancer of the bone with high metastatic potential in pediatric as well as adolescent and young adults. The survival rate for metastatic and relapsed OS patients is <30% and there is currently no effective standardized salvage therapy. Lack of efficacy is attributed to genetic complexity present in OS that is partly due to moderate levels of replication stress (RS). While high levels of RS can induce cell death, moderate RS levels may cause genomic instability that contributes to OS progression. Therefore, induction of RS to high levels that cause cell death could be a promising therapeutic strategy. Bromodomain and extra-terminal domain [BET proteins (BRD2,3, and 4)] are a family of epigenetic readers that not only regulate gene expression networks, but also regulate DNA replication and RS. Thus, we tested the hypothesis that BET inhibition will potentiate the efficacy of salvage therapy through exacerbation of RS in xenograft models of aggressive OS. The effect of the bivalent BET inhibitor (BETi), AZD5153, as a single agent and in combination with cytotoxic agents such as topotecan and ifosfamide was evaluated. Combination index and Bliss independence analyses demonstrated additive to synergistic cell growth inhibition in OS cell lines upon treatment with clinically relevant concentrations of AZD5153+topotecan/ifosfamide. Treatment with PROTAC ARV825 that degrades BET proteins, resulted in similar growth inhibitory effects. Significant increase in PARP cleavage was observed following AZD5153+topotecan treatment compared to single agent, indicating enhancement of apoptosis. In addition, western blot and comet assays showed that BETi+topotecan induces its effect, at least partly, through increased DNA damage and RS *in vitro*. *In vivo* efficacy and safety studies focused on patient-derived xenografts (PDXs) of naïve and pre-treated OS that harbor RS signatures. AZD5153 as a single agent significantly suppressed tumor growth in both naïve (PDX96) and pretreated (TT2) OS PDX models compared to vehicle ($p < 0.05$, Two-way ANOVA; Holm-Sidak). Anti-tumor effect correlated with increased γ -H2AX following AZD5153 exposure in PDX, indicative of increased RS. Moreover, RNA-seq analysis integrated with kinome profiling data from BETi-treated PDX exhibited deregulation of factors involved in RS. Combination treatments of BETi+topotecan/ifosfamide indicated that AZD5153 potentiated the anti-cancer effect of salvage therapy in TT2 OS PDX, was well tolerated, and increased the probability of survival in mice. Efficacy in an OS RS+ metastatic lesion model is in progress. These data collectively suggest that BET inhibition as a single agent and in combination with low-dose salvage therapy holds promise as novel treatment strategies for inducing RS-mediated cell death in aggressive OS.

*Dobrota E, Pandya PH, Jannu AJJ, Bijangi-Vishehsaraei K, Bailey BJ, Barghi F, Shannon HE, Riyahi N, Damayanti NP, Young C, Malko R, Justice R, Albright E, Sandusky Gem Wurtz LD, Collier CD, Petricoin E, Coy K, Trowbridge M, Sinn AL, Ferguson MJ, Huang K, Zhang J, Saadatzadeh MR, Pollok KE. "Multi-omics analytic pipeline of pediatric and AYA solid tumor patient-derived xenografts reveals biomarkers of therapeutic sensitivity and resistance." IU Simon Comprehensive Cancer Center Cancer Research Day (Oct 2022) Indianapolis, Indiana., USA

Solid tumors account for ~60% of pediatric, as well as adolescent, and young adult (AYA) cancers. Notably, prognosis remains quite poor for those patients with progressive disease underscoring the need for new therapeutic approaches that are efficacious and safe. Precision genomics has emerged as a valuable tool for revealing potential disease mechanisms and treatment options. A multi-OMICs approach that identifies and cross-validates oncogenic signaling networks in aggressive cancers is an attractive method to help prioritize therapeutic vulnerabilities that can be tested in preclinical models and ultimately in clinical trials.

Establishment of patient-derived xenograft (PDX) models provides an opportunity to expand tumor specimens from patients so that genomically-guided therapies can be tested. An integrated multi-OMICs pipeline was developed to identify therapeutic targets and understand to what extent the molecular signatures of a panel of PDX remain stable or evolve following serial passaging in immunodeficient mice.

PDX models derived from naïve and pre-treated sarcoma or Wilms tumor patients which differ in genomic complexity were established via direct implantation of tumor specimens into NOD/SCID/ γ null mice. The original tumor and corresponding PDX passages were evaluated by a multi-OMICs approach including whole genome sequencing, RNA-seq, reverse phase protein array, and pathway enrichment analyses. CDK4/6 pathway hyperactivation was prioritized as a high-risk signature for proof-of-concept in-vivo validation in an osteosarcoma PDX derived from a patient with metastatic disease.

Major tumorigenic gene and pathway alterations were validated, and potential therapeutic targets identified by integrative pathway analysis. While some divergence between original tumor and the respective PDX was evident at different levels of DNA analysis, cancer-associated genes and oncogenic pathways were retained in the majority of PDXs derived from sarcomas and Wilms tumors following serial passaging. CDK4/6 small molecule inhibitor Palbociclib decreased OS tumor growth providing validation of a targeted anti-tumor response but also highlighted the need for combination therapy.

Through multi-OMICs analyses, we not only identified actionable pathways in PDXs but also validated the fidelity of molecular signatures of PDX models following serial passaging. PDX that retain the original oncogenic molecular signature will help prioritize targets for development of efficacious and safe therapies that can ultimately be translated to the clinic.

Dobrota E, Pandya PH, Jannu AJJ, Bijangi-Vishehsaraei K, Bailey BJ, Barghi F, Shannon HE, Riyahi N, Damayanti NP, Young C, Malko R, Justice R, Albright E, Sandusky G.E. Wurtz LD, Collier CD, Petricoin E, Coy K, Trowbridge M, Sinn AL, Ferguson MJ, Huang K, Zhang J, Saadatzaheh MR, Pollok KE. "Integrative Analysis of Multi-omics Data on Pediatric and AYA solid tumor Patient-Derived Xenografts Unveils Biomarkers of Therapeutic Response." IU Simon Comprehensive Cancer Center Cancer Research Day (May 2023) Indianapolis, Indiana., USA; *2nd Place Award – Research Associate category*

Solid tumors account for ~60% of pediatric, as well as adolescent, and young adult (AYA) cancers. Notably, prognosis remains quite poor for those patients with progressive disease underscoring the need for new therapeutic approaches that are efficacious and safe. Precision genomics has emerged as a valuable tool for revealing potential disease mechanisms and treatment options. A multi-OMICs approach that identifies and cross-validates oncogenic signaling networks in aggressive cancers is an attractive method to help prioritize therapeutic vulnerabilities that can be tested in preclinical models and ultimately in clinical trials.

Establishment of patient-derived xenograft (PDX) models provides an opportunity to expand tumor specimens from patients so that genomically-guided therapies can be tested. An integrated multi-OMICs pipeline was developed to identify therapeutic targets and understand the stability or evolution of molecular signatures in a panel of PDXs following serial passaging in immunodeficient mice.

PDX models derived from naïve and pre-treated sarcomas [osteosarcoma (OS) and rhabdomyosarcoma (RMS)] or Wilms tumor patients, which differ in genomic complexity, were established via direct implantation of tumor specimens into NOD/SCID/ γ null mice. The original tumor and corresponding PDX passages were evaluated by a multi-OMICs approach including whole genome sequencing, RNA-seq, reverse phase protein array, and pathway enrichment analyses. Major tumorigenic gene and pathway alterations were validated, and potential therapeutic targets identified by integrative pathway analysis. Overall, cancer-associated genes and oncogenic pathways were retained in the majority of the PDXs analyzed following serial passaging.

CDK4/6 pathway hyperactivation was prioritized as a high-risk signature for validation in an OS PDX derived from a patient with metastatic disease. CDK4/6 inhibitor (CDK4/6i) Palbociclib decreased OS tumor growth, providing validation of a targeted anti-tumor response but also highlighted the need for combination therapy.

Similarly, activated CDK4/6 pathway has also been a molecular signature of interest in the highly aggressive embryonal RMS (ERMS) models we have developed as well. In ERMS PDX HT174, we employed a small molecule inhibitor screen based on actionable components of the molecular signature (RNA-seq and protein validation) which included inhibition of CDK4/6 pathway. While the tumor was sensitive to a CDK4/6i, the highest sensitivity was to a bromodomain and extra-terminal domain proteins inhibitor (BETi)/AZD5153. Notably, BETi resulted in a profound decrease in tumor growth during the initial phase of dosing, but the tumors eventually developed AZD5153 resistance and grew, again highlighting the need for combination therapies. The mechanism-of-resistance is under investigation. Data obtained will provide new avenues to maximize the benefit of BETi therapy in ERMS.

As such, through multi-OMICs analyses, we were established PDXs that retained the original oncogenic molecular signatures as a way to identify actionable pathways and prioritize targets for development of efficacious and safe therapies that can ultimately be translated to the clinic.