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Oncogenic *NTRK3* mutations exhibit differential sensitivity to tropomyosin receptor kinase inhibitors in patients with acute myeloid leukemia

Sunil K. Joshi^{1,2,3}, Ariane Huang², Janét Pittsenbarger², Ujwal Shinde⁴, Camilo Posso⁵, Paul D. Piehowski⁶, Sara J.C. Gosline^{7,8}, Richard D. Press^{2,9}, Marina A. Gritsenko⁷, Chelsea Hutchinson⁷, Karl K. Weitz⁷, Kevin Watanabe-Smith^{2,10}, Nicola Long², Karin D. Rodland^{2,7}, Jeffrey W. Tyner^{2,3,11}, Brian J. Druker^{2,3}, Cristina E. Tognon^{2,10,#}

¹Department of Medicine, Division of Hematology, Stanford University School of Medicine, Stanford, CA, USA

²Knight Cancer Institute, Oregon Health & Science University, Portland, OR, USA

³Division of Hematology & Medical Oncology, Department of Medicine, Oregon Health & Science University, Portland, OR, USA

⁴Department of Chemical Physiology and Biochemistry, Oregon Health & Science University, Portland, OR, USA

⁵Biological Sciences Division, Pacific Northwest National Laboratory, Richland, WA, USA

⁶Environmental and Molecular Sciences Division, Pacific Northwest National Laboratory, Richland, WA, USA

⁷Biological Sciences Division, Pacific Northwest National Laboratory, Richland, WA, USA

⁸Department of Bioengineering, Oregon Health & Science University, Portland OR, USA

⁹Department of Pathology, Oregon Health & Science University, Portland, OR, USA

¹⁰Division of Oncological Sciences, Oregon Health & Science University, Portland, OR, USA

¹¹Department of Cell, Development, & Cancer Biology, Oregon Health & Science University, Portland, OR, USA

#CORRESPONDENCE: Cristina E. Tognon, E-mail: tognon@ohsu.edu

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AUTHOR'S CONTRIBUTION:

Study supervision: S.K. Joshi, B.J. Druker, C.E. Tognon

Conception and design: S.K. Joshi, C.E. Tognon

Development of methodology: S.K. Joshi, U. Shinde, C.E. Tognon

Acquisition of data: S.K. Joshi, A. Huang, J. Pittsenbarger, U. Shinde, P. D. Piehowski, R. D. Press, Marina A. Gritsenko, Chelsea Hutchinson, Karl K. Weitz, Nicola Long

Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): S.K. Joshi, A. Huang, J. Pittsenbarger, U. Shinde, C. Posso, S.J.C. Gosline, K. Watanabe-Smith, C.E. Tognon

Writing, review, & editing of the manuscript: S.K. Joshi, U. Shinde, C. Posso, P. D. Piehowski, Sara JC Gosline, K. Rodland, J.W. Tyner, B.J. Druker, C.E. Tognon

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Genomic analyses of tumors have identified rare oncogenic aberrations in the neurotrophic receptor tyrosine kinase (NTRK) genes—*NTRK1*, *NTRK2*, *NTRK3*—encoding TrkA, TrkB, TrkC receptors, respectively¹. Characterization of Trk oncoproteins in solid malignancies led to the Food and Drug Administration (FDA) approval of first-generation Trk inhibitors, larotrectinib² and entrectinib³. Second-generation Trk inhibitors, such as repotrectinib⁴, have been granted FDA breakthrough designation for patients failing first-line Trk inhibitors. Other inhibitors, such as selitrectinib⁵ and taletrectinib⁶, are currently in clinical trials.

Beyond solid malignancies, there is a growing appreciation for Trk signaling in hematologic malignancies owing to the success of Trk inhibitors⁷. Similar to solid tumors, Trk fusions (e.g., *ETV6-NTRK3*) represent the primary genetic aberration that confer oncogenic behavior in Trk-driven liquid tumors⁷. Our laboratory previously identified four novel, *NTRK2* and *NTRK3* non-kinase domain point mutations in patients with leukemia. These mutations result in Trk activation via increased cell surface abundance and receptor dimerization and importantly are amenable to Trk inhibition⁸. Herein we characterize two additional novel *NTRK3* kinase domain point mutations, L568V and P792S, identified through exome sequencing of 805 AML patient samples via the Beat AML research program⁹ (**Figure 1A**) with variant allele frequencies (VAFs) of 26% and 32%, respectively. These specimens were collected with informed consent from patients with AML according to a protocol approved by the Oregon Health & Science University (OHSU) institutional review board (IRB #4422; NCT01728402). Both were confirmed via Sanger sequencing (**Figure 1B-C**).

Initially, both patients were diagnosed with myelodysplastic syndromes (MDS) prior to AML progression. Patient 2232, harboring *NTRK3*^{P792S}, transformed into AML after 7 months, received standard of care induction and consolidation, and remained in remission one-year post-transplant but was lost to follow up. Patient 4838 with *NTRK3*^{L568V} was initially treated with lenalidomide and azacitidine but transformed to AML with the likely co-acquisition of *NRAS* and

NTRK3 mutations and died within 79 days despite a trial of palliative chemotherapy. A DNA sample during the MDS phase was available for this patient at approximately 135 days prior to AML transformation. Sequential sequencing efforts (**Supp Table 1**) revealed that the *NTRK3*^{L568V} mutation was undetectable during MDS but arose during the evolution to AML (**Figure 1D**). Based upon the VAFs, the *NRAS*^{Q61H} and *NTRK3*^{L568V} mutations appear to arise in the same clone. No DNA was available to perform a similar clonal analysis for patient 2232. Both patients exhibited similar co-occurring mutations including: *NRAS* mutations (Q61H for L568V & G12D for P792S), identical *IKZF1*^{N159S} mutations, and *RUNX1* alterations (**Supp Table 1**; Patient Characteristics). While P792 is a conserved residue among receptor tyrosine kinases, the L568 residue is not, but is within an area of conserved residues (**Supp Figure 1A**).

Both *NTRK3* mutations conferred interleukin 3 (IL-3) independent growth in Ba/F3 cells, while overexpression of *NTRK3*^{wildtype(WT)} or *NTRK3*^{I695T} did not (identified in TCGA¹⁰; TCGA-AB-2821; **Figure 1E**). Immunoblot analysis revealed increased total and phosphorylated TrkC expression in mutant-transformed Ba/F3 cells (**Figure 1F**).

To explore how *NTRK3* mutants activate TrkC, we performed molecular dynamic simulation (MDS) studies with the available crystal structure of the TrkC kinase domain¹¹ and homology modeling to fill in missing loop regions. We developed structural models of *NTRK3*^{WT}, *NTRK3*^{L568V}, and *NTRK3*^{P792S} in their 'active' or Aspartate-Phenylalanine-Glycine (DFG)-in and 'inactive' or DFG-out states to assess how the mutants altered the structure, stability, dynamics, or interactions of TrkC. Root mean square fluctuation (RMSF) analyses suggested that both mutants enabled an increase in local flexibility relative to *NTRK3*^{WT}, which was most pronounced in the TrkC inactive conformation (**Figure 1G**) relative to the active conformation (**Figure 1H**). Three regions were evaluated: Region 1, containing L568 and encompassing the P-loop that interacts with ATP; Region 2, a loop region proximal to P792 spatially; and Region 3, encompassing the activation loop. Region 4 served as a negative control as it is mostly unperturbed in both mutants regardless of receptor conformation. MDS analyses revealed

residue hyper-mobility within regions surrounding or encompassing the activation loop (i.e., Region 2 & 3) as evident by the magnitude of fluctuation relative to the *NTRK3*^{WT} baseline (denoted by dotted line; **Figure 1G-H**). *NTRK3*^{L568V} showed an approximate 3-fold difference in local fluctuation at Regions 2 and 3 in the inactive conformation. *NTRK3*^{P792S} showed an approximate 6-fold difference in fluctuation. The enhancement in local flexibility resulted in increased pocket volume (**Figure 1I-L**) near the receptor activation site, which again was most prominent in the inactive TrkC conformation (**Figure 1J-K**). WT TrkC receptor in the inactive conformation had a pocket volume of 494.4 Å³ (**Figure 1I**). This increased to 993.7 and 638.8 Å³ in the setting of L568V and P792S, respectively (**Figure 1J-K**). L568 is juxtaposed with the substrate binding site whereas P792 is located far from the substrate binding site and yet still affects inhibitor interactions. These mutant pocket volume sizes mirrored the active conformation of the WT receptor (743.0 Å³, **Figure 1L**). Mechanistically, these structural studies implicate that both *NTRK3* mutants possibly perturb the inactive receptor to adopt the active conformation, thereby conferring increased kinase activity with resulting oncogenicity.

To further characterize the downstream impact of *NTRK3* mutations, we performed global, phospho-serine/threonine, and phospho-tyrosine enriched proteomic analyses on Ba/F3 cells expressing empty vector, *NTRK3*^{WT}, *NTRK3*^{L568V}, and *NTRK3*^{P792S}. All mutant-expressing cell lines clustered distinctly from *NTRK3*^{WT}, empty vector, Ba/F3 parental cell lines (**Figure 2A**). The number of unique genes is summarized in **Supplemental Table 1** (Global Summary). These data were used to decipher pathways uniquely up- or down-regulated in each mutant cell line (**Figure 2B**). Common pathways, such as oxidative phosphorylation and E2F targets, were highlighted for both mutants, respectively compared to *NTRK3*^{WT}. Each mutant also possessed unique pathways, such as fatty acid metabolism for *NTRK3*^{L568V} and mitotic spindle for *NTRK3*^{P792S}. Gene set enrichment analysis (GSEA) brought forth upregulated terms associated with cell division (**Supp Figure 2A & B**). Kinase substrate enrichment analysis (KSEA) highlighted significant enrichment of PI3K/MTOR pathway substrates for *NTRK3*^{L568V}. Similar

substrates were highlighted for *NTRK3*^{P792S} but did not reach statistical significance (**Figure 2C**). Together these data support activation of canonical downstream TrkC signaling and reveal novel metabolic and catabolic pathways that may inform *NTRK3*^{L568V} and *NTRK3*^{P792S} activity.

We also assessed *ex vivo* sensitivity of primary cells available from patient 4838 harboring *NTRK3*^{L568V} and compared this IC₅₀ to the median IC₅₀ calculated for all patient samples tested with entrectinib⁹. Inhibitors with IC₅₀ values <20% of the median IC₅₀ are considered “effective”¹². The *NTRK3*^{L568V} patient fell outside this threshold, demonstrating a modest sensitivity to entrectinib with an IC₅₀ of 0.503 nM (**Figure 3A**). *Ex vivo* inhibitor data was not available for the patient harboring *NTRK3*^{P792S}. *NTRK3*^{L568V}-transformed Ba/F3 cells also exhibited decreased sensitivity to entrectinib (IC₅₀=120.2 nM) and larotrectinib (IC₅₀=375.8 nM; **Figure 3B-C**). Interestingly, *NTRK3*^{P792S}-Ba/F3 cells demonstrated robust sensitivity to both Trk inhibitors (IC₅₀=5.0 nM; larotrectinib and 1.3 nM; entrectinib). Both *NTRK3*-mutant-expressing Ba/F3 cell lines were sensitive to second generation Trk inhibitors – repotrectinib, selitrectinib, and taletrectinib (**Figure 3D-F**; all data summarized in **Supp Table 1**; Inhibitor Screen Summary).

We investigated the differential sensitivity pattern of *NTRK3*^{L568V} to Trk inhibitors via molecular docking studies by generating *in silico* models. We assessed larotrectinib and repotrectinib, two prototypical examples of first- and second-generation TrkC inhibitors, docked onto the active and inactive conformations of the TrkC kinase domain. Docking studies, including the dissociation constants calculated (**Supp Table 1**; Dissoc. Constants Docking Exp), were concordant with the trends observed in the cell-based IC₅₀ data (**Figure 3C & D**). Repotrectinib had lower dissociation constants for both mutants compared to larotrectinib. We then analyzed the multiple interactions—hydrophobic, π-π, cation-π, and ionic—that facilitate binding between the WT or mutant receptors in the active conformation with larotrectinib and repotrectinib.

We focused on the active conformation of the TrkC receptor given that larotrectinib and repotrectinib are both type I Trk inhibitors and, therefore preferentially bind in the active, DFG-in conformation. A higher aggregate interaction score is reflective of better interactions against a given *NTRK3* mutant, which likely translates into increased Trk inhibition and decreased cell viability experimentally (**Supp Figure 1B-C; Figure 3G-J**). In the case of *NTRK3*^{L568V}, repotrectinib demonstrated enhanced binding with a cumulative interaction score of 75.3 compared to 59.0 observed with larotrectinib binding (**Figure 3G-H**). The higher affinity for repotrectinib is likely driven by increased hydrophobic and π - π interactions and presence of a new cation- π interaction not seen with the binding of larotrectinib to the L568V mutant (**Figure 3G-H**). Although repotrectinib binds the active site of *NTRK3*^{P792S} in a different orientation than *NTRK3*^{L568V}, a similar trend was observed with interaction scores of 107.8 and 68.2 for repotrectinib and larotrectinib, respectively (**Figure 3I-J**). Taken together, perturbation of the *NTRK3* kinase domain not only alters its dynamicity but impinges its sensitivity to first-generation Trk inhibitors, most notably in the case of L568V.

Herein we characterized two *NTRK3* kinase domain mutations in patients with AML and confirmed their oncogenicity. Importantly, experimental validation and molecular docking studies demonstrate that both mutations exhibit differential sensitivity against first-generation Trk inhibitors but robust sensitivity against second-generation inhibitors. Although specific genetic aberrations have long been known to drive the development of various cancers, our work shows that a reliance on genomics alone serves as a bottleneck and limits the deployment of precise therapeutics. To this end, the development of cell-based models enables rigorous evaluation of small molecule inhibitor sensitivity and resistance, thereby informing clinical practice.

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FIGURE LEGENDS:

Figure 1: *NTRK3* point mutations found in acute myeloid leukemia (AML) patient samples alter local flexibility of the inactive receptor, enabling TrkC activation and oncogenicity.

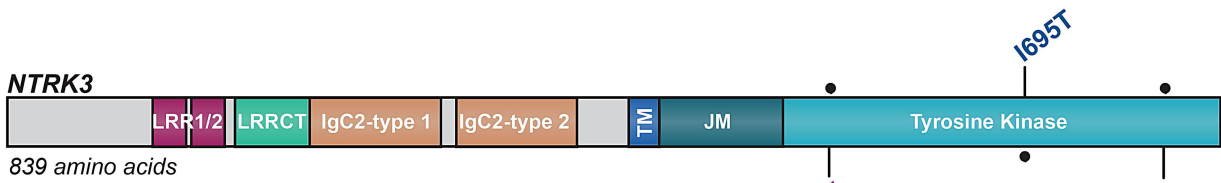
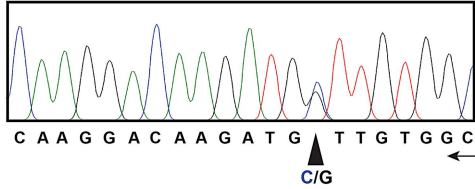
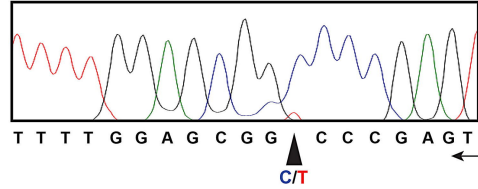
A. Lollipop diagram depicts the location of *NTRK3* point mutations (Below: Beat AML; Above: TCGA). The locations of the following domains are included: LRR 1, LRR 2, LRRCT, Ig-like C2-type 1, Ig-like C2-type 2, transmembrane (TM), juxtamembrane (JM), and tyrosine kinase. **B-C.** Electropherograms from Sanger sequencing of patient genomic DNA confirm the presence of *NTRK3*^{L568V} and *NTRK3*^{P792S} mutations. Peaks correspond to the following nucleotides: A (green), T (red), C (blue), and G (black). Arrows indicate direction of sequencing. **D.** Fish plot illustrating evolution of *NTRK3*^{L568V} from DNA isolated from bone marrow aspirate as captured by target panel DNA sequencing at myelodysplastic syndrome (MDS) stage (Day -135) and at time of transformation to AML (Day 0). Top x-axis indicates clinical time point. Clonal fractions are displayed as double the Variant Allele Fraction (VAF), consistent with the assumption of heterozygous mutations without copy number variation except for DNMT3A, which is commonly biallelic and not doubled in this depiction. ETV6 and IKZF1 are presented as a single event, using the larger VAF of the pair, as the clonal order could not be resolved with data available. Similarly, the *NTRK3* and *NRAS* VAFs are also presented as a single event. **E.** *NTRK3*^{L568V} and *NTRK3*^{P792S} mutations transform the murine Ba/F3 pro-B cell line and enable IL-3 independent growth. No growth was observed in Ba/F3 cells harboring an empty vector (pMX-IRESpuro), wildtype (WT) *NTRK3*, or *NTRK3*^{G695T}. Total viable cells are plotted over time and cell growth was measured after the withdrawal of interleukin 3 (IL-3). This experiment was repeated at least twice with consistent results. **F.** Expression of total and phosphorylated TrkC is increased in mutant-transformed Ba/F3 cells relative to WT cells. GAPDH served as a loading control. Prior to lysis, WT cells were grown in IL-3 supplemented media and all cell lines were starved overnight in 0.1% bovine serum albumin (BSA) RPMI. **G-H.** MDS studies reveal differences in root mean square fluctuation (RMSF, Angstroms (Å)) between *NTRK3* mutations and wildtype (WT) in the active (Aspartate-Phenylalanine-Glycine (DFG)-in) and inactive (DFG-out) TrkC conformations. Areas denoted in gray highlight regions of residues that exhibit fluctuation and/or served as a negative control for simulation studies. **I-L.** Pocket volume size (Å³) was measured as a surrogate to assess TrkC activation in both the inactive and active conformations. Increase in pocket volume was observed following the addition of *NTRK3* mutations in the inactive (DFG-out) conformation, denoting increased TrkC activation. Orange: ATP-binding loop; Yellow: Aspartate-Phenylalanine-Glycine (DFG) motif, DFG-in (active conformation), or DFG-out (inactive conformation); Green: cavity to assess changes in pocket-volume size. For detailed methods see: <https://doi.org/10.6084/m9.figshare.26516974.v4>.

Figure 2: Proteomic analysis of *NTRK3* mutant-expressing Ba/F3 cells highlights activation of canonical downstream signaling pathways and alterations in metabolism. **A.**

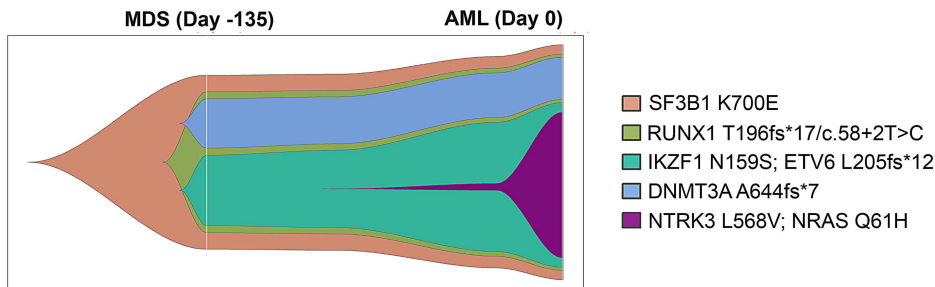
Visualization of global proteomic (left), phospho-serine/threonine (middle) and phospho-tyrosine (right) profiling by principal component analysis (PCA) shows distinct clustering amongst parental, empty vector, *NTRK3*^{WT}, and *NTRK3*-mutant expressing cells. **B.** Volcano plots depicting the significantly differentially expressed peptides identified in *NTRK3* mutant-expressing Ba/F3 cells as assessed by -log₁₀ (t-test adjusted) values. Peptides were annotated using the Hallmarks of cancer pathway analysis. Three pathways are highlighted for each mutant. **C.** Kinase substrate enrichment analysis (KSEA) analyses reveal alterations in PI3K/AKT/MTOR and MAPK pathways in *NTRK3* mutant-expressing cell lines. Red bars = significantly upregulated kinase activity. Grey bar = not significant. For detailed methods see: <https://doi.org/10.6084/m9.figshare.26516974.v4>.

Figure 3: *NTRK3* mutants are unequivocally sensitive to second-generation Trk inhibitors owing to enhanced local interactions. **A.** *Ex vivo* entrectinib sensitivity assay results from the primary acute myeloid leukemia (AML) patient sample harboring *NTRK3*^{L568V} from Beat AML data base. Freshly isolated mononuclear cells isolated from the bone marrow were incubated with increasing doses of entrectinib and assayed for viability after three days of incubation. Sensitivity was evaluated by calculating % median IC₅₀, which has historically been a marker for patient samples demonstrating sensitive to a given drug screened in the inhibitor assay⁹. **B-F.** Six replicates of wild type (WT) and mutant *NTRK3* Ba/F3 cells were plated with varying concentrations of entrectinib (**B**), larotrectinib (**C**), repotrectinib (**D**), selitrectinib (**E**), and talotrectinib (**F**) for 72 hours. *NTRK3*^{WT} cells were plated in media supplemented with interleukin 3 (IL-3). Cell viability was determined using a tetrazolamine-based viability assay. Viability is represented as a percentage of the untreated control. The average mean ± standard error of the mean (SEM) is shown. **G-J.** Larotrectinib (**G, I**) and repotrectinib (**H, J**) were docked *in silico* on the active conformation (Aspartate-Phenylalanine-Glycine (DFG)-in) of L568V- and P792S-mutated TrkC receptor to study extent of local interactions. Purple boxes at bottom right corner indicate aggregate interaction scores. A higher score correlates with enhanced interactions between TrkC receptor and respective inhibitor. Four different interactions are highlighted: hydrophobic (green), cation-π (blue), π-π (red), and ionic (pink). The cation-π interaction is formed by the proximity of a positively charged residue (Lys, Arg) with aromatic residues (Phe, Trp, and Tyr). Docking repotrectinib within the L568V- or P792S-mutated TrkC receptors, enables formation of the cation-π interaction that stems from the lysine residue at 572 and arginine at 702, respectively. In the case of both mutants, these positively-charged residues interact with a fluorinated phenyl group of repotrectinib, enabling formation of the cation-π interaction. The new cation-π interaction coupled with an increased number of hydrophobic, π-π, and ionic interactions enable repotrectinib to retain a single-digit nanomolar potency against both *NTRK3* mutations, which is not the case with larotrectinib. Notably, in the case of the *NTRK3*^{L568V} mutant, repotrectinib's potency exceeds that of entrectinib and larotrectinib by approximately 30 to 90 orders of magnitude, respectively (**Suppl Table 1**). For detailed methods see: <https://doi.org/10.6084/m9.figshare.26516974.v4>.

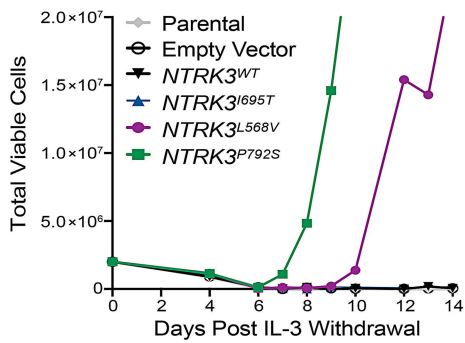
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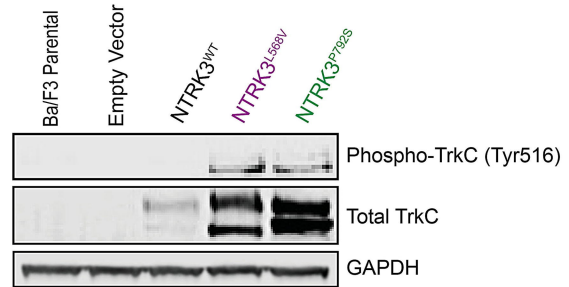
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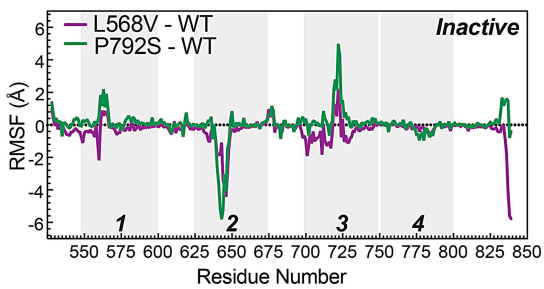
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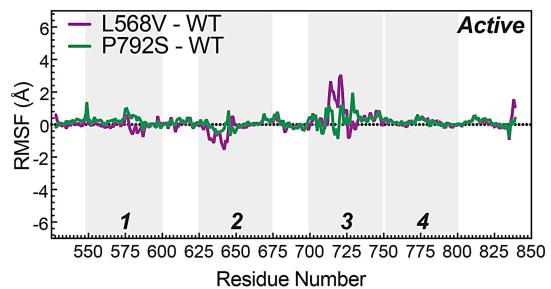
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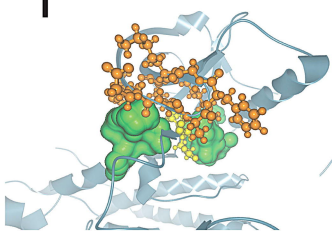
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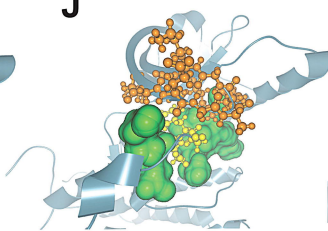
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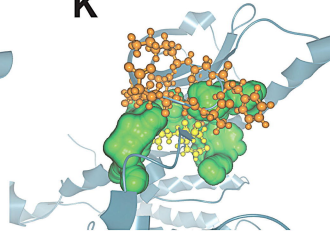
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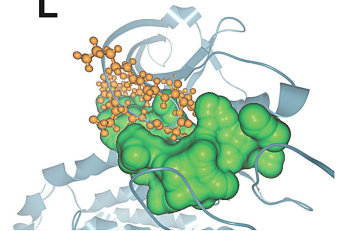
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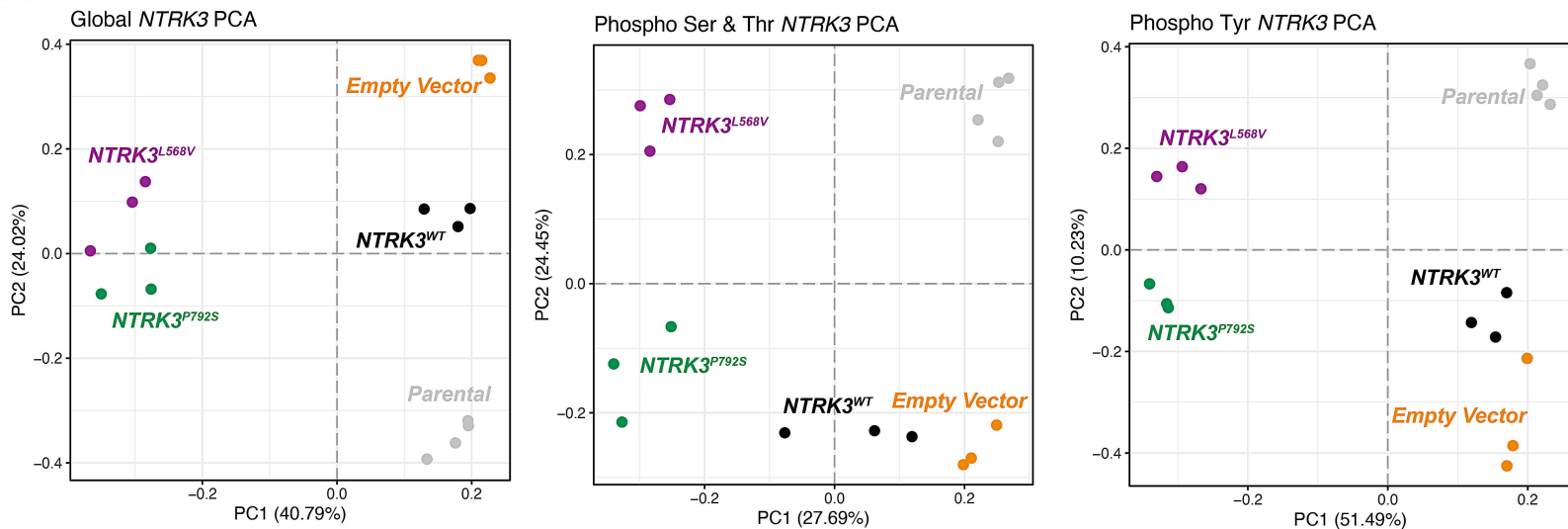
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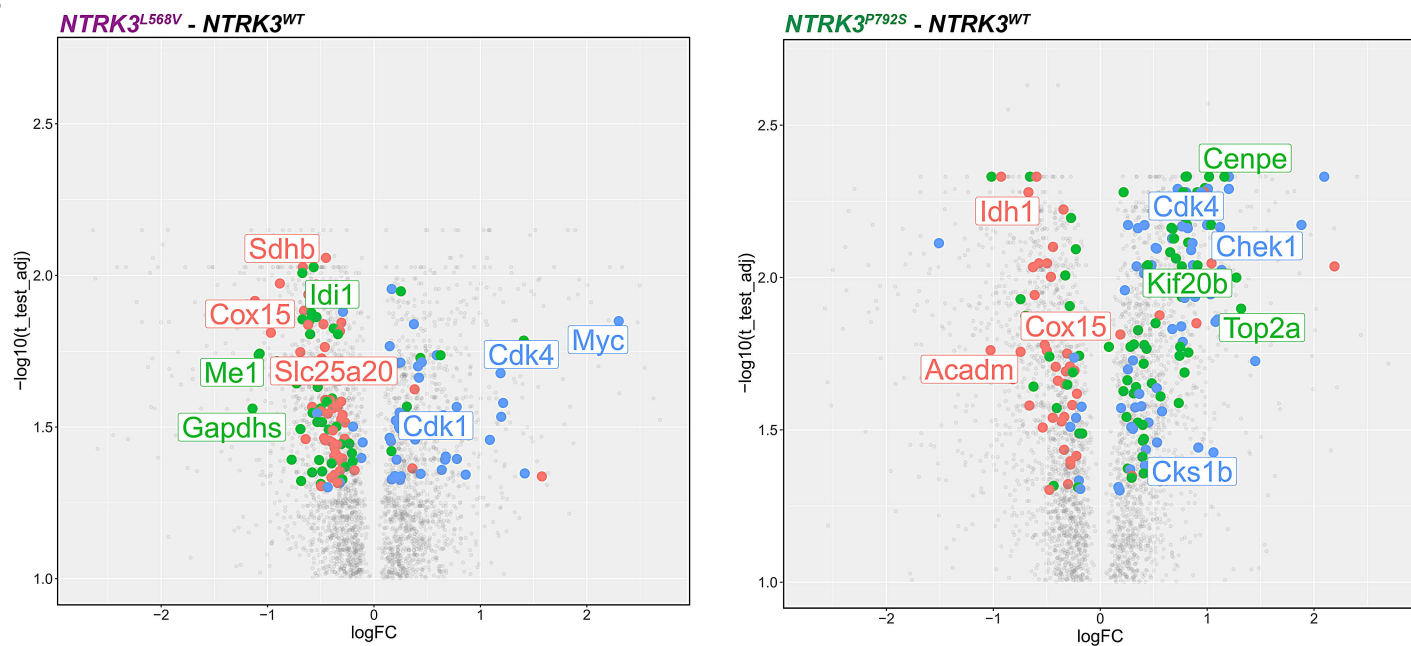
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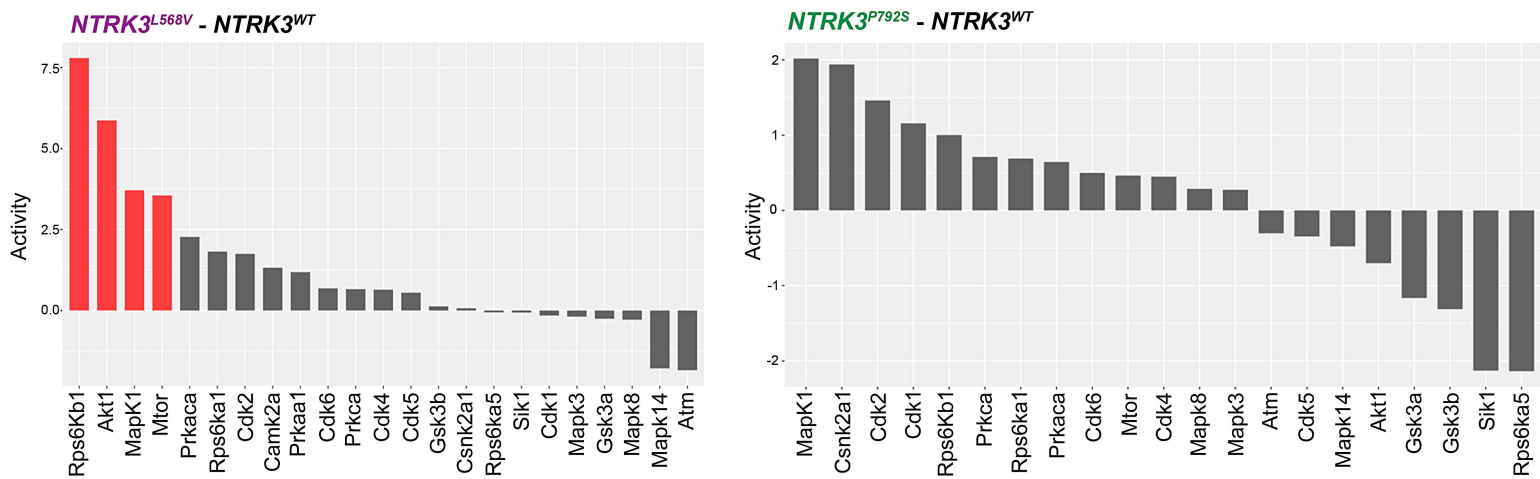
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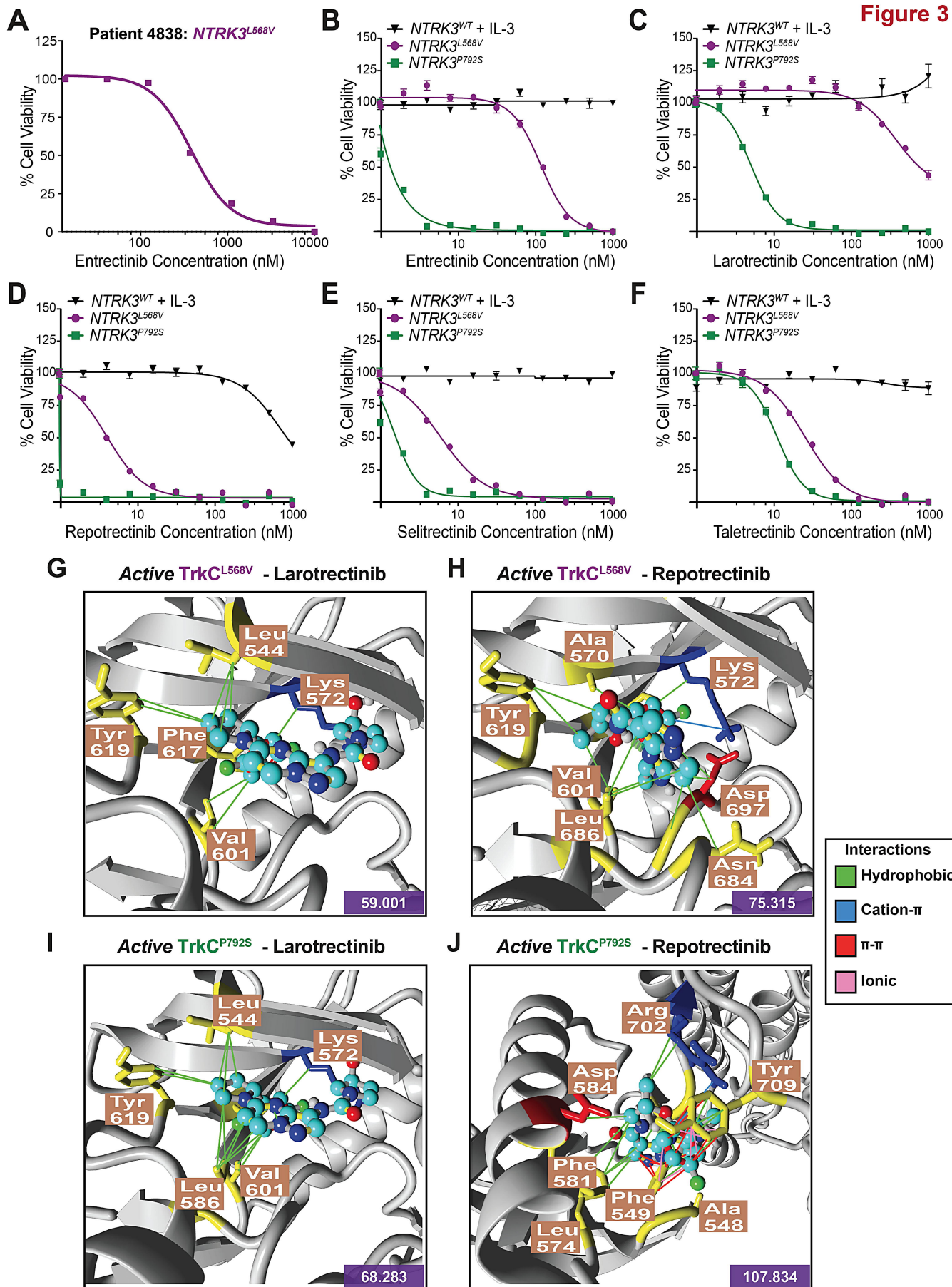


B



C





SUPPLEMENTAL DOCUMENT:

Oncogenic *NTRK3* mutations exhibit differential sensitivity to tropomyosin receptor kinase inhibitors in patients with acute myeloid leukemia

Sunil K. Joshi^{1,2,3}, Ariane Huang², Janét Pittsenbarger², Ujwal Shinde⁴, Camilo Posso⁵, Paul D. Piehowski⁶, Sara JC Gosline^{7,8}, Richard D. Press^{2,9}, Marina A. Gritsenko⁷, Chelsea Hutchinson⁷, Karl K. Weitz⁷, Kevin Watanabe-Smith^{2,10}, Nicola Long², Karin D. Rodland^{2,7}, Jeffrey W. Tyner^{2,3,11}, Brian J. Druker^{2,3}, Cristina E. Tognon^{2,10,#}

¹Department of Medicine, Division of Hematology, Stanford University School of Medicine, Stanford, CA

²Knight Cancer Institute, Oregon Health & Science University, Portland, OR

³Division of Hematology & Medical Oncology, Department of Medicine, Oregon Health & Science University, Portland, OR

⁴Department of Chemical Physiology and Biochemistry, Oregon Health & Science University, Portland, OR

⁵Biological Sciences Division, Pacific Northwest National Laboratory, Richland, WA

⁶Environmental and Molecular Sciences Division, Pacific Northwest National Laboratory, Richland, WA

⁷Biological Sciences Division, Pacific Northwest National Laboratory, Richland, WA

⁸Department of Bioengineering, Oregon Health & Science University, Portland OR

⁹Department of Pathology, Oregon Health & Science University, Portland, OR, USA

¹⁰Division of Oncological Sciences, Oregon Health & Science University, Portland, OR

¹¹Department of Cell, Development, & Cancer Biology, Oregon Health & Science University, Portland, OR

#CORRESPONDENCE: Cristina E. Tognon, E-mail: tognon@ohsu.edu

SUPPLEMENTAL FIGURE & TABLE LEGENDS:

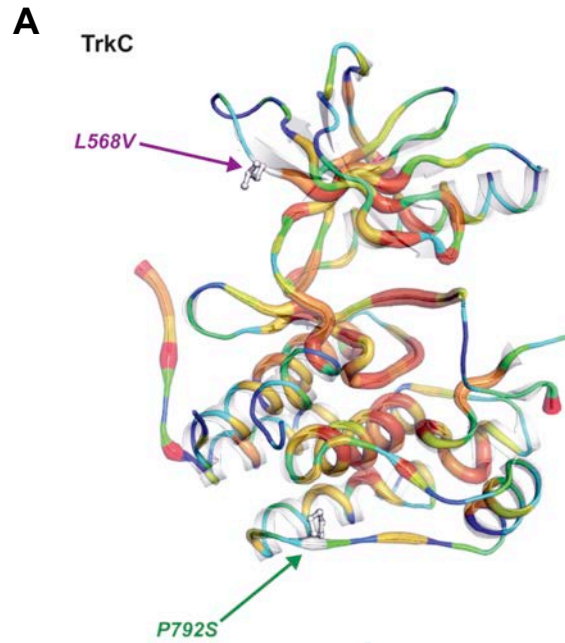
Supplemental Figure 1: *NTRK3* mutations lie in regions that are relatively conserved in receptor tyrosine kinases. **A.** The sequence conservation among receptor tyrosine kinases was ascertained using ConSurf-DB¹, a repository for the evolutionary conservation patterns of a majority of PDB proteins. The analysis was performed using the Ntrk3_human (Uniprot ID: Q16288) and was displayed on the protein structure (PDB ID: 6KZD) using PyMol. Tube thickness indicates the highest conservation score. P792 is relatively more conserved than L568. **B-C.** Larotrectinib (**B**) and repotrectinib (**C**) were docked *in silico* on the active conformation (Aspartate-Phenylalanine-Glycine (DFG)-in) of the Wild type (WT) TrkC receptor to study extent of local interactions. Purple boxes at bottom right corner indicate aggregate interaction scores. A higher score correlates with enhanced interactions between TrkC receptor and respective inhibitor. Four different interactions are highlighted: hydrophobic (green), cation- π (blue), π - π (red), and ionic (pink). The cation- π interaction is formed by the proximity of a positively charged residue (Lys, Arg) with aromatic residues (Phe, Trp, and Tyr).

Supplemental Figure 2: GO Biological Process annotation of up and down regulated pathways identified in *NTRK3* mutant expressing cells. Enrichment scores of the top eight Go Biological Processes (GOBP) identified as uniquely upregulated or downregulated in *NTRK3*^{L568V} (**A**), and *NTRK3*^{P792S} (**B**) expressing cells when peptides from *NTRK3*^{WT} expressing cells have been removed. Significance = adjusted p-value.

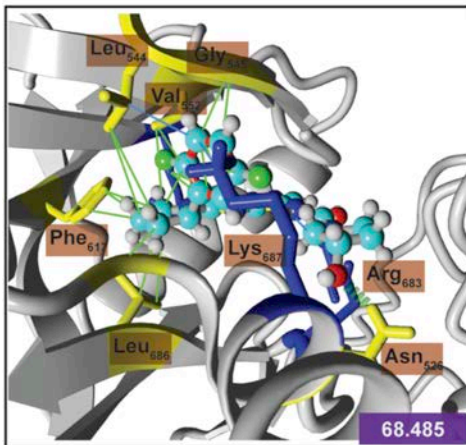
Supplemental Table 1: Tab 1: Patient characteristics. *NTRK3* mutation information, VAF, Variant Allele Frequency; CLIA, Clinical Laboratory Improvement Amendment; **Tab 2: Global Summary.** Total unique genes identified in differential protein expression analysis comparing *NTRK3* mutants to *NTRK* wildtype and empty vector control; **Tab 3: Inhibitor Summary Screen.** Summary of IC₅₀ values for inhibition of *NTRK3* mutant-transformed Ba/F3 cell growth by small-molecule inhibitors calculated from data presented in **Figures 3B-F**; **Tab 4: Dissociation Constants for Docking Experiment.** Summary of estimated dissociation constants between larotrectinib and repotrectinib for inactive and active conformations of TrkC^{WT}, TrkC^{L568V}, and TrkC^{P792S}. **Tabs 5 to 9: Full Proteomic Data from Ba/F3 Engineered Cell Lines.** Proteomic data from Ba/F3 cell lines engineer to express pMX-IRESpuro vector control, *NTRK3* wild type (*NTRK3*^{WT}), *NTRK3*^{L568V}, or *NTRK3*^{P792S}. **(Please, see Excel file)**

REFERENCES:

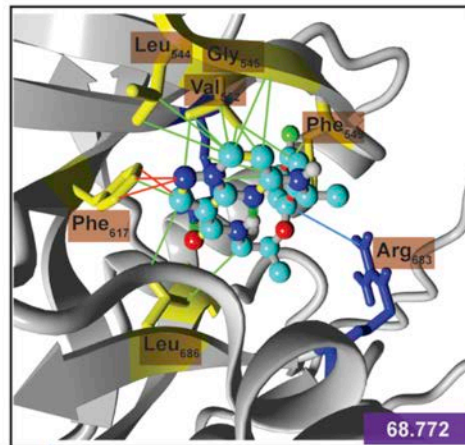
1. Ben Chorin A, Masrati G, Kessel A, et al. ConSurf-DB: An accessible repository for the evolutionary conservation patterns of the majority of PDB proteins. *Protein Sci.* 2020;29(1):258-267.



B Active TrkC^{WT} - Larotrectinib



C Active TrkC^{WT} - Repotrectinib



Interactions	
	Hydrophobic
	Cation- π
	π - π
	Ionic

Supplemental Figure 2

