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Authors

Paradis, Justine S
Acosta, Monica
Saddawi-Konefka, Robert
[et al.](#)

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Synthetic Lethal Screens Reveal Co-Targeting FAK and MEK as a Multimodal Precision Therapy for *GNAQ*-Driven Uveal Melanoma

Justine S. Paradis^{1, #}, Monica Acosta^{1, #}, Robert Saddawi-Konefka¹, Ayush Kishore¹, Frederico Gomes¹, Nadia Arang^{1, 2}, Manoela Tiago³, Silvia Coma⁴, Simone Lubrano¹, Xingyu Wu¹, Kyle Ford⁵, Chi-Ping Day⁶, Glenn Merlino⁶, Prashant Mali⁵, Jonathan A. Pachter⁴, Takami Sato^{7, 8}, Andrew E. Aplin^{3, 8}, J. Silvio Gutkind^{1, 9, *}

¹Moore's Cancer Center, University of California San Diego, La Jolla, CA, USA

²Biomedical Sciences Graduate Program, University of California San Diego, La Jolla, CA, USA

³Department of Cancer Biology, Thomas Jefferson University, Philadelphia, PA, USA

⁴Verastem Oncology, Needham, United States

⁵Department of Bioengineering, University of California San Diego, San Diego, CA, USA

⁶Laboratory of Cancer Biology and Genetics, Center for Cancer Research, NCI, NIH, MD, USA

⁷Department of Medical Oncology, Thomas Jefferson University, Philadelphia, PA, USA

⁸Sidney Kimmel Cancer Center, Thomas Jefferson University, Philadelphia, PA, USA

⁹Department of Pharmacology, University of California San Diego, La Jolla, CA, USA

Abstract

Purpose: Uveal melanoma (UM) is the most common eye cancer in adults. Approximately 50% of UM patients develop metastatic UM (mUM) in the liver, even after successful treatment of the primary lesions. mUM is refractory to current chemo- and immune-therapies, and most mUM patients die within a year. UM is characterized by gain-of-function mutations in *GNAQ/GNA11*, encoding Gαq proteins. We have recently shown that the Gαq-oncogenic signaling circuitry involves a non-canonical pathway distinct from the classical activation of PLCβ and MEK-ERK. *GNAQ* promotes the activation of YAP1, a key oncogenic driver, through focal adhesion kinase (FAK), thereby identifying FAK as a druggable signaling hub downstream from *GNAQ*. However, targeted therapies often activate compensatory resistance mechanisms leading to cancer relapse and treatment failure.

Experimental Design: We performed a kinome-wide CRISPR-Cas9 sgRNA screen to identify synthetic lethal gene interactions that can be exploited therapeutically. Candidate adaptive

*Correspondences should be addressed to J. Silvio Gutkind, Moore's Cancer Center, University of California San Diego, 3855 Health Sciences Dr, La Jolla, CA 92037. sgutkind@ucsd.edu. Phone: 858-534-5980.

[#]These authors contributed equally to this work

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resistance mechanisms were investigated by co-targeting strategies in UM and mUM *in vitro* and *in vivo* experimental systems.

Results: sgRNAs targeting the PKC and MEK-ERK signaling pathways were significantly depleted after FAK inhibition, with ERK activation representing a predominant resistance mechanism. Pharmacological inhibition of MEK and FAK showed remarkable synergistic growth-inhibitory effects in UM cells and exerted cytotoxic effects leading to tumor collapse in UM xenograft and liver mUM models *in vivo*.

Conclusions: Coupling the unique genetic landscape of UM with the power of unbiased genetic screens, our studies reveal that FAK and MEK-ERK co-targeting may provide a new network-based precision therapeutic strategy for mUM treatment.

Keywords

FAK; MEK; GNAQ; uveal melanoma; targeted therapy; precision medicine

Introduction

G protein-coupled receptors (GPCRs) are the largest family of cell surface proteins with over 800 members (1), and their dysregulation contributes to some of the most prevalent human diseases (2–4). GPCRs represent the largest family of targets for approved drugs. Strikingly, our recent analysis of human cancer genomes revealed that nearly 30% of human cancers present mutations in G proteins and GPCRs (5). In particular, uveal melanoma (UM) can be defined as a $G\alpha_q$ -driven malignancy. Indeed, Approximately 93% of UM lesions harbor activating mutations in *GNAQ* or *GNA11*, encoding for the alpha subunits $G\alpha_q$ and $G\alpha_{11}$ of the heterotrimeric G protein respectively (6,7). An additional 4% harbor mutations in the $G\alpha_q$ -linked receptor *CYSLTR2*, also acting as a driver oncogene (8).

UM is diagnosed in about 2,500 adults in the United States every year, and is the most common primary cancer of the eye in adults and the second most common melanoma subtype after skin cutaneous melanoma (SKCM) (9). While the majority of early-stage UM lesions can be treated by irradiation or enucleation, approximately 50% of the patients will metastasize, primarily to the liver, within 5–10 years after diagnosis (10). Inactivating mutations or copy loss of the *BAP1* gene, which is located on chromosome 3p21, are strongly associated with metastasis in UM patients (11), supporting that BAP1 functions as a metastasis suppressor (12). Most metastatic UM (mUM) patients are refractory to current chemotherapies and immune checkpoint blockade (13). Ultimately, the majority of advanced disease patients succumb within a year due to the suboptimal efficacy of these treatments, often combined with severe toxicities, underlying the high unmet medical need for new therapeutic strategies.

Most of the recent clinical research efforts in UM have focused on inhibiting the $G\alpha_q$ classical signaling pathway, PLC β -PKC-ERK. MEK inhibitors (MEKi) selumetinib and trametinib have been extensively evaluated for mUM. Despite encouraging results in pre-clinical studies, MEK inhibition with these agents has been shown to have nearly no impact on the overall survival of mUM patients, as single-agent or when combined

with chemotherapy (14–16). Our recent findings uncovered a non-canonical $G\alpha_q$ signaling pathway leading to the Rho-dependent activation of the Hippo/YAP pathway, which contributes to aberrant cancer cell growth (17,18). By further decoding this oncogenic signaling circuitry, we showed that the non-receptor focal adhesion kinase (FAK) is an integral node of this non-canonical $G\alpha_q$ pathway (19). Interestingly, FAK overexpression has already been associated with various cancer types including ovarian, head and neck, and breast cancers (20–22). Furthermore, multiple FAK inhibitors (FAKi) – defactinib (VS-6063), PF-562271, GSK2256098 and IN10018 – have been already tested in clinical trials, showing manageable toxicity profiles (23), and hence can be considered for mUM treatment. However, single-agent targeted therapies often activate compensatory mechanisms resulting in treatment resistance. This prompted us to perform a kinome-wide CRISPR-Cas9 sgRNA screen to identify synthetic lethal gene interactions in the context of FAKi that can be exploited therapeutically (24). Our study demonstrates that dual inhibition of MEK and FAK act synergistically to promote the conversion of cytostatic to cytotoxic inhibition of tumor growth, thereby identifying a new treatment option for mUM.

Materials and Methods

Reagents

DMEM, RPMI-1640 and antibiotic/antimycotic solution were purchased from Sigma-Aldrich. Turbofect, DMEM/F12 Glutamax, basic fibroblast growth factor (bFGF), epithelial growth factor (EGF), B-27 and N2 supplements were purchased from ThermoFisher Scientific. Fetal bovine serum (FBS, Lot 18H165) and penicillin/streptomycin from Sigma Aldrich, Polybrene from Millipore, Aquabluer from MultiTarget Pharmaceuticals.

VS-4718, VS-6063 and VS-6766 were provided by Verastem Oncology. All inhibitors used in this study were purchased from Selleck Chemicals (Houston, USA). Anhydrous DMSO, carboxymethyl cellulose (CMC), Polyethylene Glycol 400 (PEG400), Tween 80 were purchased from Sigma-Aldrich. Inhibitors were dissolved in DMSO to a stock concentration of 10 mM, aliquoted and stored at -80°C for *in vitro* experiments. For *in vivo* treatments, trametinib was prepared at a stock concentration 5mg/ml in DMSO and freshly diluted in PBS containing 5.2% PEG400 + 5.2% Tween 80 and a dose of 1mg/kg was administered once daily by intraperitoneal injection. VS-4718 was freshly diluted at 5 mg/ml in 0.5% carboxymethyl cellulose (CMC) (C5678, Sigma-Aldrich; St. Louis, MO) + 0.1% Tween 80 (P1754, Sigma Aldrich; St. Louis, MO) in sterile water (B. Braun Medical; Irvine, CA) and a dose of 10ml/kg was administered by oral gavage twice a day.

FAK, phospho-FAK, ERK, phospho-ERK, cleaved PARP1, cleaved Caspase 3, YAP, Cas9, and GAPDH antibodies were purchased from Cell Signaling. BAP1 and PARP1 were purchased from Santa Cruz Biotechnology. BrdU was purchased Abcam. All antibodies were diluted in 5% BSA at 1:500–1:1000 before use for immunoblotting and in 3% BSA at 1:100–1:400 for immunohistochemistry.

pHAGE PGK-GFP-IRES-LUC-W was from addgene (#46793).

Cell Lines

HEK 293T were cultured in High Glucose DMEM and supplemented with 10% FBS and antibiotic/antimycotic solution (100U penicillin, 0.1 mg/ml streptomycin and 0.25 µg/ml amphotericin B). UM cell lines were cultured in RPMI-1640 media supplemented with 10% FBS (92.1, OMM1.3, OMM1.5 and OMM1.5^{Cas9}) or 20% FBS (MP41, MP46, MP38 and MM28) and antibiotic/antimycotic solution. All cell lines were tested for mycoplasma and proved to be mycoplasma free using the MycoAlert™ PLUS Mycoplasma Detection Kit (Lonza).

Sphere Formation Assay

Cells were seeded in 96-well ultra-low attachment culture dishes (Corning) at 100 cells/well with the indicated concentration of 10 nM trametinib, 1 µM VS-4718 or both. Medium consisted of serum-free DMEM/F12 Glutamax supplemented bFGF (20 ng/ml), EGF (20 ng/ml), B-27 (1:50 dilution), and N2 supplement (1:100 dilution). Three weeks after seeding the number of spheres in each well and their sizes were assessed by bright-field microscopy and quantify using ImageJ (25).

Immunoblotting

UM cells were harvested at the indicated timepoints post-treatment with 10 nM trametinib and 1 µM VS-4718. Cell lysates were subjected to SDS/PAGE on 10% acrylamide gels and electroblotted to PVDF membranes. Blocking and primary and secondary antibody incubations of immunoblots were performed in Tris-buffered saline/Tween 20 [10 mM Tris (pH 7.4), 150 mM NaCl, and 0.1% Tween 20] supplemented with 5% (wt/vol) bovine serum albumin (BSA). The primary antibodies were used according to the manufacturers' instructions. HRP-conjugated donkey anti-rabbit and anti-mouse IgGs were used at a dilution of 1:5,000, and immunoreactive bands were detected using enhanced chemiluminescence. Full blots are shown in Supplementary Fig. S4

Flow cytometry

Apoptosis was determined by detecting phosphatidylserine exposure on cell plasma membranes using the fluorescent dye FITC Annexin V (BD Biosciences, San Diego, CA, USA) according to the manufacturer's protocol. Briefly, at the end of the treatment period, the cells (including floating cells) were harvested and washed twice with cold PBS. After resuspension in 100 µl 1X binding buffer, 5 µl FITC Annexin V was added and the cells were incubated for 15 min at room temperature in the dark. Finally, 400 µl 1X binding buffer was added to the cells and flow cytometric analysis was conducted. The experiments were conducted in triplicate.

Immunohistochemistry

For IHC, all tissue samples were processed and stained as previously described (26). Slides were scanned using a Zeiss Axioscan Z1 slide scanner equipped with a 20×/0.8 NA objective. All images analysis was performed using the QuPath software (27) to perform cell detection and quantify each marker as a percentage of cells stained.

Kinome-wide CRISPR screen

Generation and validation of stable Cas9 lines—For lentivirus-Cas9 production, 293T cells were plated in a poly-D-lysine-coated dish and, 16 hours later, transfected with 12 µg pLenti-CAS9-Blast, 8 µg psPAX2, 4 µg pCMV-VSV-G, using 32 µl TurboFect™ Transfection Reagent, and media was refreshed 6 hours post-transfection. 48 and 72h hours later, the virus-containing media was collected, filtered through a low protein binding filter unit (PVDF, 0.45µm, Sigma-Aldrich) and stored at 4°C up to 5 days or at –80°C.

For lentivirus-Cas9 infection, OMM 1.5 cells were plated in a 6-well plate and, 16 hours later, transduced using 1 ml virus-containing media + 1 ml complete media + 10 µg/ml polybrene. The plate was centrifuged for 15 min at 1200 rpm. The same process was repeated after 24 hours and 48 hours later the cells were selected with blasticidin (10 µg/ml).

Cas9 editing efficiency was measured by quantifying the editing frequency of the safe-harbor locus

AAVS1(TGCCTAACAGGAGGTGGGGGTTAGACCCAATATCAGGAGACTAGGAAGGAGGAGGCCTAAGGATGGGGCTTTTCTGTCACCAATCCTGTCCCTAGTGGCCCCACTGTGGGGTGGAGGGG) by next-generation sequencing using two separate sgRNA sequences (sgT1 and sgT2). Briefly, cells were transduced with either AAVS1 sgRNA and, after hygromycin-B selection, genomic DNA was isolated using the Qiagen DNeasy Blood and Tissue kit (Cat # 69504). The AAVS1 region was amplified, DNA gel-purified, and barcoded using the NEBNext Multiplex Oligos for Illumina kit (E7335) for multi-plex sequencing. Analysis of genome editing was performed using CRISPResso (28).

Pooled CRISPR library details and lentivirus preparation—In total, 3,052 unique sgRNAs targeting 763 human kinome genes for 4 guides per target were used for pooled CRISPR screens (Brunello Human Kinome CRISPR Knockout Library (29); Addgene Cat #75312). Lenti-X 293T cells were seeded in 15 cm plates and transfected the next day with: pooled CRISPR library DNA (21 ug), PAX2 (14 ug), and VSV-g (7 ug). Viral media was collected 48- and 72-hours post-transfection, pooled and concentrated using an ultracentrifugation protocol, and stored at –80°C in aliquots.

Infectious virus titer determination and transduction—Several aliquots of concentrated viral stock were used to infect separate 15 cm plates containing 4.8×10^6 OMM1.5-Cas9 cells that were seeded the day before. Non-infected plates for OMM 1.5 parental and Cas9 were also plated as antibiotic selection controls. 48 hours post-transduction, cells were treated with 2 ug/ml puromycin for 72 hours. After puromycin selection, complete cell death was achieved in uninfected control plates. The number of puromycin-resistant cells in each infected plate was counted using the Countess Automated Cell Counter to calculate the functional titer of the viral stock. Cells for screening were infected with a virus dose to achieve a MOI of 0.4–0.5 with sufficient cell numbers plated to obtain a screening depth of >1000 cells per sgRNA.

Screen set-up— 4.8×10^6 cells were seeded into 15 cm plates a day before CRISPR library transduction. 48 hours post-transduction, cells were subject to puromycin selection (2 µg/ml; 72 hours). Cells were amplified and seeded into 15 cm plates (4×10^6 cells per plate) divided

into two treatment arms: 3 replicate plates for either vehicle/DMSO or VS-4718 treatments. 4×10^6 cells from each individual plate were passaged into a new plate containing DMSO or $0.5 \mu\text{M}$ VS-4718 every 3–4 days for a total of 10 days of treatment. 5×10^6 cells were aliquoted from each plate at the beginning and end of the screen and stored at -80°C for sgRNA quantification.

sgRNA quantification—Genomic DNA was isolated using the Qiagen DNeasy Blood and Tissue kit (Cat # 69504) according to manufacturer's protocol. DNA concentration was measured using a Qubit assay (ThermoFisher Scientific; Cat #Q33226). Amplification of sgRNAs for next-generation sequencing was performed according to the Broad Institute's recommended protocol. NGS read counts were processed, aligned, and analyzed using PinAPL-Py (30).

Pre-screen evaluation of sgRNAs: sgRNAs were quantified immediately prior to the start of the screen to evaluate CRISPR library representation: sgRNA representation was 99.5% and gene representation was 100%.

Post-screen evaluation of sgRNAs: sgRNAs were quantified from each replicate plate at the end of the screen and analyzed to identify sgRNAs depleted in VS-4718 treated cells relative to DMSO treated cells.

Synergy determination

Dose-response curves and determination of IC50 values—Cells were seeded at a density of 5×10^3 to 1×10^4 cells/well in 96-well white plates. Eight different dilutions of each inhibitor were assayed in technical triplicates for 72 h in each experiment. Cell viability was measured with the Aquabluer Cell Viability Reagent on a Spark microplate reader (Tecan). Using the GraphPad Prism v8.2.0 software, the half-maximal inhibitor concentration values (GI_{50}) were determined from the curve using the nonlinear log (inhibitor) vs response-variable slope (three parameters) equation. GI_{50} values were only determined for compounds which inhibited growth by more than 50%.

Synergy determination with the Chou-Talalay method—The Chou-Talalay method (31) was used to determine possible synergistic effects of selected kinase inhibitor combinations. Briefly, cells were seeded at a density of 5×10^3 to 1×10^4 cells/well in 96-well white plates (CulturePlate; PerkinElmer Inc.). Cells were treated with either single inhibitors or combinations thereof using eight different dilutions of each inhibitor and in technical triplicates. Cell viability was measured, after 72 h treatment, with the Aquabluer Cell Viability Reagent on a Spark microplate reader (Tecan). Combination Index (CI) values showing either synergy (< 1) or antagonism (> 1) were calculated using the following equation:

$\text{CI} = (\text{D})_1 / (\text{D}_x)_1 + (\text{D})_2 / (\text{D}_x)_2$, where D_x equal the concentration of the tested substance used in the single treatment that was required to decrease the cell viability by x% and D equals the concentration of the tested substance 1 in combination with the concentration of the tested substance 2 that when combined decreased the cell number by x%.

Synergy determination with the Bliss delta score—The Bliss independence model(32) assumes a stochastic process in which two drugs elicit their effects independently, and the expected combination effect was calculated using the following equation: $IAB = IA + IB - IA \times IB$, where IA and IB are the single agent inhibition levels at fixed concentrations. If the experimentally measured effect of the drug combination was equal to, higher than or lower than the expected effect (IAB), the combination was considered to be additive (Bliss =0), synergistic (<0) or antagonistic (>0), respectively.

Human Xenograft Tumor Models

All animal studies were approved by the Institutional Animal Care and Use Committee (IACUC) of University of California, San Diego with protocol S15195. Female 4- to 6-week-old NOD.Cg-Prkdcscid Il2rgtm1Wjl/SzJ (SCID-NOD) mice were purchased from the UCSD in-house breeding program. Mice were injected subcutaneously in both flanks with either 2×10^6 or 2.5×10^6 92.1 or OMM1.3 cells, respectively. Mice were monitored twice weekly for tumor development. Tumor growth analysis was assessed as $LW^2/2$, where L and W represent length and width of the tumor. 5 mg/ml VS-4718 was prepared in 0.5% carboxymethyl cellulose (CMC) (Sigma-Aldrich; St. Louis, MO) and 0.1% Tween 80 (Sigma-Aldrich) in sterile water. 0.1 mg/ml trametinib was prepared in 5.2% polyethylene glycol 400 (Sigma-Aldrich) and 5.3% Tween 80 (Sigma-Aldrich) in DPBS. Mice were administered 50 mg/kg VS-4718 (Verastem Oncology; Needham, MA) twice daily by oral gavage and 1 mg/kg Trametinib once daily by intra-peritoneal injection (IP); control group was treated with each vehicle. Mice were euthanized at the indicated time points and tumors were isolated for sequencing, histologic, and immunohistochemical evaluation. Results of mice experiments were expressed as mean \pm SEM of a total of tumors analyzed.

Human Metastasis Tumor Model

Generation of stable GFP-Luc expressing 92.1—For lentivirus-GFP-Luc production, 293T cells were plated in a poly-D-lysine-coated 15cm dish and, 16 hours later, transfected with 30 μ g pHAGE PGK-GFP-IRES-LUC-W, 3 μ g VSV-G, 1.5 μ g Tat1b, 1.5 μ g Rev1b, 1.5 μ g Gag/Pol using 25.2 μ l p3000 buffer and 25.2 μ l of Lipofectamine 3000 transfection reagent, and media was refreshed 16 hours post-transfection. 48 and 72h hours later, the virus-containing media was collected, filtered through a low protein binding filter unit (PVDF, 0.45 μ m, Sigma-Aldrich) and stored at 4°C up to 5 days or at -80°C.

For lentivirus-Cas9 infection, 92.1 cells were plated in a 6-well plate and, 16 hours later, transduced using 1 ml virus-containing media + 1 ml complete media + 10 μ g/ml polybrene. The plate was centrifuged for 15 min at 1200 rpm. GFP expression was validated by fluorescent microscopy.

Splenic injection—Mice were injected with 1×10^6 92.1 GFP-Luc cells in the spleen, followed by removal of the spleen at 2 minutes post-injection. Tumor implantation by bioluminescence was assessed twice weekly by bioluminescence images captured using the In Vivo Imaging System (IVIS) Spectrum (Perkin Elmer, Santa Clara, CA). To this end, mice received an intraperitoneal injection of 200 mg/kg D-luciferin firefly potassium salt diluted in PBS 15 minutes before imaging (GoldBio, St Louis, MO). Total bioluminescence

was determined upon subtracting the background from the region of interest (ROI). Vehicle, trametinib, VS-4718 or trametinib/Vs-4718 were administered, starting 7 days post-surgery, with the described above dosing.

Statistical analysis.

GraphPad Prism version 8 for Windows (GraphPad Software, San Diego, CA) was used to perform data analyses, variation estimation and validation of test assumptions. Statistical analysis was performed using either a paired Student's *t* test or one-way ANOVA.

Results

Identification of conditionally lethal drug targets of FAK inhibition.

As the kinome has been the target of most drug discovery efforts, the identification of kinases whose activity are essential for UM survival in the context of FAK inhibition may facilitate the discovery of new co-targeting strategies. To perform an efficient loss-of-function screen, we used a two-vector CRISPR system, expressing a *Cas9* transgene in representative UM cells (OMM1.5, originally derived from metastasis, hereby called OMM1.5^{Cas9}), in which gene editing efficiency was validated (Fig. 1 and Supplementary Fig. S1A,B). OMM1.5^{Cas9} cells were infected with the *Human Kinome Brunello* pooled sgRNAs library (29), targeting 763 kinase genes and containing 3,052 unique sgRNAs along with 100 non-targeting controls, and subjected to puromycin selection. Surviving cells were treated with either selective FAKi (VS-4718) or vehicle for 10 days, followed by the collection of cellular DNA for gRNA analysis by NextGen sequencing (Fig. 1A). Depleted sgRNAs (dropouts) suggest that inhibition of their gene targets could sensitize cells to FAKi treatment. Our analysis of synthetic lethal interactors of FAKi revealed a significant enrichment of Gα_q-PLC and MAPK components of the classical Gα_q signaling cascade (Fig. 1B and Supplementary Fig. S1C). Using the pan-PKC inhibitor, Go-6983 (33), we found that inhibition of the canonical PLCβ-PKC-ERK pathway decreased UM cell viability and that this inhibitory effect was further enhanced by combining Go-6983 with VS-4718 (Fig. 1C). However, PKC inhibition with Go-6983 was less effective in reducing active ERK (pERK) than the MEKi trametinib (Fig. 1D). This is aligned with prior studies supporting only partial pERK reduction after prolonged PKC inhibition (34), which may explain its very modest clinical activity in mUM (35). Thus, we focused on direct MEK inhibitors for further studies. Interestingly, inhibition of FAK by VS-4718 led to a gradual increase in pERK levels (Fig. 1D, E) and UM cells stably expressing the constitutive active MEK mutant (MEK-S218/222D, MEK-DD) were more resistant to VS-4718 (Fig. 1F), suggesting a possible compensatory mechanism that may contribute to drug resistance.

Synergistic antiproliferative effect of MEK and FAK co-targeting in UM cells.

Co-targeting of MEK and FAK inhibits both canonical and non-canonical Gα_q signaling pathways, and may provide a suitable drug combination for clinical application in mUM. We used the effect-based Bliss model and the dose-effect-based Chou-Talalay combination index (CI) to assess synergistic, additive, or antagonistic drug interaction in a given combination. We confirmed that trametinib (MEKi) and VS-4718 (FAKi) decrease UM cell growth as single agents. CI and Bliss scores both converged to demonstrate a synergistic

interaction between trametinib and VS-4718 (Fig. 2A). Synergistic antiproliferative effects were observed using multiple clinically relevant MEKi (trametinib, cobimetinib, and selumetinib) as well as the second-generation RAF/MEK inhibitor VS-6766, combined with two different FAKi (VS-4718 and the clinically relevant defactinib). All combinations showed remarkable synergistic activity at relevant doses supporting a general drug-drug class pharmacodynamics interaction (Fig. 2B). Using a panel of *GNAQ* mutant UM cells lines with different expression levels of the metastasis suppressor protein BAP1 (12), we confirmed similar synergistic profiles in BAP1 wild-type or null cells (Fig. 2C & D) suggesting that this combination may also be active in mUM.

MEKi/FAKi combination increases UM apoptotic cell death and reduces melanosphere formation.

Several FAKi inhibitors have been evaluated in the clinic and demonstrated primarily a cytostatic effect as single agents (23). Given the high synergistic activity of MEKi and FAKi combination *in vitro*, we next evaluated the ability of trametinib and VS-4718 to induce cell death. As shown in Fig. 3A, FACS analysis suggests that trametinib induces an increase in the fraction of apoptotic cells [Annexin V⁺], while VS-4718 has no apoptotic activity as a single agent. Nonetheless, when both drugs were combined, at a ratio of 1:100 based on our isobologram analysis (Supplementary Fig. S2), the apoptotic response was significantly increased.

One of the most common signaling cascades involved in apoptosis is the activation of a highly specialized family of cysteinyl-aspartate proteases (caspases) (36). Caspase-mediated cell death is achieved through the cleavage of multiple key proteins that are essential to cell survival. PARP-1 is one of the substrates of caspases and a well-established marker for apoptosis (37). Aligned with our prior results, cleaved-PARP levels were strongly increased by the combination of FAKi and MEKi (Fig. 3B), while single agents were less active. We then evaluated the ability of VS-4718 and trametinib to induce apoptosis in *BAP1* null mUM cells. FAKi alone showed a minimal apoptotic response, and the combination of VS-4718 and trametinib resulted in increased cleaved-PARP compared to trametinib alone (Fig. 3B).

Cancer initiating cells are believed to play a central role in drug resistance and metastasis (38). A typical feature of these cells is their ability to form 3D-tumorspheres when cultured in suspension in a stem-cell medium. Interestingly, we found that the metastatic-derived OMM1.3 cells, but not 92.1 cells originating from a primary tumor, were able to form melanospheres. Using this model, we found that both VS-4718 and trametinib significantly reduced melanosphere formation as single agents when compared to vehicle (Fig. 3C) and this effect was further enhanced when both drugs were combined. Our results demonstrate that MEKi/FAKi combination acts by inhibiting cancer cell proliferation, including cancer initiating cells, and enhancing apoptotic cell death. This suggests that this multimodal targeted therapy could be effective on primary and metastatic UM.

Potent antitumoral and cytotoxic effects of MEKi/FAKi combination in UM xenograft and liver metastasis models.

We used UM xenograft models to evaluate the anticancer activity of the MEKi/FAKi combination *in vivo*. Tumor-bearing mice were randomly divided into four groups: control, trametinib, VS-4718 and trametinib/Vs-4718. In this regard, whereas VS-4718 has better pharmacokinetics than defactinib in mice, the latest is better able to inhibit tumor FAK activity with tolerability in cancer patients (39). Therefore, VS-4718 is used as a surrogate FAKi in murine pre-clinical models. As shown in Fig. 4A and B, in the 92.1 xenograft model, although trametinib and VS-4718 effectively induced tumor stasis as single agents, only the combination of trametinib with VS-4718 was able to induce tumor regression. No significant difference in body weight was observed between the control and any of the treated groups (Fig. 4C), suggesting that all treatments were well tolerated by the mice. These results were further confirmed using another UM cellular system harboring *GNAQ* mutations, OMM1.3, albeit in this case MEKi alone was quite potent (see Supplementary Fig. S3). We extended these studies in allograft experiments using the recently developed *GNAQ*-driven B2905 syngeneic mouse melanoma model (40). These cells are highly sensitive to the Gαq inhibitor FR900359, as we reported in a large panel UM cells (41), and to the combined inhibition of MEK and FAK (Supplementary Fig. 4). These cells exhibit limited responses to FAKi *in vivo*, likely reflecting the complexity of genetic alterations of these cells (40). MEKi limited tumor growth, but prolonged inhibition resulted in acquisition of resistance and rapid tumor regrowth, thereby compromising animal survival. Remarkably, the combination therapy was quite effective in promoting tumor regression, and no resistance was observed for the prolonged duration of the mouse experiments (>40 days).

We next evaluated MEK/ERK and FAK/YAP pathway activities in the UM xenograft tumor specimens by immunohistochemistry (Fig. 4D and E). As expected, pERK1/2 was strongly decreased only in the trametinib and trametinib/Vs-4718 treated groups. We monitored YAP nuclear exclusion as a readout for YAP inactivation upon FAKi treatment (18), and observed decreased nuclear YAP in the VS-4718 and trametinib/Vs-4718 treatment groups exclusively. The percentage of proliferating BrdU positive cells was significantly decreased in all treated groups, with the VS-4718 and trametinib/Vs-4718 treatment groups showing the most significant inhibition. Using cleaved-Caspase3 (cl-Casp3) as a marker for apoptosis, we detected a significant increase of cl-Casp3 in the trametinib/Vs-4718 group compared to the single agent groups, suggesting that the MEKi/FAKi combination induces a switch from cytostatic to cytotoxic activity, consistent with the increased cl-PARP levels observed *in vitro*.

To assess the efficacy of the MEKi/FAKi combination treatment on a murine system that better represents the advanced disease, we developed a preclinical liver metastasis model. Luciferase/GFP expressing 92.1 cells (92.1-Luc) were injected into the spleen and allowed hematogenous dissemination followed by complete splenic resection (Fig. 5A). Mice were sacrificed 8-weeks post-injection and necropsies showed macroscopic hepatic metastases with no sign of dissemination to any other organs (Fig. 5B), confirming the strong liver tropism of UM cells. Remarkably, when tumor bearing mice were treated with trametinib, VS-4718 or trametinib/Vs-4718, the metastatic burden was reduced by MEKi and FAKi

treatment and their combination (Fig. 5C, D). FAKi alone was cytostatic, whereas MEKi and MEKi/FAKi combination induced tumor regression. However, detailed analysis showed that the residual disease in the trametinib alone treated group represented 24.2 ± 0.6 % of the initial tumor compared to 7.5 ± 0.2 % in the combination treated group ($p=0.0002$), achieving none versus 40% complete responses, respectively (Fig. 5E). Resistance to trametinib in patients has led to the failure of this agent in UM clinical trial. To further investigate resistance to trametinib as a single agent or as part as a combination with FAKi in our metastatic model, we adjusted the dose of trametinib to reach maximum plasma levels similar to those achieved in humans at the recommended therapeutic dose, 2 mg/day, which is approximately 22 ng/ml upon repeated dosing (42). Mice treated with 0.1 mg/kg trametinib, achieving approximately 20 ng/ml (43) become resistant to treatment with a progressive increase of metastatic burden, whereas tumor burden remains barely undetectable in mice treated with the combination. Taken together, these findings indicate that treatment with trametinib combined with VS-4718 inhibited UM tumor growth in subcutaneous and liver metastasis models.

Discussion

The limited responses of UM to immunotherapies (40) and the lack of an FDA-approved standard of care for mUM patients poses an urgent unmet medical need that necessitates the development of new targeted treatment options. Most clinical studies in UM have focused on canonical kinases activated downstream from G_q/G_{11} with MEK as the major therapeutic target (44). MAPK/ERK pathway inhibitors have been proven to be effective treatments in various cancer types, but their effectiveness is often short-lived, with resistance developing often after the start of treatment (45–48). UM was no exception and despite encouraging results in an initial phase-II trial, with the MEK inhibitor selumetinib (AZD6244) showing an improved progression-free survival (PFS) compared to dacarbazine or temozolomide (15). The subsequent phase III double-blind trial failed to show improvement in PFS and overall survival (OS) with selumetinib + dacarbazine compared to dacarbazine alone (16). Here, we found that adaptive activation of MEK-ERK represents a compensatory resistance mechanism to FAK inhibition, and that in turn concomitant targeting FAK and MEK-ERK converts the cytostatic effects of FAKi into cytotoxic, resulting in UM cell death. These findings suggest that dual inhibition of MEK and FAK may represent a promising therapeutic option for advanced and metastatic UM patients.

Our previous studies revealed that G_{α_q} triggers the activation of the FAK/Hippo-YAP pathway, which represents a major driver in UM progression (18,19). Given that, we aimed at identifying synthetic lethal interactors that could potentiate FAKi efficiency and simultaneously reduce the risk of developing drug resistance (24), which is often observed in single-drug therapy. Our kinome-wide CRISPR/Cas9 screen unveiled potent synthetic lethality between FAKi and the MEK/ERK pathway, uncovering the therapeutic potential of a co-targeting strategy. This idea is reinforced by a recent study demonstrating that treatment of UM cells with the MEKi trametinib increased YAP activity and cell proliferation (49), and our findings that FAK/Hippo-YAP pathway inhibition leads to a gradual increase of pERK. These results suggest that both G_{α_q} -activated pathways, FAK/Hippo-YAP and MEK/ERK, can be part of compensatory feed-back processes, and that horizontal inhibition of

FAK and MEK might be necessary in order to achieve tumor regression, and prevent treatment resistance and tumor relapse (Fig. 6). In addition, concomitant inhibition of FAK and MEK triggers the activation of apoptotic cell death programs, whose full elucidation may facilitate the identification of mechanistic biomarkers predicting therapeutic response. Of interest, most UM and mUM models used in our studies are driven by the *GNAQ* oncogene. However, based on its high similarity with the *GNA11* oncogene, our findings are also expected to be applicable to *GNA11* mutant UM lesions (6,7), which should be nonetheless tested experimentally.

Several potent FAKis have been tested in the clinic in multiple cancer types, showing target inhibition and manageable toxicity profiles. Indeed, based on our findings, a Phase 1 clinical trial using IN10018 as a potent FAKi has been recently initiated in mUM (NCT04109456). The use of FAKis as single agents may act primarily as cytostatic, suggesting the possibility of identifying suitable combination strategies to reach a greater clinical efficacy. Our study supports the clinical potential of co-targeting FAK and its sensitizing pathway, MEK/ERK, as a precision therapy approach in *GNAQ*-driven UM, achieving tumor regression. In this regard, the FAKi defactinib has been recently combined with a RAF/MEKi (VS-6766) in multiple cancer types (NCT03875820). Through use of an intermittent dosing schedule, the defactinib/VS-6766 combination has shown a manageable safety profile with initial clinical activity in early clinical trial results to date (50). Altogether, we believe our study presents a strong potential for clinical translation as a multimodal signal-transduction-based precision therapy for mUM.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

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Statement of significance

Most metastatic uveal melanoma (mUM) patients are refractory to chemotherapies and immune checkpoint blockade, leading to patient death within a year of diagnosis. To date, there are no effective treatment options for mUM, highlighting an urgent need for novel therapeutic strategies. Uveal melanoma (UM) is characterized by gain-of-function mutations in *GNAQ/GNA11*, encoding Gαq proteins. Our recent studies identified focal adhesion kinase (FAK) as an integral component of the oncogenic Gαq signaling circuitry in UM, and the clinical benefits of targeting FAK in mUM are already under current investigation. However, single-agent targeted therapies often activate adaptive mechanisms resulting in drug resistance and treatment failure. Taking advantage of the unique genetic landscape of UM and the use of unbiased genetic screens, our studies reveal that horizontal inhibition of FAK and the adaptive activation of MEK-ERK results in UM cell death and tumor regression, thereby providing a novel multimodal precision therapy for mUM.

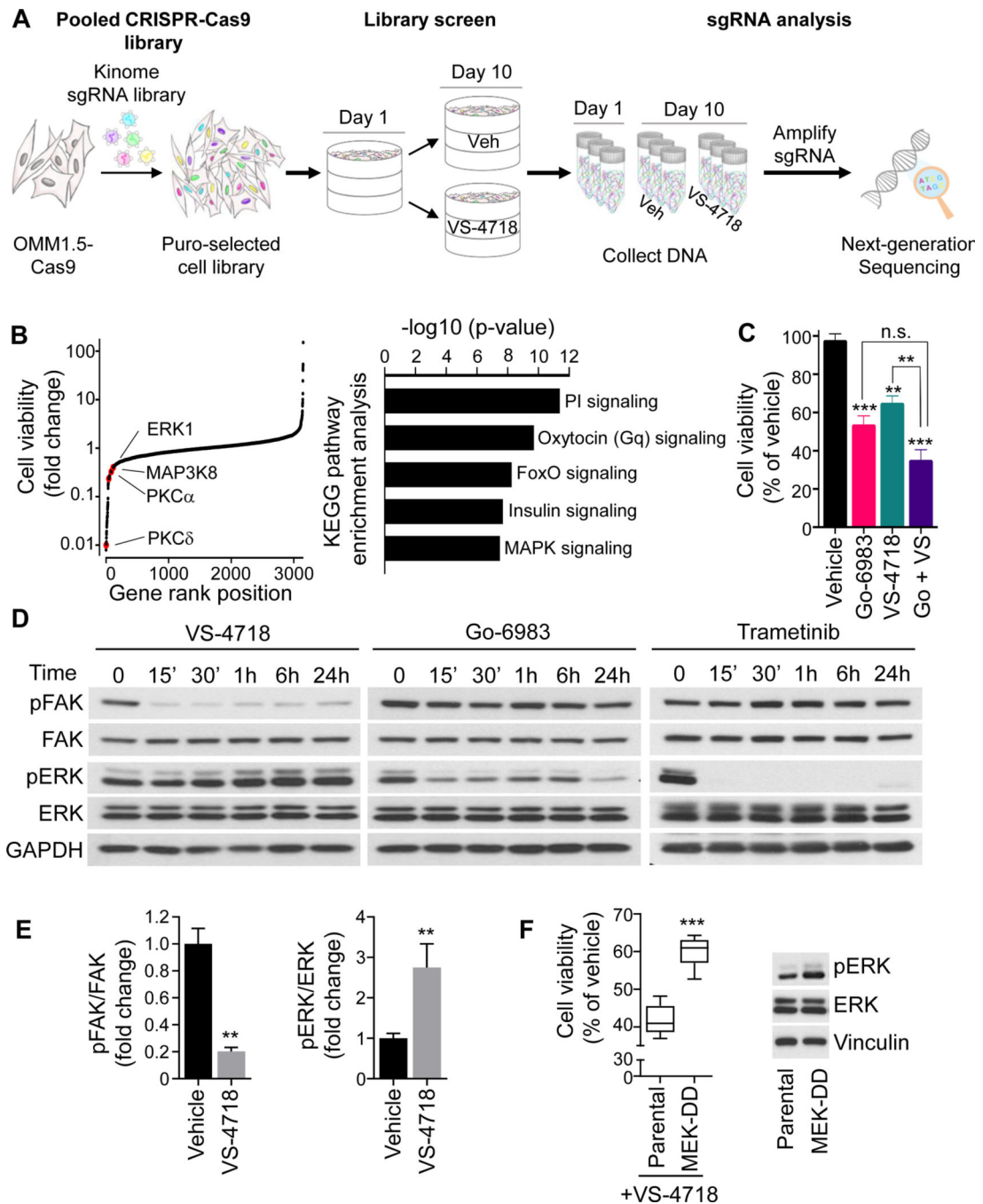


Figure 1. Kinome-wide CRISPR screen for synthetic lethal interactors of FAKi.

(A) OMM 1.5 cells expressing Cas9 were infected with the Brunello Human Kinome CRISPR sgRNA KO library at a MOI of 0.3. After selection, cells were treated with vehicle or 0.5 μ M VS-4718 (FAKi) for 10 days. (B) *Left*, Cell viability represented as fold change in FAKi-treated cells compared to control. Highlighted significant hits represent synthetic lethal genes with FAKi treatment. *Right*, KEGG pathways analysis for the top depleted sgRNAs (n=200). (C) 92.1 cell viability after 72h treatment with vehicle, 1 μ M Go-6983 (PKCi), 1 μ M VS-4718 or combination of both. (D) Time-course analysis of FAK and ERK

phosphorylation in 92.1 cells treated with VS-4718 (1 μ M), Go-6983 (1 μ M) or trametinib (MEKi, 10nM). **(E)** Quantification of pFAK/FAK and pERK/ERK ratios in 92.1 cells treated with 1 μ M VS-4718 or vehicle for 1h. **(F)** *Left*, Cell viability after 72h treatment with VS-4718 (1 μ M) in 92.1 cells expressing or not MEK-DD (S218/222D). *Right*, Immunoblot showing pERK levels in 92.1 cells expressing or not MEK-DD (S218/222D). **(C, E and F)** Data shown represent the mean \pm SEM of three independent experiments. ***p<0.001; **p<0.01; n.s. not significant.

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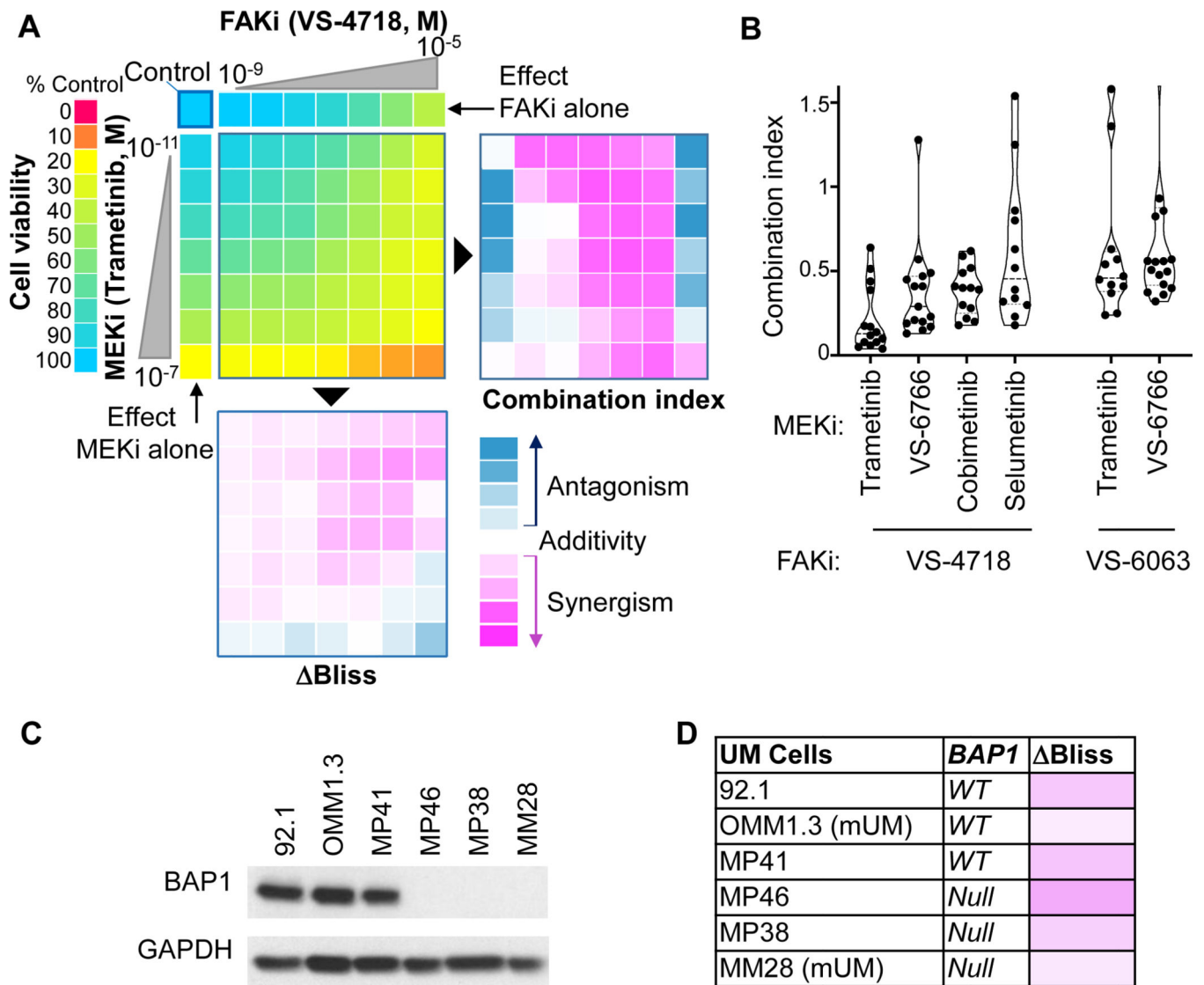


Figure 2. Synergy between FAKi and MEKi in UM and mUM cells.

(A) *Left*, 92.1 cell viability 72h after treatment. *Right*, Combination Index values (CI) determined using the Chou-Talalay method (CI<1 synergism, CI=1 additivity, CI>1 antagonism, scale from -2 to +2). *Bottom*, Bliss scores (score<0 synergism, score=0 additivity, score>0 antagonism, scale from -1 to +1). (B) CI at relevant doses (viability=50±5%) using various combinations of FAKi/MEKi. (C) Immunoblot depicting BAP1 levels in UM and mUM patient-derived cells. (D) Delta score (Bliss), assessing synergism between MEKi (trametinib, 10nM) and FAKi (VS-4718, 1µM) in a panel of UM and mUM cells with distinct BAP1 status.

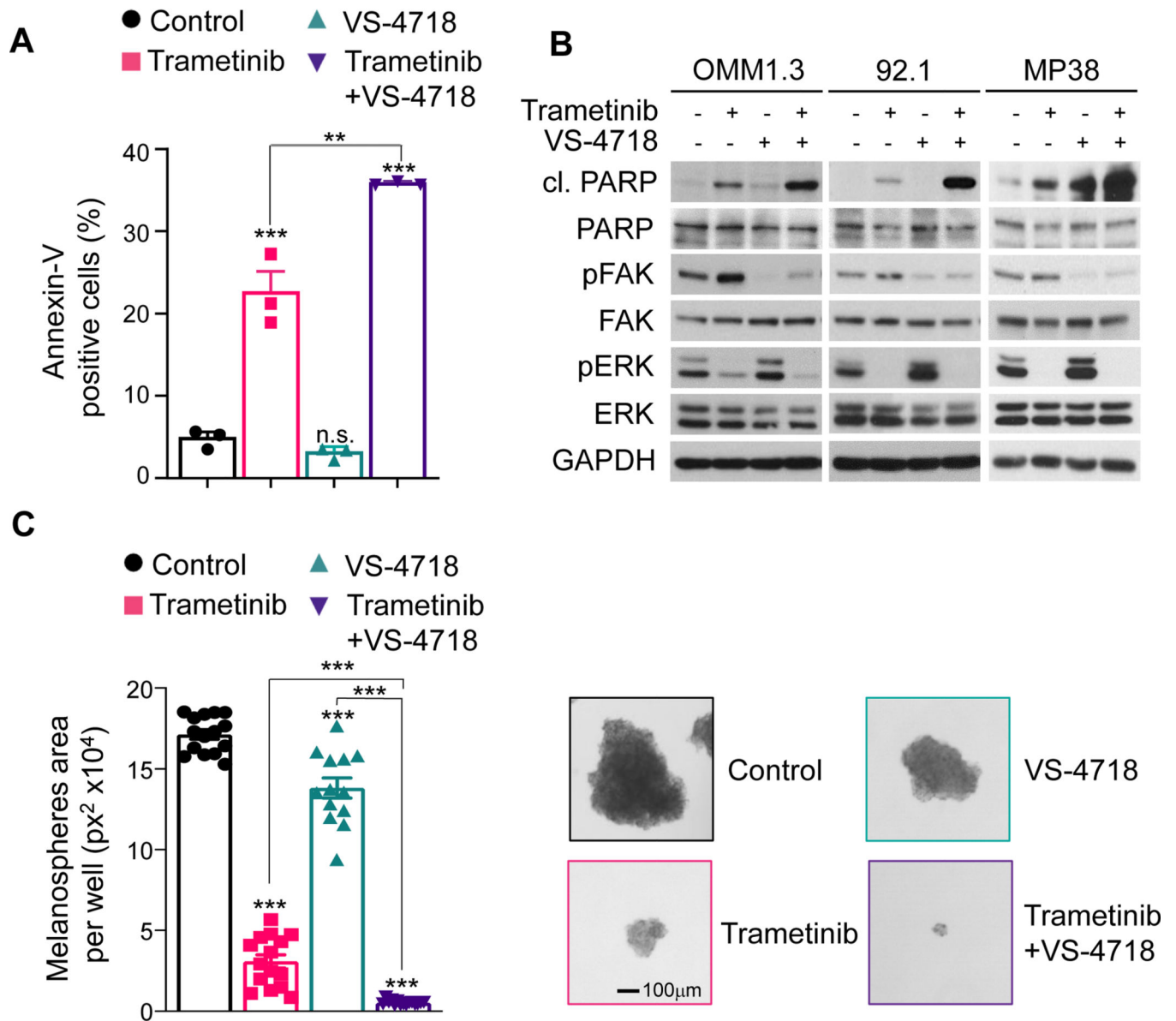


Figure 3. MEKi/FAKi combination induces apoptosis and reduces UM melanosphere formation. (A) FACS analysis of cells positive for Annexin V was used to assess the apoptotic response to trametinib (10 nM), VS-4718 (1 µM), and their combination after 24 h of treatment. (B) Immunoblot showing cleaved-PARP, pFAK (pY-397) and pERK levels upon treatment with vehicle, trametinib (10 nM), VS-4718 (1 µM) or trametinib+VS-4718 for 48 hours in UM cells. (C) *Left*, OMM1.3 melanospheres formation after treatment with vehicle (control), trametinib (10 nM), VS-4718 (1 µM) or trametinib+VS-4718 for 3 weeks. *Right*, Representative spheres (A and C) Data shown represent the mean ± SEM of three independent experiments. ***p<0.001; **p< 0.01; n.s. not significant.

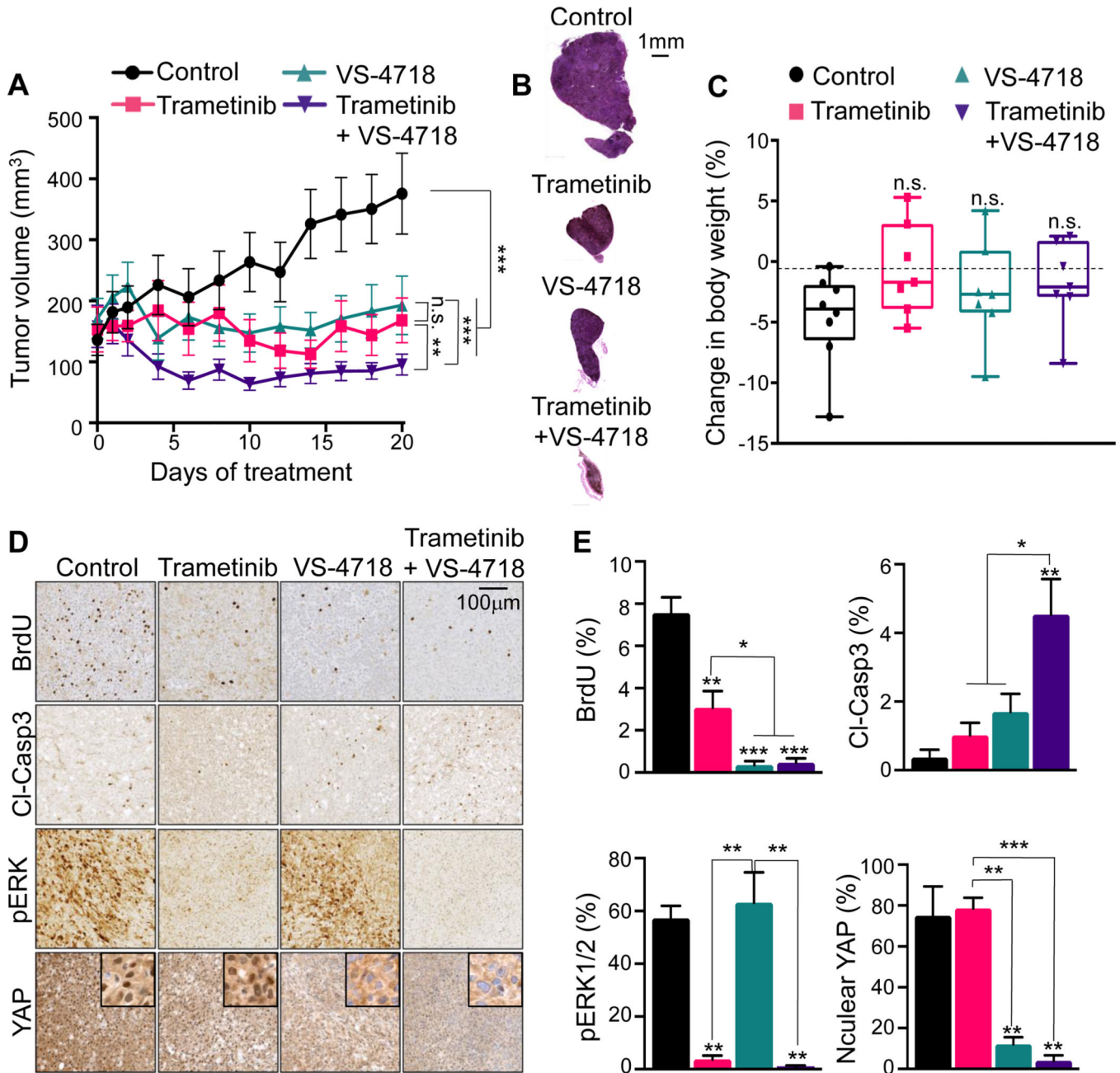


Figure 4. MEKi/FAKi combination UM growth in *in vivo* xenograft mouse models.

(A) Changes in 92.1 xenograft tumor volume in mice treated with vehicle (Control), trametinib 1 mg/kg, VS-4718 50 mg/kg or trametinib+VS-4718. (B) H&E staining of representative xenograft tumor sections after 20 days of treatment. (C) Difference in mice body weight between day 0 and day 20 of the indicated treatment in 92.1 xenografts mice. Box and whiskers plot with minimum and maximum whiskers (7 mice/group). (D) 92.1 tumor bearing mice were treated with vehicle (Control), trametinib 1 mg/kg, VS-4718 50 mg/kg or trametinib+VS-4718 for 20 days. Representative IHC staining tumor sections for BrdU, cleaved-Caspase3 (cl-Casp3), pERK and YAP. Scale bar is 100 μ m and insets are

50 μm wide. **(E)** Quantification of the IHC stained tumor sections. **(A and E)** Data are mean \pm SEM (7 mice/group). * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$; n.s. not significant.

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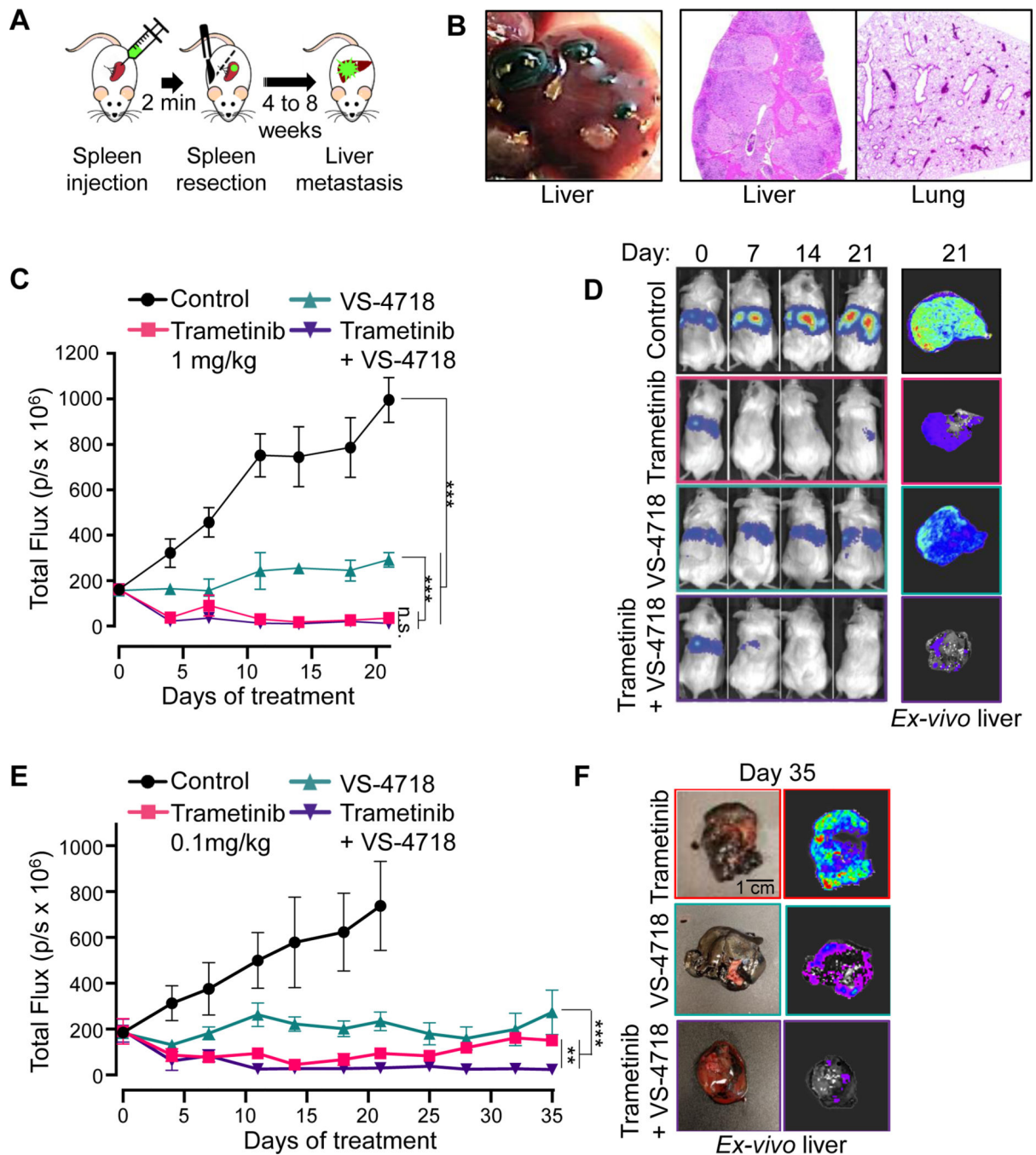


Figure 5. MEKi/FAKi combination reduces UM cells growth in an *in vivo* liver metastasis model. (A) Schematic of the hematogenous dissemination model for UM liver metastasis using 92.1 GFP-Luc cells. (B) *Left*, Macroscopic view of liver metastasis 8 weeks post-splenic injection. *Right*, H&E staining of liver and lung. (C) Hepatic tumor burden tracked by IVIS imaging after injection of 92.1 UM cells in SCID/NOD mice treated with vehicle (Control), trametinib 1 mg/kg, VS-4718 50 mg/kg or both. Data are mean±SEM (6 mice/group). ***p<0.001; n.s. not significant. (D) Representative mice treated with vehicle (Control), trametinib 1 mg/kg, VS-4718 50 mg/kg or both, at the indicated days of treatment, and

representative *ex-vivo* imaging of the liver obtained at day 21. **(E)** Hepatic tumor burden tracked by IVIS imaging after injection of 92.1 UM cells in SCID/NOD mice treated with vehicle (Control), trametinib 0.1 mg/kg, VS-4718 50 mg/kg or both. Data are mean±SEM (5 mice/group). ***p<0.001; **p<0.01. **(F)** Representative *ex-vivo* imaging of the liver from mice treated for 35 days with trametinib 0.1 mg/kg, VS-4718 50 mg/kg or both.

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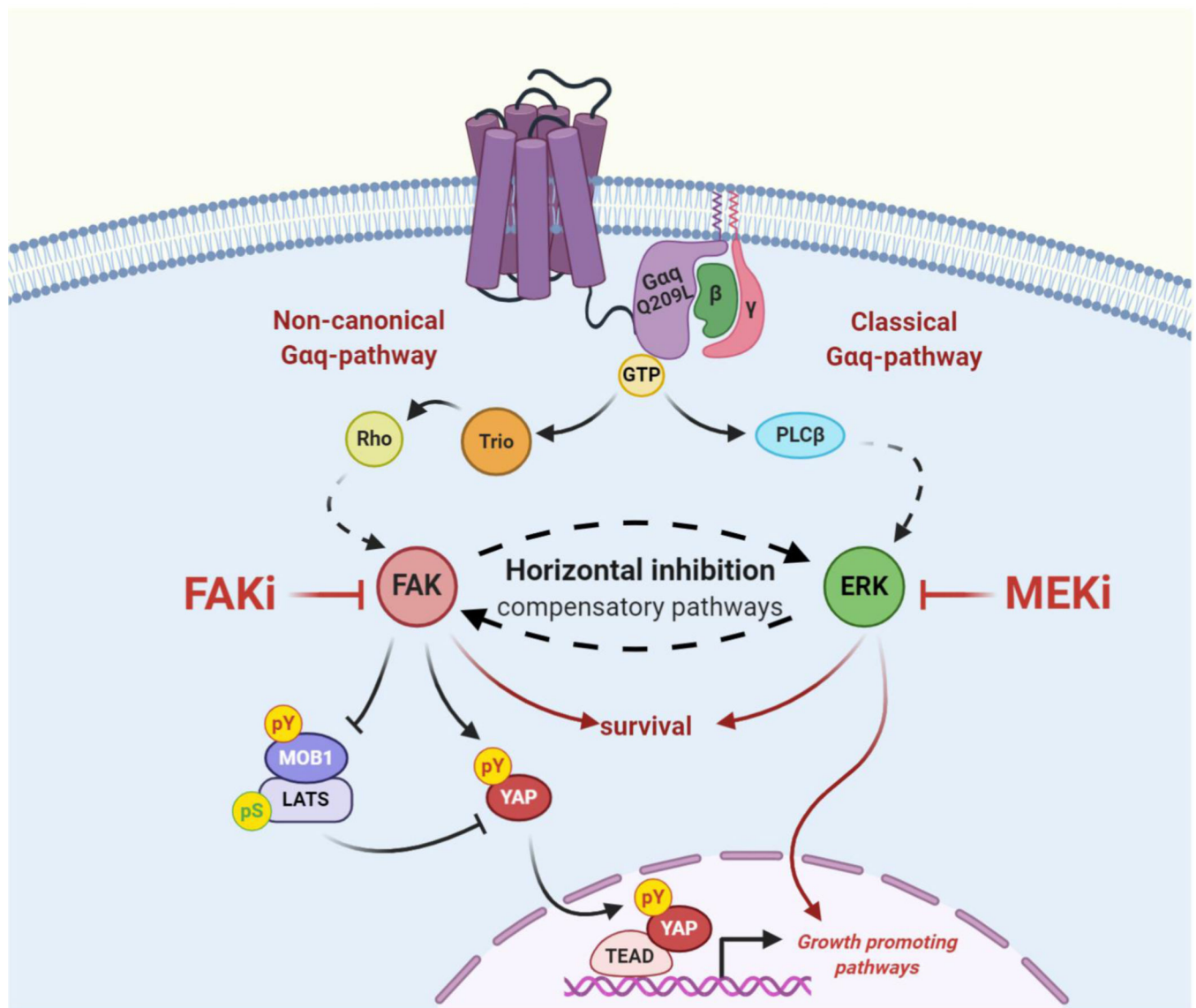


Figure 6. Horizontal inhibition of compensatory pathways in UM using MEKi/FAKi combination.

The cartoon depicts the proposed pathways by which active *GNAQ* mutant controls cell proliferation in UM cells. Horizontal inhibition of FAK and MEK likely acts by disabling growth promoting pathways regulated by YAP while concomitantly targeting parallel converging core survival mechanisms, thereby resulting in mUM regression.