

# Combined MEK and STAT3 inhibition reprograms the tumor microenvironment to overcome immunotherapy resistance in pancreatic cancer

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## ABSTRACT

Pancreatic ductal adenocarcinoma (PDAC) is characterized by immune exclusion, stromal desmoplasia, and resistance to immune checkpoint inhibition (ICI). We have previously demonstrated that reciprocally activated RAS/RAF/MEK/ERK and JAK/STAT3 pathways mediate therapeutic resistance, while combined MEK and STAT3 inhibition (MEKi/STAT3i) overcomes this resistance in PDAC. Herein, we show that combined MEKi/STAT3i alters stromal architecture and remodels the tumor-infiltrating innate and adaptive immune compartments by downregulating immunosuppressive myeloid populations and promoting T-cell enrichment and activation in *Ptf1a<sup>cre/+</sup>;LSL-Kras<sup>G12D/+</sup>;Tgfbr2<sup>flox/flox</sup>* (PKT) mice. MEKi/STAT3i-mediated reprogramming of the PDAC tumor microenvironment primes tumor-infiltrating CD8<sup>+</sup> T-cells for PD-1 blockade to augment their activation and functional cytotoxicity. As such, the addition of MEKi/STAT3i to PD-1 blockade dramatically inhibits tumor growth and prolongs survival, overcoming resistance to ICI, in PKT mice. Importantly, treatment of a patient with chemotherapy-refractory metastatic PDAC with MEKi (Trametinib), STAT3i (Ruxolitinib), and PD-1 inhibitor (Nivolumab) was not only well-tolerated but also yielded significant clinical benefit.

**Significance:** These data uncover a novel paradigm in which combined MEKi/STAT3i reprograms the stromal and immune microenvironment to overcome immunotherapy resistance in PDAC. The clinical efficacy of combined MEKi, STAT3i, and anti-PD1 treatment in chemotherapy-refractory PDAC provides encouraging signals for its translatability, and is being currently pursued in a clinical trial.

## INTRODUCTION

Pancreatic ductal adenocarcinoma (PDAC) remains a major therapeutic challenge due to its innate and acquired chemoresistance. The major contributors to therapeutic resistance that have been difficult to overcome in PDAC are mutations in the *KRAS* oncogene (1), the presence of a dense desmoplastic stroma that acts as a barrier to drug delivery and effector immune cell infiltration (2), and the immunosuppressive tumor microenvironment (TME) that renders the tumor resistant to immune checkpoint inhibition (ICI) immunotherapy (3).

Despite unprecedented success in other solid tumors (4), clinical trials utilizing single or dual ICI have largely been unsuccessful in PDAC patients (5,6). The failure of ICI in PDAC has been attributed not only to the fibroinflammatory desmoplastic stroma in PDAC that promotes T-cell exclusion, but also to the establishment of an immunosuppressive ecosystem in the TME. Cellular elements of this ecosystem comprise innate immune populations such as tumor-associated macrophages (TAM) and myeloid-derived suppressor cells (MDSC), which induce T-cell suppression through diverse mechanisms (7). A critical unmet need in the field of PDAC immunotherapy, therefore, is the discovery of novel therapeutic strategies not only to mitigate these resistance mechanisms but also to overcome ICI resistance in PDAC.

*KRAS* mutations are not only the predominant oncogenic driver in over 90% of PDAC, but also promote an inflammatory program that establishes immune privilege in the TME (1,8). Since targeting *KRAS* has remained an elusive goal, our approach has been to target downstream effectors of Ras through MEK inhibition (MEKi). However, clinical trials of MEKi have been unsuccessful in PDAC (9), likely due to the activation of resistance pathways. We have shown that MEK inhibition (MEKi) results in reciprocal activation of STAT3 signaling, which confers therapeutic resistance and continued PDAC cell growth. Combined inhibition of JAK/STAT3 (STAT3i) and MEKi overcomes this resistance mediated by parallel feedback loop activation, remodels the tumor stroma, attenuates tumor growth in patient-derived xenograft (PDX) models, and improves survival in the aggressive PDAC genetically engineered

*Ptf1a<sup>cre/+</sup>;LSL-Kras<sup>G12D/+</sup>;Tgfbr2<sup>flox/flox</sup>* (PKT) mouse model (10,11).

In the present study, we identify a novel paradigm in which combined MEKi/STAT3i remodels stromal fibrosis to enhance CD4<sup>+</sup> and CD8<sup>+</sup> T-cell recruitment while attenuating immunosuppressive TAMs and MDSCs in the PDAC TME. The addition of MEKi/STAT3i to PD-1 blockade overcomes ICI resistance by significantly augmenting anti-tumor responses and dramatically improving survival in PKT mice, compared with vehicle treatment or anti-PD-1 monotherapy. The addition of MEKi/STAT3i to PD-1 blockade not only augments the recruitment of activated and memory T-cell populations, but also improves their degranulating capacity and functional cytotoxicity. Importantly, we demonstrate the clinical efficacy and tolerability of combined Trametinib (MEKi), Ruxolitinib (STAT3i), and Nivolumab (anti-PD1) treatment in a patient with chemotherapy-refractory PDAC, providing strong evidence of the translatability of this therapeutic strategy in PDAC patients.

## RESULTS

### **Combined MEKi/STAT3i remodels stromal fibrosis and promotes an anti-tumor immune microenvironment.**

We have previously shown that combined MEKi/STAT3i, decreases tumor burden and improves survival in an aggressive GEMM of PDAC (10). We extend these results here to show that combined MEKi/STAT3i significantly remodels the tumor stroma as evidenced by decreased trichrome blue, smooth muscle actin ( $\alpha$ -SMA), and Sirius Red staining of pancreatic tumor sections in PKT mice (**Fig. 1A**). Moreover, MEKi/STAT3i significantly reduces intratumoral mucin content (Alcian Blue staining) and cellular proliferation (Ki67), as well as increases microvessel density (CD31 staining) compared with vehicle treatment (**Supplementary Fig. S1A**). To assess the immune repercussions of stromal remodeling with MEKi/STAT3i compared with vehicle treatment, flow cytometric analysis of PKT tumors (n=7-10 mice per arm) revealed a significant decrease in CD11b<sup>+</sup> myeloid cells and F4/80<sup>+</sup>

macrophages, as well as a concomitant increase in CD4<sup>+</sup> and CD8<sup>+</sup> tumor-infiltrating T-cell populations (**Fig. 1B**). These changes were confirmed with immunofluorescence (**Supplementary Fig. S1B**).

To comprehensively assess global immune changes associated with MEKi/STAT3i in PKT mice, we performed multiparametric immune profiling of pooled pancreatic tumor samples utilizing time-of-flight mass cytometry (CyTOF). Compared with vehicle treatment, MEKi/STAT3i, resulted in broad-based changes in the innate (CD11b<sup>+</sup>), adaptive (CD4<sup>+</sup> and CD8<sup>+</sup>), and humoral (CD19<sup>+</sup>) immune compartments (**Fig. 1C**). Specifically, compared with vehicle treatment, MEKi/STAT3i treatment resulted in a dramatic decrease in CD11b<sup>+</sup> myeloid cells, F4/80<sup>+</sup> macrophages, M2-like macrophages (F4/80<sup>+</sup>CD206<sup>+</sup>), and MDSCs (CD11b<sup>+</sup>F4/80<sup>-</sup>Ly6G<sup>+</sup>/Ly6C<sup>+</sup>) (**Fig. 1D**), as well as a significant increase in proportions of total (CD3<sup>+</sup>), T-helper (CD4<sup>+</sup>), cytotoxic (CD8<sup>+</sup>), and activated (CD69<sup>+</sup>) T-cell populations (**Fig. 1E**). MEKi/STAT3i also improved effector (CD44<sup>+</sup>) and central (CD62L<sup>+</sup>) memory T-cell markers, as well as TCR $\beta$  expression, in the PDAC TME compared with vehicle treatment (**Supplementary Fig. S1C**). Taken together, these data suggest that combined MEKi/STAT3i reprograms the fibroinflammatory tumor stroma to promote an anti-tumor immune microenvironment by decreasing immunosuppressive myeloid cell populations and augmenting T-cell recruitment and activation.

### **Anti-tumor effects of MEKi/STAT3i are T-cell dependent**

We have previously shown that MEKi/STAT3i results in improved survival compared with vehicle treatment or MEKi or STAT3i monotherapy in PKT mice (10). Given the aforementioned immunologic changes with MEKi/STAT3i, we sought to determine if the anti-tumor effects of this combination therapy are T-cell dependent by performing CD4<sup>+</sup> and CD8<sup>+</sup> T cell depletion in this model concurrently with MEKi/STAT3i (**Supplementary Fig. S2**). Non-T-cell depleted PKT mice treated with MEKi/STAT3i showed a significant decrease in tumor weight compared with vehicle

treatment and demonstrated 100% survival up to 80 days of treatment (**Fig. 1F&G**). In contrast, T-cell depleted mice treated with MEKi/STAT3i showed no significant decrease in tumor weight compared with vehicle-treated mice (**Fig. 1F**), despite confirmation of pERK and pSTAT3 inhibition in tumor lysates (**Fig. 1H**). Furthermore, the median survival of the T-cell depleted MEKi/STAT3i mice was significantly diminished compared with MEKi/STAT3i treatment without T-cell depletion (median 60.5 vs. not reached [NR] days, respectively; log-rank  $p=0.0001$ ; **Fig. 1G**). These results show that the decreased tumor burden and improved survival with MEKi/STAT3i in PDAC are T-cell dependent.

### **MEKi/STAT3i-mediated alterations in the PD-1/PD-L1 axis support the addition of immune checkpoint inhibition in PDAC**

To further explore the functional status of tumor-infiltrating CD8<sup>+</sup> T-cells in MEKi/STAT3i-treated versus vehicle-treated mice, we examined CD8<sup>+</sup> T-cell antigen experience (PD1<sup>+</sup>) and degranulating activity (CD107a<sup>+</sup>) in these cells. Flow cytometric (**Figure 1I**) and CyTOF analysis (**Supplementary Fig. S1B**) revealed a significant upregulation of PD1<sup>+</sup> tumor-infiltrating CD8<sup>+</sup> T-cells in mouse tumors treated with MEKi/STAT3i compared with vehicle treatment. Despite a significant increase in antigen-experienced degranulating CD8<sup>+</sup>PD1<sup>+</sup>CD107a<sup>+</sup> T-cells with MEKi/STAT3i treatment (**Fig. 1J**), we also observed an increased proportion of PD1<sup>+</sup>CD107a<sup>-</sup> non-degranulating CD8<sup>+</sup> T-cells (**Fig. 1K**). Moreover, we found no significant changes in PD-L1 expression in PKT tumors between vehicle treatment, MEKi or STAT3i monotherapy, and combined MEKi/STAT3i treatment (**Supplementary Figure S3**). Taken together, these immunologic effects associated with MEKi/STAT3i suggest that the addition of ICI to MEKi/STAT3i may overcome resistance to immunotherapy in immunologically inert PDAC.

### **Treatment with MEKi/STAT3i overcomes resistance to PD-1 blockade**

To build on these observations, PKT mice were treated with vehicle, αPD-1 monotherapy, MEKi/STAT3i, and MEKi/STAT3i combined with αPD-1 antibody (**Supplementary Fig. S4A**). Consistent with our previous data (10), MEKi/STAT3i significantly reduced tumor burden compared with vehicle treatment in PKT mice. Moreover, in keeping with prior evidence in *Kras<sup>G12D/+</sup>;Trp53<sup>fl/+</sup>;Pdx-Cre* (KPC) mice (12) as well as clinical trials of ICI in PDAC patients (5,6), αPD-1 monotherapy had no effect on tumor burden compared with vehicle treatment in PKT mice, further confirming that PDAC is refractory to ICI. Importantly, the addition of MEKi/STAT3i to αPD-1 blockade significantly reduced tumor burden compared to αPD-1-treated, as well as MEKi/STAT3i-treated, mice (**Fig. 2A&B**). In fact, in this aggressive PKT PDAC GEMM, where the pancreata of vehicle- and αPD-1-treated mice are completely replaced by tumor by 6 weeks of age, MEKi/STAT3i treated mice show only 19% tumor area, on average, after 4 weeks of treatment (~8 weeks of age), while the addition of MEKi/STAT3i to αPD-1 treatment maintained normal pancreas architecture with <6% tumor area (**Fig. 2A**).

### **Addition of MEKi/STAT3i to αPD-1 blockade significantly improves survival in PKT mice**

Having shown that MEKi/STAT3i drives stromal, as well as innate and adaptive immune, remodeling (**Fig. 1**), and that the addition of MEKi/STAT3i to αPD-1 reduced tumor burden compared with αPD-1 monotherapy, we next sought to determine if this combination therapy overcomes resistance to ICI to improve survival in PKT mice (treatment schema in **Supplementary Fig. S4B**). Confirming our previous data, MEKi/STAT3i significantly increased survival compared with vehicle treatment (median 103 vs. 44 days;  $p=0.0013$ ). Consistent with the lack of response to αPD-1 monotherapy in PDAC GEMMs (13-16), we observed no difference in survival between αPD-1 monotherapy and vehicle treatment (median 45 vs. 44 days,  $p=0.34$ ). Importantly, the addition of MEKi/STAT3i to PD-1 blockade dramatically improved PKT survival compared with αPD-1 monotherapy (median 181 vs. 45 days,  $p=0.0005$ ), as well as combined MEKi/STAT3i (median 181 vs. 103 days,  $p=0.0064$ ; **Fig. 2C**).

Furthermore, there was no difference in body weight between vehicle and treatment groups, indicating that the combination of MEKi/STAT3i with PD-1 blockade is well tolerated with no additional toxicity compared with either  $\alpha$ PD-1 monotherapy or MEKi/STAT3i *in vivo* (**Supplementary Fig. S4C**). These results highlight the significant role of MEKi/STAT3i in overcoming resistance to ICI in an immunologically inert GEMM of PDAC.

### **The addition of MEKi/STAT3i to PD-1 blockade improves recruitment, activation, and functional cytotoxicity of tumor-infiltrating T-cells**

Next, we examined the immunologic underpinnings of the dramatic improvement in survival of PKT mice treated with combined MEKi/STAT3i and  $\alpha$ PD-1 compared with  $\alpha$ PD-1 monotherapy. CyTOF-based immune profiling demonstrated that, compared with both vehicle treatment and  $\alpha$ PD-1 monotherapy, the addition of MEKi/STAT3i to PD-1 blockade resulted in a significant increase in tumor-infiltrating CD4<sup>+</sup> and CD8<sup>+</sup> T-cells, as well as CD3<sup>+</sup> T-cells demonstrating T-cell activation markers CD44 and CD69 and central memory marker CD62L (**Fig. 2D**). These broad-based changes in adaptive immune populations were then corroborated by flow cytometry in the CD4<sup>+</sup> and CD8<sup>+</sup> T-cell compartments separately. In both CD4<sup>+</sup> and CD8<sup>+</sup> T-cell compartments alike, compared to  $\alpha$ PD-1 monotherapy or vehicle treatment, the addition of MEKi/STAT3i to PD-1 blockade significantly augmented proportions of activated (CD69<sup>+</sup>), effector memory (CD44<sup>+</sup>CD62L<sup>-</sup>), central memory (CD44<sup>+</sup>CD62L<sup>+</sup>), tissue resident memory (CD44<sup>+</sup>CD62L<sup>-</sup>CD103<sup>+</sup>), and degranulating effector (CD44<sup>+</sup>CD62L<sup>-</sup>CD107a<sup>+</sup>) tumor-infiltrating T-cells (**Fig. 2E & Supplementary Fig. S5**).

Given significant improvements in the recruitment and activation profile of tumor-infiltrating T-cells, we next investigated if the addition of MEKi/STAT3i to PD-1 blockade enhances the functional cytotoxicity of CD8<sup>+</sup> T-cells compared with  $\alpha$ PD-1 monotherapy and vehicle treatment. Using *ex vivo* CD8<sup>+</sup> T-cell:tumor cell IFN- $\gamma$  release assays, PKT-derived EpCAM<sup>+</sup> tumor cells were co-cultured with splenocyte-derived CD8<sup>+</sup> T-cells from PKT mice



treated with vehicle,  $\alpha$ PD-1 alone and MEKi/STAT3i/  $\alpha$ PD-1, and the co-culture conditioned media subjected to IFN- $\gamma$  ELISA. The addition of MEKi/STAT3i to PD-1 blockade significantly increased the cytolytic activity of CD8<sup>+</sup> T-cells compared with  $\alpha$ PD-1 monotherapy or vehicle treatment (**Fig. 2F**). Further validating these functional T-cell data, the addition of MEKi/STAT3i to PD-1 blockade significantly increased tumor-specific granzyme B levels in PKT tumor lysates, compared with  $\alpha$ PD-1 monotherapy or vehicle treatment (**Fig. 2G**). These data collectively show that the addition of MEKi/STAT3i to PD-1 blockade invigorates the adaptive immune compartment to overcome T-cell dysfunction and ICI resistance in the PDAC TME. Moreover, MEKi/STAT3i, which improves tumor-infiltrating CD8<sup>+</sup> T-cell antigen experience but does not completely rescue their degranulating capacity (**Fig. 1I-K**), primes an activated CD8<sup>+</sup> T-cell compartment for ICI to further augment the cytolytic function of PDAC-infiltrating CD8<sup>+</sup> T-cells.

### **Attenuation of suppressive innate immune microenvironment in PKT mice is driven by MEKi/STAT3i but not anti-PD-1 monotherapy**

Given recent evidence that myeloid cell-specific PD-1 ablation—independent of T-cell-directed anti-PD-1 effects—may contribute to effector T-cell infiltration and antitumor immunity (17), we sought to investigate the differential effects of PD-1 blockade alone and combined MEKi/STAT3i/ $\alpha$ PD-1 on the innate immune microenvironment in PKT tumors. CyTOF-based immune profiling revealed that  $\alpha$ PD-1 monotherapy had no impact on innate immune infiltration in PKT tumors compared with vehicle treatment. The addition of MEKi/STAT3i to PD-1 blockade, however, significantly reduced the recruitment of tumor-infiltrating F4/80<sup>+</sup> macrophages, CD206<sup>+</sup> M2-like macrophages, and Ly6GC<sup>+</sup> MDSCs compared with  $\alpha$ PD-1 monotherapy or vehicle treatment (**Fig. 3A**). Tissue-based immunofluorescence (**Fig. 3B**) and flow cytometric analysis (**Supplementary Fig. S6**) confirmed a significant decrease in CD11b<sup>+</sup> myeloid, F4/80<sup>+</sup> macrophages, and Ly6G<sup>+</sup> MDSCs following MEKi/STAT3i/ $\alpha$ PD-1 compared with  $\alpha$ PD-1 monotherapy or vehicle treatment. These data reinforce that the dampening of

immunosuppressive innate populations in the PDAC TME is driven predominantly by MEKi/STAT3i without significant contributions from  $\alpha$ PD-1 monotherapy, and may underlie the adaptive immune invigoration observed with combination MEKi/STAT3i and PD-1 blockade.

### **Combination MEKi/STAT3i and PD-1 blockade demonstrates efficacy in a metastatic PDAC patient with chemotherapy-refractory disease**

A 69-year-old male with a remote history of Hodgkin's lymphoma presented with a resectable microsatellite stable, low-tumor mutation burden (TMB) PDAC in the pancreatic body. He received neoadjuvant fluorouracil (5-FU), leucovorin, irinotecan, and oxaliplatin (FOLFIRINOX) for three months before undergoing curative-intent distal pancreatectomy and splenectomy. Pathology revealed a margin-negative resection, and a 2-cm focus of residual adenocarcinoma with 0/33 involved lymph nodes. He then completed three months of adjuvant FOLFIRINOX. Two months following completion of chemotherapy, he was diagnosed with a local recurrence in the pancreatic bed as well as a metastatic lesion in segment VII of the liver. He initiated second-line chemotherapy with gemcitabine/nab-paclitaxel but had progression of disease with a significant reduction in performance status (**Fig. 4A**).

After multidisciplinary discussion, he received off-label combination treatment with MEKi Trametinib (Mekinist/Novartis, 2mg PO QD), STAT3i Ruxolitinib (Jakafi/Incyte, 5mg PO BID), and  $\alpha$ PD-1 Nivolumab (Opdivo/Bristol-Myers Squibb, 240mg IV Q2W) (**Fig. 4A**). Within one month of commencing treatment, the patient experienced a dramatic improvement in clinical and performance status. Follow-up PET/CT obtained after 3 months of treatment demonstrated significant response in the pancreatic bed mass, with nearly 50% reduction in size from 43 mm to 23 mm (**Fig. 4B**) and a metabolic response with reduction in SUV<sub>max</sub> from 8.4 to 6.5 post-treatment in the liver metastasis on PET/CT (**Fig. 4C**). The regimen was well-tolerated with no reported major adverse events while on therapy.

## DISCUSSION

The present manuscript uncovers a novel treatment strategy in PDAC that modulates the immunosuppressive tumor stroma to activate functional T-cell anti-tumor immunity to overcome ICI resistance. Resistance to ICI is a hallmark of PDAC (5,6), defined by T-cell exclusion and tissue-level immune tolerance driven by *KRAS*-mediated tumor-intrinsic programs (1,8) as well as interwoven suppressive cellular networks comprising innate immune populations and tumor-promoting cancer-associated fibroblasts (CAF) (7). The therapeutic strategy highlighted herein of disrupting oncogenic signaling via reciprocally activated *KRAS*/MEK/ERK and JAK/STAT3 pathways in PDAC tumor cells using MEKi/STAT3i (10) addresses all these major barriers to overcome ICI resistance in an aggressive and immunologically inert PDAC GEMM. In addition to remodeling of stromal architecture which may augment intratumoral T-cell trafficking and antigen experience, the attenuation of MDSCs and M2-like TAMs observed with MEKi/STAT3i treatment may drive the pro-immunogenic T-cell landscape that sensitizes PKT mice to ICI. It is also possible that *direct* effects of MEKi/STAT3i on myeloid cellular signaling and/or myelopoiesis contributes to the observed adaptive immune invigoration in the PDAC TME. While the tolerogenic role of myeloid-specific STAT3 signaling has been well established in pancreatic and other solid tumors (18), the role of myeloid-specific MEK/ERK signaling (19) and its immunologic repercussions is incompletely understood. The findings herein warrant further investigation into the differential contributions of MEK/ERK and STAT3 signaling in myelopoiesis, myeloid-specific lineage commitment, and multi-directional crosstalk with disparate immune constituents of the PDAC TME.

Emerging evidence has elucidated the signaling crosstalk between tumor cells and cancer associated fibroblasts (CAF)—the predominant constituent of the PDAC stroma—in promoting hallmarks of its aggressive biology, such as epithelial-to-mesenchymal transition (EMT) and proliferative capacity (20). In addition to our group's previous discovery of

reciprocally activated MEK/ERK (downstream of RAS) and STAT3 signaling in tumor cells promoting therapeutic resistance (10,11), other groups have reported CAF-mediated factors (such as TGF- $\beta$ ) further perpetuating tumor cell-specific MEK/ERK and STAT3 co-activation (20). The impact of abrogating MEK/STAT3-driven tumor-stromal crosstalk is exemplified in the present study, with combined MEKi/STAT3i substantially remodeling stromal composition in PKT mice. The immunologic changes associated with such stromal remodeling following MEKi/STAT3i treatment, however, are more intriguing since growing evidence underscores a complex and non-linear relationship between stromal desmoplasia and effector immune infiltration in the PDAC TME (21). The immune landscape is controlled in part by diverging signaling cues from heterogeneous CAF subtypes (22). It is plausible, therefore, that context-dependent targeting of tumor-promoting inflammatory CAFs—and not indiscriminate stromal depletion (23)—will be more effective in overcoming immune exclusion and therapeutic resistance in PDAC. To this end, MEKi/STAT3i may exert its immune potentiating effects by mitigating inflammatory CAF phenotypes or abolishing CAF-intrinsic tolerogenic signaling mechanisms (e.g., STAT3 signaling (24)) that orchestrate immune suppression and establish immune privilege in the PDAC TME. Ongoing work in our laboratory is dissecting how inhibition of cooperative MEK/ERK and STAT3 signaling within PDAC CAFs reshape the immune microenvironment to sensitize PDAC to ICI.

Beyond the scientific underpinnings of this novel therapeutic strategy to overcome ICI resistance in PDAC, the impressive clinical benefit and tolerability of Trametinib, Ruxolitinib, and Nivolumab in a patient with metastatic chemo-refractory PDAC is an encouraging signal of its translatability. While such dramatic responses are not uniformly expected in all patients enrolled in our upcoming phase I trial investigating this novel combination regimen in metastatic PDAC, the immunologic consequences of MEKi/STAT3i treatment *in vivo* will be incorporated into a

predictive biomarker discovery platform that may inform strategic selection of future patients for these immunotherapeutic combinations.

In summary, our data present novel insight into a combinatorial targeted therapy regimen that overcomes ICI resistance in PDAC by restoring elements of T-cell immunosurveillance and reprogramming the immunosuppressive TME (3), laying the foundation for its clinical application in the near future.

## METHODS

### *In vivo studies*

Tumor-bearing *Ptf1a*<sup>Cre/+</sup>*Kras*<sup>G12D/+</sup>*Tgfbr2*<sup>flox/flox</sup> (PKT) mice (provided by Dr. Harold Moses, Vanderbilt University Medical Center, Nashville, TN) were generated as previously described (25). For endpoint analysis (i.e., flow cytometry, CyTOF, histology), PKT mice were treated with vehicle (0.5% HPMC + 0.1% Tween 80), trametinib (MEKi, Novartis, 0.067 mg/ml/mouse, oral gavage three times weekly), JAK1/2 inhibitor, ruxolitinib (STAT3i, TargetMol, 0.4 mg/ml/mouse, oral gavage three times weekly), αPD-1 antibody (BioXCell, Clone #BE0273, 200 μg/mouse, intraperitoneal injection twice weekly), or combined MEKi/STAT3i or MEKi/STAT3i with αPD1 beginning at 4-4.5 weeks of age. Mice were sacrificed after four weeks of treatment or when moribund. For survival studies, MEKi (0.067 mg/ml/mouse) and STAT3i (0.4 mg/ml/mouse) were administered by oral gavage 5 days/week starting at 4.5 weeks of age; αPD-1 antibody dosing remained intraperitoneal injection twice weekly. Six weeks after treatment initiation, the frequencies of MEKi/STAT3i and αPD1 antibody dosing were reduced to three times weekly and once weekly, respectively, and continued until mice were moribund (**Supplementary Figure S4A&B**). T-cell depletion experiments are described in **Supplementary Methods**.

All animal experiments were performed in compliance with the ethical guidelines of the Institutional Animal Care and Use Committee at the University of Miami (Protocol #18-040).

### ***Flow cytometry***

Pancreatic tumors harvested from PKT mice were enzymatically digested using solution of 0.6 mg/ml of collagenase P (Roche), 0.8 mg/ml Collagenase V (Sigma Aldrich), 0.6 mg/ml soybean trypsin inhibitor (Sigma Aldrich), and 1800 U/ml DNase I (Thermo Scientific) in RPMI medium for 20-30 minutes at 37°C. Samples were then washed and resuspended in cold PEB solution followed by straining through 40 µm mesh filter to obtain single cell suspension. Spleen samples were processed using RBC lysis buffer, followed passage through 100 µm mesh filter. Samples were frozen at -80°C until further use. Prior to flow cytometry staining, samples were thawed, washed prior to incubation with FcR blocking reagent (Miltenyi Biotec), and subsequently stained with fluorescently conjugated antibodies listed in **Supplementary Table S1**. Live/dead cell discrimination was performed using Live/Dead Aqua (Life Technologies) or Ghost Red Dye 780 (TONBObiosciences) as per manufacturer's protocol. Cell fixation was performed using 1% formaldehyde solution (Thermo Fisher). Flow cytometric data acquisition was performed on CytoFLEX S (Beckman Coulter) and analyzed using FlowJo v10 software (BD Life Sciences). Gating strategies are depicted in **Supplementary Figure S7**.

### ***Mass cytometry (CyTOF)***

Single-cell suspensions from PKT tumors were flow sorted using FACS Aria III (BD Life Sciences) to remove non-cellular contaminants. Samples were resuspended and viability staining performed using 2.5 µM cisplatin (Fluidigm) working solution following by Fc receptor blocking and surface staining for 30 min at room temperature using tagged antibodies as depicted in **Supplementary Table S2**. Fixation was then performed using 1.6% formaldehyde

solution (Thermo Fisher). Cells were suspended in 125 nM Cell-ID Intercalator-Ir solution (Fluidigm) overnight at 4°C. The following day, cells were washed and resuspended in 0.1X EQ Four Element Calibration Beads (Fluidigm) before acquisition. Data were acquired by Helios Mass Cytometer (Fluidigm) and analyzed using FlowJo v10 and Cytobank (Beckman Coulter). To generate viSNE plots, individual files from each treatment group were concatenated into a single file and then a subset of equal events were selected at random through FlowJo DownSample plugin. Total number of CD45<sup>+</sup> cells from each group were used to create viSNE through Cytobank with 2000 iterations, perplexity of 30, and theta of 0.5.

### ***Histologic Analysis and Western Blotting***

See **Supplementary Methods**

### ***T-cell Functional Cytotoxicity Assays***

PKT mice were treated with vehicle, αPD-1, or MEKi/STAT3i with αPD-1 for 4 weeks before sacrifice and processing of tumor samples for flow cytometry. FACS-sorted viable EpCAM<sup>+</sup> tumor cells and total splenocytes were isolated from each mouse. Tumor cells were irradiated (4 Gy for 7 minutes) and plated in a direct co-culture system with isolated splenocytes. Condition media was collected after 72 hours of co-culture and protein content determined by BCA method. Secreted IFN-γ was measured by ELISA as per manufacturer's instruction (R&D Systems).

For measurement of intratumoral Granzyme B (GzB) levels, pancreatic tumor homogenates were prepared from PKT mice treated with vehicle, αPD-1, or MEKi/STAT3i with αPD-1 for 4 weeks. Tumor lysates were quantified and 40 μg protein loaded for analysis of GzB levels by ELISA (R&D Systems).

### ***Statistical analysis***

Descriptive statistics were calculated using Prism 9.0 (GraphPad Software Inc.). Results are shown as mean $\pm$ SEM. Multiple comparisons were performed using ANOVA followed by Tukey's multiple comparisons test. The paired two-sided Student's t test was used for two-group comparison. Survival curves were estimated using the Kaplan-Meier method and differences between groups were assessed using the log-rank test. An  $\alpha$ -cutoff  $\leq 0.05$  was used to define statistical significance.



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## FIGURE LEGENDS

### **Figure 1. Combined MEKi/STAT3i inhibits stromal fibrosis, remodels the immune microenvironment, and reduces tumor growth in a T-cell-dependent manner. (A)**

Trichrome blue,  $\alpha$ -SMA, and Sirius Red staining in tumor sections from PKT mice treated with vehicle or MEKi+STAT3i for 4 weeks; relative areas of positive staining of respective markers from tissue sections in vehicle- and MEKi/STAT3i-treated mice are indicated in adjacent histograms. **(B)** Levels of total myeloid (CD11b<sup>+</sup>), macrophage (F4/80<sup>+</sup>), and T-cell populations (CD4<sup>+</sup> and CD8<sup>+</sup>) were compared between vehicle and MEKi/STAT3i-treated PKT mice by flow cytometry (n=8-10 mice/arm). **(C)** Mass cytometry time-of-flight (CyTOF) FlowSOM plots depicting changes in total myeloid (CD11b<sup>+</sup>), T-cell (CD4<sup>+</sup> and CD8<sup>+</sup>) and B-cell (CD19<sup>+</sup>) populations in PKT mice treated with vehicle or MEKi+STAT3i (n=7-8 mice/arm).

Representative viSNE plots demonstrating changes in individual **(D)** myeloid and **(E)** T-cell subsets between vehicle- and MEKi+STAT3i-treated PKT mice as analyzed by CyTOF. Parent CD11b<sup>+</sup> **(D)** and CD3<sup>+</sup> **(E)** cell populations are denoted by dashed line. PKT mice were treated with vehicle, MEKi+STAT3i, or MEKi+STAT3i following T-cell depletion with anti-CD4 and anti-CD8 antibodies (scheme in **Supplementary Fig. S2**). Differences in **(F)** pancreas weight at sacrifice and **(G)** overall survival were compared among treatment arms. **(H)** Western blot demonstrating target inhibition of pERK1/2 and pSTAT3 in mice treated with MEKi+STAT3i  $\pm$  T-cell depletion. Flow cytometric analysis of PKT mice treated with vehicle or MEKi+STAT3i showing an increase in **(I)** PD-1<sup>+</sup> tumor-infiltrating CD8<sup>+</sup> T-cells, **(J)** CD8<sup>+</sup>PD-1<sup>+</sup>CD107a<sup>+</sup> T-cells, and **(K)** CD8<sup>+</sup>PD-1<sup>+</sup>CD107a<sup>-</sup> T-cells in MEKi+STAT3i treated mice (n=8-10 mice/arm). Data are shown as mean  $\pm$  SEM. Scale bar = 50  $\mu$ m. \*, p<0.05; \*\*, p<0.01; \*\*\*, p<0.001.

### **Figure 2: Addition of MEKi/STAT3i to PD-1 blockade overcomes immune checkpoint inhibition (ICI) resistance, improves survival, and augments recruitment, activation, and functional cytotoxicity of tumor-infiltrating T-cells. (A)** Representative images from H&E

sections from PKT mice treated with vehicle,  $\alpha$ PD-1, MEKi/STAT3i, or MEKi/STAT3i plus  $\alpha$ PD-1 for 4 weeks (*left*). Both percent tumor area (**A**, *right*) and (**B**) pancreatic weight at sacrifice were compared among treatment arms (n=5 mice/arm). (**C**) Kaplan-Meier survival plot depicting overall survival in PKT mice treated with vehicle,  $\alpha$ PD-1, MEKi/STAT3i, or MEKi/STAT3i plus  $\alpha$ PD-1 beginning at 4-4.5 weeks of age. (**D**) viSNE plots demonstrating levels of tumor-infiltrating CD3<sup>+</sup> cell subsets (CD4<sup>+</sup>, CD8<sup>+</sup>, CD69<sup>+</sup>, CD44<sup>+</sup>, and CD62L<sup>+</sup>) in vehicle,  $\alpha$ PD-1, and combined MEKi/STAT3i/ $\alpha$ PD-1 treated PKT mice as analyzed by CyTOF (n=7-8 mice/arm). (**E**) Analysis by flow cytometry showing changes in total CD3<sup>+</sup>, CD4<sup>+</sup>, and CD8<sup>+</sup> tumor-infiltrating T-cells as well as levels of activated CD69<sup>+</sup> and degranulating effector (CD4<sup>+</sup>CD62<sup>+</sup>CD107<sup>+</sup>) CD4<sup>+</sup>/CD8<sup>+</sup> T-cells in PKT mice treated with vehicle,  $\alpha$ PD-1, or MEKi/STAT3i/ $\alpha$ PD-1 (n=7-10 mice/arm). (**F**) Schematic depicting *ex vivo* tumor cell:splenocyte co-culture experiment. Splenocytes were isolated from PKT mice following treatment with vehicle,  $\alpha$ PD-1, and MEKi/STAT3i/ $\alpha$ PD-1 and co-cultured with irradiated PKT tumor cells for 72 hours. IFN- $\gamma$  release was determined by ELISA (right). (**G**) ELISA demonstrating granzyme B levels in total tumor lysate in vehicle,  $\alpha$ PD1, and MEKi/STAT3i/ $\alpha$ PD-1 treatment arms (n=3 mice/arm). Data are shown as mean $\pm$ SEM. Scale bar = 50  $\mu$ m. \*, p<0.05; \*\*, p<0.01; \*\*\*, p<0.001; \*\*\*\*, p<0.0001.

**Figure 3: Differential effects on innate immune remodeling are driven by MEKi/STAT3i with minimal contribution from anti-PD-1 monotherapy in PKT mice.** (**A**) viSNE plots depicting CyTOF analysis of CD11b<sup>+</sup> myeloid cell subsets (F4/80<sup>+</sup>, CD206<sup>+</sup>, and Ly6G<sup>+</sup>C<sup>+</sup>) in PKT mice treated with vehicle,  $\alpha$ PD-1, and MEKi/STAT3i plus  $\alpha$ PD-1 for 4 weeks (n=7-8 mice/arm). (**C**) Immunofluorescent staining and quantification of CD11b<sup>+</sup>, F4/80<sup>+</sup>, and Ly6G<sup>+</sup> levels in PKT mice among indicated treatment arms (n=3-4 mice/arm). Data are shown as mean  $\pm$  SEM. Scale bar = 50  $\mu$ m. \*, p<0.05; \*\*, p<0.01; \*\*\*, p<0.001; \*\*\*\*, p<0.0001.

**Figure 4: Combination MEKi/STAT3i and PD-1 blockade demonstrates efficacy in a metastatic PDAC patient with chemotherapy-refractory disease. (A)** Treatment timeline for patient with chemotherapy refractory PDAC prior to initiation of trametinib, ruxolitinib, and nivolumab (TRN) treatment. Pre-treatment and post-treatment PET/CT scan showing a significant reduction in size and FDG avidity of both locally recurrent tumor in pancreatic bed **(B)** and in a segment VII liver metastasis **(C)** on PET/CT imaging following 3 months of TRN treatment.

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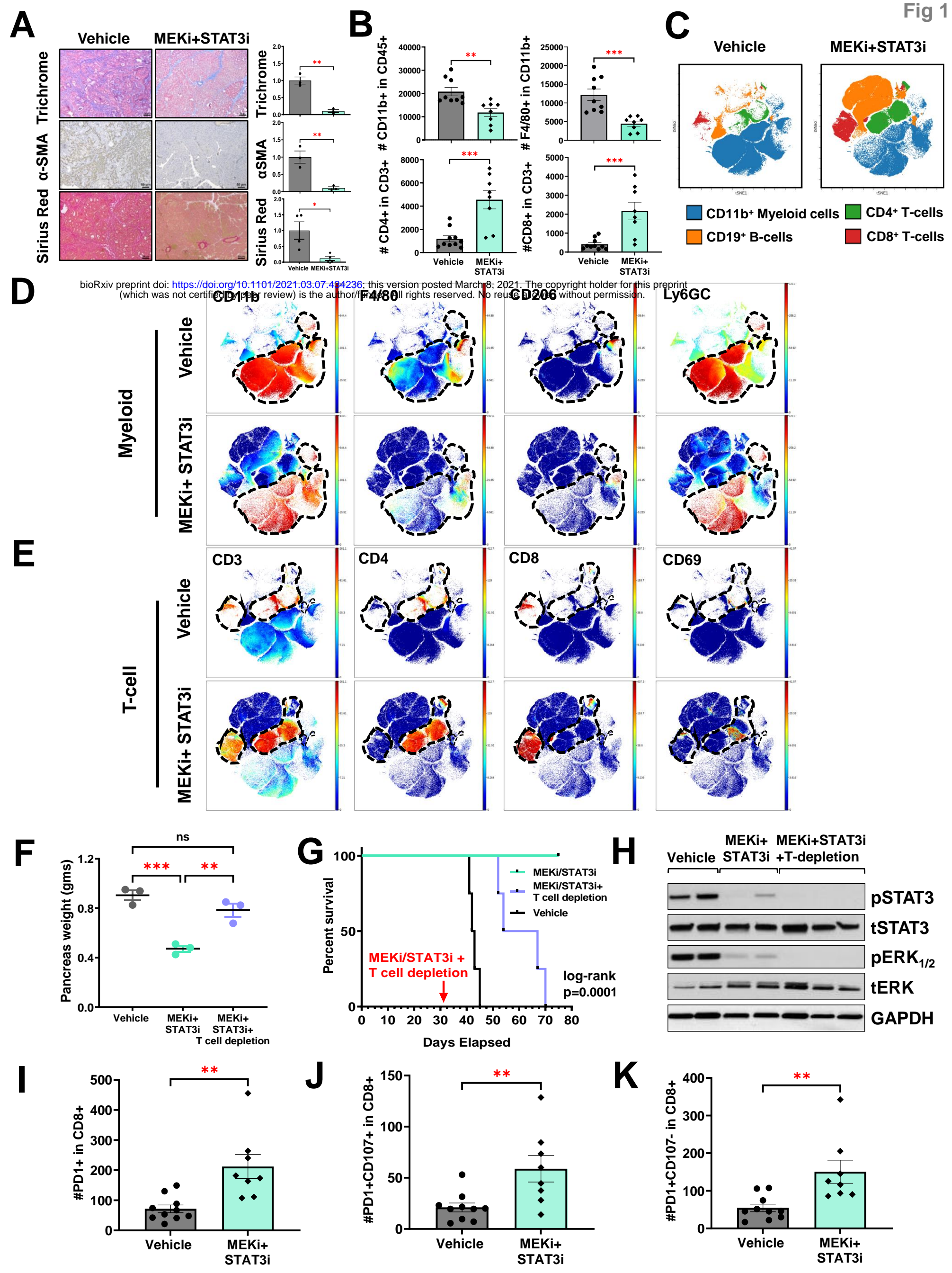
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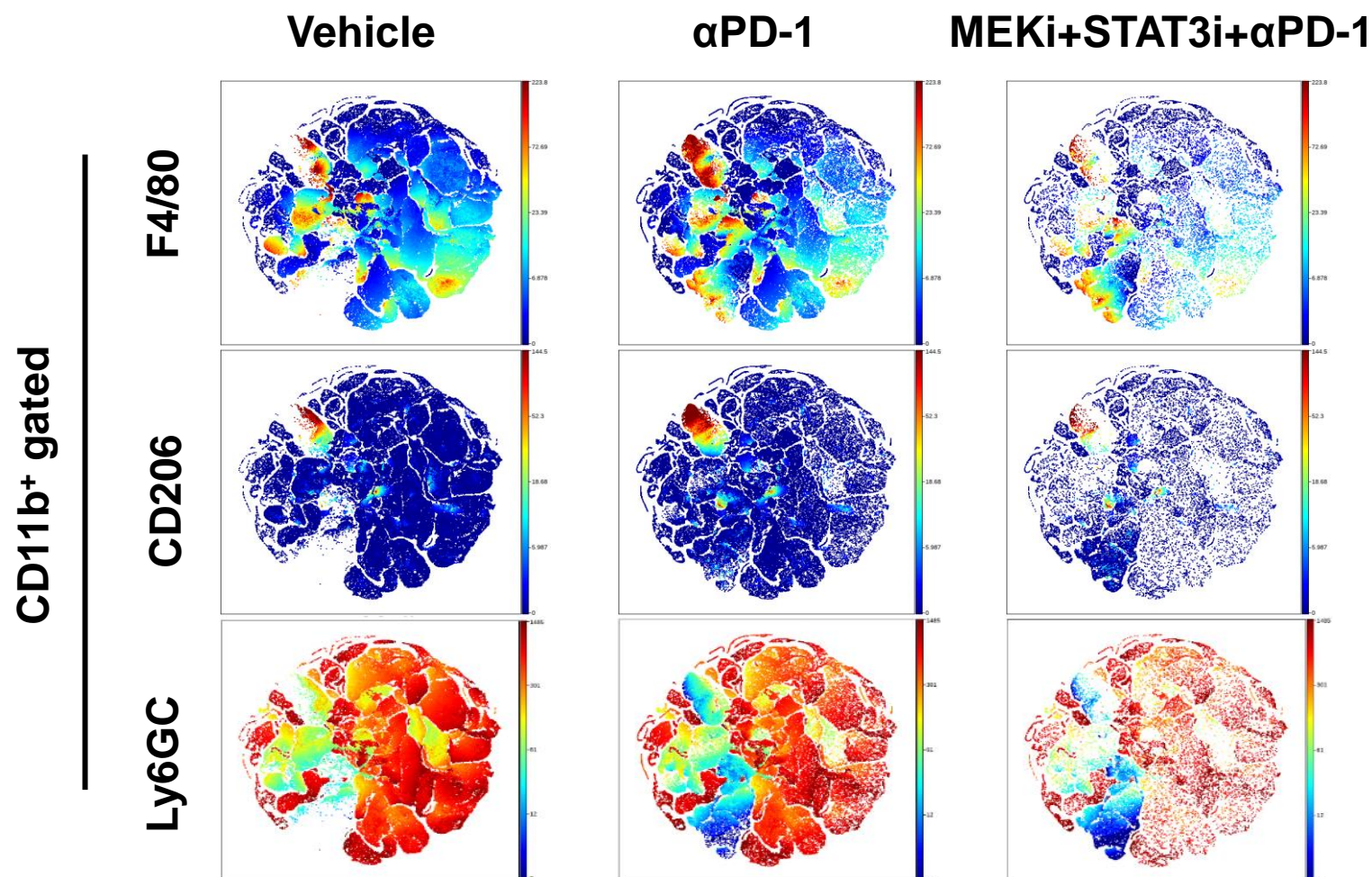










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