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An engineering approach to decode immune responses

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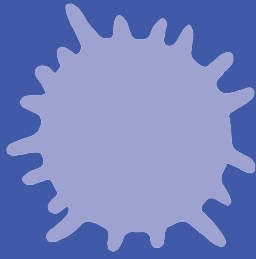
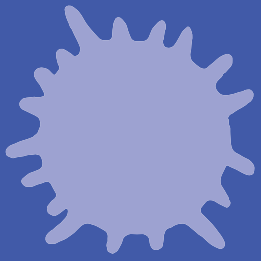
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Chapter 5.

QPCTL regulates macrophage and monocyte abundance and inflammatory signatures in the tumor microenvironment

OR: "Practising pronouncing glutaminy-peptide cyclotransferase in the bathroom mirror before every group meeting"

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Abstract

The enzyme glutamyl-peptide cyclotransferase-like protein (QPCTL) catalyzes the formation of pyroglutamate residues at the NH₂-terminus of proteins, thereby influencing their biological properties. A number of studies have implicated QPCTL in the regulation of chemokine stability. Furthermore, QPCTL activity has recently been shown to be critical for the formation of the high affinity SIRP α binding site of the CD47 “don’t-eat-me” protein. Based on the latter data, interference with QPCTL activity—and hence CD47 maturation—may be proposed as a means to promote anti-tumor immunity. However, the pleiotropic activity of QPCTL makes it difficult to predict the effects of QPCTL inhibition on the tumor microenvironment (TME). Using a syngeneic mouse melanoma model, we demonstrate that QPCTL deficiency alters the intratumoral monocyte-to-macrophage ratio, results in a profound increase in the presence of pro-inflammatory cancer-associated fibroblasts (CAFs) relative to immunosuppressive TGF- β 1-driven CAFs, and leads to an increased IFN and decreased TGF- β transcriptional response signature in tumor cells. Importantly, the functional relevance of the observed TME remodeling is demonstrated by the synergy between QPCTL deletion and anti PD-L1 therapy, sensitizing an otherwise refractory melanoma model to anti-checkpoint therapy. Collectively, these data provide support for the development of strategies to interfere with QPCTL activity as a means to promote tumor-specific immunity.

Introduction

Regulation of immune cell activity at sites of infection or cancer growth frequently occurs through a balance of signals that are received by immune activating and immune inhibitory receptors¹. For example, while activation of myeloid cells, including neutrophils, macrophages and monocytes, often occurs through Fc receptor signaling, such activation can be prevented through the simultaneous engagement of ITIM/ITSM-containing inhibitory receptors such as SIRP α . Specifically, binding of the “don’t-eat-me” signal CD47, which is widely expressed on hematopoietic and non-hematopoietic cells, to the SIRP α receptor on myeloid cells has been shown to result in decreased myeloid effector function, including suppression of target cell phagocytosis by macrophages and tumor cell killing by neutrophils²⁻⁴.

The inhibitory capacity of CD47 is dependent on the maturation of its SIRP α binding site by the ER-resident enzyme QPCTL^{5,6}. Similar to its secreted family member QPCT, QPCTL catalyzes the cyclization of N-terminal glutamine and glutamic acid residues on target proteins into a pyroglutamate residue (pGlu)^{7,8}. As shown by structural analysis, the pGlu residue at the N-terminus of CD47 contributes to the interaction surface with SIRP α ⁹; and, through genetic screening, it was shown that the activity of QPCTL is critical for the formation of this residue, making this enzyme a key regulator of the high-affinity CD47-SIRP α binding site⁶. In line with this, prevention

of pGlu formation on CD47, either by genetic knock-out or small molecule inhibition, leads to reduced SIRP α binding and increased macrophage- and neutrophil-dependent killing of antibody-opsionized target cells. Based on its role in regulating CD47-SIRP α signaling, and the possibility to develop small molecule inhibitors of enzymatic activity, QPCTL forms a potentially interesting target in cancer immunotherapy.

In addition to CD47, chemokines such as CCL2 and CX₃CL1 have been identified as QPCTL and/or QPCT substrates¹⁰⁻¹². The formation of the N-terminal pGlu on CCL2 was shown to increase its *in vivo* activity, both by conferring resistance to aminopeptidases and by increasing its capacity to induce CCR2 signaling¹⁰. Likewise, pGlu-modified CX₃CL1 appears to show an increased capacity to promote CX₃CR1 signaling *in vitro*¹². Finally, around 600 human proteins harbor a N-terminal glutamine or glutamic acid residue after the predicted signal peptide cleavage site, and it is plausible that additional QPCTL/QPCT substrates exist amongst this group of proteins¹³.

Because of its known or potential role in the post-translational modification of different immune- or tumor cell-related molecules, it is difficult to predict the overall effects of QPCTL inhibition on the tumor microenvironment (TME), and the poor pharmacokinetics of available QPCT/QPCTL inhibitors such SEN177⁶ has precluded evaluation of such effects by small-molecule inhibition. To address this question, we have generated a QPCTL-deficient mouse model and combined it with QPCTL-deficient tumor cells, to map the effects of QPCTL deficiency on either cellular compartment on the composition of the TME. The obtained data reveal that QPCTL deficiency results in a skewing of the macrophage-monocyte ratio, causes an approximately 20-fold change in the balance between TGF- β -producing myofibroblastic cancer-associated fibroblasts (myCAFs) and cytokine-secreting inflammatory CAFs (iCAFs), and shifts tumor cells from a TGF- β -responding to an IFN-responding state. Collectively, these changes convert the TME to pro-inflammatory environment that sensitizes tumors to anti-PD-L1 therapy. Together, these results suggest that therapeutic manipulation of QPCTL activity may synergize with current cancer immunotherapies.

Results

Genetic deletion of QPCTL does not significantly alter the immune compartment

Previous work has indicated that genetic deletion of QPCTL can be used to study its function *in vivo*^{10,14}; however, the possible effects of QPCTL inactivation on the TME has not been studied. To determine how blockade of QPCTL activity alters the TME, we applied CRISPR-Cas9 mediated gene-editing to generate a QPCTL-deficient (QPCTL^{-/-}) C57BL/6 mouse strain that is compatible with commonly used tumor models. To first determine whether QPCTL deficiency results in abrogation of pGlu formation on CD47, peripheral blood cells from QPCTL^{-/-} mice and WT littermates were stained with mouse SIRP α and an anti-mouse CD47 antibody that recognizes CD47 independent of pyroglutamate formation⁶. As compared to WT littermates, blood cells of

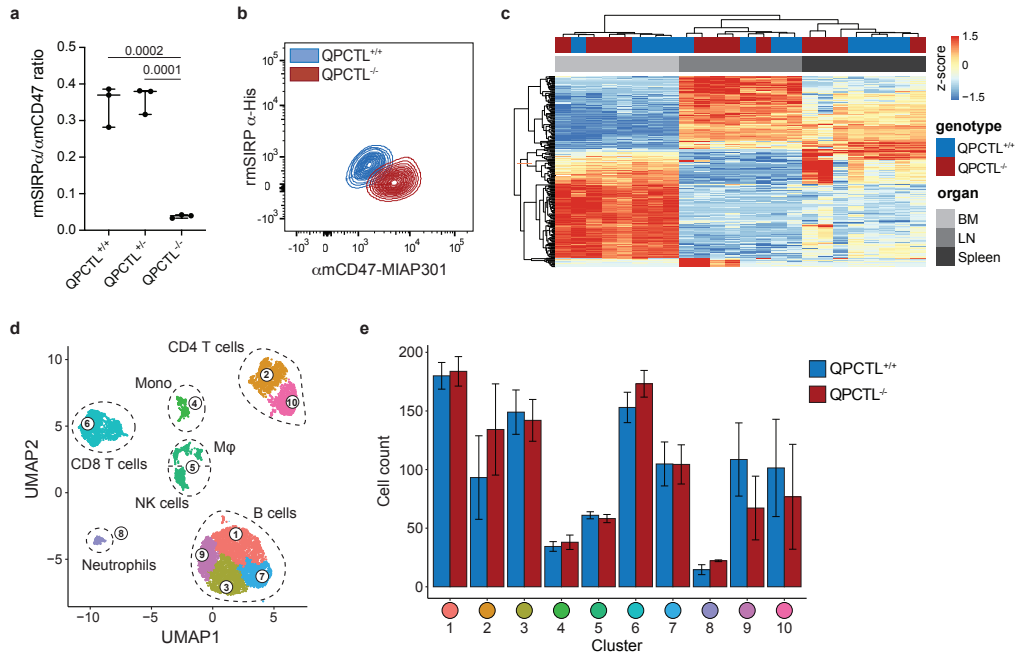


Figure 1. Generation and characterization of QPCTL-deficient mice. (a) Ratio of recombinant mouse (rm) SIRP α -His and anti-mouse (am)CD47 antibody (clone MIAP301) binding to blood cells from QPCTL^{+/+}, QPCTL^{+/-}, and QPCTL^{-/-} mice, as measured by flow cytometry. Dots depict the ratio of rmSIRP α -His/amCD47-MIAP301 mean fluorescence intensity (MFI) on blood cells from individual mice, group medians are indicated and whiskers represent min/max, $n = 3$ mice per group. (b) Flow cytometry plot depicting data described in a) for blood cells from a representative QPCTL^{+/+} and QPCTL^{-/-} mouse. (c) Heatmap depicting hierarchical clustering performed on the 1,000 most differentially expressed genes in bone marrow (BM), lymph node (LN) and spleen samples from QPCTL^{+/+} and QPCTL^{-/-} mice. (d-e) Unbiased Euclidean distance-based clustering of immune cells obtained from spleens of QPCTL^{+/+} and QPCTL^{-/-} mice. UMAP 2-dimensional projection (d) depicts the obtained clusters. Cell counts of both genotypes within each cluster are depicted (e). Bars indicate group means, error bars represent standard error of the mean. P values were determined by one-way ANOVA followed by Tukey's HSD test (a) or by two-sided Student's T test with Bonferroni correction for multiple testing (e). Significant P values (< 0.05) are indicated in the plots. Data are representative of 3 independent experiments (a-b), or were obtained in a single experiment (c-d). UMAP, Uniform Manifold Approximation and Projection.

QPCTL-deficient mice displayed significantly decreased SIRP α binding, thereby providing the first evidence that QPCTL is also a critical CD47 modifier *in vivo* (Fig. 1a-b).

To understand how QPCTL deficiency influences steady-state immune cell frequencies and gene expression, QPCTL^{-/-} and QPCTL^{+/+} littermates were subjected to histopathological, transcriptomic and flow cytometric analysis. Histopathological assessment of QPCTL^{-/-} mice revealed no significant morphological aberrations relative to littermate controls (Supplementary Data 1), and gene expression analysis of spleen, lymph nodes and bone marrow revealed no genotype-specific transcriptional changes (Fig. 1c, Supplementary Fig 1a), indicating that QPCTL deficiency does not result in major alterations in steady-state immune activity. Likewise, no substantial differences in cell counts or immune cell population frequencies were observed in blood (Supplementary Fig. 1b-c). In spleen, a modest increase in the frequency of NK cells of total non-myeloid cells and a decrease in the fraction of activated cells of total CD4⁺ T cells was observed in QPCTL^{-/-} mice, but

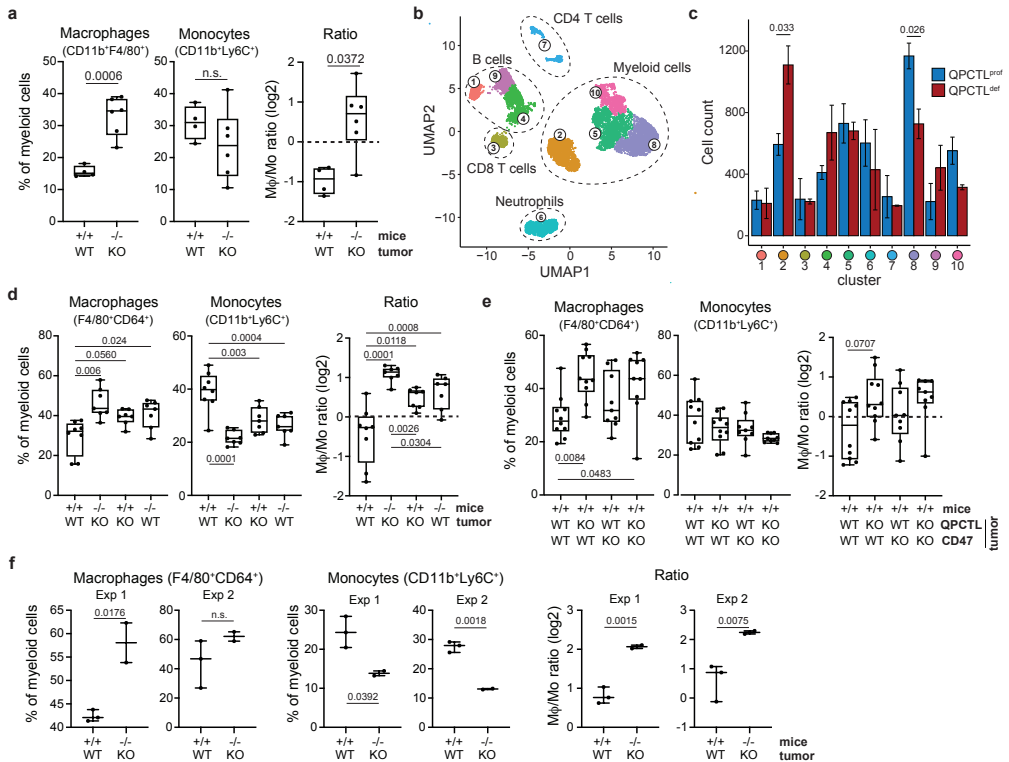


Figure 2. Tumor and host QPCTL deficiency alters the Mφ-Mo ratio in the TME. (a) Frequency of macrophages and monocytes of myeloid cells (CD11b⁺), and Mφ-Mo ratio, in the TME of QPCTL^{+/+} (*n* = 4) and QPCTL^{-/-} (*n* = 6) mice inoculated with QPCTL-WT or QPCTL-KO B16F10 melanoma cells, respectively. Tumors were analyzed between 14–16 days post inoculation. (b) UMAP visualizing 30,000 cells sampled from the data shown in a. 5,000 cells were randomly drawn from each sample (*n* = 3 mice per group) prior to analysis. Colors indicate clusters obtained by Euclidean distance-based hierarchical clustering, cluster phenotype is shown in **Supplementary Figure 2f**. (c) Contribution of cells from QPCTL^{+/+} (*n* = 3) and QPCTL^{-/-} (*n* = 3) TMEs to each cluster shown in b. Bars indicate group means, error bars represent standard error of the mean. (d) Frequency of macrophages and monocytes of myeloid cells (CD11b⁺), and Mφ-Mo ratio, in the TME of QPCTL^{+/+} and QPCTL^{-/-} mice inoculated with either QPCTL-WT or QPCTL-KO B16F10 melanoma cells (*n* = 7–8 per group). Tumors were analyzed between 12–14 days post inoculation. (e) Frequency of macrophages and monocytes of myeloid cells (CD11b⁺), and Mφ-Mo ratio, in the TME of QPCTL^{+/+} mice inoculated with QPCTL-WT, QPCTL-KO, CD47-KO, or CD47/QPCTL double-KO (dKO) B16F10 cells. Tumors were analyzed between 14–16 days post inoculation. (f) Frequency of macrophages and monocytes of myeloid cells (CD11b⁺), and Mφ-Mo ratio, in the TME of QPCTL^{+/+} and QPCTL^{-/-} mice inoculated with QPCTL-WT and QPCTL-KO MC38 cells, respectively. Data from 2 independent experiments are shown (*n* = 5 per experiment). Tumors were analyzed at 22 (experiment 1) or 29 (experiment 2) days post inoculation. Dots indicate measurements from individual mice, group medians are indicated and whiskers represent min/max. *P* values were determined by two-sided Student's T test without (a, f) or with Bonferroni correction for multiple testing (c), or by one-way ANOVA followed by Tukey's HSD test (d, e). Significant *P* values (< 0.05) are indicated in the plots. For all boxplots, dots represent individual mice, group median and 25th/75th percentiles are indicated by the box, whiskers indicate min/max. Data are representative of at least 2 independent experiments (a, d, f), or were obtained in a single experiment (b, c, e). UMAP, Uniform Manifold Approximation and Projection.

no significant changes in other immune cell type frequencies were identified (**Supplementary Fig. 1d**). The absence of substantial differences in immune cell frequencies was corroborated by unbiased hierarchical clustering of cells obtained from QPCTL^{+/+} and QPCTL^{-/-} spleen samples (**Fig. 1d** and **Supplementary Fig. 1e**).

QPCTL-deficiency alters macrophage-monocyte-ratios in the TME

To test whether systemic QPCTL deficiency influences immune cell infiltration in the TME, QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with wild-type (QPCTL-WT) and QPCTL knock-out (QPCTL-KO) B16F10 melanoma cells (**Supplementary Fig. 2a**), respectively, and TMEs were analyzed by flow cytometry 14–16 days post inoculation. QPCTL-proficient and -deficient tumors grew with similar kinetics and were similarly infiltrated by large numbers of myeloid cells (**Supplementary Fig. 2b-c**). Importantly, within the myeloid subset, QPCTL-deficient TMEs exhibited a significant higher frequency of macrophages and a substantially increased macrophage-monocyte (M ϕ -Mo) ratio (**Fig. 2a**). Although no other significant changes within the immune infiltrate could be detected (**Supplementary Fig. 2d**), an increase in the frequency of B cells, a trend toward a decrease in monocytes, and a decrease in the frequency of CD4⁺ T cells within the non-myeloid immune cell subset was observed in peripheral blood samples from QPCTL-deficient tumor-bearing mice (**Supplementary Fig. 2e**). Also, when immune infiltrates of QPCTL-deficient and -proficient TMEs were analyzed through unbiased hierarchical clustering, an increase in F4/80⁺ cells (macrophages) and a decrease in Ly6C^{high} cells (monocytes) in QPCTL-deficient TMEs was observed (**Fig. 2b-c, Supplementary Fig. 2f**).

To determine whether the increased M ϕ -Mo-ratio could be attributed to a lack of QPCTL activity in either host or tumor cells, QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with either QPCTL-WT or QPCTL-KO melanoma tumor cells (**Supplementary Fig. 3a**). Both tumor and host QPCTL deficiency led to an increased M ϕ -Mo-ratio, but the most profound increase in M ϕ -Mo-ratios was observed when QPCTL activity was lacking in both cell compartments (**Fig. 2d, Supplementary Fig. 3b-c**). In blood, the most pronounced differences in immune cell frequencies were found when comparing tumor-bearing versus non-tumor-bearing animals—independent of QPCTL activity—emphasizing that QPCTL deficiency does not impact the systemic immune compartment in a major way (**Supplementary Fig. 3d**). To explore to what extent loss of pGlu-modified CD47 contributed to the altered intra-tumoral M ϕ -Mo balance, wild-type mice were inoculated with QPCTL-, CD47- or double-KO tumor cells. Absence of CD47 resulted in M ϕ frequencies that were numerically higher than observed in recipients of WT B16 tumor cells, but to a lower extent than observed in recipients of QPCTL-KO cells, and did not significantly alter M ϕ -Mo-ratios (**Fig. 2e**). This suggests that, if the CD47-modifying activity of QPCTL contributes to the altered myeloid cell composition in these tumors, it likely plays a minor role.

To test whether the role of QPCTL as a modifier of the TME extended to other tumor models, QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with QPCTL-WT and QPCTL-KO MC38 colon carcinoma cells, respectively. Following tumor outgrowth, a profound increase in M ϕ -Mo-ratio was observed in QPCTL-deficient TMEs (**Fig. 2f, Supplementary Fig. 4a-b**), while no significant changes were detected in other immune cell subsets (data not shown). Together, these data indicate that QPCTL affects the TME composition in at least two different tumor models, and that combined tumor and host QPCTL deficiency leads to a significant increase in M ϕ -Mo-ratio.

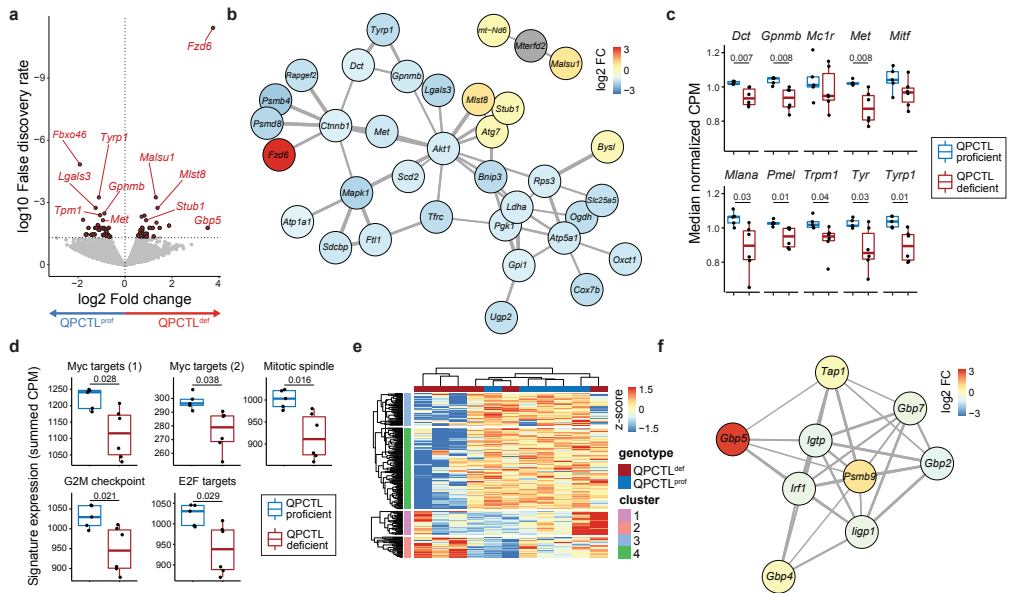


Figure 3. QPCTL deficiency results in suppression of melanogenesis and cell metabolism. mRNA sequencing was performed on sorted CD45-negative cells from QPCTL-proficient ($n = 5$) and QPCTL-deficient ($n = 6$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. **(a)** Differential gene expression analysis comparing CD45-negative cells obtained from QPCTL deficient versus QPCTL-proficient TMEs. Genes with a false discovery rate (FDR) < 0.05 are indicated in red. Selected genes are indicated in the plot. **(b)** Network analysis (StringDB) performed on all significantly (FDR < 0.05) differentially expressed genes. Genes with a medium interaction strength (> 0.4) are included. Line thickness indicates interaction strength. Nodes are colored based on log₂ fold differences obtained in **a**. **(c)** Transcript abundance of selected genes in the melanogenesis pathway. Boxplots indicate group median and 25th/75th percentiles, whiskers indicate the interquartile range multiplied by 1.5, dots signify individual samples. **(d)** Signature expression of cell cycle-associated hallmark signatures from MSigDB, calculated as the summed CPM of all genes within each signature. Boxplots indicate group median and 25th/75th percentiles, whiskers indicate the interquartile range multiplied by 1.5, dots signify individual samples. **(e)** Hierarchical clustering of the 1,000 most differentially expressed genes across all samples, depicted as a row-normalized heatmap. **(f)** Network analysis (StringDB) performed on genes from cluster 2 **(e)**. Genes with a medium interaction strength (> 0.4) are included. Line thickness indicates interaction strength. Nodes are colored based on log₂ fold differences obtained in **a**. P values were determined by one-way ANOVA followed by Tukey's HSD test **(c, d)**. Significant P values (< 0.05) are indicated in the plots. Data are representative of 2 independent experiments. CPM, counts per million; MSigDB, Molecular Signatures Database.

QPCTL deficiency is associated with suppressed intra-tumoral melanogenesis and cell metabolism

Having established that QPCTL deficiency is associated with an alteration in intra-tumoral immune cell composition, we set out to investigate the effect of QPCTL deficiency on the tumor cell and stromal cell compartment of the TME. RNA sequencing of CD45-negative cell fractions from QPCTL-deficient and -proficient TMEs showed that QPCTL deficiency resulted in differential expression of a substantial set of genes (**Fig. 3a**, **Supplementary Fig. 5a**), and network analysis revealed that expression of multiple genes involved in melanogenesis (*Dct*, *Tyrp1*, *Gpnmb*) was reduced in QPCTL-deficient melanomas (**Fig. 3b**). Assessment of the expression level of a broader set of genes involved in melanogenesis likewise showed dampening of this pathway (**Fig. 3c**). Interestingly, these transcriptional changes coincided with the functional abrogation of melanin production, as evidenced by a loss of pigmentation of QPCTL-deficient tumors (**Supplementary**

Fig. 5b). Expression of a network of genes involved in cell cycle (*Mapk1*, *Akt1*) and cell metabolism (*Pgk1*, *Atp5a1*, *Oxct1*) was additionally found to be decreased in CD45-negative cells in the QPCTL-deficient setting (**Fig. 3b, d, Supplementary Fig. 5c**). While dampened expression of these gene sets was consistently observed, effect sizes were small and had no discernible effect on tumor outgrowth (**Supplementary Fig. 2b-c**).

To further explore putative transcriptional alterations in the CD45-negative compartment as a result of QPCTL deletion, obtained transcriptomes were clustered based on the top 1,000 most variable genes, revealing a cluster of genes that was enriched in 3 out of 6 QPCTL-deficient samples (gene cluster 2, **Fig. 3e**). Network analysis performed on this cluster showed that this gene set contained a small network comprised of IFN induced genes (*Gbp* family members, *Tap1*, *Irf1*; **Fig. 3f**). Interestingly, previous studies have shown that IFN γ can act as a suppressor of melanogenesis in B16F10 melanoma cells^{15–17}, a notion that potentially links this IFN signature to the observed decrease in tumor pigmentation.

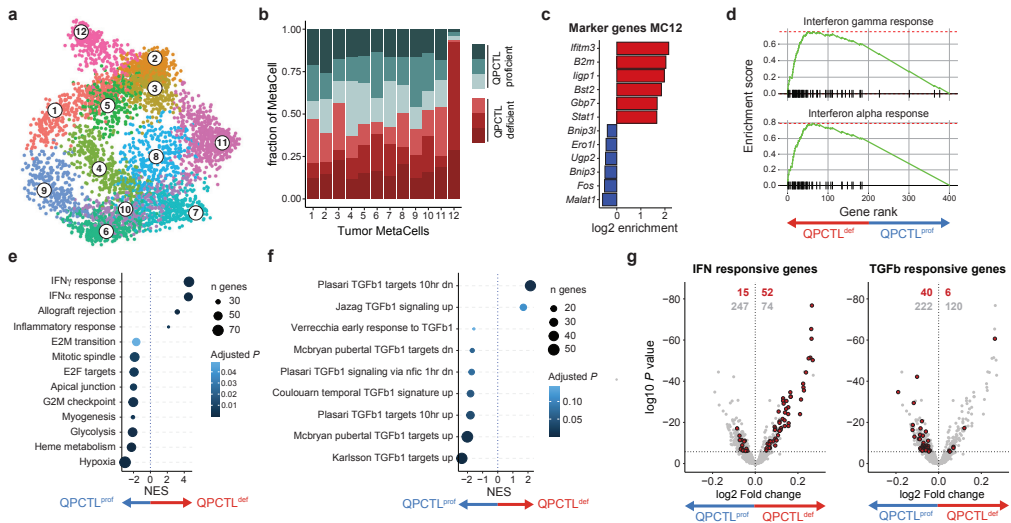


Figure 4. QPCTL deficiency leads to an increased IFN- and decreased TGF-b-response signature in tumor cells. scRNA sequencing was performed on sorted live cells from QPCTL-proficient ($n = 3$) and QPCTL-deficient ($n = 3$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. (a) 2-dimensional MetaCell projection of the tumor cell compartment. Single cells are colored by MetaCell. (b) Stacked barchart depicting the sample composition of each tumor cell MetaCell. Cell counts from each sample were normalized to 1,000 cells. (c) Enrichment of marker genes (6 highest and lowest expressed) in tumor cell MetaCell 12. (d) Gene set enrichment analysis performed on the top and bottom 200 genes expressed by MC12 (see **Supplementary Fig. 7b**). Gene-enrichment plots for the IFN γ and IFN α response gene-sets are depicted. (e-f) Differential gene expression analysis comparing tumor cells derived from QPCTL-proficient and QPCTL-deficient TMEs, followed by gene set enrichment analysis using either hallmark (e) or immunologic (f) gene sets from MSigDB. Results obtained from the immunologic signature gene sets were filtered for those containing "TGFb". Gene sets with a $P < 0.05$ are shown. (g) Volcano plots depicting differential gene expression analysis. Horizontal line indicates an adjusted P value cutoff of 0.05. IFN (left) or TGF-b (right) signature genes are highlighted in red (see **Supplementary Table 1** for signature genes). Red numbers denote quantity of significant differentially expressed genes within the signature, grey numbers denote the quantity of remaining differentially expressed genes. Depicted data were obtained in a single experiment, consisting of 6 mice. NES, normalized enrichment score; MSigDB, Molecular Signatures Database.

Single cell transcriptomic profiling reveals remodeling of the tumor micro-environment by QPCTL deficiency

The above data indicate that QPCTL inactivation affects both immune cell composition of the TME and the transcriptome of non-immune cells at the tumor site. With the aim to potentially link these two observations we inoculated QPCTL^{+/+} and QPCTL^{-/-} mice with QPCTL-WT and QPCTL-KO B16F10 cells, respectively, and performed scRNAseq of both immune cells and non-immune cells from the resulting QPCTL-deficient and -proficient TMEs. Applying the MetaCell algorithm¹⁸ on 13,093 transcriptomes derived from 6 TMEs (3 QPCTL-proficient, 3 QPCTL-deficient), showed 3 transcriptionally divergent cell supertypes, reflecting immune cells (*Ptprc*, *Irgam*), fibroblasts (*Col1a1*, *Acta2*), and tumor cells (*Mlana*, *Pmel*) (**Supplementary Fig. 6**). To identify cell type-specific changes that accompany QPCTL deficiency, each of these supertypes was subsequently re-clustered and analyzed separately.

To investigate which of the transcriptional changes observed in the CD45-negative compartment could be mapped to the tumor cell compartment, this supertype was re-grouped into transcriptionally disparate MCs (**Fig. 4a, Supplementary figure 7a**), and the relative contribution of cells from either QPCTL-deficient mice or QPCTL-proficient samples to the different MCs was examined. Strikingly, 1 MC (MC12) was nearly exclusively observed in QPCTL-deficient samples (**Fig. 4b**). Examination of the marker genes of MC12 showed a prominent presence of IFN induced transcripts *Ifitm3*, *B2m*, *Bst2* and *H2-D1* (**Fig. 4c**), and gene-set enrichment analysis performed on MC12 marker genes identified both IFN γ and IFN α response as the strongest enriched gene-sets (**Fig. 4d, Supplementary figure 7b-c**). IFN γ and IFN α/β response signatures show a high level of overlap¹⁹⁻²¹, making it difficult to assign the observed response to either cytokine. To potentially deconvolute these signatures, and diagnose other putative cytokine response signatures, the CytoSig model¹⁹ was applied to MC12. In line with the GSEA, this analysis detected transcription response signatures of all IFN types, with the highest score being observed for IFN γ (**Supplementary figure 7d**). CytoSig additionally identified a reduction in TGF- β signaling in MC12, and retrospective analysis of the bulk RNAseq data revealed reduced expression of TGF- β responsive genes in QPCTL-deficient TMEs (**Supplementary figure 7e**). Furthermore, differential gene expression analysis showed that QPCTL deficiency was associated with a reduced TGF- β and an increased IFN responsive signature (**Fig. 4e-g**) across all MCs, potentially indicating either an altered abundance of—or sensitivity to—these cytokines. Notably, in vitro sensitivity of B16F10 cells to both TGF- β and IFN γ was unaltered by QPCTL deficiency (**Supplementary figure 7f-i**), arguing in favor of an altered abundance of these cytokines in the TME.

Next, the immune cell compartment was grouped into 11 MetaCells (MCs), classified as either CD3⁺ lymphocytes (*Cd8a*, *Cd3e*), dendritic cells (*Ccr7*, *H2-Aa*), or macrophages/monocytes (*Adgre1*, *Fcgr1*; **Fig. 5a-b, Supplementary Fig. 8a**). The macrophage/monocyte (M ϕ /Mo) MCs could be further subdivided into 2 groups that were marked by high expression of either *Ccr2* and *Irga4* (M ϕ /Mo-1 subgroup) or *Ms4a7* and *Pf4*^{fl} (M ϕ /Mo-2 subgroup; **Fig. 5b-c**), suggestive of a blood- versus tissue-derived origin²²⁻²⁵. Analysis of the contribution of cells from QPCTL-deficient and -proficient TMEs to individual MCs indicated that QPCTL deficiency changed the relative abundance of the

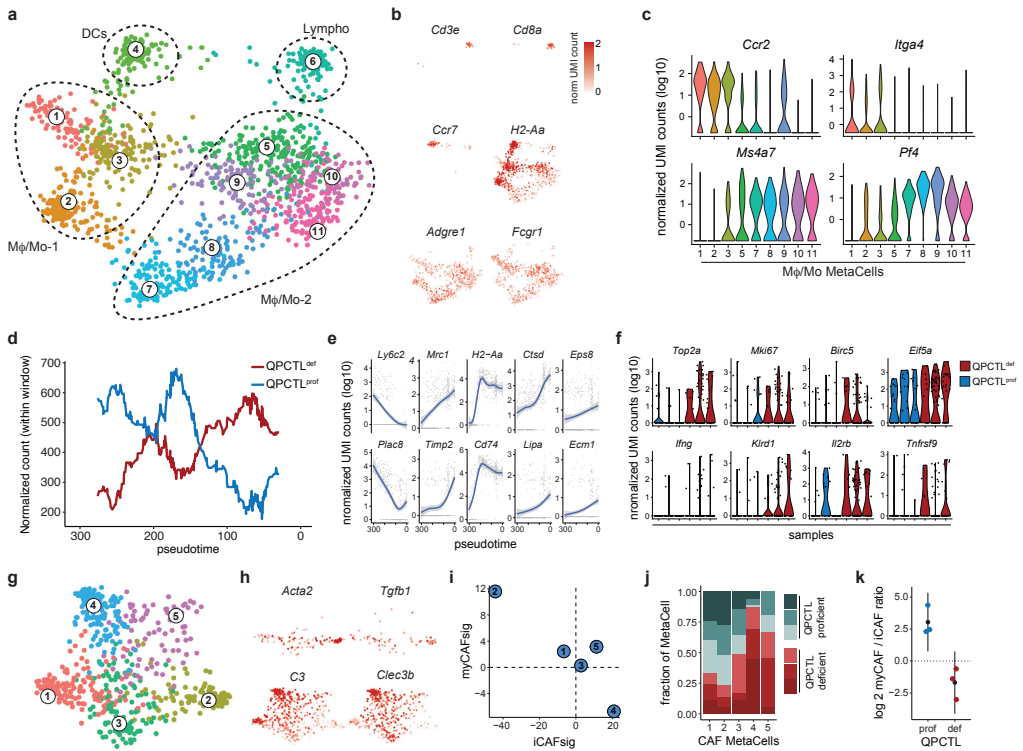


Figure 5. QPCTL deficiency alters the immune cell compartment and CAF polarization in the TME. scRNA sequencing was performed on sorted live cells from QPCTL-proficient ($n = 3$) and QPCTL-deficient ($n = 3$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. **(a, b)** 2-dimensional MetaCell projection of the immune cell compartment. Single cells are colored by MetaCell **(a)**, or normalized UMI count **(b)** of selected genes. **(c)** Violin plots depicting normalized UMI counts of selected genes across M ϕ /Mo MCs. **(d, e)** SlingShot Trajectory analysis performed on M ϕ /Mo subset 1 (MC1, 2 and 3). **(d)** QPCTL-deficient or QPCTL-proficient TMEs replicates were pooled, and normalized cell counts were tallied within windows of 60 cell wide, sliding 1 cell per frame. Lines indicate normalized cell counts within each window. **(e)** Normalized UMI counts of selected genes that are significantly associated with pseudotime. Blue lines indicate general additive linear models, grey areas indicate confidence intervals, grey dots represent single cells. **(f)** Violin plots depicting normalized UMI counts of selected genes within the CD3⁺ lymphoid cell MetaCell (MC6). **(g, h)** 2-dimensional MetaCell projection of the fibroblast compartment. Single cells are colored by MetaCell **(g)**, or normalized UMI count **(h)** of selected genes. **(i)** Enrichment of iCAF and myCAF signatures (**Supplementary Table 2**) in each CAF MetaCell. Signature values represent summed log₂ transformed enrichment values, calculated using the MetaCell algorithm. **(j)** Stacked barchart depicting sample composition of each CAF MetaCell. Cell counts from each sample were normalized to 1,000 cells. **(k)** myCAF/iCAF ratio detected in QPCTL-proficient and -deficient TMEs. Colored dots indicate individual mice, black dots indicate means, whiskers indicate the standard deviation. Depicted data were obtained in a single experiment, consisting of 6 mice. iCAF, inflammatory cancer-associated fibroblast; myCAF, myofibroblastic cancer-associated fibroblast; UMI, unique molecular identifier.

different cell states that jointly comprised the M ϕ /Mo-1 subgroup (**Supplementary Fig. 8b**). As intra-tumoral M ϕ /Mo cells can exist within a continuum of transcriptional cell states^{26,27}, pseudotime analysis was performed on M ϕ /Mo subgroup 1 to examine if such a continuum could be observed, and whether this was linked to QPCTL deficiency. This analysis demonstrated a strong continuous association between pseudotime (i.e. cell state) and sample-origin (**Fig. 5d**). To investigate the transcriptional changes underlying this association, genes were clustered based on their expression kinetics across pseudotime (**Supplementary Fig. 8c**), revealing a gradual loss of expression of Mo-related genes *Ly6c2* and *Plac8* across pseudotime (**Fig. 5e**), with the lowest expression levels

found in the area that contained the highest fraction of cells from QPCTL TMEs. At the same time, an increase in transcripts linked to ‘inflammation-resolatory’ Mf’s (*Mrc1*, *Timp2*), antigen-presentation (*H2-Aa*, *Cd74*) and Mf effector function (*Eps8*, *Ctsd*, *Ecm1*, *Lipa*)^{28–31} was observed. Consistent with these findings, Mf/Mo cells that were more dominant in QPCTL-deficient samples had reduced expression of monocyte-associated transcripts, and displayed transcriptional similarity to previously identified tumor-associated macrophage subsets³² (**Supplementary Fig. 8d**). Together with the observed Mf-Mo skewing (**Figure 2**), these data argue in favor of a model in which QPCTL deficiency in the TME leads to transcriptional changes that drive Mo-to-Mf conversion. Moreover, the intra-tumoral macrophages identified by flow cytometry (**Figure 2**) exhibited high-level surface expression of MHC class II (*H2-Aa*), CD206 (*Mrc1*) and CCR2 (*Ccr2*), linking these cells to the Mf/Mo population identified through scRNAseq (**Supplementary Fig. 8e**).

Congruent with the flow cytometric analysis, no differences in the frequencies of CD3⁺ lymphoid cells derived from QPCTL-deficient versus QPCTL-proficient TMEs could be observed in the scRNAseq dataset. However, as activated lymphoid cells are potent producers of IFN, we queried whether transcriptional features associated with lymphocyte activation were detected more frequently in lymphocytes from QPCTL-deficient TMEs. Interestingly, expression of genes associated with TCR-triggering (*Ifng*, *Il2rb* and *Tnfrsf9*) and cell cycle activity (*Top2a*, *Mki67*, *Birc5*) was detected more frequently in cells derived from the QPCTL-deficient tumors (**Fig. 5f**). Likewise, lymphocytes derived from QPCTL-deficient TMEs showed increased expression of the *Ccl3*, *Ccl4*, and *Ccl5* chemokines, and of the T cell activation-related genes *Ly6a*, *Nkg7*, and *Gzmb* (**Supplementary Fig. 8f**). While the increase in *Ifng* gene expression in lymphocytes in QPCTL-deficient samples was only modest, the parallel observation of other aspects of lymphocyte activation in these samples suggests that these cells may, at least in part, be responsible for the IFN responsive signature that is observed in the tumor cell compartment.

Diverse subsets of cancer-associated fibroblasts (CAFs) that possess distinct immunomodulatory functions have been reported in the TME of different cancer types³³, and two highly distinct populations— termed TGF- β -producing myfibroblastic CAFs (myCAFs) and IL-1-driven inflammatory CAFs (iCAFs)—have been identified in a recent set of cross-species studies^{34,35}. As a reduced TGF- β response signature was identified as one of the characteristics of tumor cells in QPCTL-deficient tumors, we next asked whether QPCTL deficiency affected CAF polarization. MetaCell-based clustering within the fibroblast cell supertype resulted in 5 transcriptionally distinct MCs (**Fig. 5g**, **Supplementary Fig. 8g**). Assessment of transcripts known to be involved in CAF function showed that MC2 was enriched for *Tgfb1* transcripts and several myCAF markers (e.g. *Acta2*, *Itgb1*), whereas MC4 exhibited more pronounced expression of genes involved in functional inhibition of TGF- β (*Ltbp1*, *Dcn*) and multiple iCAF markers (*C3*, *Clec3b*; **Fig. 5g**, **Supplementary Fig. 8h**). In line with this, analysis of signature enrichment-scores pertaining to these two subsets showed that MC2 and MC4 scored the highest for either the myCAF or iCAF signature, respectively (**Fig. 5h**). Moreover, increased surface expression of the myCAF and iCAF markers CD29/ITGB1 and Ly6C was detected on MC2 and MC4, respectively (**Supplementary Fig. 8i**). Strikingly, MC2 and MC4 displayed the highest depletion and enrichment in QPCTL-deficient and -proficient

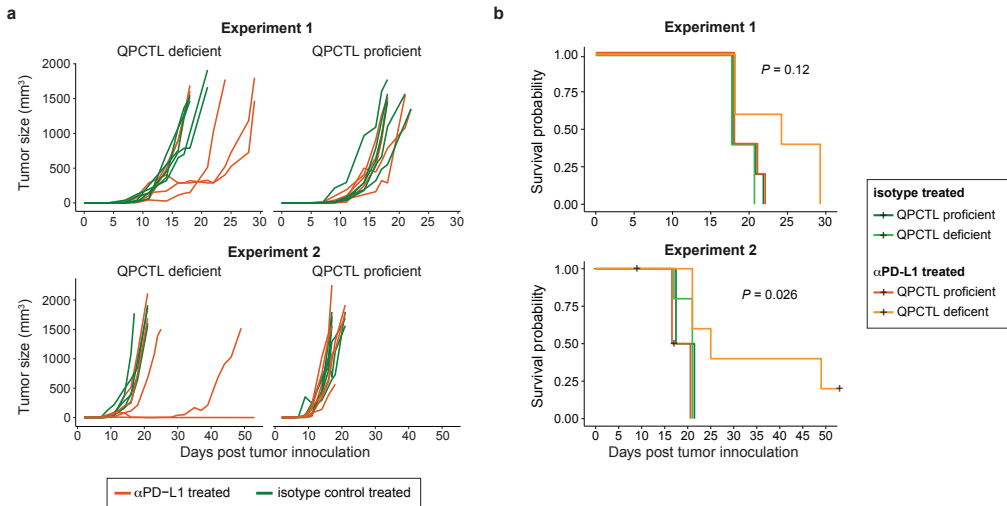


Figure 6. QPCTL deficiency sensitizes the tumor microenvironment to anti-PD-L1 treatment. QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with QPCTL-WT and QPCTL-KO B16F10 melanoma cells, respectively. Each group subsequently received either αPD-L1 or isotype control antibody treatment at day 7, 9 and 11 post tumor inoculation. **(a)** Tumor growth curves, assessed until day 50 post tumor inoculation. Lines represent individual mice. Data from two experiments are depicted ($n = 5$ per group). **(b)** Survival probabilities of mice treated with αPD-L1 or isotype control antibody in a QPCTL-proficient and -deficient setting. Black plus-signs indicate censored events. Data from two independent experiments are depicted. Global P values were determined by log-rank test **(b)**. Data from 2 independent experiments are depicted.

TMEs (Fig. 5i), resulting in a 20-fold increase in the iCAF/myCAF ratio in QPCTL-deficient TMEs (Fig. 5j). Thus, in the absence of QPCTL activity, polarization of fibroblasts toward TGF- β -producing myCAFs is reduced in favor of the more pro-inflammatory iCAFs.

QPCTL deficiency enhances susceptibility of B16F10 tumors to anti-PD-L1 treatment

The B16F10 melanoma commonly shows a poor response to single agent PD1/PD-L1 checkpoint blockade^{36,37}. Having observed that QPCTL deficiency alters the TME to a more proinflammatory state, we hypothesized that QPCTL deficiency may modulate the sensitivity of B16F10 tumors to such PD-1/PD-L1 blocking therapies. To test this, QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with QPCTL-WT and QPCTL-KO B16F10 cells, respectively. Upon tumor formation (6 days post-inoculation) mice were treated with either an anti-PD-L1 or isotype control antibody. In QPCTL-proficient animals, tumor growth progressed rapidly and was not influenced by anti-PD-L1 therapy (Fig. 6a). In contrast, anti-PD-L1 therapy did result in improved tumor control in QPCTL-deficient mice (Fig. 6a), and led to a prolonged survival in approximately 50% of animals (Fig. 6b), providing direct evidence that the TME modulatory effect of QPCTL removal has functional consequences.

Discussion

QPCTL activity is known to influence the properties of a number of molecules that are active in the TME and may potentially influence additional—as of yet unidentified—substrates. To obtain a global view of the cumulative effects of QPCTL activity on the host's immune response to tumor growth, we made use of a QPCTL^{-/-} mouse model in combination with syngeneic QPCTL-KO tumor cell lines. We conclude that inactivation of QPCTL alters M ϕ -Mo abundance, increases IFN pathway activity relative to TGF- β pathway activity, and leads to a profound increase in iCAFs relative to myCAF in the TME (**Supplementary Fig. 9a**). In line with the observed skewing of the TME to a pro-inflammatory state that is induced by QPCTL deficiency, we demonstrate that such deficiency leads to the sensitization of B16F10 melanomas to anti-PD-L1 treatment.

The current study has the following limitations: 1) Germline deletion of QPCTL may potentially lead to developmental alterations that influence the host's response to tumor challenge, e.g. affecting the capacity of certain CAF or immune subsets to differentiate, independent of QPCTL activity during tumor outgrowth. However, the absence of clear phenotypic alterations at baseline, and the fact that an increased M ϕ -Mo-ratio was observed in wild-type mice challenged with QPCTL-deficient tumor cells, argue against this possibility. 2) In the present study we have aimed to model the effects of depletion of QPCTL activity on the tumor micro-environment, whereas glutaminyl cyclase inhibitors will, based on the similarity of their active sites, likely inhibit both QPCTL and QPCT activity^{38,39}. Such inhibition of QPCT may be relevant as siRNA-mediated suppression of QPCT has been shown to reduce expression of CCL2, CX₃CL1 and CD54/ICAM¹². In future work, dual inactivation of QPCTL and QPCT may form a means to test this.

The depletion of monocytes that we observe in QPCTL-deficient TMEs may potentially be explained by a decreased functionality of the CCL2-CCR2 signaling axis. In pre-clinical models of breast cancer, the CCL2-CCR2 axis has been shown to influence the abundance of monocytes in primary tumors⁴⁰ and metastatic lesions⁴¹. Furthermore, monocyte recruitment was found to be reduced after thioglycolate challenge of mice that were either QPCTL-deficient or treated with QPCT/QPCTL inhibitors¹⁰. However, it is important to note that at high concentrations, pyroGlu-CCL2 and unmodified CCL2 demonstrate similar chemotactic activity¹⁰, and the effect of impaired pyro-glutamylated CCL2 will therefore depend on local concentrations.

Contrary to expectations, we observed a relative increase in macrophage frequencies in QPCTL-deficient TMEs. Transcriptomic profiling of intra-tumoral M ϕ /Mo cells revealed that these macrophages expressed monocyte-associated molecules (e.g. *Ccr2*, *Itga4*) and pseudotime analysis suggests the existence of intermediate M ϕ /Mo cell states, together arguing in favor of their monocytic origin. In prior work, abrogation of the CCR2-CCL2 signaling axis in monocytes has been shown to strongly reduce the accumulation of intra-tumoral macrophages⁴². Based on these data, we propose that the boosting of intra-tumoral macrophages by QPCTL inactivation occurs through a mechanism that is independent of CCL2, and is potentially driven by an accelerated monocyte-to-macrophage differentiation program.

Having observed a number of independent alterations in the TME that are induced by QPCTL deficiency, it is of interest to speculate on the possible causal relationship between these individual changes. One possible scenario (**Supplementary Fig. 9b**) is that suppression of the CCR2-CCL2 axis due to the lack of QPCTL activity causes an early reduction in the influx of monocytes, which have been shown to form a major source of TGF- β ⁴³. Such an initial deficit in TGF- β abundance could potentially limit myCAF polarization and favor differentiation toward iCAFs, relieving TGF- β -driven suppressive effects on myeloid and lymphoid effector cells. Notably, TGF- β can act as a suppressor of IFN γ production by NK cells^{44,45} and CD8⁺ T cells^{46,47}, and increased abundance of activation-associated transcripts was observed in lymphocytes from QPCTL-deficient TMEs, together suggesting that this cell pool may be the source of the observed IFN-response signature. Genetic ablation experiments (**Figure 2**) indicated that CD47 does not play a significant mechanistic role in the TME-modulatory effects of QPCTL in the B16 melanoma model. However, as the CD47/SIRP α axis acts primarily through inhibition of activating signals of ITAM-containing receptors, such as activating Fc receptors on myeloid cells, it is plausible that the observed synergy between QPCTL deficiency and treatment with an opsonizing anti-PD-L1 antibody is at least partially mediated through its effect on the CD47 pathway.

In summary, our data provide evidence that removal of QPCTL activity can shift the TME from an immunosuppressive (monocyte skewed, myCAF, TGF- β) towards a pro-inflammatory (macrophage skewed, iCAF, IFN) milieu, and acts synergistically with anti-PD-L1 therapy to enhance tumor control and survival. If this TME-remodeling effect can also be achieved through pharmacological inhibition of QPCTL activity in human cancers, such inhibitors may offer potential in combination treatment strategies that include checkpoint blocking antibodies and/or tumor-opsonizing antibodies.

Methods

Generation of transgenic mice

C57/Bl6JR mice were obtained from Janvier. QPCTL^{-/-} mice carrying an 811bp deletion in exon 2 of the *Qpctl* gene were generated on the C57BL/6JRj background using pronuclear microinjection in mice zygotes with a CRISPR/Cas9 mixture (50 ng/ml Cas9 RNA and 25 ng/ml sgRNA, in water). The sgRNA was targeted to the second exon of the *Qpctl* gene (5'-GCACAATCAATAAGGGACGC-3'). QPCTL^{-/-} mice and QPCTL^{+/+} mice were identified by PCR using the following primers: Fwd_KO (5'-GTTTTAGGGATGGATGCCGC-3'), located before the 811bp deletion, Fwd_WT (5'-GGACTCCTAGTAGGCAACGG-3'), located in the 811bp deletion, and Rev (5'-GGCTGTTTTGGGATCTTCGG-3'), located after the 811bp deletion.

Evaluation of mouse blood cell counts

Whole blood of mice was collected by heart puncture and total cell counts were determined using a

DxH500 Hematology Analyzer (Beckman Coulter).

Peripheral blood collection and preparation

Whole blood of mice was collected into heparin-coated tubes by heart puncture or tail vein puncture at indicated time points. Samples were incubated twice for 5 minutes in erythrocyte-lysis buffer (0.15M NH_4Cl , 10mM KHCO_3 , 0.1 mM EDTA, pH 7.4), and washed once in staining buffer (0.5% BSA in PBS). Cells were then used for antibody staining, as described below.

Cell lines

B16F10 cells and MC38-AMS cells were kindly provided by D. Peeper (Division of Molecular Oncology & Immunology, Oncode Institute, The Netherlands Cancer Institute, Amsterdam, The Netherlands). The MC38-AMS cell line is a variant of the MC38 cell line available from Kerfast. Whole exome sequencing was performed to compare the MC38-AMS and MC38-Kerfast line, and data have been uploaded to the Sequence Read Archive. B16F10 and MC38-AMS cells were cultured in DMEM (Gibco) supplemented with 10% FCS and penicillin-streptomycin. Cells were cultured at 37 °C and 5% CO_2 .

CRISPR/Cas9-mediated generation of CD47 and QPCTL knockout cells

To generate QPCTL- knockout (KO), CD47-KO, and WT control B16F10 cell lines, cells were transfected with pLentiCRISPR v.2 vector encoding sgRNA targeting the murine QPCTL (5'-TATTGATTGTGCGACCCCCG-3') or CD47 (5'-AGCAACAGCGCCGCCGCAA-3') gene, or left untransfected. Culture medium of transfected cells was supplemented the next day with 2 $\mu\text{g ml}^{-1}$ puromycin for at least 2 days. Selected cells were expanded, and subsequently sorted on the basis of $\alpha\text{mCD47-MIAP301}^{\text{LO}}$ mSIRP α -Fc $^{\text{LO}}$ phenotype (in case of CD47 knockout), or $\alpha\text{mCD47-MIAP301}^{\text{HI}}$ mSIRP α -Fc $^{\text{LO}}$ phenotype (in case of QPCTL knockout), in order to obtain bulk knockout populations. WT control B16F10 cells were sorted based on morphology gating only. Next, single cells were isolated and expanded, and approximately 50 knockout or wild-type clones were pooled to obtain pure knockout or wild-type populations. To generate CD47/QPCTL double KO (dKO) cell lines, B16F10 QPCTL KO cell lines were transfected with pLentiCRISPR v.2 vector encoding sgRNA targeting the murine CD47 gene. 1 day after transfection, culture medium was supplemented with 2 $\mu\text{g ml}^{-1}$ puromycin for at least 2 days. Single cells were isolated and expanded, and 12 clones were pooled to obtain knockout populations. To generate QPCTL-KO and control MC38-AMS cell lines, cells were transduced with pLentiCRISPR v.2 vector encoding sgRNA targeting the murine QPCTL gene or a non-targeting control gRNA. 2 days after transduction, culture medium was supplemented with 2 $\mu\text{g ml}^{-1}$ puromycin for at least 4 days. Next, single cells were isolated and expanded, and 12 knock-out or control clones were pooled to obtain cell populations for further use. Gene disruption was validated by sequence analysis of the relevant gene locus by TIDE⁴⁸ analysis and, in case of CD47, by flow cytometry.

Tumor challenge

To analyze the effect of QPCTL deficiency in both host and tumor cells or in host cells only, 8- to 25-week-old male and female QPCTL^{-/-} or wild-type QPCTL^{+/+} littermate controls were injected with 2×10^5 of the indicated B16F10 tumor cell line in a 100 μ L solution of PBS (Lonza) and Matrigel (Corning) (1:1) in the right flank on day 0. To analyze the effect of tumor cell CD47 deficiency or CD47 and QPCTL-double deficiency, 9–12-week-old C57/Bl6JR (female; Janvier) were injected with 2×10^5 of the indicated B16F10 tumor cell line. To analyze the effect of QPCTL deficiency in both host and tumor cells in MC38-AMS tumors, 8–25-week-old QPCTL^{-/-} or wild-type QPCTL^{+/+} littermate controls were injected with 5×10^5 of the indicated MC38-AMS tumor cell line. Tumors were measured 3 times a week, and mice were sacrificed 13–17 days (B16F10 tumors) or 21–29 days (MC38 tumors) after tumor challenge. Mice with a tumor volume equal or below 40mm³ were excluded and tumors used for subsequent flow cytometry analyses ranged from 75–1436 mm³ (B16F10 tumors) or 112.5 - 786.5mm³ (MC38 tumors).

TME single-cell preparation

Tumors were fragmented on ice and were subsequently digested in DMEM (10 ml per tumor) supplemented with collagenase IV (2 mg ml⁻¹, Sigma Aldrich) and DNase I (50 μ g ml⁻¹, Sigma Aldrich) for 30 min at 37 C. Subsequently, 40 ml DMEM supplemented with 10% FCS was added per tumor, and cell suspensions were passed through 100 μ m filters. Next, samples were incubated for 5 minutes in erythrocyte-lysis buffer (0.15M NH₄Cl, 10mM KHCO₃, 0.1 mM EDTA, pH 7.4), and washed once in staining buffer (0.5% BSA in PBS). Tumor single cell suspensions were then counted and used for antibody staining.

Flow cytometry

Cell surface CD47 was assessed by staining of blood immune cells with the anti-mouse CD47 antibody MIAP301 at a dilution of 1:100 or 1:200 plus His-tagged recombinant mouse SIRP α (rmSIRP α -His) (4, 12 or 36 μ g ml⁻¹), in PBS containing 0.5% (w/v) BSA (Sigma) and 0.2% (w/v) sodium azide (Sigma) (FACS buffer) for 30 min at room temperature, protected from light. After two washes with FACS buffer, cells were stained with a fluorochrome-labeled anti-His antibody at a dilution of 1:100 or 1:200 for 30 min at 4 °C in FACS buffer, while protected from light. Cells were then washed with FACS buffer, and DAPI, propidium iodide, or 7-AAD Viability Staining Solution (eBioscience) was added to allow dead cell exclusion. Antibodies used to analyze immune cells in tumor single cell suspensions are listed in **Supplementary Table 3**. Measurements were performed on an LSRII, LSRFortessa, or FACSCantoII (BD Biosciences). Data were analyzed using FACS Diva software (BD Biosciences) and FlowJo software.

Unbiased flow cytometry data analysis

Samples were preprocessed using FlowJo software, compensating for spectral overlap, selecting IR-Dye⁻CD45⁺ single cells and removing outlier cells. Further analysis was performed in R, implementing the FlowCore package⁴⁹. Samples were subsampled to obtain 10,000 or 30,000 total

cells for spleen or tumor analysis, respectively. Next, a logicle (biexponential) transformation was applied to the measured fluorescence intensities. Uniform Manifold Approximation and Projection (UMAP) was used for dimension reduction, and subsequently used for hierarchical clustering by Euclidean distance (Ward's method). Relative contributions of cells derived from QPCTL^{-/-} and QPCTL^{+/+} samples to each of the clusters was then assessed.

IFN γ and TGF- β sensitivity of B16F10 cell lines

For IFN γ sensitivity testing, QPCTL-WT or QPCTL-KO cell lines were seeded on 6-well plates at 50,000 cells per well, incubated at 37 °C for 3 hours to allow cells to adhere, and subsequently treated with indicated amounts of IFN γ (Mouse IFN-gamma Recombinant Protein, ThermoFisher Scientific) for 21 hours. Cells were then harvested with trypsin-EDTA (Gibco), washed twice with PBS, and examined either through flow-cytometry or western blotting. For TGF- β sensitivity testing, QPCTL-WT or QPCTL-KO cell lines were seeded on 6-well plates at 50,000 cells per well and incubated at 37 °C for 16 hours. Next, cells were pre-incubated at 37 °C for 4 hours in culture medium containing 0.2% FCS, and subsequently incubated at 37 °C with indicated quantities of recombinant mouse TGF- β 1 protein (R&D systems) for 1 hour in culture medium containing 0.2% FCS. Cells were then harvested with trypsin-EDTA (Gibco), washed twice with PBS, and examined through western blotting. For flow-cytometry cells were stained with anti-PD-L1-BV421 and anti-H2-Kb-PE (both 1:100 dilution, see Supplementary Table 3) for 15 minutes at room temperature, washed twice with FACS buffer, and analyzed on an LSRFortessa (BD Biosciences). For western blot analyses, cells were incubated on ice for 30 minutes in 200 ml RIPA buffer (1% Triton X100, 0.1% Sodium deoxycholate, 0.1% SDS, 1 mM EDTA, 10 mM Tris pH 8, 140 mM NaCl) supplemented with Halt™ Protease and Phosphatase Inhibitor Cocktail (ThermoFisher Scientific), followed by pulse mixing on a Vortex Genie (Scientific Industries). Lysates were then centrifuged at 20,000x *g* for 20 minutes at 4 °C and protein concentrations in the resulting supernatants were determined using Pierce™ BCA Protein Assay Kit (ThermoFisher Scientific) according to manufacturer's protocol. Next, equal amounts of protein were processed using the Novex NuPAGE Electrophoresis system (Thermo Fisher Scientific) and Trans-Blot Turbo Transfer system (Bio-Rad) according to the manufacturers' instructions. Membranes were blocked using Western Blocking Reagent (Roche) for 1 hour, and subsequently stained overnight at 4 °C with indicated primary antibodies diluted in Western Blocking Reagent (see Supplementary Table 3 for antibody information) followed by 2 hours at 4 °C with either anti-rabbit or anti-mouse secondary antibodies conjugated to HRP (see Supplementary Table 3 for antibody information). Western blots were imaged using the ChemiDoc MP Imaging System (Bio-Rad).

RNA sequencing

RNA was extracted from the indicated frozen tissues using the RNeasy Mini Kit (Qiagen). Cell populations isolated by FACS were washed once in PBS, and subsequently lysed in RLT buffer (Qiagen). Whole transcriptome sequencing samples were prepared with the TruSeq Stranded

mRNA Kit (Illumina). Paired-end 50bp sequencing was performed on a NovaSeq 6000 system (S1 flowcell, Illumina), obtaining an average of 18×10^6 reads per sample. Reads were aligned to the pre-built GRCm38 genome_snp_tran reference using HISAT2⁵⁰, and transcript counts were obtained using an in-house generated pipeline (GenSum, <https://github.com/NKI-GCF/gensum>). Differential gene expression analysis was performed using the edgeR package⁵¹. Network analysis was performed using the stringDB database, applying the igraph package for visualization.

Single cell RNA sequencing analyses

Single-cell digests of QPCTL^{-/-} and QPCTL^{+/+} TMEs were generated as outlined above. Cells were stained with IR-Dye for dead cell exclusion and with anti-mouse TotalSeqä Hashtag antibodies (TotalSeq-A0301-06, Biolegend), pooled in equal numbers, and were single-cell sorted on a BD Fusion cell sorter. Single-cell RNA isolation and library preparation was performed according to the manufacturer's protocol of the 10X Genomics Chromiumä Single Cell 3' kit, and the cDNA library was sequenced on the NextSeq™550 Sequencing System (Illumina). A total of $\pm 3.7 \times 10^8$ reads resulted in the detection of 14,888 cells with a median of 3,344 detected genes per cell. Feature-barcode matrices were generated using the Cell Ranger software of the 10X Genomics Chromium pipeline. Further processing was subsequently performed using the MetaCell¹⁸ and Seurat R packages⁵². Cells that contained less than 500 UMIs or had a mitochondrial transcript fraction of >0.2 were removed. Next, variable genes across the dataset were identified with a normalized variance/mean threshold at 0.1 and a down-sampled coverage threshold at 80, yielding 1,021 genes. These genes were subsequently used as anchors to search for gene-gene correlations across the dataset, and genes with correlations of >0.1 were included. The obtained genes were then clustered into 50 separate gene-modules, and each was annotated manually (**Supplementary Table 4**).

To identify the major cell types, a feature-gene list was compiled of gene-modules that contained marker genes for various cell types (modules 10, 17, 19, 21, 24, 26, 29, 30, 33, 39, 40, 44, 47, 48, 49; Supplementary Table 3), and these feature-genes were used to generate MetaCells. The obtained MetaCells were then classified as either immune cells, fibroblasts, or tumor cells, as shown in **Supplementary figure 6**. MetaCells that contained significant expression of marker genes from multiple cell types were identified as 'doublet MetaCells', and excluded from further analysis.

Subsequent analysis was performed within each individual cell type. In brief, doublet detection was performed using the HTODemux function of Seurat, setting the positive quantile at 0.99. Cells containing either a high amount of UMIs (UMI-thresholds: Immune [10,000], fibroblast [11,000], tumor cell [30,000]) or gene-counts (gene-count-thresholds: Immune [3,000], fibroblast [4,000], tumor cell [5,700]) were considered doublets and excluded. Feature genes used for cell type-specific MetaCell generation were obtained using the mcell_gset_filter_varmean and mcell_gset_filter_cov functions implemented in the Metacell package. These features genes were filtered for genes involved in cell cycle (gene-module 7 and 20) and ribosomal proteins (gene-module 2).

For all plots showing normalized UMI counts, a center log ratio normalization was applied, as implemented in the Seurat package⁵². To calculate sample fractions within MetaCells, cell counts

were first normalized to 10,000 cells within each sample-hashtag to allow comparisons.

Pseudotime analysis was performed using the Slingshot algorithm⁵³. Gene-level general additive models were fitted to feature-genes used for MetaCell generation applying the fitGAM function from the TradeSeq R package⁵⁴, setting knots at 5. Only genes that associated significantly (adjusted P value < 0.05) with pseudotime were used in subsequent analysis. Genes were then clustered based on expression kinetics across pseudotime based on Euclidean distance (**Supplementary Table 5**). To assess sample composition across pseudotime, the 3 replicates from QPCTL^{-/-} or QPCTL^{+/+} TMEs were analyzed together. To allow pooling of replicates, cell counts were normalized to the total number of cell counts within each sample. Normalized cell counts were then tallied within windows of 60 cell-codes wide, sliding 1 cell-code per frame. Differential gene expression analysis was performed using the FindMarkers function implemented in Seurat. Wilcoxon Rank Sum test was used to obtain log₂ fold changes.

For comparison of Immune MetaCell 1–3 to the external monocyte/macrophage cell clusters, the scRNAseq dataset from Gubin *et al.*³² was retrieved from the Gene Expression Omnibus (GSE119352). The external data was subsequently filtered for monocyte/macrophage cell clusters (Mac_s1–5), and normalized through centered log-ratio transformation. Differentially expressed genes within each cluster was then determined by comparing each cluster to all others applying Wilcoxon Rank Sum test (FindAllMarks function, Seurat R package). 25 marker genes were then selected for each MetaCell, defined as the 25 most enriched genes within that MetaCell. Each of these gene-sets was then used to compute a similarity score with each of the Mac_s clusters. Each similarity score was calculated by filtering the Wilcoxon Rank Sum test results of a given Mac_s cluster for a marker gene-set, followed by a weighted sampling of the log₂ transformed fold change values (sampling 10,000 times, with replacement, weighted by the MetaCell gene-enrichment value), and finally averaging (median) the obtained values.

Pathology

For histopathological analyses, 2 μm-thick hematoxylin-eosin-stained sections were prepared from formalin-fixed, paraffin-embedded murine tissues, including skin, spleen, thymus, lymph nodes, liver, pancreas, gastrointestinal tract, heart, lung, kidneys, testes, ovaries, accessory sex glands, bone marrow (sternum and extremity), and muscles. Sections were evaluated and scored by an animal pathologist blinded to animal genotype.

Statistical analysis

All statistical analyses were performed either with R (V4.0.5, ‘Shake and Throw’) or Graphpad (V8.4.1, Prism software). All statistical test were two-sided, unless otherwise indicated. Differences were considered statistically significant if $P < 0.05$. The n values used to calculate statistics, the type of replicates and the relevant significant P values are noted in the figure legends.

Ethical compliance

All animal experiments were approved by the Animal Welfare Committee of the Netherlands Cancer Institute (NKI), in accordance with national guidelines. Animals were maintained in the animal department of the NKI, housed in individually ventilated cage systems under specific-pathogen-free conditions, and received food and water freely. Mice were used at 8-25 weeks of age.

Data availability

Transcriptomic data presented in the manuscript have been deposited to the Gene Expression Omnibus, and can be accessed as series GSE180201. Exome data for MC38-Kerafast and MC38-AMS have been deposited to the Sequence Read Archive, and can be accessed as project PRJNA753254. R scripts used to produce key figures in the manuscript have been submitted to GitHub (https://github.com/kasbress/QPCTL_Project).

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Author contributions

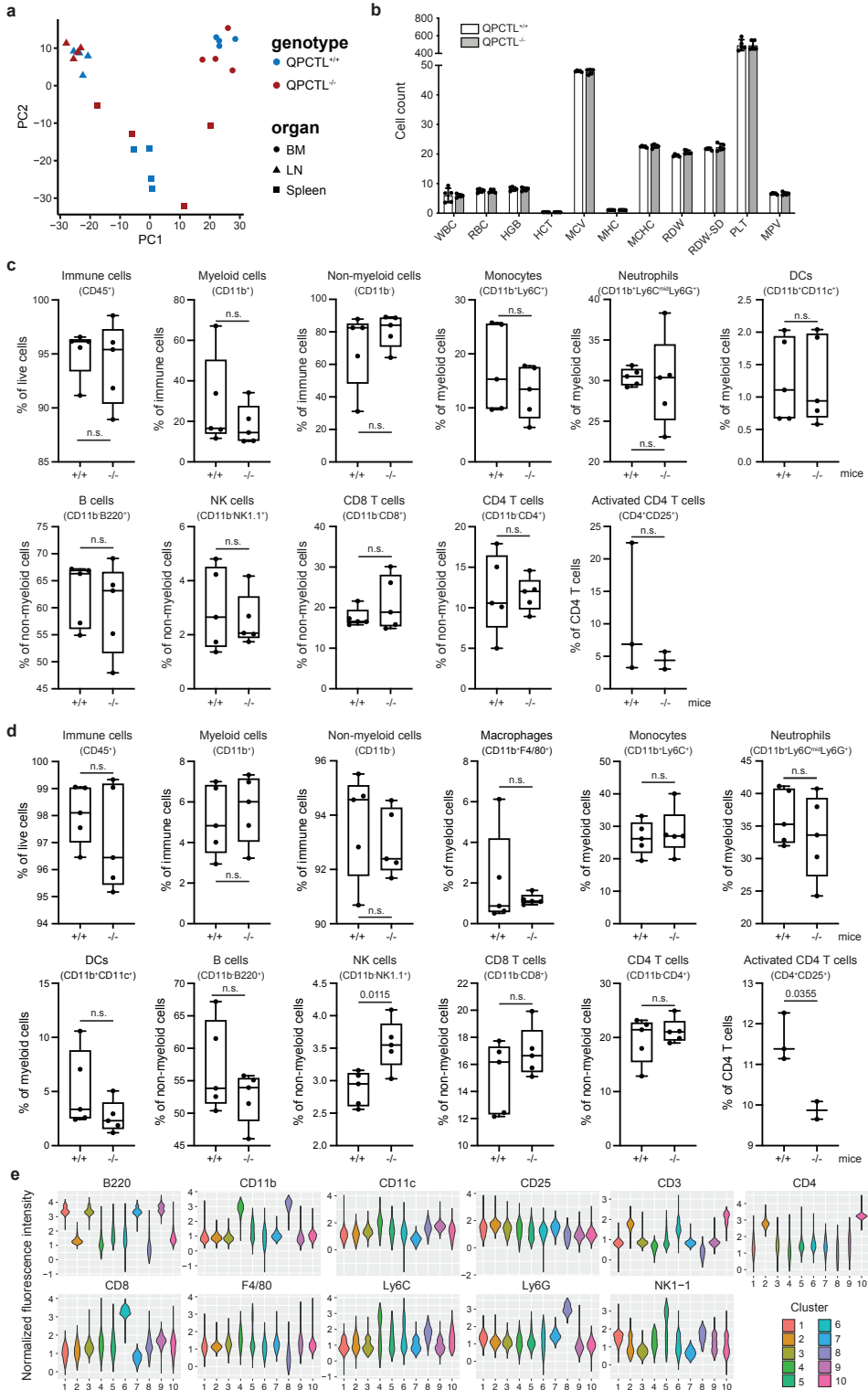
M.E.W.L. and K.B. conceived the project, designed and performed experiments, interpreted data, curated data and co-wrote the manuscript. M.T. designed and performed experiments. N.P, J.S, B.S and M.B. performed experiments. L.K established the QPCTL^{-/-} strain. T.N.S. conceived the project, designed experiments, interpreted data and co-wrote the manuscript.

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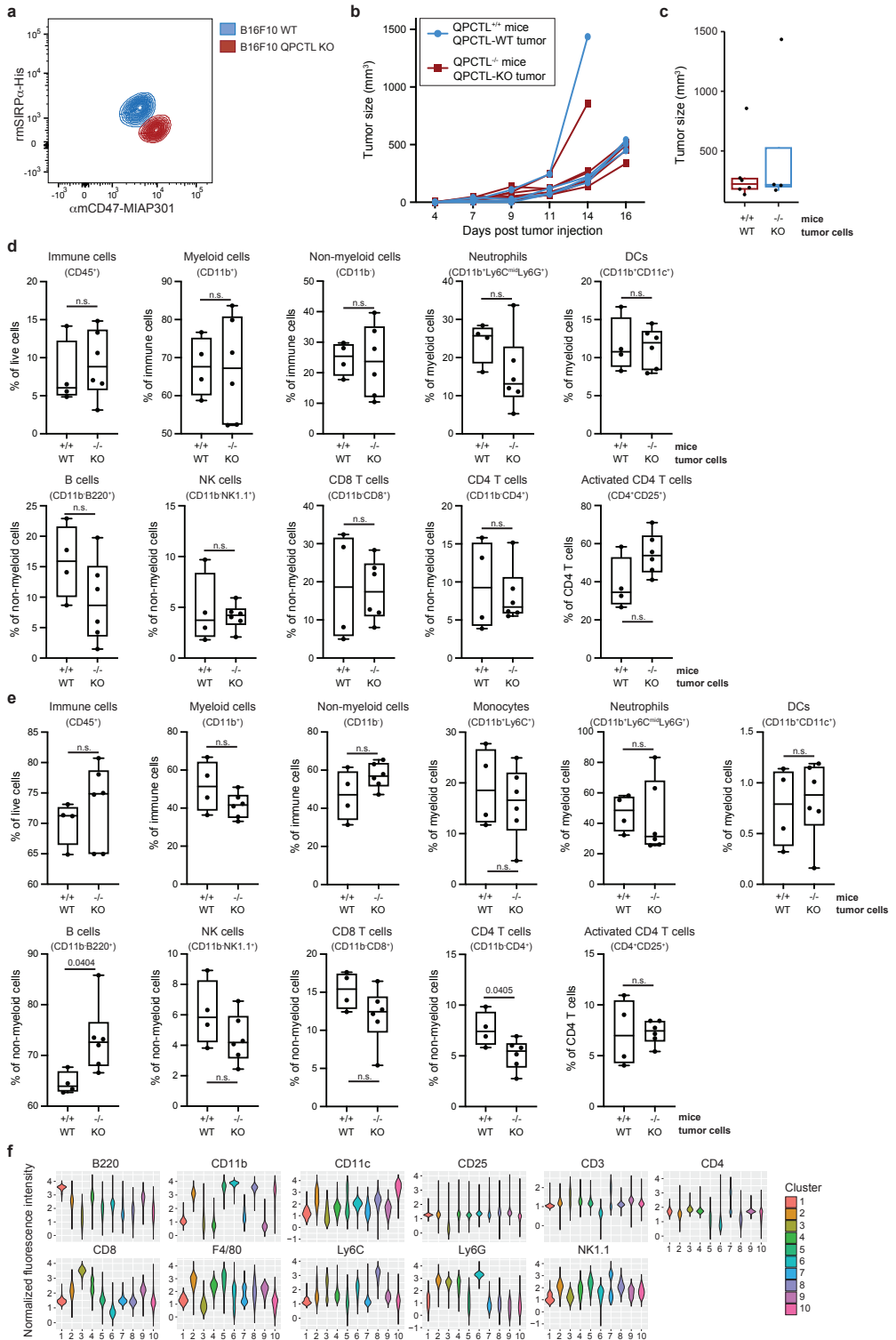
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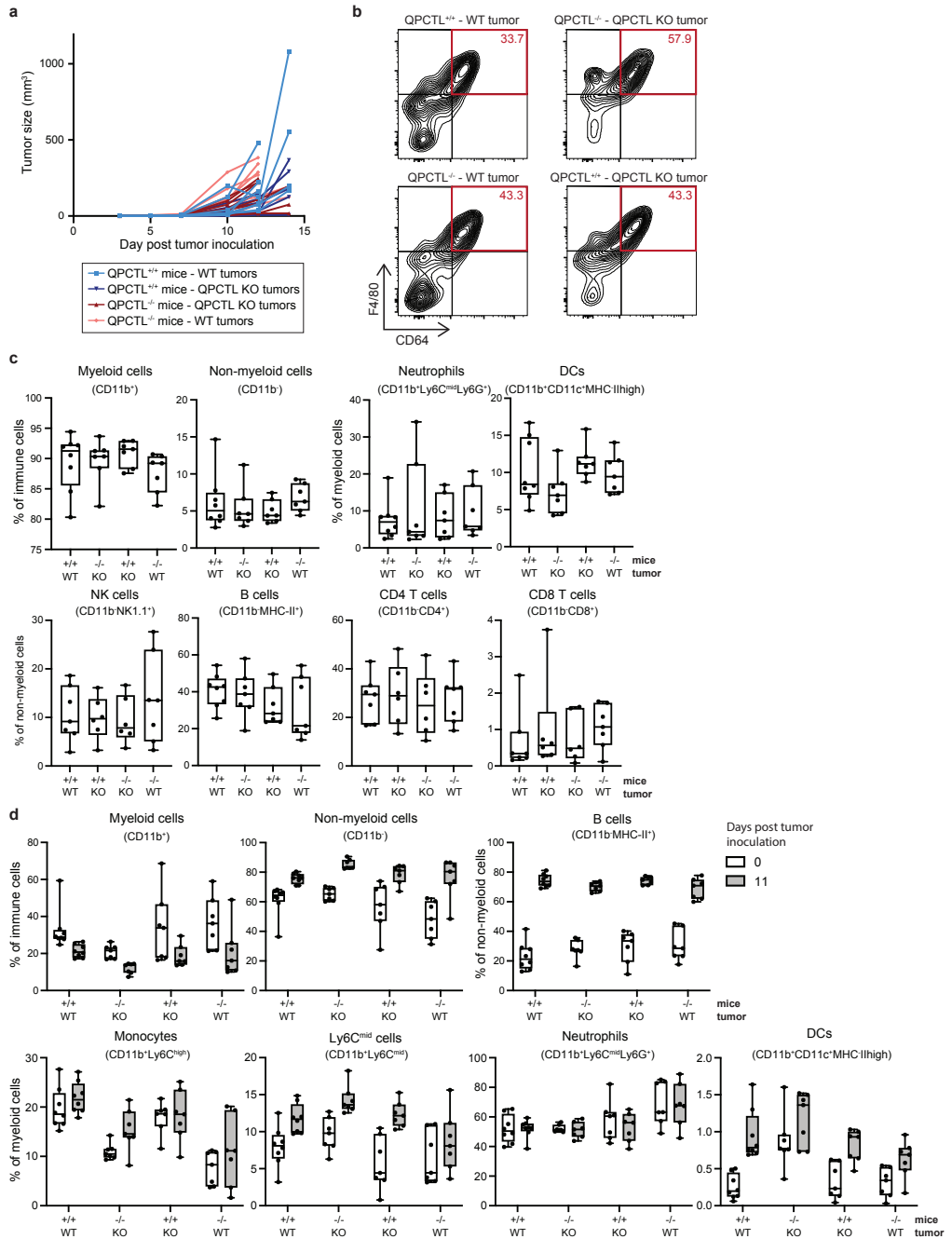
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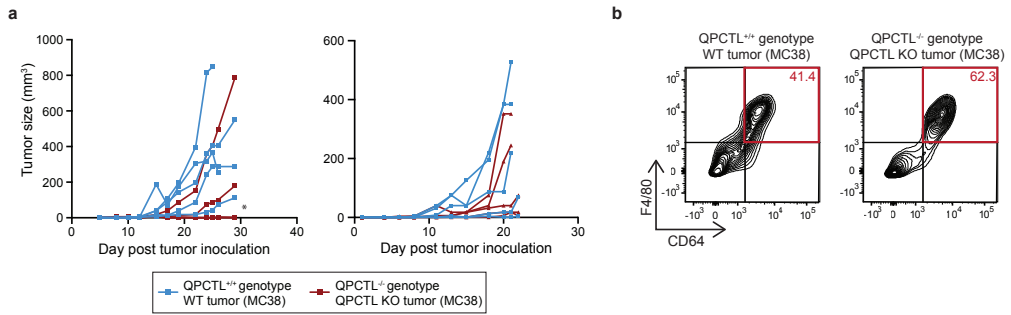
Supplementary Figure 1 (on previous page). Phenotypic analysis of QPCTL KO mice (related to Figure 1). (a) Principal component analysis performed on the 1,000 most differentially expressed genes in bone marrow (BM), lymph node (LN) and spleen samples from QPCTL^{+/+} and QPCTL^{-/-} mice. The first two components are plotted. (b) Quantification of indicated parameters in peripheral blood ($n = 5$ mice per group). Dots depict data from individual mice, bars represent group means, error bars indicate standard deviation. (c) Quantification of indicated immune cell populations as a frequency of total live, myeloid (CD11b⁺), non-myeloid (CD11b⁻) or total CD4 T cells (CD3⁺CD4⁺), in blood of QPCTL^{+/+} and QPCTL^{-/-} mice. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. $n = 3$ (QPCTL^{+/+}) or $n = 2$ (QPCTL^{-/-}) mice for activated CD4 T cells; $n = 5$ mice per group for all other immune cell subtypes. (d) Quantification of indicated immune cell populations as a frequency of total live, myeloid (CD11b⁺), non-myeloid (CD11b⁻), or total CD4 T cells (CD3⁺CD4⁺), in spleen of QPCTL^{+/+} and QPCTL^{-/-} mice. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. $n = 3$ (QPCTL^{+/+}) or $n = 2$ (QPCTL^{-/-}) mice for activated CD4 T cells; $n = 5$ mice per group for all other immune cell subtypes. (e) Violin plots depicting expression of indicated markers by the cell clusters described in Fig. 1d. *P* values were determined by two-sided Student's *T* test (c, d). Data were obtained in single experiments. WBC, white blood cell; RBC, red blood cell; HGB, hemoglobin; HCT, hematocrit; MCV, mean corpuscular volume; MCHC, mean corpuscular hemoglobin concentration; RDW, red cell distribution width; RDW-SD, red cell distribution width - size distribution; PLT, platelet; PV, mean platelet volume.



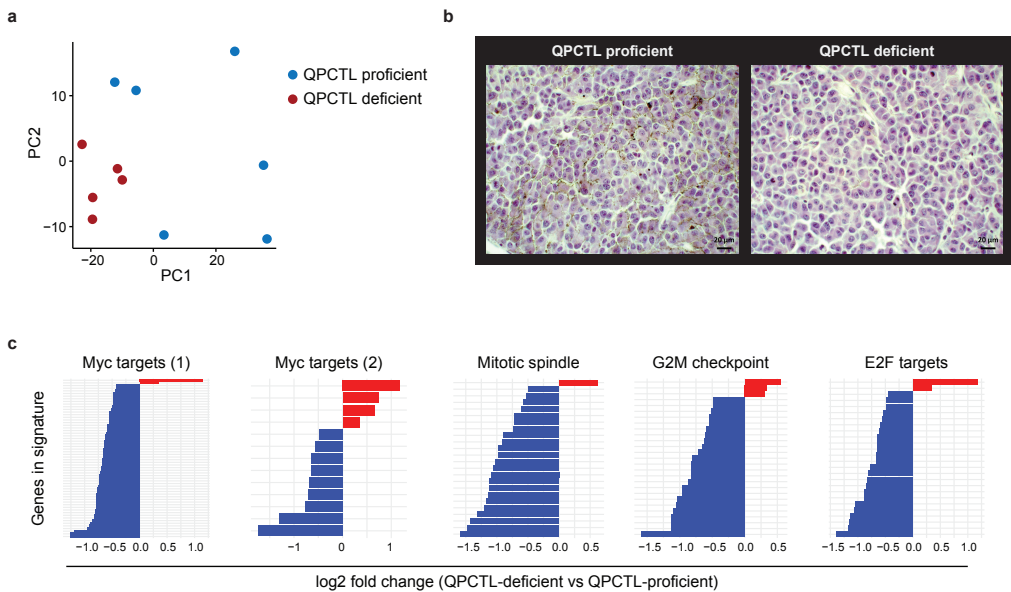
Supplementary Figure 2 (on previous page). Effect of QPCTL deficiency on the TME and blood cell compartment in tumor-bearing animals (related to Figure 2). (a) Representative flow cytometry plot depicting rmSIRP α -His and α mCD47 antibody (clone MIAP301) binding to QPCTL-WT and QPCTL-KO B16F10 cells. (b-f) QPCTL^{+/+} ($n = 4$) or QPCTL^{-/-} ($n = 6$) mice were inoculated with B16F10 WT and B16F10 QPCTL KO tumor cells, respectively. Mice were sacrificed between day 14–16 post inoculation. (b, c) Tumor growth curves and tumor sizes at day 14 post inoculation. Boxplots indicate group median and 25th/75th percentiles, whiskers indicate the interquartile range multiplied by 1.5, dots signify individual samples. (d) Quantification of indicated immune cell populations as a frequency of total live, total immune (CD45⁺), myeloid (CD11b⁺), non-myeloid (CD11b⁻) or total CD4 T cells (CD3⁺CD4⁺), in QPCTL-proficient or -deficient TMEs. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. (e) Quantification of indicated immune cell populations as a frequency of total live, total immune (CD45⁺), myeloid (CD11b⁺), non-myeloid (CD11b⁻) or total CD4 T cells (CD3⁺CD4⁺), in blood of tumor bearing mice in a QPCTL-proficient or -deficient setting. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. (f) Violin plots depicting marker expression within the obtained clusters described in **Fig. 2b**. P values were determined by two-sided Student's T test (d, e). Data are representative of at least 2 independent experiments.



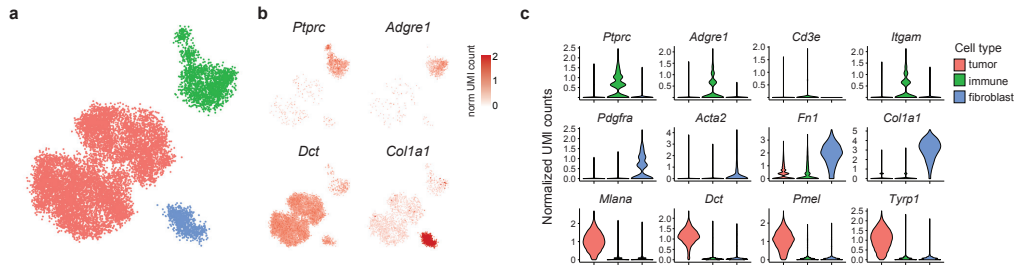
Supplementary Figure 3 (on previous page). Effect of QPCTL deficiency in tumor and host cell compartments (related to Figure 2). QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with either QPCTL-WT or QPCTL-KO B16F10 melanoma cells ($n = 7-8$ per group). Mice were sacrificed between 12–14 days post tumor inoculation. **(a)** Tumor growth curves. **(b)** Representative flow cytometry plots of data described in **Fig. 2d**, depicting macrophages (F4/80⁺CD64⁺ cells) amongst total myeloid (CD11b⁺) cells in the TME. Numbers depict the percentage macrophages within the myeloid cell gate. **(c)** Quantification of indicated immune cell populations as a frequency of total immune (CD45⁺), myeloid (CD11b⁺) or non-myeloid (CD11b⁻) cells, in the indicated TMEs. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. **(d)** Quantification of indicated immune cell populations as a frequency of total immune (CD45⁺), myeloid (CD11b⁺) or non-myeloid (CD11b⁻) cells, in blood before tumor inoculation (0 days), or 11 days after tumor inoculation. Dots depict individual mice, boxplots indicate group median and 25th/75th percentiles, whiskers indicate min/max. *P* values were determined by one-way ANOVA followed by Tukey's HSD test **(c)**. Significant *P* values (< 0.05) are indicated in the plots. Data are representative of at least 2 independent experiments **(a-c)**, or were obtained in a single experiment **(d)**.



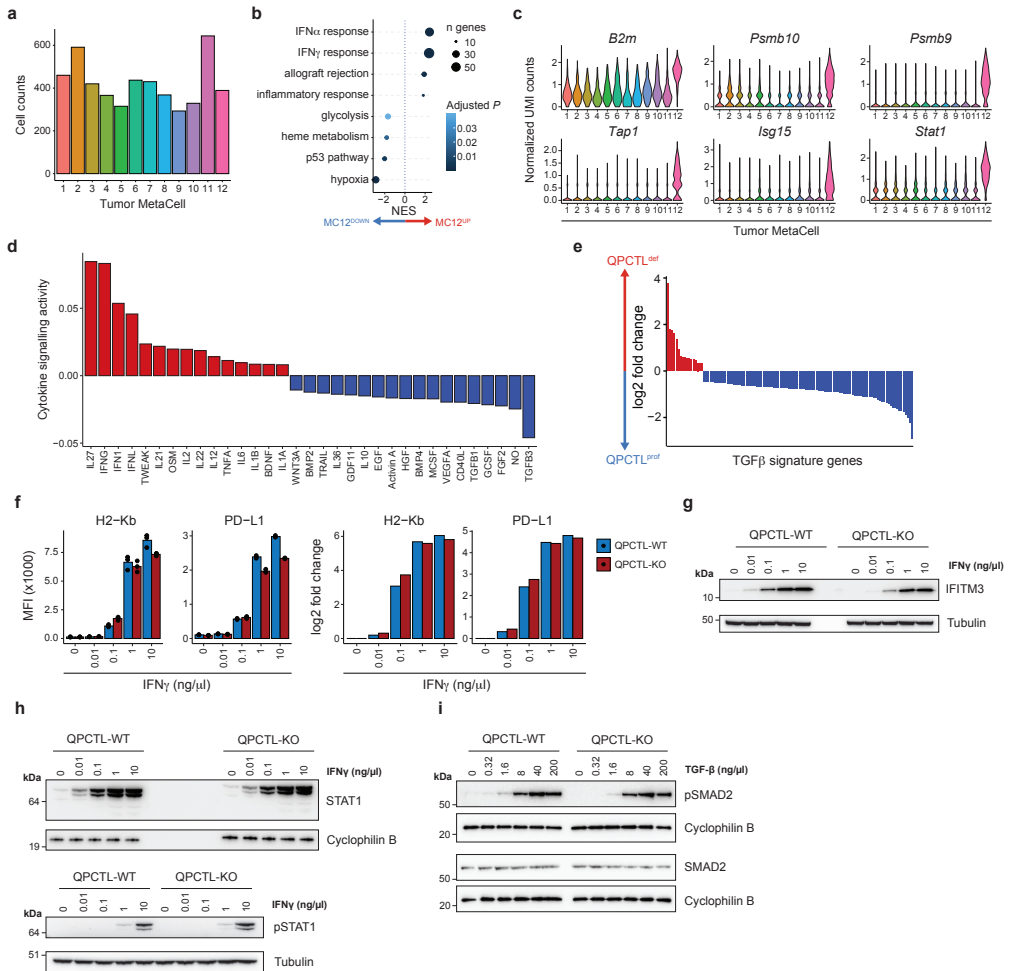
Supplementary Figure 4. Effect of QPCTL deficiency on TME composition (related to Figure 2). QPCTL^{+/+} and QPCTL^{-/-} mice were inoculated with QPCTL-WT and QPCTL-KO MC38 cells, respectively. Data from 2 independent experiments are shown ($n = 5$ per experiment). Mice were sacrificed at 22 (experiment 1) or 29 (experiment 2) days post tumor inoculation. (a) Tumor growth curves in two independent experiments. Asterisk (*) indicates 3 overlapping lines. (b) Representative flow cytometry plots of data described in Fig. 2f, depicting macrophages (F4/80⁺CD64⁺) amongst total myeloid (CD11b⁺) cells in the TME. Numbers depict the percentage macrophages within the myeloid cell gate. Data were obtained in 2 independent experiments.



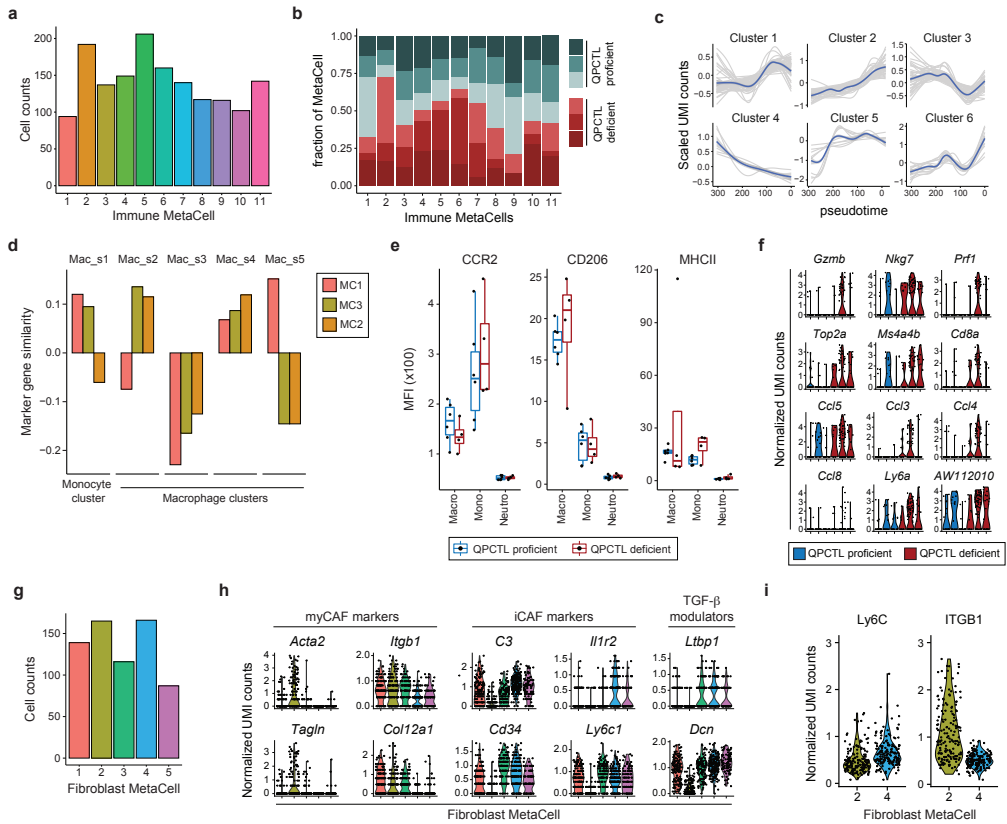
Supplementary Figure 5. Effect of QPCTL deficiency on tumor melanogenesis and cell cycle-related gene expression (related to Figure 3). mRNA sequencing was performed on sorted CD45-negative cells from QPCTL-proficient ($n = 5$) and QPCTL-deficient ($n = 6$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. (a) Principal component analysis performed on the 1,000 most differentially expressed genes across all samples. (b) Representative hematoxylin-eosin stained sections from QPCTL-proficient and -deficient TMEs. Note the presence of melanin signal (brown) in the QPCTL-proficient, but not the QPCTL-deficient sample. (c) Waterfall plots depicting log₂ fold change values of genes from indicated hallmark pathways obtained from MSigDB. Only significantly ($P < 0.05$) differentially expressed genes are included. Data are representative of 2 independent experiments. PC, principal component; MSigDB, Molecular Signatures Database.



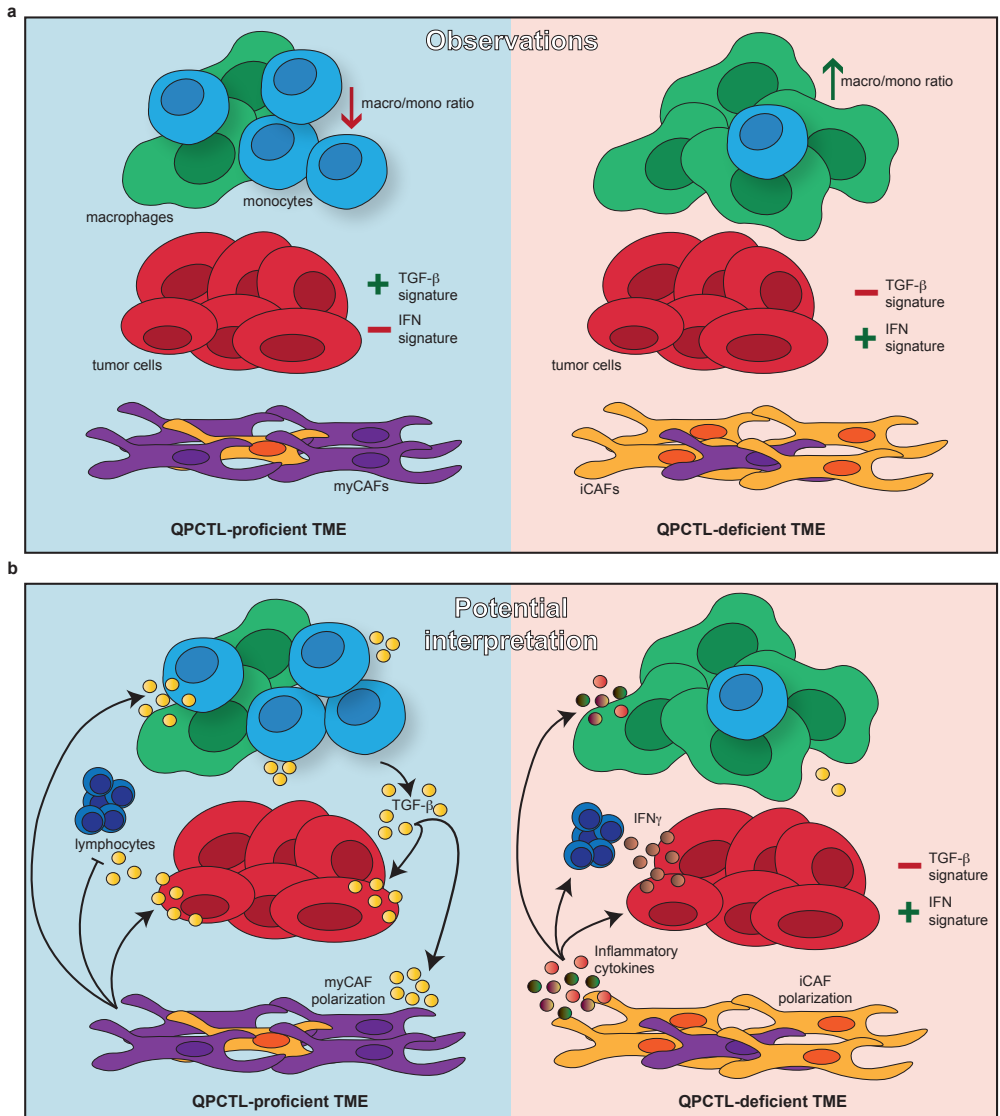
Supplementary Figure 6. Characteristics of the TME cell supertypes identified by single cell RNA sequencing (related to Figure 4 and 5). scRNA sequencing was performed on sorted live cells from QPCTL-proficient ($n = 3$) and QPCTL-deficient ($n = 3$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. **(a, b)** 2-dimensional MetaCell projection of all cells analyzed by scRNAseq. Single cells are colored by MetaCell **(a)**, or normalized UMI count **(b)** of selected genes. **(c)** Violin plots depicting gene expression of selected cell type-specific genes within the major clusters obtained. Depicted data were obtained in a single experiment, consisting of 6 mice.



Supplementary Figure 7. QPCTL deficiency leads to an increased IFN- and decreased TGF- β -response signature in tumor cells (related to Figure 4). (a-d) scRNA sequencing was performed on sorted live cells from QPCTL-proficient ($n = 3$) and QPCTL-deficient ($n = 3$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. (a) Absolute cell counts per tumor cell MetaCell included in the analysis. (b) Gene set enrichment analysis performed on the top and bottom 200 genes expressed by MC12. Hallmark gene sets from MSigDB were used in the analysis, and only significant gene sets ($P < 0.05$) were included in the plot. (c) Normalized UMI counts of selected IFN responsive genes within each tumor MetaCell. (d) Cytokine signaling activity in tumor cell MC12 as predicted by the CytoSig algorithm (Jiang *et al.*, Nature Methods, 2021). (e) Analysis of bulk RNAseq data (Fig. 3). Log₂ fold changes of TGF- β associated transcripts between QPCTL-deficient and -proficient samples. (f) H2-Kb and PD-L1 cell surface expression of QPCTL-WT and QPCTL-KO B16F10 lines after a 20-hour incubation with the indicated amounts of IFN γ . For fold-change plots, group means were calculated and normalized to the MFI detected in the control treated samples. (g-h) Protein levels of IFITM3 (g), STAT1 and phosphorylated STAT1 (h) detected in cell lysates obtained from QPCTL-WT and QPCTL-KO B16F10 lines after a 20-hour incubation with indicated amounts of IFN γ . (i) Phosphorylation state of SMAD2 detected in cell lysates obtained from QPCTL-WT and QPCTL-KO B16F10 lines after a 1-hour incubation with indicated amounts of TGF- β . Data were obtained in a single experiment, consisting of 6 mice (a-e), or are representative of 2 independent experiments (f-i).



Supplementary Figure 8 QPCTL deficiency alters the immune cell compartment and CAF polarization in the TME (related to Figure 5). (a-d and f-i) scRNA sequencing was performed on sorted live cells from QPCTL-proficient ($n = 3$) and QPCTL-deficient ($n = 3$) B16F10 TMEs. Tumors were harvested at day 14 post inoculation. (a) Absolute cell counts per immune MetaCell included in the analysis. (b) Stacked barchart depicting the sample composition of each immune MetaCell. Cell counts from each sample were normalized to 1,000 cells. (c) Gene clusters obtained through hierarchical clustering of gene expression kinetics across pseudotime. Graphs depict general additive models fitted for each gene in grey. Blue lines represent average trends for each cluster. Note that gene cluster 2 and 4 exhibit a strong positive and negative association with pseudotime, respectively. Single gene examples from these clusters are depicted in **Figure 4e**. (d) Comparison of marker-gene expression between the MetaCells comprising M ϕ /Mo subset 1 (MC1, 2 and 3) and the M ϕ /Mo cell clusters described by Gubin and colleagues (Mac_s_1–5; Gubin *et al.*, Cell, 2018). The 20 most enriched genes of each MetaCell were selected, and similarity to the Mac_s clusters was calculated as the sum of log₂ transformed enrichment of those selected genes within each Mac_s cluster. See methods for details on analysis. (e) Comparison of cell surface expression levels of indicated proteins on macrophages, monocytes and neutrophils assessed in QPCTL-proficient ($n = 6$) or -deficient ($n = 4$) TMEs, analyzed at day 14 post tumor inoculation. (f) Top 12 genes with the highest relative expression in CD3⁺ lymphocytes (MC6) from QPCTL-deficient compared to QPCTL-proficient TMEs. Violin plots depicting normalized UMI counts. (g) Absolute cell counts per CAF MetaCell included in the analysis. (h) Violin plots depicting normalized UMI counts of selected genes across CAF MetaCells. (i) cell surface expression levels of indicated proteins on CAF MC2 and MC4, as assessed using barcoded antibodies. Data were obtained in a single experiment, consisting of 6 mice (a-d and f-i), or are representative of 2 independent experiments (e).



Supplementary Figure 9. Graphical abstract. (a) Cartoon depiction of the findings in the study. **(b)** Potential scenario that connects the reported findings.