PIKfyve controls dendritic cell function and tumor immunity

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Abstract

The modern armamentarium for cancer treatment includes immunotherapy and targeted therapy, such as protein kinase inhibitors. However, the mechanisms that allow cancer-targeting drugs to effectively mobilize dendritic cells (DCs) and affect immunotherapy are poorly understood. Here, we report that among shared gene targets of clinically relevant protein kinase inhibitors, high *PIKFYVE* expression was least predictive of complete response in patients who received immune checkpoint blockade (ICB). In immune cells, high *PIKFYVE* expression in DCs was associated with worse response to ICB. Genetic and pharmacological studies demonstrated that PIKfyve ablation enhanced DC function via selectively altering the alternate/non-canonical NF-κB pathway. Both loss of *Pikfyve* in DCs and treatment with apilimod, a potent and specific PIKfyve inhibitor, restrained tumor growth, enhanced DC-dependent T cell immunity, and potentiated ICB efficacy in tumor-bearing mouse models. Furthermore, the combination of a vaccine adjuvant and apilimod reduced tumor progression *in vivo*. Thus, PIKfyve negatively controls DCs, and PIKfyve inhibition has promise for cancer immunotherapy and vaccine treatment strategies.

Keywords

PIKfyve, dendritic cell, kinase inhibitor, apilimod, immune checkpoint blockade, NF-κB, tumor, antigen-specific T cell, cancer vaccine

Main

The success of immunotherapy has fundamentally altered our understanding of cancer and changed the standard of care for cancer treatment.[1,2] Understanding the mechanisms of immunotherapy efficacy and resistance are needed to further improve cancer outcomes.[3,4] Attention has centered on the countless ways in which tumor-associated antigen (TAA)-specific T cells are directly or indirectly suppressed.[4-8] However, fewer studies have investigated the contribution of professional antigen presenting cells (APCs), such as dendritic cells (DCs), to antitumor immunity. DCs prime, activate, and sustain T cell responses across cancer types and are required for durable immunity against tumors.[9,10] Moreover, they hold a dominant role in extinguishing ongoing immune responses, making them among the most powerful regulators of antitumoral T cell responses.

Though there are many elements in the tumor microenvironment that favor a pro or anti-cancer response, we now understand that the success of many approved cancer therapies ultimately depends on the sustained activation of TAA-specific CD8+ T cells through DCs.[9,11,12] For example, the "DC-T cell axis" is required for response to immune checkpoint blockade (ICB) therapy. Loss of either cell type reduces ICB efficacy in preclinical models.[13] DC or T cell signatures are associated with ICB treatment response in cancer patients.[3,14-16] Studies have shown that certain systemic chemotherapy agents may enhance DC antigen presentation by enhancing tumor cell antigenicity.[17,18] However, studies reporting a direct effect on DCs are rare,[19,20] with a few agents promoting a more "DC-like" phenotype in suppressive myeloid-derived cells.[21] Studies have focused on chemotherapy-induced activation of DCs as a mechanism to potentiate immunotherapy strategies, such as DC vaccines or ICB.[13,21-24]

The contribution of DCs to targeted therapy efficacy, such as protein kinase inhibitors, is largely unknown. Unlike ICB or chemotherapy, these drugs target DNA damage responses or oncogenic signaling pathways preferentially utilized by cancer cells.[25,26] Targeting BRAF, EGFR, PDGRA, KIT, PIK3CA, ALK, MET, ROS1, ERBB, CDK12, and others has improved cancer care, however durable responses are uncommon.[27,28] Historically, the ability of the FDA-approved kinase inhibitors to interact with the immune system was ignored. More recently, preclinical studies have included characterization of the immune system, focusing on associations between treatment response and T cell profiles.[29-34] Efficacy of a receptor kinase inhibitor was shown to be dependent on T cells which served as a rationale for combination therapy with ICB.[35,36] However, these studies have not explored the contribution of DCs to those outcomes.

As the number of clinical trials combining immunotherapy and protein kinase inhibitors continue to increase, it is imperative to understand whether they rely on DCs and modulate the DC-T cell axis. Therefore, we aimed to identify DC-associated kinases and their inhibitors in the context of ICB and characterize their ability to alter DC phenotypes and DC-dependent therapies in the tumor microenvironment.

Results

DC *PIKFYVE* expression is associated with ICB efficacy

Protein kinases are important targets for cancer therapy. To explore the relevance of their gene targets in ICB-induced tumor immunity, we first identified the 25 most common and unique gene targets of Phase I/Phase II/FDA-approved kinase inhibitors offered in a commercial screening library[37–39] (*ALK, AURKA, BTK, CSF1R, EGFR, FGFR1, FLT1, FLT3, IGF1, IGFR1, IKBKB, JAK1, JAK2, KDR, KIT, MET, MTOR, NTRK1, PDGFRA, PIKFYVE, PTK2, RAF1, RET, SRC, and SYK*). We then explored the potential involvement of these 25 genes in treatment response in a clinical cohort of patients who received ICB and clinical bulk RNA-sequencing (RNA-seq) of their tumors at the University of Michigan as a part of their cancer treatment (**Table 1**).[40]

Seventeen percent of our cohort exhibited RECIST-defined[41] complete response (CR) to ICB (**Fig. 1a**). As expected, the CR group had the best overall survival in our cohort (**Fig. 1b**). Of the kinase inhibitor gene targets evaluated, high pre-treatment *PIKFYVE, KDR, NTRK1, IKBKB, FLT1, or FGFR1* expression was each associated with lower odds of achieving CR when controlling for cancer type, biopsy site, age at the time of treatment initiation, and ICB agent (**Fig. 1c**). Amongst these targets, we found that only high *PIKFYVE* expression was independently associated with worse progression-free survival (PFS) on treatment (**Fig. 1d**). Furthermore, a high pre-treatment CD8+ T cell activation score[42,43] was associated with better overall survival, whereas a high PIKfyve score had the opposite effect (**Fig. 1e**). In a published prospective study,[44] we found that pre-treatment PIKfyve scores were not different between patients with melanoma who were ICB-treatment-naïve ("NAÏVE") or had previously progressed on treatment ("PROG") (**Supp. 1a**). Interestingly, a high PIKfyve score was associated with worse overall survival in the PROG cohort when controlling for mutational subtype, stage of disease, and neoantigen load (**Supp. 1b-c**).Therefore, *PIKFYVE,* encoding a cytosolic lipid kinase,[45–47] may be a therapeutically targetable gene with relevance in patients receiving ICB.

We then postulated that the correlation between *PIKFYVE* and ICB-associated outcomes in cancer patients may depend on cell type. We utilized published single cell RNA-seq (scRNA-seq) datasets to assess this hypothesis. We found that *PIKFYVE* was expressed in immune cells across multiple cancer types[48] (**Supp. 1d-h**). Importantly, in a study of patients with melanoma treated with ICB,[49] pre-treatment expression of *PIKFYVE* in conventional DCs (cDCs) was lower in responders (R) when compared to non-responders (NR) (**Fig. 1f-g**).There were no differences in *PIKFYVE* expression in any other immune cell type. In addition, a patient with endometrial cancer with CR[50] had lower expression of *PIKFYVE* in their cDCs compared to nonresponders (**Supp. 1i-j**). Together, these data nominate DCs as an immune cell in which *PIKFYVE* may play a significant role in cancer immunity and ICB-associated outcomes.

PIKfyve suppresses DC function

Given these unexpected findings, we sought to understand why patients with lower DC *PIKFYVE* expression in tumors were inclined towards better clinical outcomes following

ICB therapy. We hypothesized that PIKfyve may have a direct and functional role in modulating DCs and DC-mediated immune response against tumors. To this end, we performed genetic and pharmacologic manipulation of PIKfyve in DCs to gain a comprehensive and accurate portrayal of its functional role in this context.

First, we treated non-tumor-bearing, wild-type mice with apilimod, a potent and specific PIKfyve inhibitor that has been studied in many cell types and evaluated in Phase II clinical trials.[35,51–60] Next, we performed a global assessment of immune cells enriched from spleens. We discovered increased proportions of CD69+CD8+ T cells in apilimod-treated mice compared to vehicle (**Supp. 2a-b**). There were no changes in the percentage of terminally differentiated, memory, or exhausted CD8+ T cells between groups (**Supp. 2c-g**). We also assessed myeloid lineages (**Supp. 2h**). We found no change in total surface expression of CD11c, F4/80, or CD11b (**Supp. 2i-k**). Interestingly, there was an increase in the total surface expression of XCR1 with apilimod treatment (**Fig. 2a**). Within the cDC subset, there was an increase in the relative percentage of cDC1s (**Fig. 2b**) versus cDC2 cells (**Supp. 2l**) with apilimod treatment. This warranted further study as cDC1s are a cDC subtype which expresses XCR1 and is involved in cross-presentation and CD8+ T cell-mediated anti-tumor responses.[61–63] An increase in total surface expression of XCR1 was also observed in cultured cDCs treated with apilimod compared to dimethyl sulfoxide (DMSO) (**Supp. 3a-b**). To confirm that these changes were due to loss of functional PIKfyve, we bred *Pikfyve*[f/f] ("WT") mice with *Itgax*[Tg/0] ("CD11c-Cre") mice to specifically and conditionally delete *Pikfyve* in CD11c+ DCs ("KO"). We cultured cDCs isolated from WT and KO mice *in vitro* and validated loss of PIKfyve expression (**Supp. 3c**). Intriguingly, we found that surface (**Fig. 2c**) and total protein (**Fig. 2d**) expression of XCR1 was also increased in cDCs from *Pikfyve* KO versus WT mice.

Given this evidence that bonafide *Pikfyve* loss could alter cDC state, we broadly assessed phenotypic changes in cDCs with PIKfyve ablation. We found that surface (**Fig. 2e**) and total protein (**Fig. 2f**) expression of MHC-I (H2-kb, H2-kd) were significantly increased in KO cDCs when compared to those from WT mice. To determine if these findings were phenocopied by pharmacologic PIKfyve inhibition, we treated cDCs from WT mice with apilimod. Apilimod-treated cDCs showed increased surface and total protein expression of MHC-I when compared to control (**Fig. 2g-h**). Similarly, surface (**Fig. 2i**) and total protein expression (**Fig. 2j**) of MHC-II were increased in the KO versus WT cDCs. Furthermore, apilimod-treated cDCs had increased surface and total protein expression of MHC-II (**Fig. 2k-l**).

To understand whether this increase in MHCI/II expression occurred in isolation or represented a change in overall DC maturation, we examined the expression of co-stimulatory molecules. Total protein levels of CD80 and CD86 expression were increased in *Pikfyve* KO versus WT cDCs (**Fig. 2m**). Apilimod treatment induced CD80 and CD86 expression in WT DCs in a dose-dependent manner (**Fig. 2n**). The percentage of CD80+CD86+ cDCs was also increased with *Pikfyve* KO (**Fig. 2o, Supp 3d**) compared to WT. This increase in CD80+CD86+ cDCs was mirrored in apilimod-treated cells alone and in combination with PolyI:C and LPS (**Fig. 2p**). Furthermore, the

total expression of CD40 was increased in *Pikfyve* KO versus WT cDCs (**Fig. 2q**). Addition of apilimod to LPS treatment also increased total expression of CD40 when compared to DMSO-treated cDCs (**Fig. 2r**).

As genetic and pharmacologic ablation of PIKfyve affected MHC and costimulatory molecules, we hypothesized that PIKfyve could, consequently, alter the ability of cDCs to present antigens and subsequently activate antigen-specific T cells. To test this possibility, we utilized the ovalbumin (OVA) and OT-I/OT-II cell systems.[43,64,65] When cultured with SIINFEKL peptide (pOVA), *Pikfyve* KO cDCs had greater total intensity and percentage of cells with H2-kb-SIINFEKL surface expression compared to WT cDCs (**Supp. 3e-f**). Similarly, apilimod-treated cDCs had greater total intensity and percentage of cells with H2-kb-SIINFEKL surface expression on cDCs compared to DMSO-treated cDCs (**Supp. 3g-h**). In addition, we observed that peptide antigen-loaded *Pikfyve* KO cDCs induced a greater percentage of IFNγ^+ and granzyme B$^+$ OT-I cells compared to WT (**Supp. 4a-c**). This increase in T cell activation was also seen with apilimod treatment compared to DMSO (**Supp. 4d-e**). Finally, we loaded cDCs with soluble OVA protein (sOVA) which were then co-cultured with OT-I and OT-II cells. There were higher percentages of Ki67$^+$ (**Supp. 4f**) and IFNγ^+ (**Fig. 2s**) OT-I cells following co-culture with sOVA-loaded *Pikfyve* KO cDCs compared to WT cDCs. These increases were also demonstrated in OT-I cells co-cultured with apilimod-treated cDCs compared to DMSO (**Fig. 2t, Supp. 4g**). In addition, there were higher percentages of Ki67$^+$ and IFNγ^+ OT-II cells following co-culture with *Pikfyve* KO and apilimod-treated cDCs when compared to their respective controls (**Supp. 4h-i, Fig. 2u-v**). Together, these data indicate that PIKfyve alters DC state and negatively controls DC maturation and function.

PIKfyve suppresses NF-κB activation in DCs

To explore the mechanism through which PIKfyve suppresses DCs, we conducted RNA-seq studies. First, we performed gene set enrichment analysis on genes differentially expressed between *Pikfyve* KO versus WT cDCs (**Supp. Table 1**). Review of MSigDB curated gene sets ("C2") revealed that a gene signature of enhanced DC maturation was positively enriched in *Pikfyve* KO cDCs (**Supp. 5a, Supp. Table 2**) consistent with our phenotypic and functional characterization of these cells.

Furthermore, examination of MSigDB hallmark gene sets ("H") revealed positive enrichment of the "TNF_SIGNALING_VIA_NFκB" gene set in *Pikfyve* KO versus WT cDCs (**Supp. 5b, Supp. Table 3**). This was intriguing as NF-κB is known to be an essential transcription factor for driving the overall maturation and acute activation of DCs.[66–69] Given our findings, we hypothesized that a more general signature of NF-κB downstream gene targets would be affected in *Pikfyve* KO cDCs. Thus, we curated a list of validated NF-κB gene targets from commercial assays (**Supp. Table 4**). In a posteriori analysis, we observed that this signature representing direct activation of downstream NF-κB genes was positively enriched in *Pikfyve* KO versus WT cDCs (**Fig. 3a**).

To corroborate these data suggesting NF-κB regulation by PIKfyve, we treated WT cDCs with apilimod to identify differentially expressed genes at timepoints coinciding with early DC activation (**Supp. Table 5-6**). Examination of MSigDB hallmark gene sets revealed positive enrichment of "TNF_SIGNALING_VIA_NFκB" in apilimod-treated cDCs at 3 and 8 hours when compared to DMSO (**Supp. 5c-d, Supp. Table 7-8**). Importantly, direct NF-κB gene targets were positively enriched in apilimod-treated cDCs at these early timepoints, thus substantiating our findings from the genetic model (**Fig. 3b-c**). Interestingly, apilimod treatment increased *Il12b* transcripts (**Supp. 5e, Supp. Table 5-6**) in addition to the secretion of IL-12p40 (**Supp. 5f**) and IL-12p70 (**Supp. 5g**). Finally, we confirmed that both genetic loss and pharmacological inhibition of PIKfyve increased relative protein levels of p-NF-κB to NF-κB (**Fig. 3d**).

NF-κB is regulated by dynamic, cascading changes in a well-characterized system of upstream cytosolic regulatory proteins that culminate in an altered transcriptional landscape.[70,71] Interestingly, neither genetic nor pharmacological PIKfyve ablation changed levels of IκB-β, but PIKfyve loss or inhibition did increase levels of p-IκB-α relative to IκB-α, which would allow for increased NF-κB phosphorylation and activation (**Fig. 3e**). To understand if this regulation extended further upstream, we also investigated the canonical and alternate/non-canonical regulators of the IκB kinase complex. Abundance of IKK-α and IKK-β were unchanged with PIKfyve ablation (**Fig. 3f**). Interestingly, the abundance of IKK-ε, p-TBK-1, and TBK-1 were increased with genetic and pharmacological PIKfyve ablation (**Fig. 3g**).

These novel findings motivated further exploration of how PIKfyve may alter NF-κB activation. Intriguingly, *Sqstm1* emerged as the most significantly upregulated gene in apilimod-treated cDCs when compared to DMSO (**Fig. 3h, Supp. Table 6**). It was also differentially expressed in KO versus WT cDCs (**Supp. Table 1**). *Sqstm1* is a promising mechanistic candidate in PIKfyve regulation as it is involved in vesicular trafficking, autophagy, and lysosome and proteasome degradation.[72,73] Importantly, *Sqstm1 can* regulate NF-κB activation under various conditions[72,74–76] and is known to interact with TBK-1.[72,73,77–80]

Utilizing the TCGA Pan-Cancer data, we found that high *SQSTM1* was associated with better overall survival in patients (**Supp. 6a**). We also found that high *SQSTM1* negatively correlated with PIKfyve scores (**Fig. 3i**). Furthermore, patients with high NF-κB gene target scores were more likely to have high *SQSTM1* expression (**Fig. 3j**). Recently, Ghislat *et al*[67] demonstrated that NF-κB pathway activation may direct maturation states within the mouse cDC1 subset. We discovered *Sqstm1* expression was higher in mouse cDC1[81] when compared to most other myeloid lineages, including cDC2 (**Supp. 6b**). We investigated this further in the patient data. As expected, a published[67] "cDC1 maturation score" positively correlated with NF-κB gene targets in TCGA Pan-Cancer patient samples (**Supp. 6c**). Moreover, patients who had higher cDC1 maturation scores were more likely to have high *SQSTM1* expression (**Fig. 3k**). Collectively, these data suggest that PIKfyve mediates suppression of DC transcriptional maturation programs through the alternate/non-canonical NF-κB regulatory pathway.

Pikfyve loss in DCs enhances anti-tumor immunity *in vivo*

Since the above results showed that PIKfyve could fundamentally transform cDCs, we examined whether these alterations had any effect on a disease setting. We hypothesized that the loss of *Pikfyve* in DCs could attenuate tumor growth in syngeneic mouse models of cancer. Subcutaneous MC38 tumors injected into *Pikfyve* KO mice manifested reduced tumor growth (**Fig. 4a**) and tumor weights (**Supp. 7a**) compared to WT mice. Furthermore, this inhibitory effect on tumor growth and terminal tumor weights was also seen in KO mice bearing MCA-205 (**Fig. 4b, Supp. 7b**) and B16F10 (**Fig. 4c, Supp. 7c**) tumors.

We then investigated whether the immune response was altered with *Pikfyve* loss in DCs in the B16F10-OVA model. As expected, the growth of tumors (**Supp. 7d-f**) was decreased when inoculated in KO versus WT mice. There was no difference in the percentage of CD11c$^+$MHC-II$^+$ DCs isolated from tumor-draining lymph nodes (TDLNs) compared to all CD45$^+$ cells (**Supp. 7g-h**). Importantly, overall H2-kb-SIINFEKL expression on the surface of DCs from the KO mice was higher than those from the WT (**Fig. 4d**). Concordantly, the percentage of H2kb-SIINFEKL$^+$ DCs was increased in the KO mice (**Supp. 7i**). When comparing the littermate pairs of WT and KO mice, there was an increase in the percentage of SIINFEKL tetramer$^+$ CD8$^+$ T cells isolated from tumors from each pair (**Fig. 4e-f, Supp. 7j**). Therefore, *Pikfyve* loss enhanced DC-mediated antigen presentation and priming of antigen-specific CD8$^+$ T cells in an *in vivo* tumor model.

We additionally explored whether loss of *Pikfyve* in DCs could alter response to anti-PD-1 therapy in the ICB-resistant B16F10 model. Tumor growth inhibition with anti-PD-1 therapy was enhanced when comparing endpoint tumor volume in *Pikfyve* KO mice to wild type mice (**Fig. 4g**).

Finally, we assessed the effects of PIKfyve inhibitor treatment on tumor-infiltrating DCs isolated from DC-selective *Pikfyve* KO versus WT mice. There was an increase in the percentage of cDC1s in KO versus WT mice (**Fig. 4h**). Importantly, treatment with apilimod increased the percentage of cDC1s amongst the WT mice but resulted in no additional increase in the KO mice. Notably, these differences were the same when evaluating total surface XCR1 expression between groups (**Fig. 4i**). There were no changes in the percentage of cDC2s (**Supp. 8a**) or total surface SIRP1α expression (**Supp. 8b**) between any groups. Combined, these data show that PIKfyve in DCs can modulate tumor outcomes *in vivo*.

Immune effects of PIKfyve are DC-dependent

Though PIKfyve inhibitors have shown anti-tumor effects in various preclinical cancer models,[35,56,60,82–84] it is unclear if the activity of DCs directly contributes to its therapeutic effect. Our combined genetic and pharmagologic *in vivo* experiments showed no additional change in *Pikfyve* KO DCs with the addition of PIKfyve inhibitor treatment. Therefore, we further hypothesized that therapeutic PIKfyve inhibition required the

presence of DCs to exert its full anti-tumor effect *in vivo*. To this end, we inoculated WT mice with subcutaneous MC38 tumors. *In vivo* treatment with apilimod reduced MC38 tumor growth compared to vehicle (**Fig. 5a**) and increased the percentage of intratumoral IFNγ⁺CD8⁺ T cells (**Fig. 5b, Supp. 9a**). The efficacy of apilimod was lost in immune-deficient NSG mice (**Supp. 9b**). To test whether drug efficacy required DCs, we used *Batf3*⁻/⁻ mice[61–63] which have loss of cDC1s. Importantly, loss of apilimod efficacy remained when comparing *Batf3*⁻/⁻ mice to immune-competent, WT mice (**Fig. 5c**).

We also investigated the importance of functional DC and T cell signaling pathways in apilimod treatment efficacy. In MC38 tumor-bearing mice, we observed that the efficacy of apilimod was reduced when mice were treated with neutralizing monoclonal antibodies against IL-12[13] (**Supp. 9c**) or IFNγ[35,42] (**Supp. 9d**) compared to isotype controls. As IL-12 is often expressed by DCs and IFNγ by T cells, the data suggests that *in vivo* efficacy of PIKfyve inhibition requires the presence of functional DC signaling and intact DC and T cell signaling pathways. To further validate this possibility in an additional tumor model, we inoculated WT, non-transgenic mice with subcutaneous B16F10 tumors. Treatment with apilimod *in vivo* reduced tumor growth compared to vehicle in this model (**Supp. 9e**). Again, efficacy of apilimod was lost when B16F10 tumors were inoculated into NSG (**Supp. 9f**) or *Batf3*⁻/⁻ mice (**Fig. 5d**).

As vaccines are a DC-dependent immunotherapy strategy, we explored whether Apilimod, as a DC-modulating and DC-dependent agent could potentially modulate cancer vaccine strategies. PolyI:C and TLR agonists are commonly used as vaccine adjuvants in different tumor vaccine models.[85] We treated DCs with PolyI:C or lipopolysaccharide (LPS) in the presence of apilimod *in vitro*. We found that PolyI:C, LPS, or apilimod stimulated MHC-I and MHC-II expression, and apilimod further enhanced this effect (**Supp. 9g-h**). In a proof-of-concept experiment, we explored whether PIKfyve inhibition, as a DC-maturing and DC-dependent therapy, could potentiate the effects of PolyI:C against tumor growth *in vivo*. First, we treated mice with apilimod or vehicle in addition to subcutaneous PolyI:C or water (control) for 21 days prior to inoculation with B16F10-OVA tumors (**Fig. 5e**). The combination of pre-treatment with apilimod and PolyI:C decreased subsequent tumor growth compared to control or either agent alone (**Fig. 5f, Supp. 9i**).

We then inoculated mice with B16F10-OVA tumors first and followed by combination therapy with vehicle or apilimod in addition to PolyI:C or water (**Fig. 5g**). Apilimod and PolyI:C alone resulted in modest tumor growth inhibition which was further potentiated by combination with both agents (**Fig. 5h, Supp. 9j**). These data suggest that PIKfyve inhibitors are clinically viable DC-dependent drugs and are promising candidates for novel combination therapy strategies with human-relevant DC-stimulating agents and adjuvants in cancer and other diseases.

Discussion

In this work, we explored whether therapeutically actionable molecular signaling pathways regulate DCs. We revealed a previously unknown fundamental mechanism for PIKfyve in controlling DC state, maturation, and function through NF-κB regulation and demonstrated its ability to potentiate immunotherapy strategies through DCs. Thus, we generated key translational insight into the potential of PIKfyve inhibitors for enhancing DC-dependent therapies in cancer, such as ICB and vaccines.

Our data are the first to comprehensively characterize a clinically viable protein kinase inhibitor strategy to enhance DC function. Though the investment into these cancer-targeted drugs has transformed cancer therapy outcomes for many patients, it is clear that further insight is required to understand how to achieve universally durable and curative responses.[25,26,28,86] With the historically recent acceptance of the essential role of immunity in cancer, new studies have shown that these drugs may modulate CD8^{+} T cell responses and the efficacy of immunotherapy[29,31–36,87–89]. Notably, their impact on DCs remains largely unexplored. DCs have long been a desired target for cancer therapy given their essential role in T cell immunity.[9–11] At present, it is challenging to manipulate DCs directly due to their small numbers, diversity of roles across subsets, and lack of specific targets. Our data identified a PIKfyve as a clinically-druggable molecular target in DCs. This work highlights the potential of targeting PIKfyve in DCs to improve T cell responses and immunotherapy.

Our study revealed a previously unexplored mechanistic association between PIKfyve, a lipid kinase that synthesizes phosphatidylinositol and regulates autophagy,[45,47,51,52,90] and NF-κB, a transcription factor that is functionally critical for DC activation and maturation.[66–70] We demonstrate that NF-κB signaling activation through PIKfyve ablation selectively occurred through the alternate or non-canonical regulators of the IκB inhibitor complex, such as TBK-1.[71,91,92] Furthermore, we identified *SQSTM1,* which has been shown to regulate NF-κB signaling under a wide range of conditions, as a possible mechanistic linchpin.[72,73,77–80] As *SQSTM1* is a critical component of autophagic regulation through TBK-1,[72,74–76] vesicle trafficking[72,73], and transcriptional regulation,[73] it is a potential nexus between PIKfyve and alternate or non-canonical NF-κB regulation. In addition, we discovered that PIKfyve may preferentially alter cDC1s in tumor models. These results are in line with a recent study demonstrating that NF-κB signaling may be crucial for cDC1 maturation.[67] Our findings provide a rationale for further investigation of the potential for PIKfyve to regulate DC identity and lineage determination.

Ultimately, these data provide a strategy to transcriptionally and functionally overcome DC suppression in the tumor microenvironment through a lipid kinase. Given that PIKfyve suppresses DC function, we reason that PIKfyve inhibitors may be selected to treat cancers with high PIKfyve expression in tumors and DCs, thereby effectively targeting both tumors and the immune system. Conversely, caution may be required when considering using PIKfyve inhibitors in cancers driven by NF-κB signaling.[93] Thus, these fundamental insights into DC regulation may inform optimal, "precision medicine" treatment strategies for further evaluation of PIKfyve inhibitors in specific cancers.

To our knowledge, our results are the first to distinguish PIKfyve inhibition as a DC-dependent cancer therapy. Our preclinical studies demonstrate that PIKfyve inhibitors target DCs directly, require DCs and DC/T cell signaling to prevent tumor progression, and potentiate ICB therapy. This DC-dependent nature of PIKfyve inhibitor efficacy is rooted in its ability to mediate antigen presentation to T cells. Thus, our data provides a mechanistic rationale for PIKfyve inhibitor therapy in combination with ICB and invites further inquiry into ways to potentiate other DC-dependent strategies.

In addition to ICB, tumor vaccination is an approach for cancer prevention and therapy.[94–99] Many vaccination strategies involve adjuvants that directly activate DCs.[85,98] In proof-of-principle experiments, we found that PIKfyve inhibition could potentiate the anti-tumor effect of Poly I:C, a cancer vaccine adjuvant. As the results of cancer vaccine therapy have remained underwhelming,[100] our data provides a rationale for further investigation of PIKfyve inhibitors to vaccinate against cancers and infectious diseases. As PIKfyve inhibition is now being clinically evaluated for efficacy against COVID-19,[59,101] our results substantiate the broad potential of PIKfyve inhibition as a DC-enhancing strategy across disease states.

Acknowledgements

Author contributions

J.E.C., W.Z., and A.M.C. conceived the idea, designed the experiments, and composed the paper; J.E.C. conducted all experiments with assistance from J.Y., J.G., T.M., and S.Y.; Y.Q. and Y. Zheng assisted with vaccine experiments; I.K., J.Y., Y.B., and T.M. assisted with flow cytometry experiments and analysis; I.K., J.Y., Y.B., T.M., L.V., H.L., and M.D.G. assisted with immunological experimental techniques; I.K., J.Y., T.M., A.P., H.X., J.Z., H.L., G.L., and M.D.G. assisted with interpretation of results; Y.Q., J.G., Y.B., J.C.T., S.W., S.G., L.V., Y. Zheng, and T.H. assisted with animal experiments. M.G., Y. Zhang, X.C., F.S., R.W., and M.C. assisted with RNA-seq datasets analysis; M.C. and M.D.G. assisted with clinical analysis and the interpretation of the results. A.M.C. supervised the project.

Competing interest declaration

A.M.C. is co-founder and SAB member of LynxDx, Esanik, Medsyn, and Flamingo Therapeutics. A.M.C. serves as a scientific advisor or consultant to EdenRoc, Proteovant, Rebus, and Tempus. W.Z. has served as a scientific advisor or consultant for NGM, CrownBio, Cstone, ProteoVant, Hengenix, NextCure, and Intergalactic. All other authors declare no competing interests.

Methods

Human studies
All clinical records in this study were obtained and utilized with the approval of Institutional Review Boards. Patients who received ICB therapy were recruited through the University of Michigan Hospital, Ann Arbor, MI, USA. Those who were enrolled in the continuous comprehensive clinical sequencing program at the Michigan Center for Translational Pathology (MCTP MI-Oncoseq program)[40,42,43,102] and had bulk RNA-sequencing libraries of pre-treatment tumor were included in this analysis. Overall survival times were determined from the initiation of therapy. Response to treatment was determined using RECIST1.1[41] criteria with patients meeting criteria for pseudoprogression (imRECIST criteria[103]) being excluded from analysis. Integrative clinical sequencing was performed using standard approved protocols in the MCTP Clinical Laboratory Improvement Amendments-compliant sequencing laboratory as previously described.[40,102,104] Total RNA purified using the AllPrep DNA/RNA/miRNA kit (Qiagen) was then sequenced using the exome-capture transcriptome platform on an Illumina HiSeq 2000 or HiSeq 2500 in paired-end mode. CRISP, the standard clinical RNA-Seq pipeline, was used to perform quality control, alignment, and expression quantification.[105] Tables of read counts were then transformed into fragments per kilobase of transcript per million mapped reads using the Bioconductor *edgeR* package in R.[106]

Gene signature score computation
We used normalized expression of genes to define CD8$^+$ T cell infiltration and activation (*CD8A, CXCL10, CXCL9, GZMA, GZMB, IFNG, PRF1, TBX21*), PIKfyve (*PIKFYVE, PIP4K2A*)[42,43], NF-κB gene targets and cDC1 maturation scores.[67] Scores were calculated by inverse-normal transformation of individual gene expression levels across the cohort followed by summation of inverse-normal values for each sample.[42,43]

Kinase inhibitor gene target selection
A catalog of kinase inhibitors of a commonly utilized commercial screening assay (Selleck Chem; Catalog L1800: https://www.selleckchem.com/screening/tyrosine-kinase-inhibitor-library.html)[37–39,107] was utilized to identify Phase I, Phase II, and FDA-approved drugs as previously described.[37]

Reagents
Apilimod was purchased from Selleck Chemicals and resuspended in dimethyl sulfoxide (DMSO) for *in vitro* studies and in 0.5% methylcellulose (vehicle) for *in vivo* experiments. Lipopolysaccharides (LPS) were purchased from Millipore Sigma (L2654) and resuspended in water. EndoFit Ovalbumin (vac-pova), H-2K^b-restricted ovalbumin

MHC class I epitope (257-264) peptide (vac-sin), and polyI:C (HMW) VacciGradeTM (vac-pic) were purchased from InvivoGen. EasySepTM Mouse CD8+ T cell isolation kit was purchased from Stemcell Technologies (19853).

Bone marrow-derived conventional dendritic cells
Bone marrow was collected from the femurs and tibias of mice. Dendritic cells were generated with bone marrow cells cultured with Flt3L (200 ng/ml) in IMDM (Gibco: 12440-053) supplemented with 10% FBS. For genetic knock-out studies, cells were collected on days 8-10. For pharmacologic studies, cells were collected and treated on day 6. Primary cell cultures were tested to be mycoplasma-free in accordance with standard laboratory procedures at MCTP.

Antigen presentation assays
For peptide-based antigen presentation assays, H-2Kb-restricted ovalbumin (OVA) peptide (100 ng/ml) was added to cDCs *in vitro*. CD8$^+$ T cells were enriched from the lymph node and spleens of OT-I mice. 1×10^5 OT-I T cells were then co-cultured with 1×10^5 cDCs previously cultured with H-2Kb-restricted OVA peptide for 12 hours in 10% FBS medium with 55 µM β-ME and 5 ng/ml IL-2 in a 96-well plate. After 2 days, OT-I cells were collected and analyzed for cytokine expression.

For protein-based antigen presentation assays, CD3$^+$ T cells were enriched from the lymph nodes and spleen of OT-I or OT-II transgenic mice. 2×10^5 OT-I or OT-II cells were then co-cultured with 2×10^5 cDCs together with soluble ovalbumin (sOVA) (10 µg/ml) for 3 days in 10% FBS medium with 55 µM β-ME and 5 ng/ml IL-2 in a 96-well plate. After 3 days, OT-I or OT-II cells were collected and analyzed for cytokine expression. In experiments with drug treatment, cDCs were pre-treated with DMSO or Apilimod (50 nM) for 4 hours and washed prior to co-culture with T cells.

Cell culture
MC38 cell line was acquired from Walter Storkus as previously described.[65,108,109] B16F10 and MCA-205 were purchased from ATCC. Ovalbumin-expressing B16F10 (B16F10-OVA) was established with pCI-neo-mOVA plasmid (Addgene plasmid #25099) and selected with 1 mg/ml of G418 for 2 weeks as previously described.[42,43] All cell lines were maintained at < 70% in culture and tested for mycoplasma contamination every 2 weeks according to MCTP standard protocols.

Flow cytometry analysis (FACS)
Cultured dendritic cells were collected and prepared as single cell suspensions. Single cell suspensions of mononuclear cells were isolated from bilateral tumors combined for each mouse with mechanical disassociation followed by Ficoll separation. Single cell suspensions of mononuclear cells were mechanically disassociated from bilateral tumor-draining lymph nodes.

Surface staining was performed by adding antibodies to single cell suspensions in MACS buffer (PBS, 2% FBS, 1 mM EDTA) for 30 minutes. For cytokine staining, cells were stimulated with phorbol myristate acetate (5 ng/ml), ionomycin (500 ng/ml),

brefeldin A, and monensin at 37°C for 4 hours followed by surface and intracellular staining with Foxp3 /Transcription Factor Staining Buffer Set (eBioscience) per the manufacturer's protocol. For tetramer staining, cells were first incubated with tetramer for 30 minutes prior to the addition of antibodies for surface staining. All data were acquired through LSR Fortessa (BD) and analyzed with FlowJoTM or FACS DIVA (BD Biosciences) software.

For flow cytometry analysis, the following antibodies were used: H-2Db (ThermoFisher Scientific: 28-14-8), H-2Kb (BD Biosciences: AF6-88.5), MHC-IA/IE (BD Biosciences: M5/114.15.2), CD11c (ThermoFisher Scientific: N418), CD80 (BD Biosciences: 16-10A1), CD86 (Biolegend: GL-1), XCR1 (Biolegend: ZET), and SIRP1α/CD172a (BD Biosciences: P84). H-2kb-SIINFEKL (BioLegend: 25-D1.16), CD90.2 (BD Biosciences: 53-2.1), CD8 (BD Biosciences: 53-6.7), CD4 (BD Biosciences: RM4.5), CD3 (BD Biosciences: 17A2), CD45 (BD Biosciences: 30-F11), CD45R (BD Biosciences: RA3-6B2), F4/80 (BD Biosciences: T45-2342), CD11b (ThermoFisher Scientific: M1/70), IFNγ (BD Biosciences: XMG1.2), and granzyme B (BD Biosciences: GB11). CD44 (BD Biosciences: IM7), CD62L (BD Biosciences: MEL-14), KLRG1 (BD Biosciences: 2F1). CD49a (BD Biosciences: Hα31/8), TIM-1 (BD Biosciences: 5D12), PD-1/CD279 (BD Biosciences: J43), Ki67 (BD Biosciences: B56), and CD69 (BD Biosciences: H1.2F3). For tetramer staining, iTAg Tetramer/PE against H-2 Kb OVA SIINFEKL (MBL TB-5001-1) was used.

Immunoblotting
Whole cell lysates were prepared in RIPA lysis buffer (ThermoFisher Scientific: 89900) with Halt™ Protease Inhibitor Cocktail (ThermoFisher Scientific: 78429). Protein concentrations were quantified with the Pierce BSA Standard Pre-Diluted Set (ThermoFisher: 23208). Samples were denatured in NuPage 1x LDS/reducing agent buffer for 10 minutes at 95°C. 30-60 µg protein samples were loaded into 4-12% Bis-Tris gels and transferred onto nitrocellulose membrane (ThermoFisher Scientific: 88018) using the BioRad Trans-blot Turbo System. Membranes were blocked with 5% non-fat dry milk and incubated with primary antibodies overnight at 4 °C and then incubated with host species-matched HRP-conjugated secondary antibodies (BioRad) for 1 hour at room temperature. Membranes were developed using chemiluminescence (PierceTM ECL Western Blotting Substrates, SuperSignalTM West Femto Maximum Sensitivity Substrate, Amersham ECL Prime) and detected using Li-Cor.

For immunoblot analysis, the following primary antibodies were used: MHC-I (ThermoFisher Scientific: PA5-115363), MHC-IA/IE (ThermoFisher Scientific: M5/114.15.2), CD86 (Cell Signaling: E5W6H), CD80 (Cell Signaling: E6J6N), CD40 (Cell Signaling: E2Z7), p-NF-κB p65 Ser536 (Cell Signaling: 93H1), NF-κB p65 (Cell Signaling: D14E12), p- IκB-α Ser32/36 (Cell Signaling: 5A5), IκB-α (Cell Signaling: 9242), IκB-β (Cell Signaling: D1T3Z), IKK-α (Cell Signaling: D3W6N), IKK-β (Cell Signaling: 8943), IKK-γ (Cell Signaling: 2685), IKK-ε (Cell Signaling: D61F9), p-TBK-1 (Cell Signaling: D52C2), TBK-1 (Cell Signaling: D1B4), vinculin (Sigma Aldrich: V9131), and total histone 3 (Cell Signaling: 96C10).

Preparation and analysis of bulk RNA-sequencing data

Total RNA was extracted from cDCs using the miRNeasy mini kit with the inclusion of the genomic DNA digestion step with the RNase-free DNase Kit (Qiagen). RNA quality was assessed by the Bioanalyzer RNA Nano Chip and depletion of rRNA prior to library generation was performed using RiboErase selection kit (Cat.# KK8561, Kapa Biosystems). Then, the KAPA RNA HyperPrep Kit (Cat.# KK8541, Roche Sequencing Solutions) was used to generate libraries, and sequencing was performed on the Illumina HiSeq™ 2500. Reads were aligned with the Spliced Transcripts Alignment to a Reference (STAR) to the mouse reference genome mm1068. Tables of read counts were then transformed into fragments per kilobase of transcript per million mapped reads using the Bioconductor *edgeR*[92] package in R for further downstream analysis. Differential expression analyses were performed using the Bioconductor *limma*[110] package in R. Gene Set Enrichment Analysis was performed using the *fgsea*[111] package.

Analysis of published bulk RNA-sequencing data

The Riaz *et al* dataset[44] was downloaded as FPKM expression from Gene Expression Omnibus (GEO) (GSE91061) and github (https://github.com/riazn/bms038_analysis). The TCGA Pan-Cancer dataset was downloaded as FPKM expression from the publications summary website (https://gdc.cancer.gov/about-data/publications/pancanatlas). The Murakami et al dataset[81] was downloaded as TPM expression from GEO (GSE149761).

Analysis of published single cell RNA-sequencing data

We identified human cDCs in all scRNA-seq datasets as previously validated[16] (*C1orf54, CPVL, LGALS2, CA2, PAK1, CLEC10A, HLA-DMA, HLA-DQB1, HLA-DRA, HLA-DRB1, LYZ, FSCN1*). Single-cell analysis was performed using the Seurat package (v4.1.1).[112]

The Qian *et al* dataset[48] (breast, colorectal, lung, and ovarian cancers) was downloaded as raw counts from the authors' website (https://lambrechtslab.sites.vib.be/en). Log1p data normalization and clustering was performed using the unsupervised graph-based clustering approach. The *AverageExpression* function was employed to extract the cell type-specific cluster gene expression.

The Sade-Feldman *et al* melanoma dataset[49] was downloaded as log2(TPM+1) expression from the Human Cell Atlas website (https://www.humancellatlas.org/). The data was subsetted for "pre-treated" samples only. The *FetchData* function was utilized to evaluate gene expression per cell type.

The Chow *et al* endometrial cancer dataset[50] was downloaded as raw counts from GEO (GSE212217). Log1p data normalization and clustering was performed using the unsupervised graph-based clustering approach. The data was subsetted for "pre-treated" samples only.

Animal experiments

All animal studies were conducted with the approval of the Institutional Animal Care and Use Committee at the University of Michigan prior to initiation of procedures and data collection. Female or male wild type C57BL/6J mice (Strain #: 000664), *Pikfyve^{f/f}* mice (Strain #: 029331), *Itgax^{Tg/0}* (Strain #: 007567) mice, OT-I TCR transgenic mice (Strain #:003831), OT-II TCR transgenic mice (Strain #:004194), *Batf3^{-/-}* mice (Strain #: 013755), and NSG^{TM} (Strain #: 005557) were purchased from The Jackson Laboratory. *Itgax^{Tg/0} Pikfyve^{f/f}* C57BL/6 mice were bred internally and genotyped according to the standard protocol provided by The Jackson Laboratory. All mice were housed under specific pathogen-free conditions. All studies were compliant with all relevant ethical regulations regarding animal research. Tumor volumes were measured by caliper every 2-3 days, and tumors did not exceed 2 centimeters in any direction. All mice were maintained under pathogen-free conditions.

For *in vivo* studies in genetic conditional knock-out models, eight- to twelve-week-old male and female sex-matched littermate pairs of *Pikfyve^{f/f}* and *Itgax^{Tg/0} Pikfyve^{f/f}* were used for all tumor studies. For the MC38 tumor model, 1.5×10^6 tumor cells were subcutaneously injected into both flanks of male mice. For the B16F10 and B16F10-OVA tumor models, 0.5×10^6 tumor cells were subcutaneously injected into both flanks of female mice. For the MCA-205 tumor model, 1×10^6 tumor cells were subcutaneously injected into both flanks of male mice. For the anti-PD-1 therapy study, on day 5 following inoculation with B16F10 tumor, 200 µg isotype control antibody (Bio X Cell: BE0089, clone 2A3) or 200 µg anti-PD-1 (Bio X Cell: BE0146, clone RMP1-14) was administered intraperitoneally to each mouse every 3 days throughout the experiment.

For drug therapy studies, six- to eight -week-old male and female wild type C57BL/6 mice were inoculated with MC38 or B16F10 syngeneic cancer cell lines. For the anti-IFNγ blockade study, on day 7 following inoculation with MC38 tumor, 100 µg isotype control antibody (Bio X Cell: BE0088, clone HRPN) or 100 µg mouse anti-IFNγ (Bio X Cell: BE0055, Clone XMG1.2) was administered intraperitoneally in 4 doses every other day to each mouse with either vehicle or apilimod 30 mg/kg given either 5 days per week (MC38) or daily (B16F10) administered by oral gavage. For the anti-IL-12 blockade study, on day 7 following inoculation with MC38 tumor, a single dose of 1 mg isotype control antibody (Bio X Cell: BE0089, clone 2A3) or 1 mg mouse anti-IL-12p40 (Bio X Cell: BE0051, Clone C17.8) was administered intraperitoneally to each mouse followed by three additional doses of 500 µg of either antibody given every other day +/- either vehicle or apilimod (30 mg/kg, 5 days per week) administered by oral gavage.

For vaccine strategy experiments, six- to eight -week-old female wild type C57BL/6 mice were inoculated with B16F10-OVA tumors. For the vaccination study, mice were pretreated with vehicle or apilimod (30 mg/kg, 5 days per week) administered by oral gavage +/- water or PolyI:C subcutaneously (100 µg on Day 1 and Day 14) for 21 days. On Day 22, B16F10-OVA tumors were inoculated and monitored for an additional 14 days in the absence of treatment. For the vaccine therapy study, B16F10-OVA tumors were inoculated. On Day 5, mice began treatment with vehicle or apilimod (30mg/kg, 5 days per week) administered by oral gavage +/- water or PolyI:C subcutaneously (100 µg on Day 5 and Day 12) for a total of eleven days of treatment.

Statistical analysis

Sample sizes for clinical statistical analyses were not predetermined. Overall survival was estimated by Kaplan-Meier methods and compared with log-rank tests. Cox proportional hazard models were used for multivariate survival analysis. Multivariate logistic regression models were used to assess binary outcomes of response to treatment. Pearson's correlation coefficient was used to assess linear correlations between variables. Chi-square test was performed to assess independence of groups. All statistical analyses were performed using R.

In vitro experiments were carried out in duplicate or triplicate independent experiments as indicated. Student t-tests were performed to assess differences between groups. All *in vivo* studies included a minimum of three littermate pairs of WT or KO mice as biological replicates for genetic conditional knock-out studies and a minimum of four mice per group for drug therapy studies. Mann-Whitney U or ANOVA with post-hoc Tukey adjustment for multiple comparisons were performed to assess differences between groups as indicated. All statistical analyses were performed using GraphPad Prism software.

Data availability

Raw data are included in the Source Data or Supplementary Information. All materials are available from the corresponding authors upon reasonable request. Clinical sequencing data are publicly available.[40,43] RNA-seq data newly generated in this study for *in vitro* analysis have been deposited in the Gene Expression Omnibus repository at NCBI (accession codes GSE235596 and GSE235599).

	Percent (*n*)	Hazard Ratio [95% CI]	*P* value
Age (time of sequencing)			
Below median	50.0% (*n*=46)	1 (*reference level*)	-
Above median	50.0% (*n*=46)	0.55 [0.29, 1.03]	0.06
Sex			
Female	44.5% (*n*=41)	1 (*reference level*)	-
Male	55.5% (*n*=51)	0.92 [0.46, 1.86]	0.82
Immune checkpoint blockade agent			
Atezolizumab	9.8% (*n*=9)	1 (*reference level*)	-
Nivolumab	27.2% (*n*=25)	0.43 [0.13, 1.40]	0.16
Pembrolizumab	59.7% (*n*=55)	0.58 [0.23, 1.44]	0.24
Combination	3.3% (*n*=3)	0.43 [0.065, 2.92]	0.39
Cancer Type			
Melanoma	9.8% (*n*=9)	1 (*reference level*)	-
Bladder	16.3% (*n*=15)	1.24 [0.39, 3.86]	0.70
Breast	14.1% (n=13)	0.73 [0.22, 2.34]	0.60
Gastrointestinal	3.3% (n=3)	1.20 [0.20, 7.02]	0.83
Head and Neck	20.7% (n=19)	0.76 [0.28, 2.05]	0.60
Kidney	6.5% (n=6)	0.27 [0.053, 1.38]	0.11
Lung	5.4% (n=5)	0.10 [0.011, 0.92]	0.042
Lymphoma	3.3% (n=3)	2.41 [0.53, 10.99]	0.25
Prostate	12.0% (n=11)	1.16 [0.39, 3.49]	0.77
Sarcoma	4.3% (n=4)	0.57 [0.13, 2.40]	0.44
Other	4.3% (n=4)	1.97 [0.53, 7.30]	0.30

Table 1: Demographics of a clinical cohort of patients treated with ICB.

Pre-treatment tumor bulk RNA-seq libraries from 92 patients who received ICB treatment and clinical sequencing at the University of Michigan were included in the analysis. Hazard ratios for overall survival are included with 95% confidence intervals. *P* values were calculated from a multivariate cox proportional hazards model.

Figure Legends

Figure 1: DC *PIKFYVE* expression is associated with ICB efficacy.
a. Pie chart of number and percentages of RECIST-defined response to treatment of patients treated with immune checkpoint blockade (ICB) at the University of Michigan, Ann Arbor (n = 92). CR, complete response; PR, partial response; SD, stable disease; PD, progressive disease.

b. Kaplan-Meier curves of overall survival of patients treated with ICB at the University of Michigan, Ann Arbor, by RECIST-defined treatment response. *P* value is determined by log-rank test.

c. Forest plot of log-odds of having complete response (CR) vs. not CR for 25 common Phase I/Phase II/FDA-approved drug target genes. Data from bulk RNA-seq plotted are log-odds with 95% confidence intervals. *P* values are determined by multivariate logistic regression controlling for cancer type, biopsy, ICB agent, and age at initiation of ICB treatment.

d. Forest plot of hazard ratios for progression-free survival of patients treated with ICB at the University of Michigan, Ann Arbor, by high vs. low gene expression for candidate drug target genes. Data from bulk RNA-seq plotted are hazard ratios with 95% confidence intervals. *P* values are determined by multivariate cox proportional hazards model controlling for cancer type, ICB agent, and age at initiation of ICB treatment.

e. Forest plot of hazard ratios for overall survival of patients treated with ICB at the University of Michigan, Ann Arbor, by high vs. low gene expression for PIKfyve score and CD8^{+} T cell activation score. Data from bulk RNA-seq plotted are hazard ratios with 95% confidence intervals. *P* values are determined by multivariate cox proportional hazards model controlling for cancer type, ICB agent, and age at initiation of ICB treatment.

f. t-SNE plot of immune cells in pre-treatment tumors from patients with melanoma from scRNA-seq data.[49] The black arrow indicates the cDC population.

g. *PIKFYVE* expression (log2 (TPM+1)) across immune cell types in nonresponders ("NR" including SD and PD response) or responders ("R" including PR and CR response) to ICB treatment. Data plotted are mean ± s.d. from scRNA-seq data.[49] *P* value determined by student t-test with Welch's correction.

Figure 2: PIKfyve suppresses DC function.
a, b. Splenic DCs in PIKfyve inhibitor-treated non-tumor-bearing mice. **(a)** Relative median fluorescent intensity of surface XCR1 in CD45^{+} cells and (**b**) percentage of cDC1 cells from spleens of mice treated with vehicle or apilimod (30 mg/kg daily) on day 5 of treatment (n = 5 per group). Data plotted are mean ± s.d. *P* values generated from student t-test.

c, d. XCR1 expression in genetic models of *Pikfyve* loss. (**c**) Relative median fluorescent intensity surface XCR1 on *Pikfyve*$^{f/f}$ ("WT") or *Itgax*$^{Tg/0}$ *Pikfyve*$^{f/f}$ ("KO") cDCs on culture day 9. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test. (**d**) Immunoblot of total XCR1 in *Pikfyve* WT vs. KO cDC lysates on culture day 9. Total histone h3 serves as loading control. Images are representative of two experiments.

e-h. MHC-I expression in genetic models of *Pikfyve* loss. (**e**) Relative median fluorescent intensity surface MHC-I (H-2kb, H-2kd) on WT or KO cDCs on culture day 9. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test. (**f**) Immunoblot of total MHC-I in *Pikfyve* WT vs. KO cDC lysates on culture day 9. Vinculin serves as loading control. Images are representative of two experiments. (**g**) Relative median fluorescent intensity of surface MHC-I on DMSO or apilimod-treated cDCs treated for 20 hours on culture day 6. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test. (**h**) Immunoblot of total MHC-I in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Vinculin serves as loading control. Images are representative of two experiments.

i-l. MHC-II expression in PIKfyve inhibition. (**i**) Relative median fluorescent intensity of surface MHC-II (MHC-IA-IE) on *Pikfyve* WT or KO cDCs on culture day 9. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test. (**j**) Immunoblots of total MHC-II (MHC-IA-IE) in *Pikfyve* WT vs. KO cDC lysates on culture day 9. Vinculin serves as loading control. Images are representative of two experiments. (**k**) Relative median fluorescent intensity of surface MHC-II on DMSO or apilimod-treated cDCs treated for 20 hours on culture day 6. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test. (**l**) Immunoblots of total MHC-II in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Vinculin serves as loading control. Images are representative of two experiments.

m, n. Immunoblots of total CD80 and CD86 (**m**) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (**n**) in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Total histone H3 serves as loading control. Images are representative of two experiments.

o, p. Representative dot plots of surface CD80 and CD86 on (**o**) *Pikfyve* WT vs. KO cDC lysates on culture day 9. Images are representative of three experiments. (**p**) Representative dot plots of surface CD80 and CD86 on DMSO or apilimod-treated cDCs +/- PolyI:C (50 μg/ml) or lipopolysaccharide (LPS, 50 ng/ml) treated for 20 hours on culture day 6. Images are representative of two experiments.

q, r. Immunoblots of total CD40 (**q**) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (**r**) in DMSO or apilimod-treated cDC lysates treated with LPS (50 ng/ml) after 20 hours on culture day 6. Vinculin serves as loading control. Images are representative of two experiments.

s-v. OT-I/OT-II cell activation. Percentage of IFNγ^+ OT-I cells after 72 hours of co-

culture with (**s**) *Pikfyve* WT vs. KO or (**t**) DMSO or apilimod pre-treated cDCs +/- sOVA (10 µg/ml) and IFNγ⁺ OT-II cells co-cultured with (**u**) WT vs. KO or (**v**) DMSO or apilimod pre-treated cDCs +/- sOVA. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test.

Figure 3: PIKfyve suppresses NF-κB activation in DCs.
a. Enrichment plots of NF-κB target genes in *Pikfyve* KO vs. WT cells on culture day 9.

b, c. Enrichment plots of NF-κB target genes in DMSO or apilimod-treated cDCs treated for (**b**) 3 hours or (**c**) 8 hours on culture day 6.

d. Immunoblots of p-NF-κB-p65 and total NF-κB-p65 (left) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (right) in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Total histone H3 serves as loading control. Images are representative of two experiments.

e. Immunoblots of p-IκB-α, IκB-α, and IκB-β (left) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (right) in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Total histone H3 serves as loading control. Images are representative of two experiments.

f. Immunoblots of IKK-α, IKK-β, and IKK-γ (left) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (right) in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Total histone H3 serves as loading control. Images are representative of two experiments.

g. Immunoblots of IKK-ε, p-TBK-1, and TBK-1 (left) in *Pikfyve* WT vs. KO cDC lysates on culture day 9 and (right) in DMSO or apilimod-treated cDC lysates after 20 hours on culture day 6. Total histone H3 serves as loading control. Images are representative of two experiments.

h. *Sqstm1* shown on volcano plot of differentially expressed genes in apilimod versus DMSO-treated cDCs treated for 8 hours on culture day 6 by log(fold change) and -log(adjusted *P* value).

i-k. TCGA Pan-Cancer analysis, (**i**) Correlation between *SQSTM1* normalized gene expression and PIKfyve score. Correlation coefficient and *P* value calculated using Pearson's product-moment correlation. (**j**) Forest plot of log-odds of having high vs. low *SQSTM1* for patients with high vs. low NF-κB gene targets expression or (**k**) high vs. low cDC1 maturation score expression.[67] Data from bulk RNA-seq plotted are log-odds with 95% confidence intervals. *P* values are determined by multivariate logistic regression controlling for cancer type.

Figure 4: *Pikfyve* loss in DCs enhances anti-tumor immunity *in vivo*.
a-c. Syngeneic tumor models. (**a**) Subcutaneous tumor volume of MC38 tumors (mm³) in *Pikfyve* KO mice (*n* = 8) or WT mice (*n* = 7). Data plotted are mean ± s.e.m. *P* value

determined by Mann-Whitney U test on Day 24. (**b**) Subcutaneous tumor volume of MCA-205 tumors (mm^3) in *Pikfyve* KO mice (*n* = 6) or WT mice (*n* = 6). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on Day 15. (**c**) Subcutaneous tumor volume of B16F10 tumors (mm^3) in *Pikfyve* KO mice (*n* = 8) or WT mice (*n* = 8). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on Day 15.

d-f. B16F10-OVA tumor model. (**d**) Relative median fluorescent intensity of surface H2-kb-SIINFEKL in WT (*n* = 6) vs. KO (*n* = 6) cDCs from tumor-draining lymph nodes of B16F10-OVA tumors. Data plotted are mean ± s.d. *P* value determined by Mann-Whitney U test. (**e**) Percentage and (**f**) dot plots of SIINFEKL Tetramer$^+$ CD8$^+$ T cells isolated from B16F10-OVA tumors. Bilateral subcutaneous tumors from each WT (*n* = 6) or KO (*n* = 6) mouse were combined for analysis. WT/KO littermate pairs #1-3 (Experiment 1) and pairs #4-6 (Experiment 2) are from two independent experiments. Data plotted are mean ± s.d. *P* value determined by paired Mann-Whitney U test.

g. Subcutaneous tumor volume of B16F10 tumors (mm^3) in *Pikfyve* KO mice or WT mice treated with anti-PD1 antibody (200μg every 3 days) or isotype control (*n* = 8 per group) on Day 15. Data plotted are mean ± s.e.m. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons.

h, i. B16F10 tumor model. (**h**) % cDC1 of all DC cells and (**i**) XCR1 median fluorescent intensity by all DC, cDC1 or cDC2 subset in *Pikfyve* KO mice or WT mice treated with vehicle or apilimod (30 mg/kg daily). Bilateral subcutaneous tumors from each mouse (*n* = 4 per group) were combined for analysis. Data plotted are mean ± s.d. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 6 of treatment.

Figure 5: Immune effects of PIKfyve are DC-dependent.
a, b. MC38 tumor model. (**a**) Subcutaneous tumor volume of MC38 tumors (mm^3) in mice treated with vehicle or apilimod (30 mg/kg x 5 days/week) (*n* = 10 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on day 15 of treatment. (**b**) Representative dot plots of percent IFNγ$^+$ of CD8$^+$ T cells in vehicle or apilimod-treated MC38 tumors. Bilateral subcutaneous tumors from each mouse were combined for analysis.

c. Subcutaneous tumor volume of MC38 tumors (mm^3) in mice treated with vehicle versus apilimod (30 mg/kg x 5 days/week) in wild-type (*n* = 10 tumors per group) or *Batf3*$^{-/-}$ mice (*n* = 4 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 15 of treatment.

d. Subcutaneous tumor volume of B16F10 tumors (mm^3) in mice treated with vehicle or apilimod (30 mg/kg daily) in wild-type or *Batf3*$^{-/-}$ mice (*n* = 10 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 9 of treatment.

e. Schematic of "vaccination" experiment demonstrating pre-treatment with oral (vehicle or apilimod) or subcutaneous (water or PolyI:C) reagents followed by B16F10-OVA tumor inoculation.

f. Individual growth curves of subcutaneous tumor volume of B16F10-OVA tumors (mm^3) following 21 days of pre-treatment with vehicle versus apilimod (30 mg/kg daily) +/- subcutaneous injection of water versus PolyI:C (100 μg on Day 1 and Day 14) (n = 8 tumors per group).

g. Schematic of "vaccine therapy" experiment demonstrating combination therapy treatment with oral (vehicle or apilimod) or subcutaneous (water or PolyI:C) reagents following B16F10-OVA tumor inoculation.

h. Individual growth curves of subcutaneous tumor volume of B16F10-OVA tumors (mm^3) in mice treated with vehicle or apilimod (30 mg/kg daily) +/- subcutaneous injection of water versus PolyI:C (100 μg once weekly) (n = 6 tumors per group).

Supplemental Figure Legends

Extended Data Fig. 1
Supp 1a. Comparison of PIKfyve scores in patients with melanoma who were ICB treatment-naïve ("NAÏVE") versus those who had previously progressed on ICB treatment ("PROG"). Data plotted are mean ± s.d. from bulk RNA-seq data.[44] *P* value is determined by Mann Whitney U test.

Supp 1b. Forest plot of hazard ratios of NAÏVE cohort by high vs. low PIKfyve score, high vs. low neoantigen load, disease stage (reference is M0), and mutation subtype (reference is None). Data plotted are hazard ratios with 95% confidence intervals from bulk RNA-seq data.[44] *P* values are determined by a multivariate cox proportional hazards model.

Supp 1c. Forest plot of hazard ratios of PROG cohort by high vs. low PIKfyve score, high vs. low neoantigen load, disease stage (reference is M1A), and mutation subtype (reference is None). Data plotted are hazard ratios with 95% confidence intervals from bulk RNA-seq data.[44] *P* values are determined by a multivariate cox proportional hazards model.

Supp 1d-g. t-SNE plots of immune cells in pre-treatment tumors from patients with (**d**) breast, (**e**) colorectal, (**f**) lung or (**g**) ovarian cancer from scRNA-seq data.[48]

Supp 1h. Barplots of *PIKFYVE* cluster average expression by cancer, stroma, or immune cell type from scRNA-seq data.[48]

Supp 1i. t-SNE plot of immune cells in pre-treatment tumors from patients with endometrial cancer from scRNA-seq data.[50]

Supp 1j. Comparison of log (*PIKFYVE*) expression in cDCs in patients with endometrial cancer who were nonresponders ("NR" including SD and PD) versus CR to ICB treatment. Data plotted are mean ± s.d. from scRNA-seq data.[50] *P* value is determined by student t-test with Welch's correction.

Extended Data Fig. 2
Supp 2a. Gating strategy for Supp 2 panels b-g for CD8^{+} T cells.

Supp 2b-g. Comparison of the percentage of (**b**) CD69^{+} effector, (**c**) naïve, (**d**) terminally differentiated, (**e**) resident memory, (**f**) central memory, and (**g**) exhausted CD8^{+} T cells from spleens of non-tumor-bearing mice treated with vehicle or apilimod (30 mg/kg daily) on day 5 of treatment (*n* = 5 spleens per group). Data plotted are mean ± s.d. *P* values generated from student t-test.

Supp 2h. Gating strategy for Figure 2 panels a-b and Supp 2 panels i-l for myeloid cells and cDCs.

Supp 2i-l. Comparison of the percentage of total (**i**) CD11c, (**j**) F4/80, and (**k**) CD11b in CD45^{+} cells and (**l**) percentage of cDC2 cells from spleens of non-tumor-bearing mice treated with vehicle or apilimod (30 mg/kg daily) on day 5 of treatment (*n* = 5 per group). Data plotted are mean ± s.d. *P* values generated from student t-test.

Extended Data Fig. 3
Supp 3a. Gating strategy for Figure 2 panels c-r and Supp panels b-h (CD11c^{+} gate).

Supp 3b. Immunoblots of PIKfyve in *Pikfyve* WT vs. KO cDC lysates on culture day 9. Total histone H3 serves as loading control. Image is representative of two experiments.

Supp 3c. Representative density plot of median fluorescent intensity of surface XCR1 in DMSO or apilimod-treated cDCs after 20 hours on culture day 6.

Supp 3d. Comparison of the percentage of CD80^{+}CD86[+-] cDCs in *Pikfyve* WT vs KO models. Data plotted are mean ± s.d. *P* values generated from student t-test across 3 replicates.

Supp 3e, f. Relative median fluorescent intensity of surface H-2kb-SIINFEKL (**e**) and percent H-2kb-SIINFEKL^{+} cells (**f**) in *Pikfyve* WT vs. KO cDC +/- pOVA (100 ng/ml) after 12 hours on culture day 9. Data plotted are mean ± s.d. *P* value determined by student t-test across 3 replicates. Representative plots shown.

Supp 3g, h. Relative median fluorescent intensity of surface H-2kb-SIINFEKL (**g**) and percent H2kb-SIINFEKL^{+} cells (**h**) in DMSO or apilimod-treated cDCs +/- pOVA (100 ng/ml) after 12 hours. Data plotted are mean ± s.d. *P* value determined by student t-test across 3 replicates. Representative plots shown.

Extended Data Fig. 4

Supp 4a. Gating strategy for Supp 4b-e (CD90$^+$ CD8$^+$ gate for OT-I cells).

Supp 4b, c. Representative dot plot and percentage of (**b**) IFNγ^+ and (**c**) granzyme B$^+$ OT-I cells after 48 hours of co-culture with *Pikfyve* WT vs. KO cDCs +/- pOVA 100ng/ml. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test.

Supp 4d, e. Representative dot plot and percentage of (**d**) IFNγ^+ and (**e**) granzyme B$^+$ OT-I cells after 48 hours of co-culture with DMSO or apilimod pre-treated cDCs +/- pOVA 100 ng/ml. *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test.

Supp 4f, g. Percentage of Ki67$^+$ OT-I (**f**) or OT-II (**g**) cells after 72 hours of co-culture with (*Pikfyve* WT vs. KO pre-treated cDCs +/- sOVA (10 μg/ml). *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test.

Supp 4h, i. Percent of Ki67$^+$ OT-I (**h**) or OT-II (**i**) cells after 72 hours of co-culture with DMSO or apilimod pre-treated cDCs +/- sOVA (10 μg/ml). *n* = 3 biological replicates. Data plotted are mean ± s.d. *P* value determined by student t-test.

Extended Data Fig. 5

Supp 5a. Enrichment plot of Dendritic Cell Maturation Up gene signature (M4562: LENAOUR_DENDRITIC_CELL_MATURATION_UP) in *Pikfyve* KO vs. WT cDCs on culture day 9.

Supp 5b. Enrichment plot of MSigDB Hallmark "TNF_SIGNALING_VIA_NFκB" in *Pikfyve* KO vs. WT cDCs on culture day 9.

Supp 5c, d. Enrichment plots of MSigDB Hallmark "TNF_SIGNALING_VIA_NFκB" in apilimod vs. DMSO-treated cDCs treated for (**c**) 3 hours or (**d**) 8 hours on culture day 6.

Supp 5e. Comparison of *Il12b* expression fold change by RT-qPCR in apilimod vs. DMSO-treated cDCs treated for 3 or 12 hours on culture day 6. Data plotted are mean ± s.d. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons across 3 replicates.

Supp 5f. Comparison of IL-12p40 protein levels by ELISA in media from apilimod vs. DMSO-treated cDCs treated for 12 hours on culture day 6. Data plotted are mean ± s.d. *P* values generated from student t-test across 3 replicates.

Supp 5g. Comparison of IL-12p70 protein levels by ELISA in media from apilimod vs. DMSO-treated cDCs treated for 24 hours on culture day 6. Data plotted are mean ± s.d. *P* values generated from student t-test across 3 replicates.

Extended Data Fig. 6

Supp 6a. Kaplan-Meier curve of overall survival of patients in the TCGA Pan-Cancer bulk RNA-seq dataset, by high or low (median) *SQSTM1* normalized gene expression. *P* value is determined by log-rank test.

Supp 6b. Comparison of *Sqstm1* normalized gene expression across myeloid lineage cell types. Data plotted are mean ± s.d. from bulk RNA-seq data.[81] *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons across all pairwise comparisons. *P* value for cDC1 versus cDC2 shown on plot.

Supp 6c. Correlation between *SQSTM1* normalized gene expression and NF-κB target genes expression in the TCGA Pan-Cancer bulk RNA-seq dataset. Correlation coefficient and *P* value calculated using Pearson's product-moment correlation.

Extended Data Fig. 7
Supp 7a-c. Tumor weights for (**a**) MC38 tumor measured on Day 26, (**b**) MCA-205 tumor measured on Day 16, and (**c**) B16F10 tumor measured on Day 16. Data plotted are mean ± s.d. *P* values determined by Mann-Whitney U test.

Supp 7d-f. Subcutaneous tumor volume from (**d**) Experiment 1 and (**e**) Experiment 2 of B16F10-OVA tumors (mm^3) in *Pikfyve* KO mice or WT mice. Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on Day 15. (**f**) Image of B16F10-OVA tumors on Day 16. Bilateral subcutaneous tumors were excised from 3 WT/KO littermate pairs per experiment in two independent experiments.

Supp 7g. Gating strategy for Fig. 4d, Supp. 7h-i ($CD45^+$ Lin(-) $CD11c^+$ MHC-II^+ gate for dendritic cells from tumor-draining lymph nodes).

Supp 7h. Percent dendritic cells of $CD45^+$ cells in *Pikfyve* WT ($n = 6$) vs. KO ($n = 6$) tumor-draining lymph nodes of B16F10-OVA tumors. Data plotted are mean ± s.d. *P* value determined by Mann-Whitney U test.

Supp 7i. Percent H2-kb-SIINFEKL$^+$ dendritic cells in WT ($n = 6$) vs. KO ($n = 6$) tumor-draining lymph nodes of B16F10-OVA tumors. Data plotted are mean ± s.d. *P* value determined by Mann-Whitney U test.

Supp 7j. Gating strategy for Fig. 4e-f (7-AAD$^-$ $CD90^+$ $CD3^+$ $CD8^+$ gate for intratumoral CD8+ T cells).

Extended Data Fig. 8
Supp 8a, b. B16F10 tumor model. (**a**) % cDC2 of all DC cells and (**b**) SIRP1α median fluorescent intensity by all DC, cDC1, or cDC2 subset in *Pikfyve* KO mice or WT mice treated with vehicle or apilimod (30 mg/kg daily). Bilateral subcutaneous tumors from each mouse ($n = 4$ per group) were combined for analysis. Data plotted are mean ± s.d. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 6 of treatment.

Extended Data Fig. 9

Supp 9a. Percent IFNγ^+ of CD8$^+$ T cells in vehicle or apilimod-treated (30 mg/kg x 5 days/week) MC38 tumors. Bilateral subcutaneous tumors from each vehicle or apilimod-treated (n = 5 per group) mouse were combined for analysis. Data plotted are mean ± s.d. *P* value determined by Mann-Whitney U test.

Supp 9b. Subcutaneous tumor volume of MC38 tumors (mm^3) treated with vehicle versus apilimod (30 mg/kg x 5 days/week) (n = 6 tumors per group) in NSG^{TM} mice. Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on day 15 of treatment.

Supp 9c. Subcutaneous tumor volume of MC38 tumors (mm^3) in mice treated with vehicle versus apilimod (30 mg/kg x 5 days/week) (n = 6 tumors per group) treated with IgG2a or αIL-12p40 blocking antibody. Data plotted are mean ± s.e.m. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 15 treatment.

Supp 9d. Subcutaneous tumor volume of MC38 tumors (mm^3) in mice treated with vehicle versus apilimod (30 mg/kg x 5 days/week) (n = 6 tumors per group) treated with IgG1 or αIFNγ blocking antibody. Data plotted are mean ± s.e.m. *P* value determined by ANOVA after post-hoc Tukey adjustment for multiple comparisons on day 15 of treatment.

Supp 9e. Subcutaneous tumor volume of B16F10 tumors (mm^3) in mice treated with vehicle or apilimod (30 mg/kg daily) (n = 10 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on day 9 of treatment.

Supp 9f. Subcutaneous tumor volume of B16F10 tumors (mm^3) in mice treated with vehicle versus apilimod (30 mg/kg daily) (n = 10 tumors per group) in NSG^{TM} mice. Data plotted are mean ± s.e.m. *P* value determined by Mann Whitney U test on day 11 of treatment.

Supp 9g, h. Relative median fluorescent intensity of (**g**) surface MHC-I (H2-kb, H2-kd) and (**h**) surface MHC-II (MHC-IA-IE) on DMSO or apilimod-treated cDCs +/- PolyI:C (50 μg/ml) or LPS (50 ng/ml) treated for 20 hours on culture day 6. Images are representative of two experiments.

Supp 9i. Subcutaneous tumor volume of B16F10-OVA tumors (mm^3) following 21 days of pre-treatment with vehicle versus apilimod (30 mg/kg x 5 days/week) +/- subcutaneous injection of water versus PolyI:C (100μg on Day 1 and Day 14) (n = 8 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on Day 14 after tumor inoculation.

Supp 9j. Subcutaneous tumor volume of B16F10-OVA tumors (mm^3) in mice treated with vehicle or apilimod (30 mg/kg x 5 days/week) +/- subcutaneous injection of water

versus PolyI:C (100µg once weekly) (n = 6 tumors per group). Data plotted are mean ± s.e.m. *P* value determined by Mann-Whitney U test on Day 11 of treatment.

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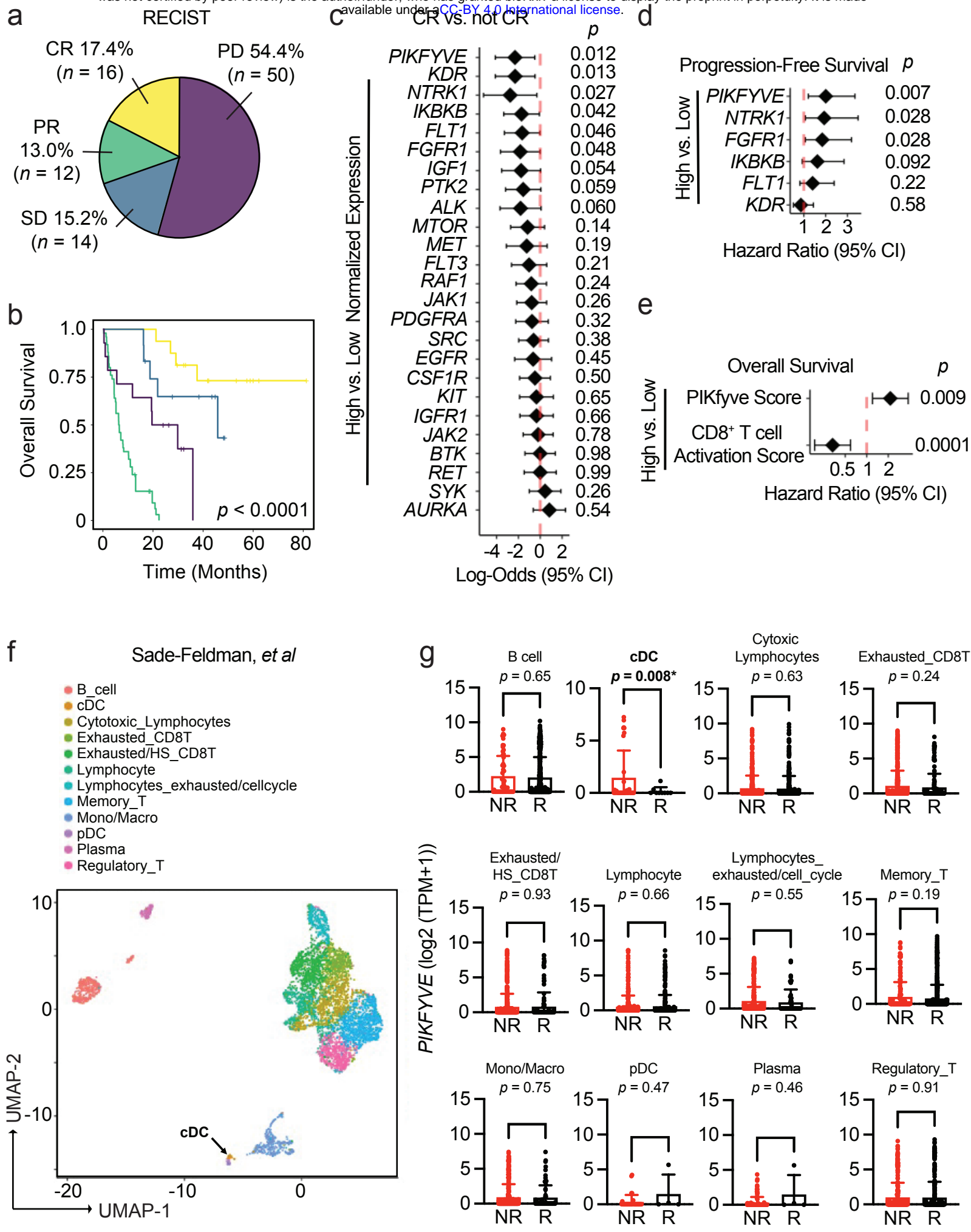
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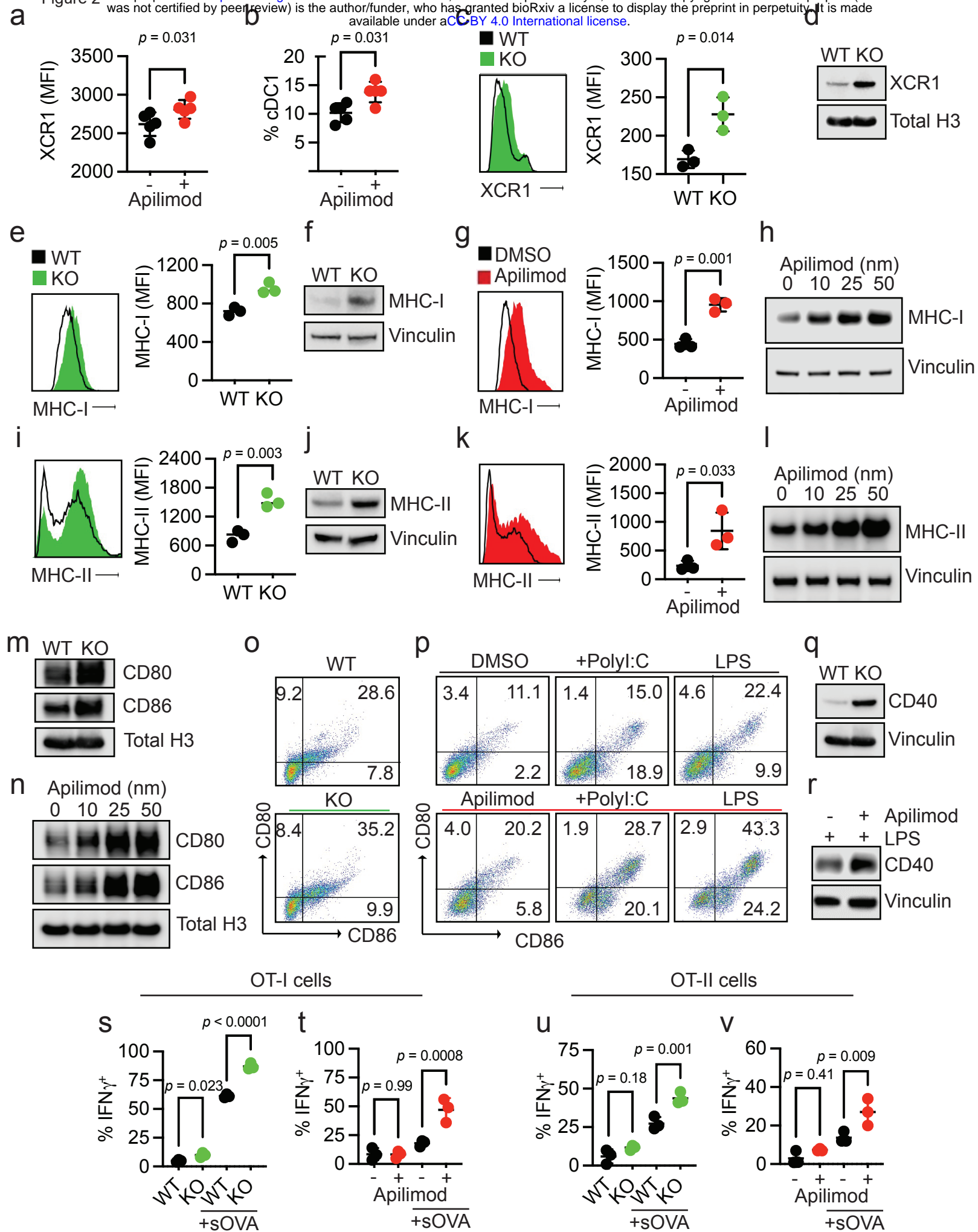
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Figure 1

Figure 2

Figure 3

Figure 5