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Macrophage IL-1 β and TNF- α create an immune-metabolic loop regulating Arginase2 in neuroblastoma

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Abstract

Neuroblastoma is the most common solid tumour of childhood, yet the prognosis for high-risk disease remains poor. We demonstrate that arginine metabolism via Arginase 2 (ARG2) drives neuroblastoma cell proliferation. Targeting arginine metabolism by blocking Cationic Amino Acid Transporter 1 (CAT-1) dependent arginine uptake in vitro or therapeutic depletion of arginine by pegylated-recombinant arginase BCT-100 significantly delays tumour development and prolongs murine survival. Tumour cells polarise infiltrating-monocytes to a M1-macrophage phenotype, which release IL-1 β and TNF- α in an RAC- α serine/threonine-protein kinase (AKT)-dependent manner. IL-1 β and TNF- α signal-back to upregulate ARG2 expression via p38 and Extracellular regulated kinases 1/2 (ERK1/2) signalling in neuroblastoma and neural crest-derived cells. Proteomic analysis reveal Stage IV human tumour microenvironments are enriched in IL-1 β and TNF- α , which is associated with a worse prognosis. Thus we describe an immune-metabolic regulatory loop between tumour cells and infiltrating myeloid cells regulating ARG2, which could clinically exploited.

Statement of significance: Neuroblastoma polarised macrophages released IL-1 β and TNF- α which signal back to regulate Arginase2 in tumour cells and drive their proliferation

Introduction

The consumption and metabolism of diverse nutrients by cancer cells is recognised as a key regulator of immunity. Glucose metabolism by cancer cells generates a tumour microenvironment that has low levels of glucose, leading to inhibition of T cell cytotoxicity through the accumulation of lactate, microenvironment acidification, and reduced aerobic glycolysis (1-3). Tumour infiltrating monocyte differentiation and cytokine release may be similarly affected, leading to perturbation of their role in coordinating the surrounding immune response (4, 5). Amino acid metabolism also plays a critical role in the function of both normal and malignant cells. Although whole body amino acid homeostasis is regulated through restricted inter-organ enzyme expression, at the cellular level enzyme expression is controlled in the intracellular compartment to maintain metabolic precursor supplies and regulate the wider tissue microenvironment (6).

Arginine is a semi-essential amino acid which is metabolised into ornithine and urea by the expression of cytoplasmic Arginase 1 (ARG1) and mitochondrial Arginase 2 (ARG2), or Nitric Oxide Synthase (NOS) enzymes into reactive nitric oxide species (7). These metabolites feed forward into diverse roles in cell signalling, proliferation and protein synthesis. Cellular breakdown of arginine also plays a critical role in regulating the immune response, a process which has been capitalised on by malignant cells to contribute to their immune escape (8). We recently identified that Acute Myeloid Leukaemias (AML) and neuroblastoma, two of the most common and devastating cancers of childhood create a potent immunosuppressive microenvironment through the expression of ARG2 enzyme which suppresses T-cell immunity (9, 10).

Although the metabolic effect of cancer cells on shaping the responsiveness of surrounding immune populations is increasingly well described, the reciprocal effects of immune cell populations on modulating cancer cell amino acid metabolism have not previously been reported. In particular the role of arginine metabolism in this process is unknown and the signals which regulate ARG2 in cancer are not well understood. Here we demonstrate how myeloid cells within the tumour microenvironment and tumour cells engage in reciprocal cross-talk to regulate the expression of ARG2 in neuroblastoma cells, and how this arginine metabolism plays a central role in neuroblastoma pathogenesis. Importantly, this study identifies arginine metabolism as a clinically relevant therapeutic target.

Materials and Methods

Patient Samples

Heparinised blood and tumour samples were obtained from 50 patients with neuroblastoma treated at the Birmingham Children's Hospital and Children's Hospital Oxford. Samples were obtained from patients with newly diagnosed neuroblastoma, at the time of diagnostic biopsy or before the start of treatment. GD2+ neuroblastoma cells were isolated from bone marrow aspirates taken from patients with stage IV disease.

Neuroblastoma murine model

Transgenic Tg(*TH-MYCN*)^{41Waw} mice were genotyped to detect the presence of human *MYCN* transgene or the Chromosome 18 insertion site, using an allelic discrimination methodology (11, 12). Specific assays were designed to measure the presence of the *MYCN* transgene (forward primer 5'-CGACCACAAGGCCCTCAGTA; reverse primer 5'-CAGCCTTGGTGTGGAGGAG; probe 6FAM-CGCTTCTCCACAGTGACCACGTCG TAMRA; Eurofins) or to the site of the transgene on chromosome 18 which is disrupted during insertion (forward primer 5'-CCACAAAATATGACTTCCTAAAAGATTT; reverse primer 5'-CATGGGACTTCCTCCTTATATGCT; probe VIC-5'-AACAATTATAACACCATTAGATATG TAMRA). After weaning, *TH-MYCN* mice were palpated for intra-abdominal tumours twice weekly. Mice with palpable tumours ranging in size between 5-20mm in diameter were then humanely sacrificed. At sacrifice, unheparinised and heparinised whole blood, as well as tumour tissue were obtained for further ex vivo analyses. Tumour tissue was processed as

above. Tumour tissues were stained with anti-mouse GD2 (BioLegend) on ice for 30 minutes. The expression of these markers was then assessed by flow cytometry.

For treatment with BCT-100, mice were treated with 60mg/kg BCT-100 or saline, twice a week, *ip* either from weaning in the prophylaxis setting or upon the development of a 5mm tumour in the treatment setting. Mice were treated until the experimental endpoint of a 10mm abdominal tumour. In the prophylaxis experiment, mice were bled before the start of treatment, midway through the treatment, 24 hours after the fifth dose of either saline or BCT-100, and at tumour endpoint. All experimental protocols were monitored and approved by either The Institute of Cancer Research Animal Welfare and Ethical Review Body, in compliance with guidelines specified by the UK Home Office Animals (Scientific Procedures) Act 1986 and the United Kingdom National Cancer Research Institute guidelines for the welfare of animals in cancer research or the University of New South Wales Animal Care and Ethics Committee and conducted according to the Animal Research Act, 1985 (New South Wales, Australia) and the Australian Code of Practice for Care and Use of Animals for Scientific Purposes (2013).

GD2+ tumour cell and myeloid cell isolation

For isolation of GD2+ tumour cells from human and murine tumours were digested using Type II collagenase, labelled with anti-GD2-PE antibody (BioLegend) and bound to anti-PE coated magnetic beads (Miltenyi Biotec, Bisley, UK). Cells were enriched according to manufacturer's instructions to be >98% GD2+ cells as confirmed by flow cytometry using a PE conjugated anti-human GD2 antibody. For isolation of primary GD2+ cells from the bone marrow of diagnosed stage IV patients, bone marrow aspirates were collected in RPMI 1640

media containing 10% FCS. Cells were lysed using erythrocyte lysis buffer (Qiagen) and the white cell fraction isolated by centrifugation. The neuroblastoma cells were labelled with purified mouse anti-human GD2 Clone 14.G2a (BD Pharmingen) and bound to anti-mouse IgG2a/b microbeads (Miltenyi Biotec). Cells were enriched according to manufacturer's instructions (Miltenyi Biotec). For isolation of monocytes, peripheral blood was collected from healthy donors. Monocytes were separated using a Lymphoprep gradient (STEMCELL Technologies) and enriched by positive selection using anti-human CD14 MicroBeads (Miltenyi Biotec).

Cell lines and cultures

Human primary, untransformed, embryonic neural crest (R1113T) or dorsal root and/or sympathetic ganglion-derived stem cells (SZ16) were obtained and cultured as previously described (13-15). Neuroblastoma cell lines (SKNAS, KELLY, IMR-32, LAN-1), the Ewing's sarcoma cell line SKNMC which has high ARG2 expression, and primary GD2+ neuroblastoma cells were cultured in RPMI 1640 medium (Sigma) supplemented with 10% v/v foetal bovine serum (FBS, Sigma), 100 U/mL penicillin and streptomycin (Gibco), 1mM sodium pyruvate (Gibco) and 2mM L-Glutamine (Gibco). All cell lines were originally obtained from ATCC and validated for authenticity by DNA short tandem repeats in line with American National Standards Institute ASN-0002-2011 (Northgene). All experiments were performed between passages 3-9, and cells were confirmed as Mycoplasma negative by PCR analysis (LookOut, SIGMA. Latest testing date September 2018). The effects of arginine deprivation were tested on cells cultured in arginine-free RPMI 1640 for SILAC

(ThermoFisher Scientific) supplemented with 10% v/v arginine-free dialysed FBS (ThermoFisher Scientific). Cells were maintained in an incubator at 5% CO₂ in air and at 37°C.

Arginase activity assays

The activity of arginase 2 present within cells was determined by measuring the conversion of arginine into urea, as previously described (10).

Antibody microarray analysis

Human stage I neuroblastoma tissue samples (n=13), human stage IV neuroblastoma biopsies (n=9) were analysed using scioDiscover antibody microarrays (Sciomics) which targets 900 cancer-related proteins (16). After sample homogenisation, proteins were extracted with scioExtract buffer (Sciomics) and labelled at an adjusted concentration with scioDye 2 (Sciomics) according to the manufacturer's instructions. A pool of all protein samples was labelled with scioDye1 and used as a reference for all experiments, allowing competitive dual-colour measurements. Array production, blocking and sample incubation were performed in compliance with strict quality control procedures as reported previously. The arrays were scanned with identical instrument laser power and adjusted PMT setting using a Powerscanner (Tecan). Spot segmentation was performed with the software GenePix Pro 6.0 (Molecular Devices).

Enzyme-linked Immunosorbent Assays (ELISA)

The concentrations of cytokines IFN- γ , IL-1 β , TNF- α , TGF- β , IL-6, IL-4, IL-13 and GM-CSF in plasma and cell culture media were measured by sandwich-ELISA kit according to specific manufacturer's instructions.

Monocyte-driven proliferation assays

Neuroblastoma cells suspended at a density of 1×10^6 cells/mL in PBS were labelled with 1 μ M CellTrace™ FarRed staining solution (Molecular Probes, ThermoFisher Scientific) at 37°C for 20 minutes. Stained cells were washed three times in RPMI-1640 and rested for 10 minutes in complete media. Labelled neuroblastoma cells were then cultured in supernatants from neuroblastoma-induced macrophages (75% final volume), with or without 1 μ g/mL anti-IL1 β (R&D Systems, Catalog #MAB201) and 1ng/mL anti-TNF α (Cell Signalling, Catalog #7321s) neutralising antibodies. Cells were harvested 5 days later and analysed on a CytoFLEX Flow Cytometer (Beckman Coulter). Histograms representing distinct generations of proliferation cells were generated using the FlowJo Software (TreeStar Inc.).

Reverse transcriptase polymerase chain reactions

Total RNA was extracted from cells using either the RNEasy Kit (Qiagen) according to the manufacturer's specifications. Extracted RNA was quantified on a NanoDrop ND-1000 spectrophotometer (ThermoScientific). First strand complimentary DNA (cDNA) was generated by incubating 1 μ g of extracted RNA with 500ng of random primers (Promega),

0.5mM dNTP (Promega), 1x reverse transcriptase buffer (Promega), 40U RNase inhibitors (RNasin, Promega) and either 100U MMLV RNase H⁺ or 15U AMV reverse transcriptase (Promega). For endpoint PCR reactions, up to 100ng of sample cDNA was incubated in 5 μ L of 10X PCR reaction buffer (Invitrogen), 0.5mM dNTPs, one unit of *Taq* polymerase (Invitrogen), 1.5mM MgCl₂, 0.5 μ M of each forward and reverse primer and nuclease free water up to a final reaction volume of 50 μ L. Human primer sequences are listed in Table S1. All quantitative PCR (RT-qPCR) reactions were conducted on a Fast 7500 real-time PCR thermal cycler (Applied Biosystems).

TaqMan Assays

RNA was isolated using the miRNeasy Mini Kit (Qiagen) and RNA concentration determined by spectrophotometry (NanoDrop 1000, ThermoFisher Scientific). RNA (10 ng per replicate) was reverse transcribed using Superscript™ III Reverse Transcriptase (ThermoFisher Scientific) according to manufacturer's instructions with random hexamer primers (0.3 μ g, ThermoFisher Scientific) and RNasin Plus RNase Inhibitor (20 units, Promega). Samples were analysed in triplicate. Samples analysed in the absence of RT enzyme or without RNA were included as negative controls. cDNA was amplified using TaqMan Gene Expression Assays for each target (Table S2, ThermoFisher Scientific) according to manufacturer's instructions. Expression of the housekeeping gene PPIA was determined for each sample using sequence specific reverse and forward primers (200nM forward primer GGACCCAACACAAATGGTTCC, 200nM reverse primer CTTTCACTTTGCCAAACACCA, 100nM FAM labelled probe ATGCTTGCCATCCAACCACTCAGTCTTG). mRNA expression was calculated using the

comparative Ct method relative to PPIA. RNA from Cell lines known to express genes of interest were included as control (Table S2)

Study approval

In accordance with the Declaration of Helsinki, patient samples were obtained after written informed consent prior to inclusion in the study. Primary human neural crest-derived stem cell lines were obtained under ethical committee approval PFS14-011 from the French Biomedical Agency for the use of embryonic material. Regional Ethics Committee (REC 10/H0501/39) and local hospital trust research approval for the study was granted for United Kingdom hospitals. The Institute of Cancer Research Ethics Committee approved all animal protocols in this study. Collection of diagnostic bone marrow aspirates from Stage IV patients was performed under ethical approval of the Medical Research and Ethics committee (MREC/98/4/023). Procedures were carried out in accordance with UK Home Office Guidelines.

Antibody microarray normalisation and statistical analysis

The acquired raw data were analysed using the linear models for microarray data (LIMMA) package of R-Bioconductor after uploading the median signal intensities. As described previously, a specialised invariant Lowess method was applied for normalisation.⁽¹⁷⁾ For the differential analysis of protein expression, a one-factorial linear model was fitted with LIMMA resulting in a two-sided t-test or F-test based on moderated statistics. Differences in protein abundance between sample groups are presented as log-fold changes (logFC) calculated for the basis 2. The presented p-values were adjusted for multiple testing by

controlling the false discovery rate according to Benjamini and Hochberg. In all comparisons, proteins were defined as significantly differential with a log-fold change above 0.5 or below -0.5 and an adjusted p-value below 0.05. Functional enrichment analyses were conducted with the STRING software (<https://string-db.org>) for the proteins with significantly differential abundance between groups, whereby up- and downregulated proteins were analysed separately.

Arginase 2 Fluorescence Intensity

Quantification of cell-by-cell fluorescence intensity for Arginase 2 expression across treatment conditions were performed using ImageJ software (National Institute of Health, USA). Briefly confocal image stacks were converted to single channel images. Pixel intensity measurements were determined from single channel images representing Arginase 2 staining with Image thresholds set to match positive structures within defined cell boundaries.

Statistical analysis

Parametric student t-tests were used to determine the statistical significance of the difference in paired observations between groups (GraphPad Prism, USA). All p values are two-tailed and p values <0.05 were considered to represent statistically significant events. Significance was recorded as * p<0.05, ** p<0.01, ***p<0.001, ****p<0.0001.

Results

Neuroblastoma conditioned macrophages release IL-1 β and TNF- α in the tumour microenvironment

Myeloid cells are a major orchestrator of cancer-related inflammation with the potential to support tumour growth, invasion and metastasis. In neuroblastoma we have previously shown a significant increase of immunosuppressive myeloid cells in peripheral blood of patients and in the tumour tissue of the transgenic neuroblastoma TH-MYCN murine model. However the role of intratumoral myeloid cells in human neuroblastoma tumours is not well understood. To understand the landscape inside human tissue at diagnosis, we first investigated the proteomic profile of 23 human neuroblastoma tumours (9 Stage I and 14 Stage IV) using a novel antibody array (18). Non-metric multidimensional scaling from protein array for all analysed samples based on the complete protein expression data revealed separate clustering of Stage I and Stage IV tumours (Figure 1A). Analysis of human neuroblastoma proteomes showed increased expression of the monocyte/macrophage marker CD14 and the granulocytic cell marker CD15 in high stage disease (Figure 1B). Immunohistochemistry of tissue microarrays of 27 tumours, revealed CD14⁺ cells infiltrated the tumour tissue (Figure 1C upper panels with histoscore Fig 1D, Supp Fig 1A) whilst CD15 staining localised around vasculature (Figure 1C lower panels with histoscore Fig 1D, Supp Fig 1A). Together these findings highlight the potential role of myeloid cells in tumourigenesis.

Monocyte function may be modulated by their environment. To investigate the influence of neuroblastoma tumour cells on monocytes, monocytes enriched from healthy donor blood

were co-cultured with sorted Ganglioside G2 (GD2)+ neuroblastoma cells from patients or cell lines. We observed that neuroblastoma conditioning led to upregulation of the macrophage marker CD68 and only a minority of cells upregulated CD206 (M2 marker) (Figure 1E and Supp Figures 1B and 1C). In addition, myeloid ARG1 activity was down-regulated (Figure 1F) consistent with polarisation to a M1-phenotype. Importantly immunohistochemistry of neuroblastomas at diagnosis confirmed the infiltration of these CD68+ macrophages within the tumour tissue (Figure 1G and Supp Figure 1D).

Tumour infiltrating myeloid cells can shape the immune response through cytokine release within tumour tissue (19). To investigate the cytokine profile of neuroblastoma induced macrophages, a broad panel of cytokines was analysed in culture supernatants. Tumour-conditioning led to an increased release of IL-1 β and TNF- α , with undetectable levels of IL-13, IL-6, IFN- γ , IL-4, TGF- β and GM-CSF consistent with a M1-phenotype (Figure 2A and B, Supp Figure 2A). Tumour cells alone released minimal cytokines (<8 pg/ml, Supp Figure 2B). Although CD15+ granulocytes released IL-8, they did not release either IL-1 β or TNF- α following tumour co-culture (Supp Figure 2C). To prove that the release of IL-1 β and TNF- α was from the macrophages, intracellular staining for cytokines was performed. Neuroblastoma conditioning led to an increased frequency of IL-1 β and TNF- α positive macrophages at 24hours and 48hours (Figure 2C, Supp Figure 3A and 3B). Using confocal microscopy we confirmed CD14+ cells sorted from patients expressed IL-1 β and TNF- α (Supp Figure 3C) and immunohistochemistry of tissue microarrays of 27 tumours confirmed the expression of IL-1 β and TNF- α in the tumour-infiltrating macrophages. (Fig 2D; Supp. Figure 4A).

IL-1 β and TNF- α secretion from myeloid cells may be regulated by AKT signalling (20). Co-culture of healthy donor monocytes with neuroblastoma led to AKT phosphorylation (Figure 2E) and AKT inhibition with MK-2206 prevented IL-1 β and TNF- α release (Fig 2F and G). No evidence for STAT3, NF- κ B, or PI3K pathway activation was identified (Supp Fig 4B). Therefore neuroblastoma cells polarise surrounding monocytes to M1-macrophages which release IL-1 β and TNF- α .

Neuroblastoma cell proliferation is dependent on arginine metabolism

Previously we established that neuroblastoma cells consume arginine from the microenvironment and catabolise this amino acid by ARG2 to create an immunosuppressive microenvironment contributing to immune escape and suboptimal immunotherapy responses (9). However, the role of ARG2 in neuroblastoma development and more widely in human cancers has only received limited study. Arginine metabolism can contribute to cell proliferation. To investigate the role of ARG2 in tumour cell proliferation, we first performed shRNA knock-down for ARG2. ARG2 knock-down led to a significant reduction in cell proliferation (Figure 3A, Supp Figure 4C) confirming the key role of this enzyme. We next blocked uptake of arginine from the microenvironment via Cationic Amino Acid Transporter-1 (CAT1), which we showed is expressed in the majority of neuroblastoma cell lines (Supp Figure 4D). N-nitro-L-arginine (L-NAME) inhibitor led to a significant decrease in tumour cell proliferation (Figure 3B). Culture of tumour cells in the absence of arginine similarly inhibited tumour cell metabolic activity (Figure 3C). BCT-100 is a PEGylated

recombinant human arginase that can deplete arginine to undetectable levels in cancer patients leading to clinical responses in adult trials (21). Culture of neuroblastoma with BCT-100 led to a rapid inhibition of cell proliferation (Supp Figure 4E), and tumour cell death characterised by PARP cleavage (Supp Figure 5A). Electron microscopy of sorted tumour cells from cell lines and patients confirms loss of cell membrane integrity, and cellular fragmentation (Figure 3D).

To investigate the *in vivo* dependence of tumour growth on arginine we used the immunocompetent *TH-MYCN* transgenic mouse model which spontaneously develop neuroblastoma tumours (11). These murine tumour cells also express ARG2 (Supp Figure 5B). We first demonstrated that *ex vivo* treatment of murine GD2+ tumour cells with BCT-100 led to a significant reduction in viable cells (Figure 3E). Treatment of *TH-MYCN* mice with twice-weekly BCT-100 led to a sustained drop in plasma arginine to almost undetectable levels (Figure 3F). To understand if tumour initiation could be delayed or prevented in the absence of arginine, mice were treated prophylactically from the time of weaning at 3 weeks of age, when the tumours were 1-2 mm in size. Neuroblastoma development was significantly delayed and mice survived for significantly longer in the BCT-100 treated group compared to control ($p=0.0001$, Figure 3G). Following this, we investigated the effect of BCT-100 on established tumours. Here we showed that murine tumour progression was significantly delayed compared to the saline control and overall survival was significantly extended ($p=0.0181$, Figure 3H). Arginine re-synthesis pathway enzymes Argininosuccinate Synthase (ASS) and Ornithine Transcarbamylase (OTC) were not

upregulated in GD2+ cells from murine tumours as mechanisms of resistance (Supp Fig 5C). No evidence for BCT-100 drug toxicity in terms of weight or clinical features were identified.

Macrophage IL-1 β and TNF- α drive tumour ARG2 expression via p38/ERK signalling

As ARG2 contributes to tumour cell proliferation, we hypothesised that these macrophage-derived cytokines may reciprocally regulate ARG2 expression. We first showed that the treatment of neuroblastoma cells with low basal expression of ARG2 (SKNAS and IMR32) with IL-1 β and TNF- α , either alone or in combination, resulted in upregulated ARG2 expression (Figure 4A and B). Sorted human GD2+ neuroblastoma cells similarly upregulated ARG2 in response to cytokines (Figure 4C). Consistent with this finding, supernatant from tumour-induced macrophages upregulated ARG2 in neuroblastoma cells (Figure 4D, Supp Figure 5D). To investigate whether the M1-macrophages would therefore enhance neuroblastoma cell proliferation via ARG2, we cultured neuroblastoma cells with induced-macrophage supernatants. Supernatants led to increased cell proliferation in neuroblastoma cells (Figure 4E – red) compared to the untreated cells (Figure 4E-black). The phenotype was partially reversed by the addition of IL- β and TNF- α neutralising antibodies (Figure 4E-green, and Supp Fig 5E).

Neuroblastoma is a pathological derivative of trunk-level neural crest cells, which normally develop into diverse populations including catecholamine-secreting cells of the adrenal medulla, sympathetic, parasympathetic and sensory neurons, and multipotent Schwann cell precursors (22, 23). We hypothesised that a microenvironment containing similar factors to postnatal inflammation may contribute to tumour initiation by upregulating ARG2 in these

embryological cells. Analogous to neuroblastoma, treating cultures of normal human embryonic ganglion precursors with IL-1 β and TNF- α led to a significant upregulation of ARG2 protein expression (Figure 4F), demonstrating the inherent responsiveness of neural crest progenitors to these signals before oncogenic transformation.

We determined that neuroblastoma express the receptors for IL-1 β and TNF- α (Figure 5A). The Interleukin 1 Receptor 1 (IL1R1) and Tumour-Necrosis Factor Receptor 1 (TNFR1) receptors can induce a signalling cascade that both converge on a common final effector pathway through ERK1/2 and p38 activation, and Ribosomal Protein S6 Kinase A5 (MSK1) activation (Figure 5B) (24-26). Treatment of neuroblastoma cells with IL-1 β and TNF- α leads to NF κ B phosphorylation by 0.5 hours and subsequent phosphorylation of ERK1/2 from 1 hour onwards (Figure 5C). Simultaneously, the cytokines also induced p38 phosphorylation at 0.5 hours (Figure 5C). PD98059 binds inactive ERK and prevents phosphorylation and activation by upstream mediators, while SB20308 inhibits p38 catalytic activity but does not affect phosphorylation. The resulting inhibition of p38 or ERK1/2 signalling leads to subsequent downregulation of ARG2 expression in SKNAS neuroblastoma cells (Figure 5D). Blockade of either p38 or ERK1/2 in isolation is insufficient to prevent ARG2 upregulation by cytokines due to compensation by the other side of the pathway (Fig 7B and Supp Fig 5F). MSK1 is at crossroads of the common downstream cascade and can be auto-regulated by kinases including ERK1/2 and p38. SB747651A blockade of MSK1 activity, which is phosphorylated from 0.5 hours onwards (Figure 5C), similarly prevented cytokine-induced ARG2 upregulation (Figure 5E).

The 1β and TNF- α enriched intra-tumoural microenvironment is associated with high-stage disease

We previously showed that ARG2 expression is highest in Stage IV tumours and is associated with a worse overall survival (9). Cytokines may be functional either within the tumour microenvironment or released into the blood to induce systemic effects. Analysis of blood from 25 neuroblastoma patients at diagnosis revealed that the majority of patients did not have significantly increased TNF- α and IFN- γ compared to healthy controls, although in 9 cases circulating levels of IL-1 β and IL-6 were significantly higher ($p=0.042$) (Figure 6A).

We hypothesised that the intratumoral cytokines driving arginine metabolism in neuroblastoma would promote high stage human neuroblastoma development. To investigate this we further analysed the proteomic profile inside 23 human neuroblastoma tumours. Heat-map representation of protein signals reveal that Stage I and Stage IV tumours show distinct molecular proteomic subgroups, with 7 Stage IV tumours (P21-27) forming a distinct group, while 3 others (P10,P15,P16), had proteomes more similar to Stage I tumours (Figure 6B). Consistent with our in vitro findings, characterisation of the Stage IV tumours identified significantly higher levels of the M1-macrophage derived cytokines IL-1 β and TNF- α than Stage I tumours (Figure 6C). In contrast, Stage I tumours had increased expression of the M2-related cytokines TGF- β , IL-10, and IL-4 (Figure 6D). No significant differences in IL-6 and IL-13 expression were identified. Consistent with this, analysis of the expression profile of 88 neuroblastomas (GEOID: GSE16476) revealed high expression of *IL-1 β* or *TNF- α* within tumours is associated with a significantly worse overall survival for neuroblastoma patients ($p=0.012$ and $p=0.027$ respectively, Figure 7A and B).

Discussion

Although it is well established that amino acid metabolism can regulate anti-cancer immunity, the capacity of the immune system to regulate cancer amino acid metabolism has rarely been characterised. In this study we identify a key reciprocal regulation between tumour cell arginine metabolism and intra-tumoural macrophages in neuroblastoma. The regulators of ARG2 expression in cancer are poorly understood despite abundant data on its cytoplasmic counterpart ARG1. ARG2 can be upregulated by hypoxia in osteosarcoma cells and non-malignant cells, whilst in pancreatic ductal adenocarcinoma models obesity correlated with increased ARG2 levels and enhanced tumour growth (27, 28)(29, 30). Studies of cytokine regulation of Arginase 2, are limited to non-malignant cells with reports that Th1 or Th2 cytokines have no effects on murine myeloid cells or can modulate ARG2 expression in murine neural stem cells.(31) In humans IL-10 may regulate ARG2 in combination with isoproterenol in macrophages.(32)

We and others have previously reported the ability of neuroblastoma to modulate circulating monocytes into an immunosuppressive phenotype on T cells and NKT cells. (33) Here we demonstrate that the tumour cells also polarise intratumoral monocytes to M1-macrophages, which express and release IL-1 β and TNF- α after AKT signal transduction. AKT inhibitors, such as Perifosine, have recently undergone early phase clinical trial development including evaluation in refractory neuroblastoma, with initial results suggesting that targeting this pathway could prolong progression-free survival (34). We show that tumour-polarised macrophages act back to regulate cancer cell arginine metabolism through IL-1 β and TNF- α , and drive tumour cell proliferation. Recently murine macrophages were shown

to increase neuroblastoma proliferation in association with STAT3 phosphorylation, although the factor responsible was not identified.(35) We demonstrate that ARG2 expression is under the control of both p38 and ERK1/2 in human neuroblastoma cells, which lie downstream of the receptors for IL-1 β and TNF- α (IL1R1 and TNFR1 respectively). The role of these cytokine pathways in cancer cell expression of ARG2 has not previously been reported. Some redundancy in the signalling cascade is evident, such that inhibition of both receptor pathways, or of their common effect on MSK1, is required to inhibit enzyme expression.

We showed that the Stage IV intratumoural microenvironment is enriched in the expression of IL-1 β and TNF- α whilst the converse is true for Stage I tumours. To date, the role of IL-1 β and TNF- α in neuroblastoma has primarily centred around the effects of these cytokines on neuroblastoma cell lines used as models of neurodegenerative disease, such as Alzheimer's disease. In terms of its effects on the malignant phenotype, recombinant TNF- α has been shown to be a growth factor for neuroblastoma cell lines, although the mechanism of action was unknown (36). A minor subset of neuroblastoma cells within tumours, may themselves express TNF- α intracellularly or on the cell membrane, but they do not release the cytokine into the microenvironment (33). For IL-1 β , little is known in the context of neuroblastoma, although it is reported to drive cyclo-oxygenase (COX-2) expression in neuroblastoma Alzheimer's disease cell line models (37). Importantly, we identified that levels of IL-1 β and TNF- α proteins in the plasma are not significantly greater than in healthy donors, indicating that it is the intra-tumoural interactions that are key.

Clinically it is clear that Stage I and Stage IV neuroblastomas are distinct at the levels of tumour dissemination, responses to chemotherapy, and patient outcome. Inter-cellular signalling within tumours remains difficult to characterise, although much has been learned from transcriptomic and epigenetic profiling of these tumours (38, 39). The functional interaction of proteins within the cellular ecosystem must be contributing to variation in tumour aggressiveness, although analysis of multiple proteins inside tumours is challenging. To our knowledge, this study is also the first proteomic characterisation of human neuroblastomas at diagnosis and the findings suggest that array-based proteomic profiling can lead to new insights into tumour immunobiology. It has been hypothesised that an immune-stimulatory event, such as infection in early childhood could contribute to the development of childhood cancers either through a normal or aberrant response. Indeed a 'delayed infection' hypothesis had been suggested for childhood acute lymphoblastic leukaemia (40). Although specific infections like Epstein-Barr Virus are directly linked to malignant transformation of cells in Hodgkin's Lymphoma or nasopharyngeal carcinoma, in the majority of paediatric malignancies no evidence of clearly defined cause and effect have been found (41, 42). It is possible that an isolated pro-inflammatory response within a tissue microenvironment could potentially lead to a cytokine profile that drives metabolism in malignant or pre-malignant cells, giving them a survival advantage, allowing development into a frank malignancy. The inflammation could be secondary to very specific infectious agents or an abnormal, pathological response due to immune defects.

Neural crest cells are highly multipotent stem cells in the embryo which give rise to diverse cell types such as melanocytes, odontoblasts, peripheral neurons and support cells,

including those of the dorsal root, sympathetic and parasympathetic ganglia, and specific endocrine cells in the thyroid and parathyroid glands and the adrenal medulla(22). We found that neural crest-derived primary cells are enriched in ARG2 protein, relative to ARG1. The role of ARG2 in embryological processes is not well understood. Neonatal CD71+ erythroid cells express ARG2, which may affect the response to commensal bacteria in the developing baby, while dendritic cells in the developing foetus also express ARG2 to modulate immune responses *in utero* (43, 44). That the expression of ARG2 in untransformed neural crest-derived stem cells, can be upregulated by immune cytokines IL-1 β and TNF- α points to the potential for metabolic changes to occur during malignant transformation or expansion. We have previously shown that AML blasts have similarly upregulated ARG2 in comparison to their non-malignant haematopoietic counterparts (10). Indeed, knock-down of ARG2 significantly reduces the ability of both types of tumour cells to proliferate, suggesting this enzyme provides an advantage to cancer growth and dissemination.

Although arginine metabolism under cytokine control can drive neuroblastoma proliferation, this axis also provides a potential therapeutic target. Targeting tumour-associated myeloid cells has received significant attention to date. Although depletions of myeloid cells can be achieved *in vivo* using anti-GR1 or anti-CR2 antibodies, the effects are very short-lived in mice and no human equivalent exists for clinical translation. One approach to target the feedback loop we have described is to inhibit IL-1 β and TNF- α cytokine activity. Anti-TNF- α therapy was the paradigm for anti-cytokine therapies with the

development of anti-TNF- α antibody (infliximab) and a decoy anti-TNF-A receptor (etanercept). Although these antibodies have demonstrated remarkable activity in autoimmune conditions, they have only undergone limited study in the setting of cancer therapy. Infliximab has been used as a single agent in patients with advanced cancer, with some patients experiencing disease stabilisation (45). The drug has also been trialled to treat renal cell carcinoma and although improvements in immune profiles were noted, there were also significant increases in adverse events (46, 47). Similar antibodies against IL-1 β (canakinumab) and its receptor IL-1R1 (anakinra) also exist. Although canakinumab has not been formally tested in patients with an existing cancer, administration of this drug has been shown to significantly reduce incidences of lung cancer and its mortality in patients with atherosclerosis (48). Future combination clinical trials of these agents could represent a novel and potential approach in children with neuroblastoma.

It is now possible to successfully target cancer arginine metabolism through therapeutic arginine depletion with BCT-100, a PEGylated recombinant arginase that induces sustained arginine depletion for months in human trials (49, 50). The drug has completed Phase I/II trials in adult malignancies with an excellent safety profile (21). In this study we demonstrated that BCT-100 not only leads to a decrease in neuroblastoma proliferation with accompanying cell death *in vitro*, but also to delayed progression and prolonged survival in neuroblastoma-bearing mice. These findings support the testing of BCT-100 in an international Phase I/II clinical trial (PARC, NCT03455140) in children with relapsed/refractory malignancies including neuroblastoma. The targeting of both immune

and metabolic drivers of tumorigenesis as presented in this study, is rational and clinically achievable, and could be a new paradigm in the treatment of neuroblastoma.

Acknowledgements

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Figure Legends

Figure 1: Neuroblastoma induces M1-macrophages

A) Non-metric multi-dimension scaling of Stage I (RED) and Stage IV (BLUE) tumours at diagnosis shows distinct proteomic profiles for these two stages of tumours B) Proteomic analysis of Stage I and IV tumours identifies significantly higher expression of CD14 and CD15 in Stage IV tumours C) Immunohistochemical staining of sections from neuroblastomas showing infiltration of CD14+ (upper) and CD15+(lower) myeloid cells. Representative sections shown of n=27, tissue microarray D) Histoscores of CD14 and CD15 staining in neuroblastoma tissue microarrays of n=27 tumours E) CD14+ monocytes from healthy donors co-cultured with sorted GD2+ tumour cells from patients upregulate CD68 expression (upper). Minimal CD206 upregulation was seen. Representative flow cytometry shown (n=5) F) CD14+ monocytes from healthy donors co-cultured with neuroblastoma have decreased arginase activity, as assessed by conversion of ornithine to urea in a colorimetric assay. (n=3) G) Immunohistochemical staining of sections from neuroblastomas showing infiltration of CD68+ macrophages. Representative sections shown of n=27, tissue microarray

Figure 2: Tumour-induced macrophages cells release IL-1 β and TNF- α through p-AKT signalling

ELISA of supernatants following co-culture of healthy donor monocytes with neuroblastoma cell lines, showing increased IL-1 β (A) and TNF- α (B) (n=7) C) Co-culture of monocytes from healthy donors with tumour cell lines for 48hours leads to upregulation of IL-1 β and TNF- α expression, compared to those cultured in RPMI10% media. Flow cytometry staining

shown, gated on CD14⁺ cells. Representative staining from 3 independent experiments D) Immunohistochemical staining of sections from neuroblastomas showing infiltration of CD33+IL-1 β + and CD33+TNF- α + macrophages. Representative sections from n=27 TMA shown E) CD14⁺ myeloid cells from healthy donors were sorted following co-culture with neuroblastoma cell lines. Co-culture leads to increased expression of p-AKT, as shown by Western blotting (n=3) Addition of AKT inhibitor MK2206 to co-cultures of CD14⁺ cells and neuroblastoma cell lines leads to inhibition of IL-1 β (F) and TNF- α (G) release (n=3) protein expression.

Figure 3: Neuroblastoma proliferation is dependent on arginine metabolism

A) shRNA knock-out of ARG2 in SKNMC (high baseline ARG2 expression) decreases cell proliferation. Fold change in cell number after 72h compared to baseline. Experiment performed in duplicate. Corresponding Western blots for ARG2 in wild-type and knock-down cell lines shown below, with actin as a loading control B) Proliferation of tumour cell lines is inhibited by CAT1 inhibition with L-NAME, measured by ³H-thymidine incorporation after 72 hours C) Cell lines were cultured with RPMI+10%FBS (R10%) or arginine-free RPMI+10%FBS (R10%-arginine). Metabolic activity was measured by MTT after 72h. n=7 replicates D) Sorted GD2⁺ neuroblastoma cells from patients were treated with BCT-100 (600ng/mL). Analysis of cell death was performed by transmission electron microscopy (Representative micrographs of 2 out of 6 patients shown). Upper panel show untreated cells. Lower panels show post treatment with 600ng/mL BCT-100. Features consistent with organelle enlargement, cell membrane permeabilisation, and cellular fragmentation with 600ng/mL BCT-100. Experiments performed on 3 separate occasions E) Sorted GD2⁺ cells

from TH-MYCN murine neuroblastomas were cultured with BCT-100 (600ng/mL) for 72 hours. The percentage of viable cells relative to untreated controls was determined by flow cytometry, using propidium iodide staining. BCT-100 leads to a decrease in murine neuroblastoma cell viability *ex vivo* F) Plasma from control (saline) and BCT-100 treated TH-MYCN mice was collected at the start (PRE), 16 days after (MID), and at tumour end-point (END). The concentration of arginine was determined by ELISA. BCT-100 maintains a significant reduction in the plasma arginine concentration *in vivo*. n=6 G) TH-MYCN mice were treated with BCT-100 (60mg/kg) twice weekly intraperitoneally (*ip*) from the time of weaning at 3 weeks of age before overt tumour formations (Prophylaxis). Kaplan-Meier curves show that the development of tumours is significantly delayed, and that survival is increased in BCT-100 treated mice H) TH-MYCN mice were treated with BCT-100 (60mg/kg) twice weekly *ip* once 5 mm tumours were palpable (Treatment). Kaplan-Meier curves show a significant prolongation of survival in BCT-100 treated mice.

Figure 4: IL-1 β and TNF- α upregulate Arginase 2 expression and tumour cell proliferation

Treatment of neuroblastoma cells SKNAS (A) and IMR32 (B) with recombinant cytokines alone or in combination leads to upregulation of ARG2, measured by western blot. Actin is shown as a loading control. Corresponding densitometry of ARG2 relative to actin shown. Representative of n=6 replicates C) Treatment of sorted GD2+ primary neuroblastoma cells (Patient 53 and Patient 54) with cytokines leads to upregulation of ARG2, measured by Western blot. Actin is shown as a loading control. Corresponding densitometry of ARG2 relative to actin shown D) Representative confocal microscopy of neuroblastoma cell line SKNAS shows expression of arginase 2 is increased following culture with the supernatants

of neuroblastoma-induced macrophages. DAPI – blue, ARG2 – green, MitoTracker – red (n=3) E) Neuroblastoma cell proliferation is enhanced in the presence of neuroblastoma-induced macrophage conditioned supernatants (MCM). The addition of anti-TNF α and IL-1 β antibodies (inhibitors) reversed the proliferative effects of MCM. Cell proliferation of neuroblastoma shown by dilution of Cell Trace reagent, measured by flow cytometry F) Treatment of embryonic dorsal root ganglion stem cell line SZ16 with recombinant cytokines alone or in combination leads to upregulation of ARG2, as measured by Western blot. Actin is shown as a loading control. Corresponding densitometry of ARG2 relative to actin shown. Representative of n=3 replicates.

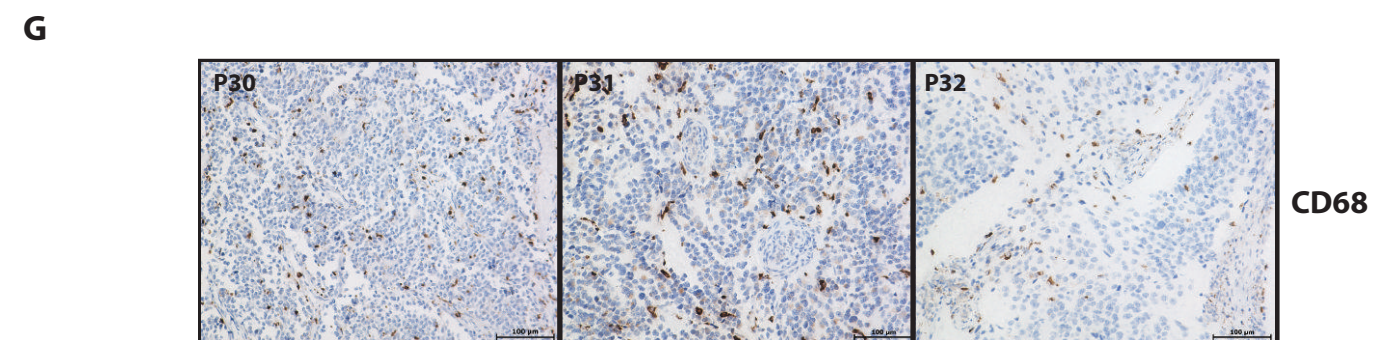
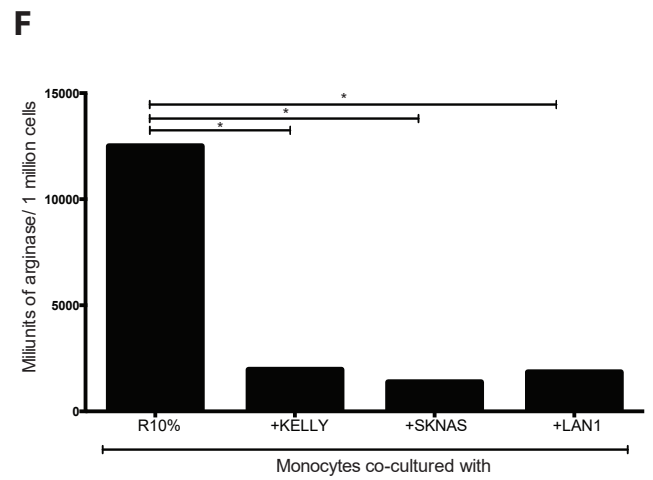
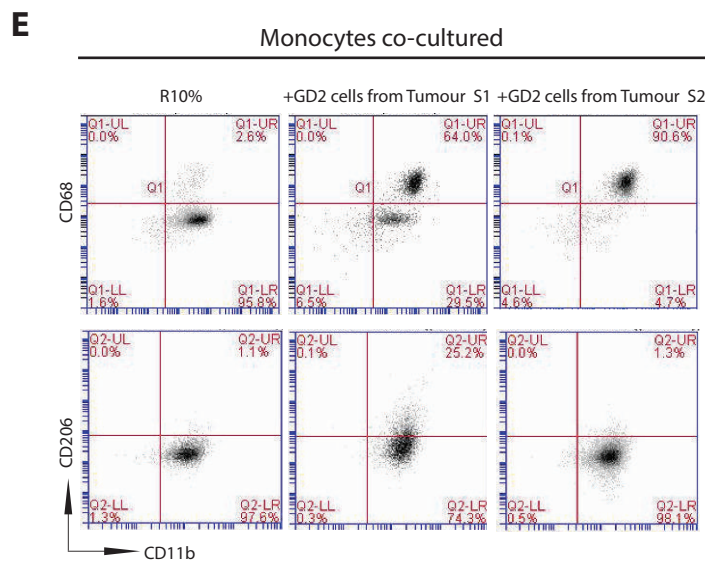
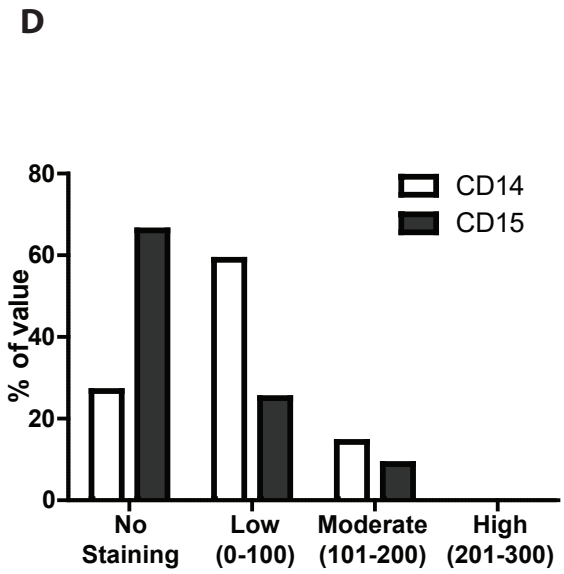
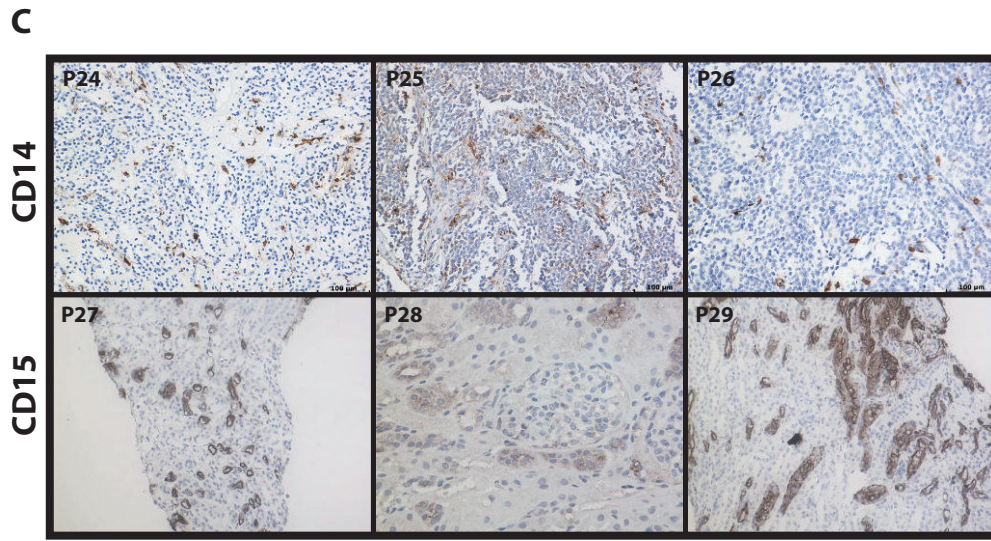
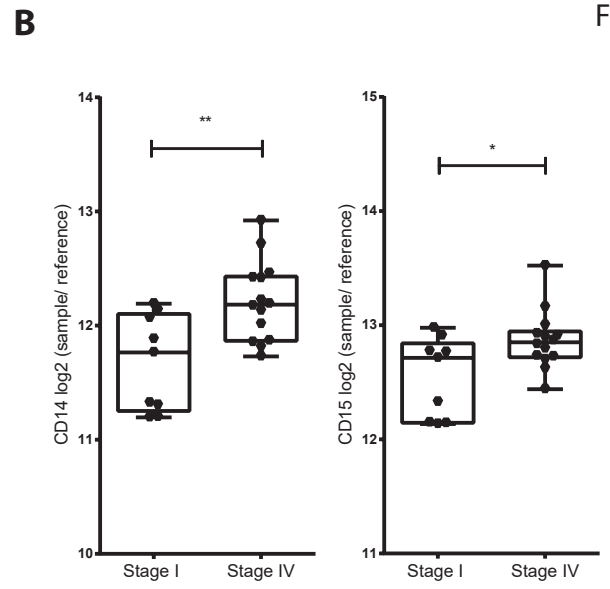
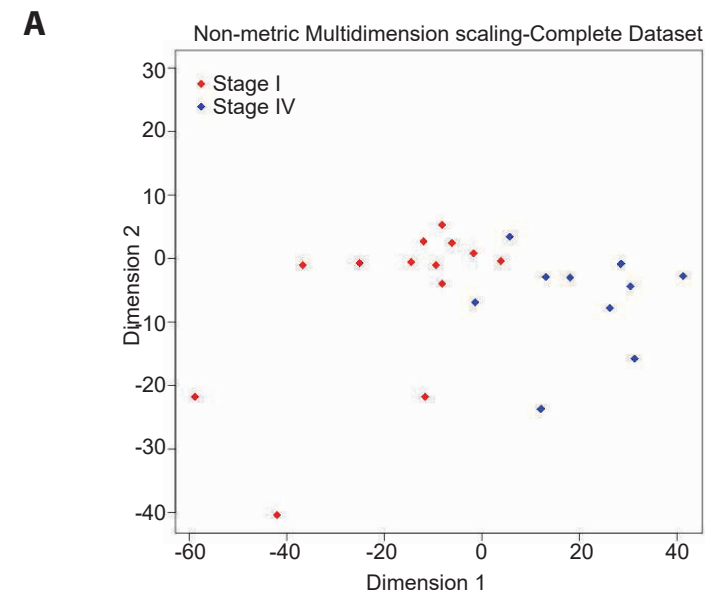
Figure 5: IL-1 β and TNF- α drive Arginase 2 expression in a p38/ERK dependent manner

A) Neuroblastoma cell lines express the IL-1 β and TNF- α receptors on the cell surface as assessed by flow cytometry. (n=3) B) Schematic showing the signalling pathway for IL-1 β and TNF- α cytokines, via ERK1/2, p38 and MSK1 C) Time course (hours) in which IL-1 β and TNF- α lead to increased p-NF κ B (0.5 hours), p-ERK1/2 (from 1 hour onwards), p-p38 (0.5 hours) and p-MSK1 (0.5 hours onwards). ERK1/2, p38, and MSK1 activity are inhibited by PD90859, SB203508, and SB747651A respectively. Western blot shown. Representative of n=3 replicates D) Treatment of SKNAS neuroblastoma cells with recombinant cytokines leads to upregulation of ARG2, which is inhibited by ERK1/2 and p38 inhibition. Western blot shown with actin as a loading control. Corresponding densitometry of ARG2 relative to actin shown N=3 replicates E) Treatment of SKNAS neuroblastoma cells with recombinant cytokines leads to upregulation of ARG2, which is inhibited by MSK1 inhibition. Western blot

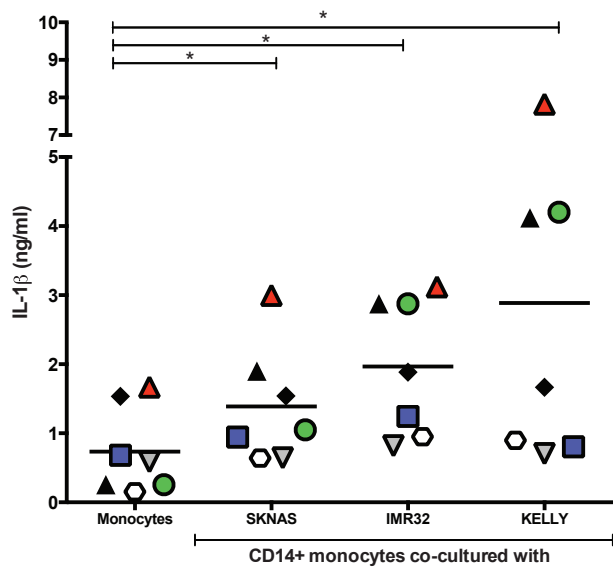
shown with actin as a loading control. Corresponding densitometry of ARG2 relative to actin shown n=3 replicates

Figure 6: The Stage IV neuroblastoma intratumoural microenvironment is enriched in IL-1 β and TNF α A) ELISA Quantification of cytokine titres in neuroblastoma patient plasma (n=26) at diagnosis identifies no significant differences in circulating levels of TNF- α and IFN- γ . Circulating IL-1 β concentrations were significantly higher in some patients at diagnosis. B) Heatmap of Stage I (RED) and Stage IV (BLUE) tumours at diagnosis shows distinct proteomic profiles for these two stages of tumours C) Proteomic analysis of Stage I and IV tumours at diagnosis identifies significantly higher expression of the IL-1 β , TNF- α , as well as IFN- γ in Stage IV tumours D) Stage I tumours express significantly higher Th2 cytokines TGF- β , IL-10, and IL-4 by proteomic analysis

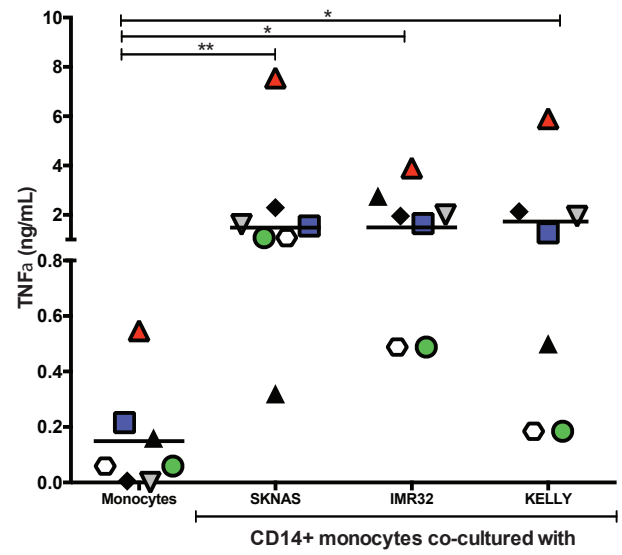
Figure 7: High IL-1 β or TNF- α expression in tumours correlate with a worse overall survival for patients Kaplan-Meier curves of n=88 neuroblastoma patients at diagnosis identifying high IL-1 β (A) or TNF- α (B) expression in tumours is associated with a worse overall survival. All data are analysed in accordance with the public Versteeg database 'R2: microarray analysis and visualization platform' (<http://r2.aml.nl>).



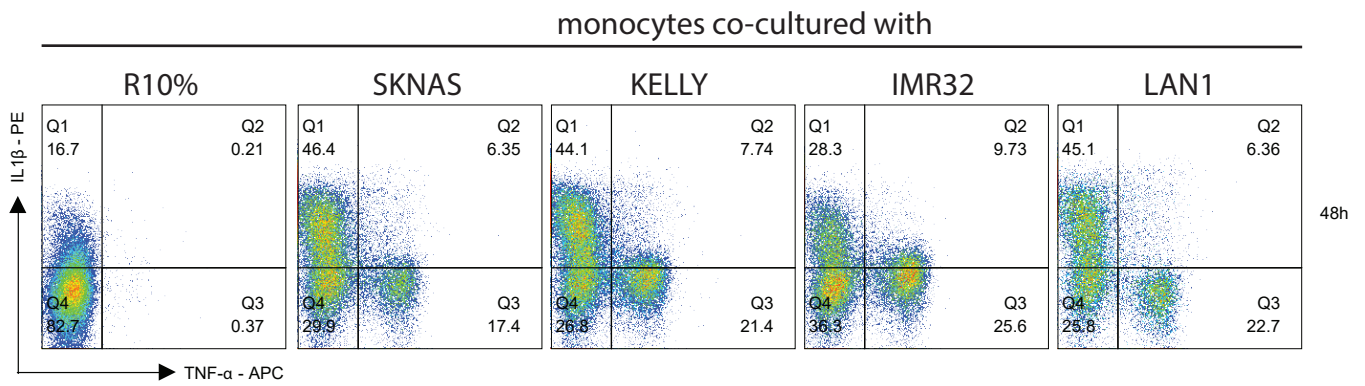
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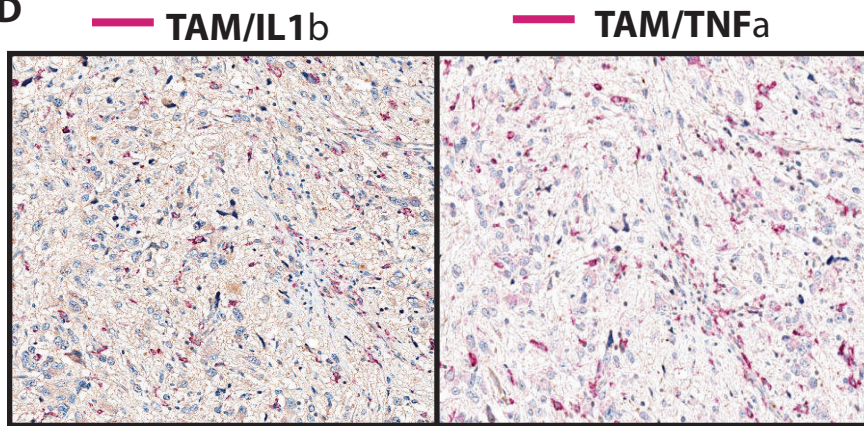
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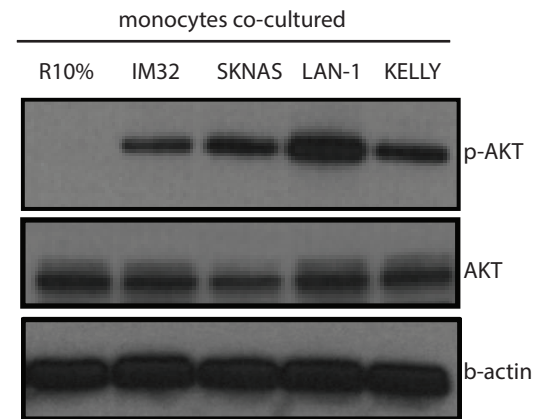
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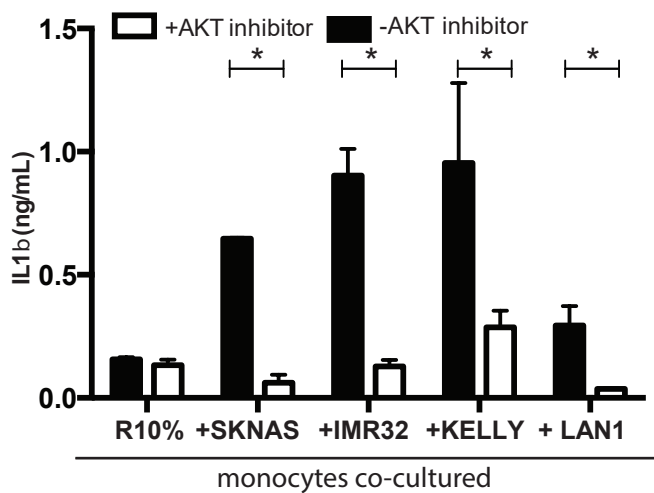
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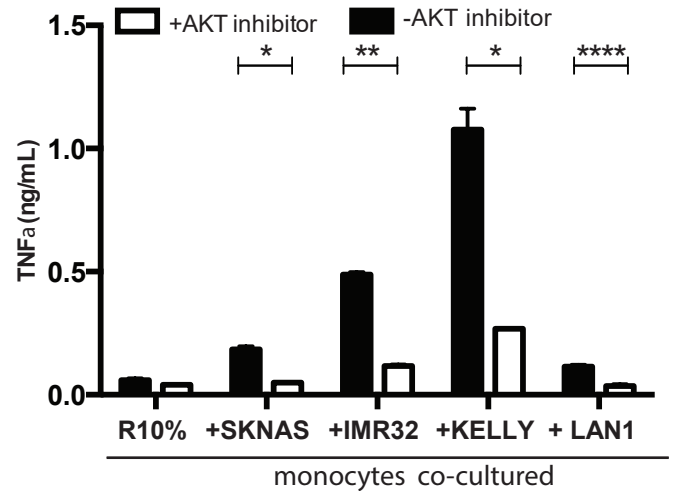
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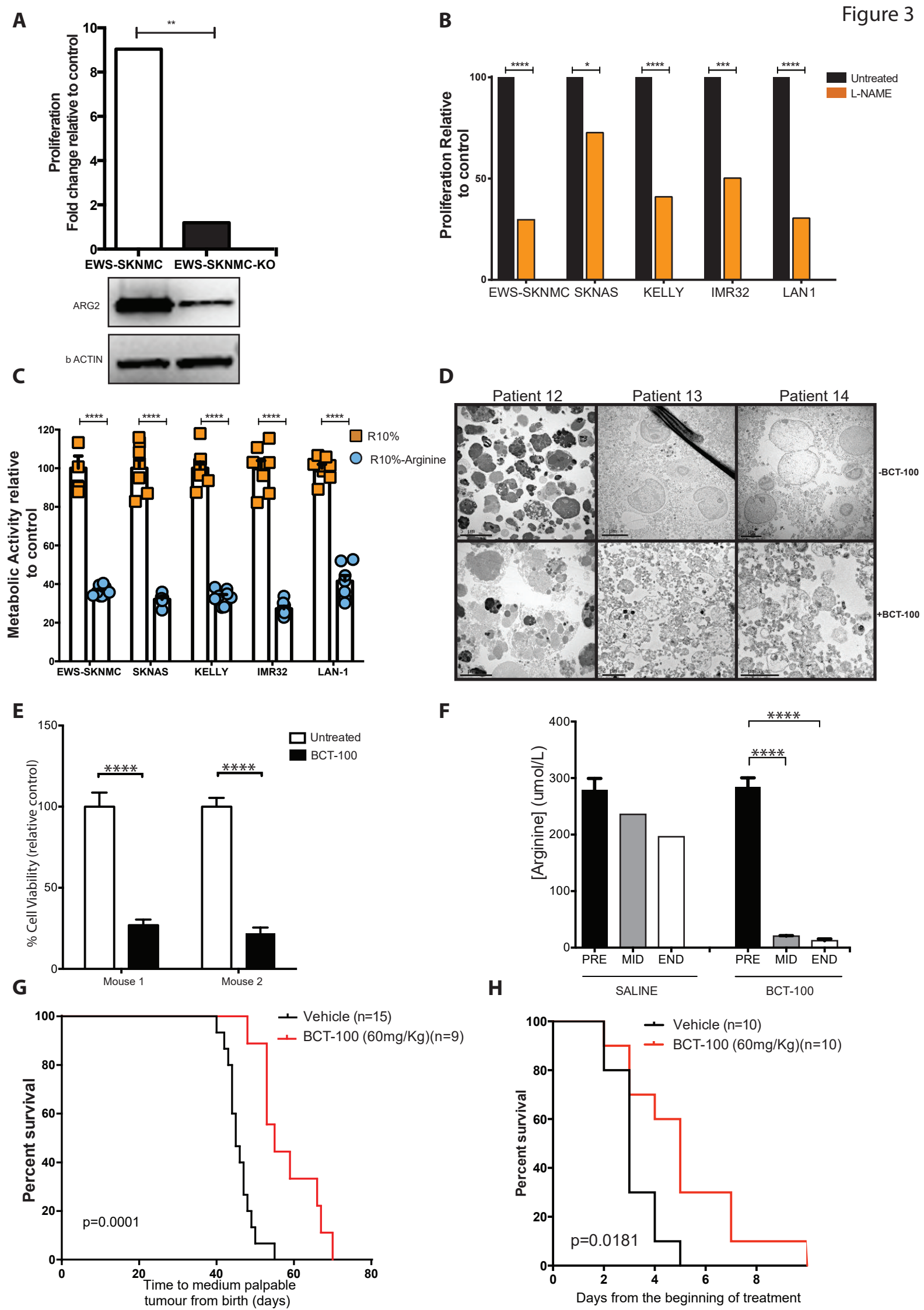


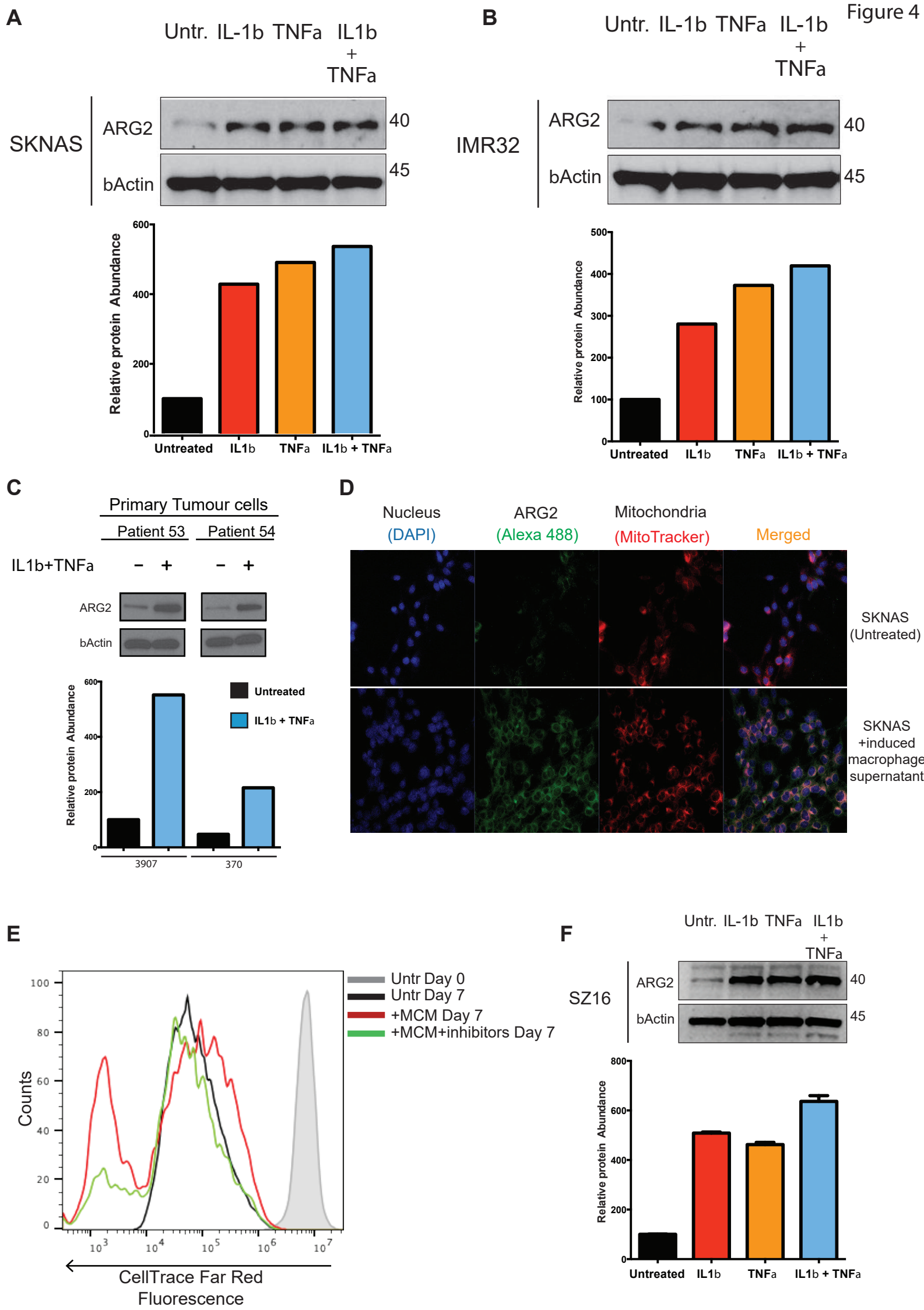
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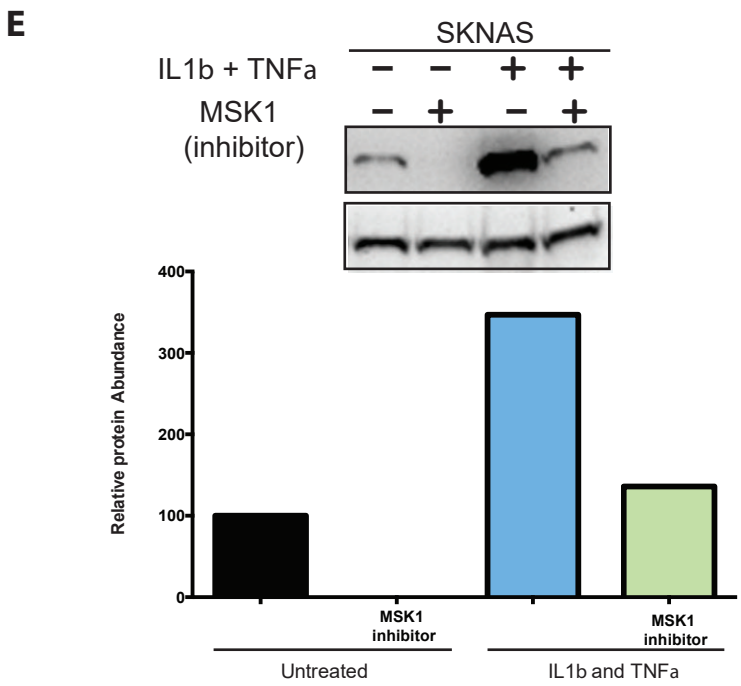
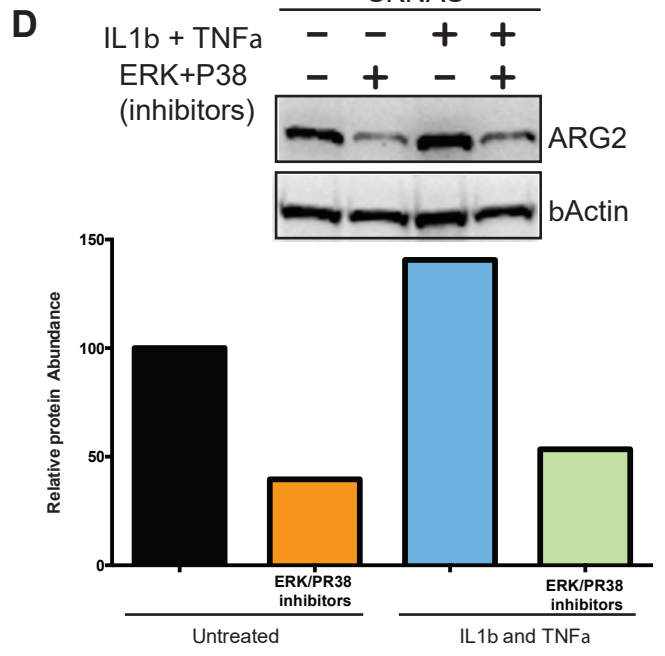
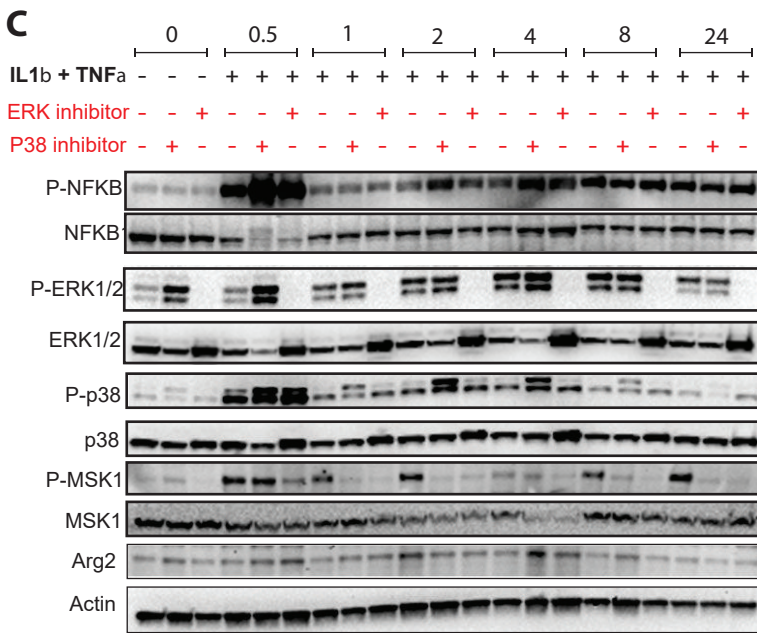
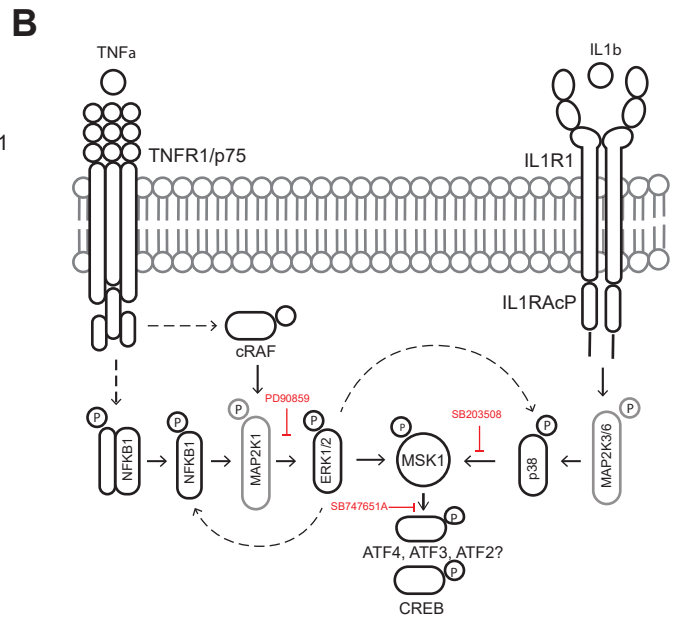
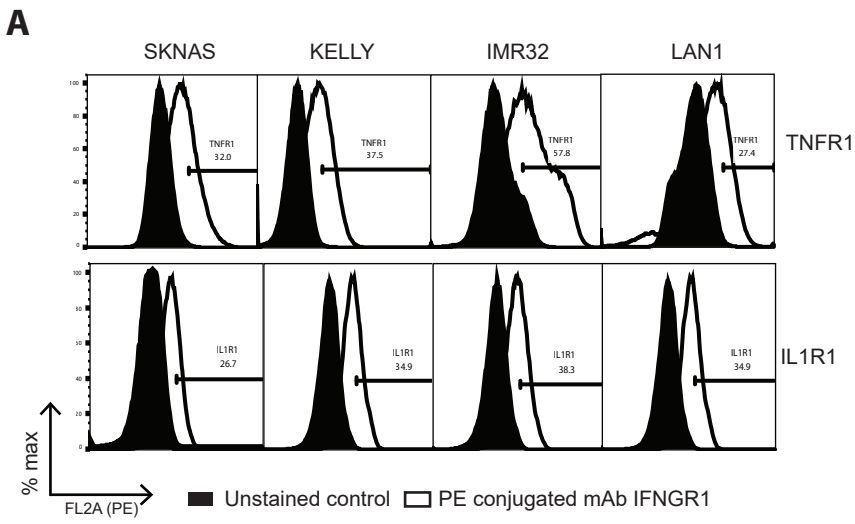


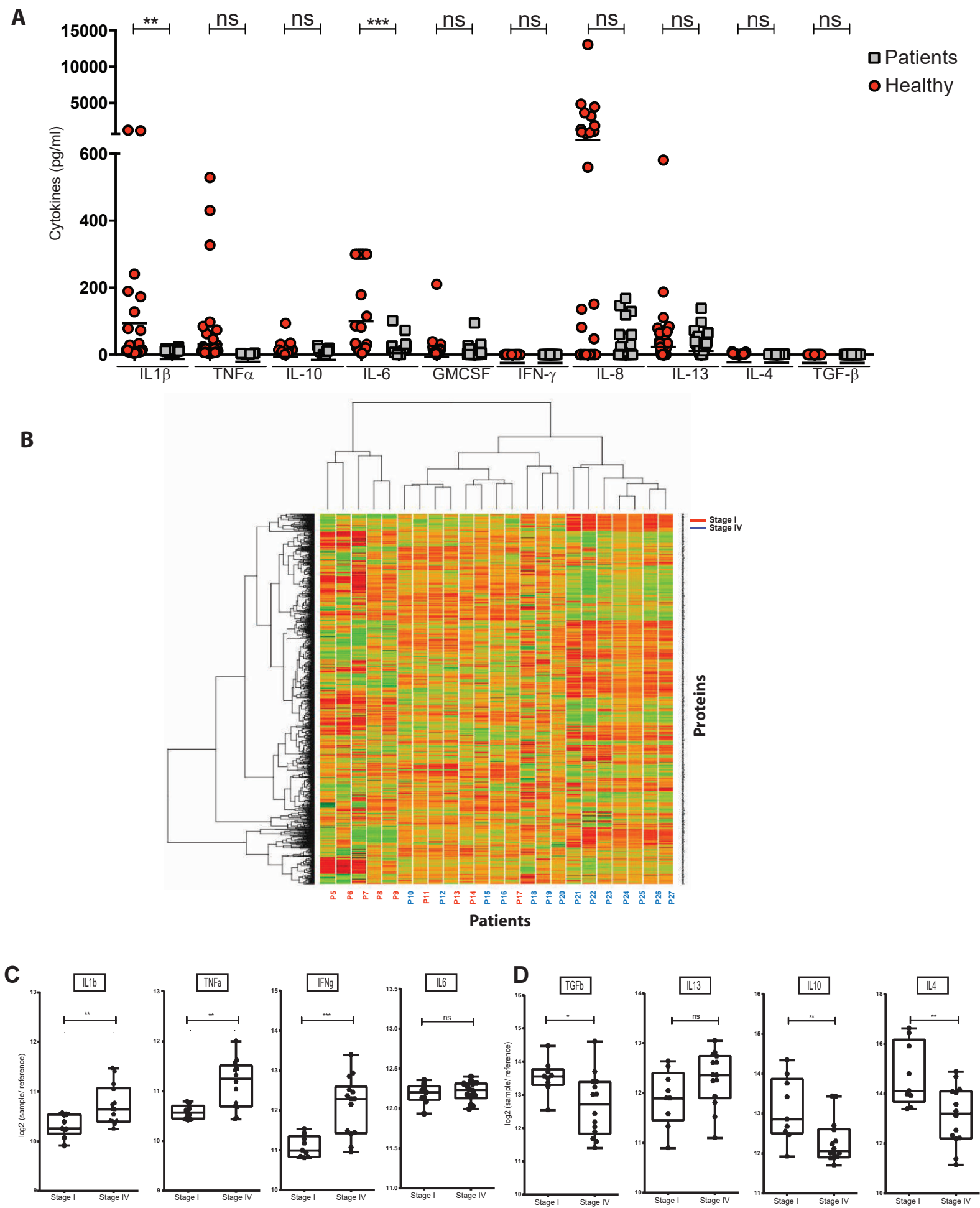
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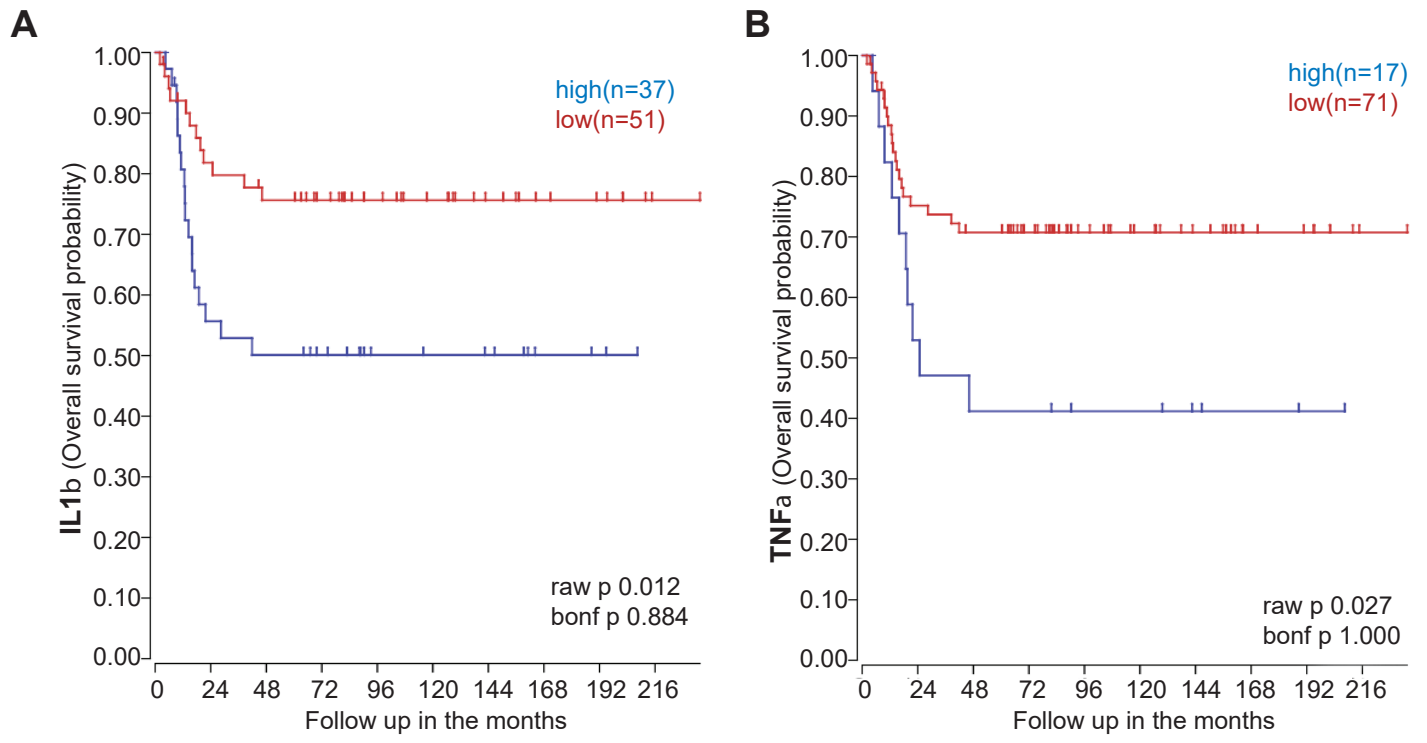












Supplementary Materials and Methods

Cell Treatment conditions

Where indicated recombinant human cytokines IL-1 β (10 ng/ml, PeproTech), and/or TNF- α (10 ng/mL, PeproTech), were added to cell cultures for up to 48 hours. For competitive inhibition of arginine 2.5 μ M L-N^G-Nitroarginine methyl ester (L-NAME, Cayman Chemicals) was added to cultures for 72 hours. ERK1/2 inhibitor PD98059 (20 μ M, CST), p38 inhibitor SB203508 (20 μ M CST), or MSK1 inhibitor SB747651A (10 μ M, Torcris) were pre-incubated with cells for 1 hour in relevant experiments. Cellular mitochondrial metabolic activity was determined using CCK-8 (Sigma) according to manufacturer's specification.

Arginase 2 knock-down

Lentiviral plasmids encoding four unique 29-mer shRNA constructs against human arginase 2, ligated upstream to a GFP reporter in the pGFP-C-shLenti vector (#TL314692, OriGene technologies) were used for the knockdown of Arg-2 gene expression in neuroblastoma cells. An additional plasmid containing a non-effective 29-mer scrambled shRNA (#TR30021, OriGene Technologies) was used as control. Recombinant lentiviral particles carrying either the scrambled shRNA or anti Arg-2 shRNA were produced by transient co-transfection with the Lenti-V-pack packaging plasmid mix (#TR30037, OriGene Technologies) into HEK293T using the MegaTran transfection protocol (#TT20003, OriGene Technologies). Particles were harvested at 48 and 72 hours post transfection, filtered through a 0.45 μ m cellulose acetate filter, and re-concentrated using the Lenti Concentrator kit (#TR30025, OriGene Technologies). Cell lines were infected with re-concentrated particles at Multiplicity of Infection (MOI) of 30 for SKNMC or 40 for KELLY. GFP-positive cells were selected using flow

assisted cell sorting and tests performed using cell populations with >95% GFP expression. Arginase2 protein expression knockdown was confirmed by immunoblotting.

Immunoblotting

Cells were harvested, washed in PBS and lysed with RIPA lysis buffer (20nM Tris-HCl pH7.5, 150nM NaCl, 2mM EDTA, 1.0% Triton X-100) containing cOmplete™ EDTA-free protease inhibitors (Roche) and PhosSTOP™ phosphatase inhibitors (Sigma). For subcellular fractionation experiments, mitochondrial and cytosolic fractions were isolated from neuroblastoma cells using the Mitochondrial Isolation Kit for Cultured Cells (Pierce) according to the manufacturer's protocol. Following a Bradford assay, equal amounts of total protein per treatment condition were electrophoresed on a 10% Tris-Glycine SDS-PAGE gels (BioRad) at 150V for 1 hour. Separated proteins were transferred to PVDF membranes using the TransBlot system (BioRad). Protein bands were detected using primary antibodies to ARG2 (Santa Cruz, sc-20151), p-AKT (CST, 9271S), AKT (CST, 2938S), p-p38 (CST, 9215S), p38 (CST, 9212S), p-ERK1/2 (CST, 9102S), ERK1/2 (CST, 9102S), MSK (CST, 3489), P-MSK1 (CST, 9595), NFkappaB (CST, 8242S), P-NFkappaB (CST, 3033S) or beta-actin (CST, 4967S). HRP-conjugated secondary antibodies of either goat anti-rabbit (CST, 7074S) or horse anti-mouse (CST, 7076S) were used for primary antibody detection. Blots were developed using ECL substrate (BioRad), exposed to CL-Xposure X-ray film (ThermoFisher Scientific) or documented using the ChemidocMP system (BioRad). Protein bands were quantified by densitometry analysis using ImageLab software (BioRad).

Immunofluorescence and confocal microscopy

Cells were seeded onto sterile poly-L-lysine coated 13mm diameter glass coverslip-inserts (VWR) at 1×10^4 per coverslip placed in a 24-well plate. Seeded cells were maintained at routine culture conditions. Live staining of mitochondria was performed by adding 100nM of MitoTracker® Red CMXRos (ThermoFisher Scientific) to culture media for 15 minutes at 37°C prior fixing of cells. Cells were washed in ice-cold PBS then fixed in 2% paraformaldehyde for 20 minutes at room temperature, followed by permeabilisation in 0.1% Triton-X for 10 minutes. Coverslips were incubated in blocking buffer consisting of 5% heat inactivated goat serum (HiNGS) in 1x PBS for 1 hour. Cells were then incubated overnight with a mouse monoclonal anti-human arginase 2 antibody (Clone OTI 3G5, OriGene) diluted 1:100 in blocking buffer. The coverslips were washed twice in PBS containing 0.005% Tween-20 (PBS-T) followed by incubation with a 1:1000 dilution of Alexa Fluor-488 conjugated anti-mouse antibody (ThermoFisher scientific) for 2 hours. After three washes with PBS-T, coverslips were air dried then mounted in SlowFade gold anti-fade mountant with DAPI (ThermoFisher Scientific) to stain the cell nuclei. Cells were examined by fluorescence microscopy using a Zeiss LSM780 fluorescence confocal microscope and co-localisation analysis performed on acquired images using ZEN software suite (Carl Zeiss Microscopy).

Immunohistochemistry and scoring

Paraffin-embedded neuroblastoma sections and tissue micro-arrays (TMA) of 27 human neuroblastomas and 5 human peripheral nerve control tissue (all cores in duplicate, US Biomax) were deparaffinised and rehydrated following quality control to confirm diagnosis

and antigen preservation. Tumours were stained on a Ventana Discovery Ultra automated system, according to manufacturer's protocol. Heat-induced antigen retrieval was performed with cell conditioning 1 buffer (CC1), pH 8.5 (Ventana). Protein blocking was applied with Background Sniper (Biocare Medical, Concord, CA).

Staining with anti-human TNF- α 1:150, Abcam) and anti-human IL-1 β (1:200, Abcam) were performed at 37°C, followed by the addition of secondary antibodies (Discovery anti-Rabbit HQ and anti-HQ HRP). DAB chromogen was applied (Discovery ChromoMap DAB, Ventana) and slides were counterstained with haematoxylin II (Ventana). Where indicated, detection was performed with anti-human CD68 (1:300, DAKO), anti-human CD33 (Abcam) or anti-human CD14 (1:400, Abcam) using the Novolink Polymer Detection System (RE7280-K, Leica). Primary antibody incubations were carried out overnight at 4°C and tissue sections were counterstained with haematoxylin and mounted in DPX (VWR). To assess nonspecific staining, samples were similarly treated but the primary antibodies omitted and replaced with isotype specific rabbit or mouse IgG (Vector Labs Peterborough UK).

Antigen expression in immunohistochemistry sections were assigned as described by Nenutil et al. 2005. Briefly, to evaluate the immunostaining intensity each slide was examined on an Olympus BX51 microscope. Representative 400x magnification fields of at least 100 tumor cells were selected and photographed with an Olympus DP70 camera and accompanying image software. Fields were assigned an antigen staining intensity score of 0 = negative, 1= weak, 2 = moderate, 3 = strong. The product of the percent positive cells and staining intensity was then derived to create a histoscore of 0-300 for each high power field. A final histoscore was then given to each specimen for each antigen.

Enzyme-linked Immunosorbent Assays (ELISA)

The concentrations of cytokines IFN- γ , IL-1 β , TNF- α , TGF- β , IL-6, IL-4, IL-13 and GM-CSF in plasma and cell culture media were measured by sandwich-ELISA kit according to specific manufacturer's instructions.

Flow cytometry

Neuroblastoma cells and myeloid cells were isolated as described above. Cell surface staining was performed with anti-human CD14, CD15, CD206, CD11b, CD68, PE anti-human IL1R1 (eBioscience) and PE anti-human TNFRSF1A(also known as TNFR1) (LSBio) antibodies. For intracellular staining, myeloid cells were stained with surface markers for 20 min at 4°C followed by fixation and permeabilisation using the Fixation/Permeabilisation Concentrate and Diluent Kit (eBioscience). Cells were washed twice then stained with PE conjugated anti-human IL-1 β and APC anti-human TNF α (eBiosciences) according to manufacturer's instructions. Cell staining was assayed on a Cyan-ADP flow cytometer (Beckman-Coulter) and analysed using FlowJo software (TreeStar Inc.). To determine cell viability after

treatment cells were washed in ice-cold PBS, stained with propidium iodide (PI) and assayed on a BD Accuri C6 flow cytometer.

Monocyte Polarisation assays

Monocytes were cultured in the presence or absence of recombinant human cytokines, neuroblastoma or neuroblastoma culture supernatants (50% of final volume), with or without 0.5mM N_G-hydroxy-L-arginine (NOHA, Cayman Chemicals), and in arginine complete or arginine-free conditions for 48 hours. Where indicated 5 µM of an AKT inhibitor (MK2206, Cayman Chemicals) was added. The culture supernatants were harvested and analysed for cytokine content.

Supplementary Figure Legends

Figure S1: M1-Macrophage infiltrate neuroblastoma

A) Immunohistochemical staining of tissue microarrays of neuroblastoma and healthy peripheral nerve tissue showing background staining (Mouse IgG), CD14+ and CD15+ staining. Tissue microarray sample key for all TMAs used in the manuscript, lower right panel B) CD14+ monocytes from healthy donors co-cultured with sorted neuroblastoma tumour cells from patients upregulate CD68 expression (upper). Minimal CD206 upregulation was seen. (n=3) C) CD14+ monocytes from healthy donors co-cultured with neuroblastoma cell lines upregulate CD68 expression (upper). Minimal CD206 upregulation was seen. GM-CSF treatment is used as a positive control for CD206 staining D) Immunohistochemical staining of tissue microarrays of neuroblastoma and healthy peripheral nerve tissue showing CD68+ staining and sample key.

Figure S2: Cytokine profile of tumour induced macrophages or granulocytes

A) Concentrations of cytokines in the supernatants of healthy donor monocytes cocultured with neuroblastoma cell lines measured by ELISA after 72hours culture B) Concentrations of cytokines in the supernatants neuroblastoma cell lines measured by ELISA after 72hours culture C) Concentrations of cytokines in the supernatants of healthy donor granulocytes cocultured with neuroblastoma cell lines measured by ELISA after 72hours culture

Figure S3: Neuroblastoma conditioning upregulates IL-1 β and TNF- α expression in macrophages

A) Co-culture of monocytes from healthy donors with tumour cell lines for 24hours leads to upregulation of IL-1 β and TNF- α expression, compared to those cultured in RPMI10% media. Flow cytometry staining shown, gated on CD14+ cells. Representative staining from 3 independent experiments shown in A. (B) Percentages of cytokine positive cells shown for 3 further healthy donors, after 48hour co-culture with neuroblastoma cell lines C) Confocal microscopy of CD14+ cells sorted from a neuroblastoma patient, showing expression of TNF- α (upper panels) and IL-1 β (middle panel). Background antibody staining control (lower panel).

Figure S4: Macrophage infiltration and downstream effects on neuroblastoma tumours

A) Immunohistochemistry staining of Tissue Microarrays of neuroblastoma tumours (n=27) and healthy control tissue, for CD33 alone, and for CD33 in combination with IL-1 β and TNF- α . B) Co-culture of healthy donor derived monocytes with neuroblastoma cell lines does not alter STAT3, NF- κ B, or PI-3K activation. Western blot shown with actin as a loading control. C) Densitometry of ARG2 expression in SKNMC cells before and after shRNA knock-out of ARG2 D) CAT1 expression in neuroblastoma cell lines and untransformed, embryonic neural crest (R1113T) or dorsal root and/or sympathetic ganglion-derived stem cells (SZ16). Actin shown as a loading control E) Neuroblastoma cell lines were treated with 60-4800ng/ml BCT-100 for 72hours. Cell proliferation after 72hours was assessed by 3H-thymidine incorporation

Figure S5: IL-1 β and TNF- α drive neuroblastoma cell proliferation

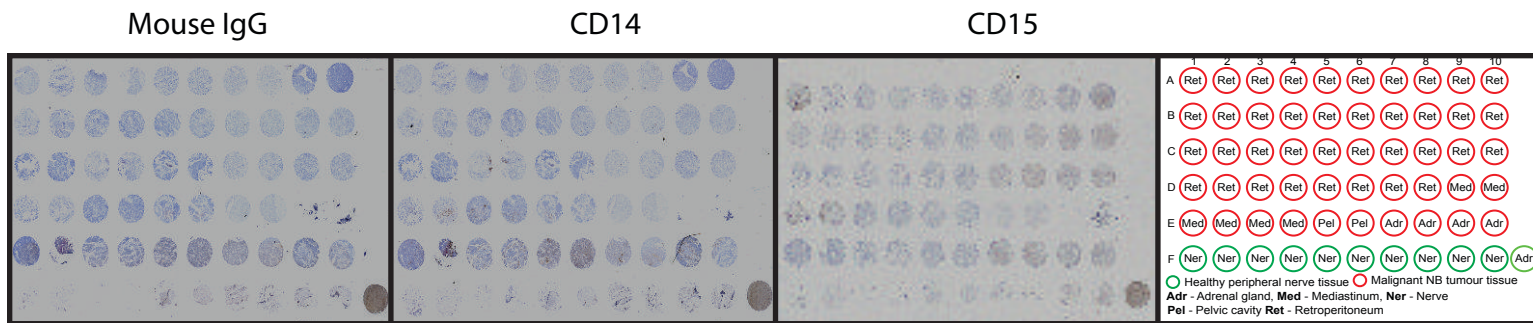
A) Neuroblastoma cell lines were treated with 600ng/ml BCT-100 or PEG alone for 72hours. Whole cell lysates were tested for PARP cleavage, which is induced by BCT-100. Actin shown as a loading control. B) ARG2 expression in GD2+ sorted neuroblastoma cells from Th-MYCN mice. Actin shown as a loading control C) ARG2, ASS, and OTC expression in GD2+ sorted neuroblastoma cells from Th-MYCN mice after BCT-100 treatment. Actin shown as a loading control D) Quantification of the upregulation of ARG2 by tumour-induced macrophage supernatants measured by confocal microscopy E) Neuroblastoma cell proliferation is enhanced in the presence of neuroblastoma-induced macrophage conditioned supernatants (MCM) or recombinant TNF α and IL-1 β . The addition of anti-TNF α and IL-1 β antibodies (inhibitors) reversed the proliferative effects of MCM. Cell proliferation of neuroblastoma shown by dilution of Cell Trace reagent, measured by flow cytometry for n=3 donors F) PD90859 (PD) or SB203508 (SB) inhibitors used as single agents do not block the combined effects of TNF α and IL-1 β on ARG2. Western blot shown with actin as a loading control.

Supplementary Table

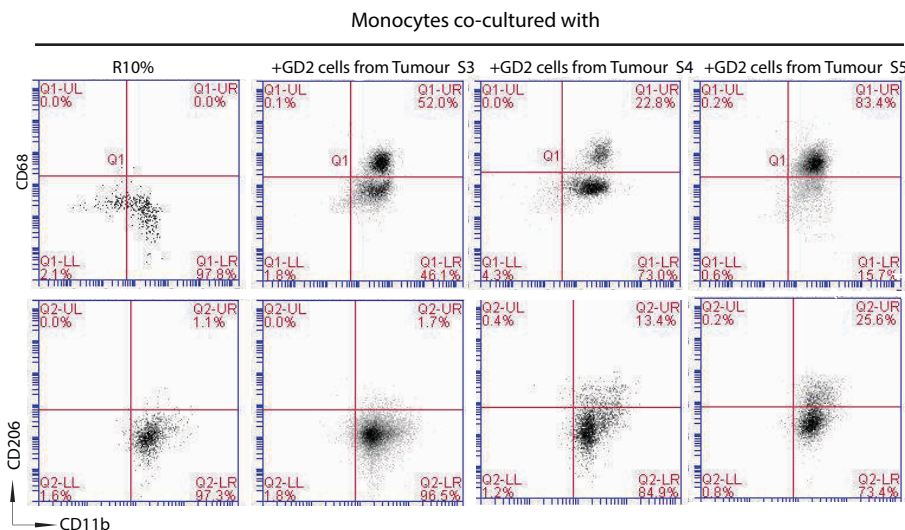
TABLE S1: Oligonucleotide primers and probes used in this study

Oligonucleotide & Probes	Source
MYCN transgene primer, Forward: CGACCACAAGGCCCTCAGTA	Hadjidaniel and Reynolds (2010)
MYCN transgene primer, Reverse: - CAGCCTTGGTGTGGAGGAG	Hadjidaniel and Reynolds (2010)
MYCN transgene probe: 6FAM TM - CGCTTCTCCACAGTGACCACGTCG-TAMRA TM	Hadjidaniel and Reynolds (2010)
Murine: Chromosome 18 insertion site Forward: CCACAAAATATGACTTCCTAAAAGATTT	This Paper
Murine: Chromosome 18 insertion site Reverse: CATGGGACTTCCTCCTTATATGCT	This Paper
Murine: Chromosome 18 insertion site probe: VIC TM -5'- AACAATTATAACACCATTAGATATG-TAMRA TM	This Paper
Human: Arginase 2 primer, Forward: ATGTCCCTAAGGGGCAGCCTCTCGCGT	Mussai et al. (2015b)
Human: Arginase 2 primer, Reverse: ATGTCCCTAAGGGGCAGCCTCTCGCGT	Mussai et al. (2015b)
Human: GAPDH primer, Forward: CCAGCCGAGCCACATCGCTC	Mussai et al. (2015b)
Human: GAPDH primer, Reverse: ATGAGCCCCAGCCTTCTC	Mussai et al. (2015b)
Human: PPIA primer, Forward: GGACCCAACACAAATGGTTCC	This Paper
Human: PPIA primer, Reverse: CTTTCACTTTGCCAAACACCA	This Paper
Human: PPIA probe: FAM TM -5'- ATGCTTGCCATCCAACCACTCAGTCTTG-TAMRA TM	This Paper

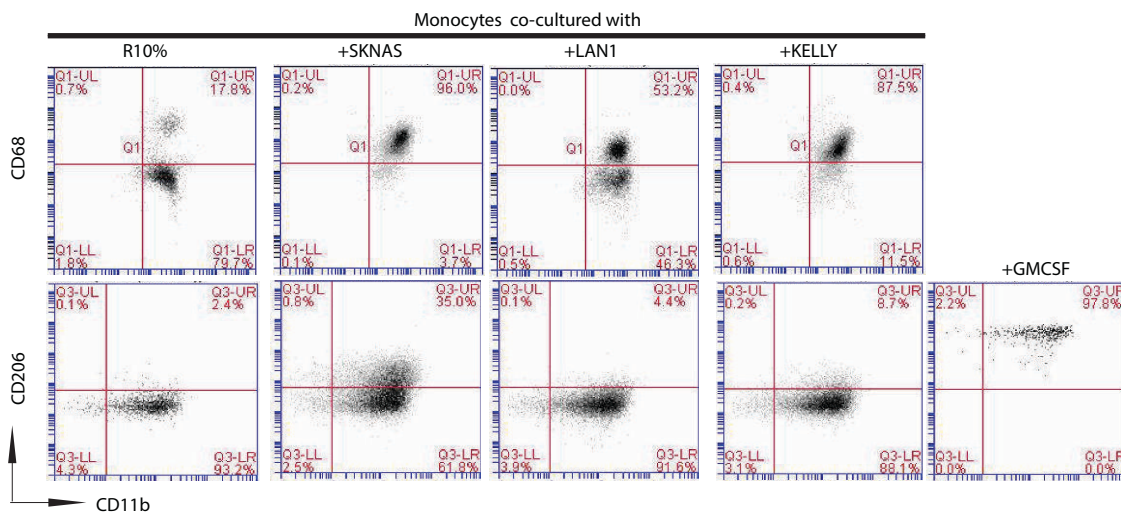
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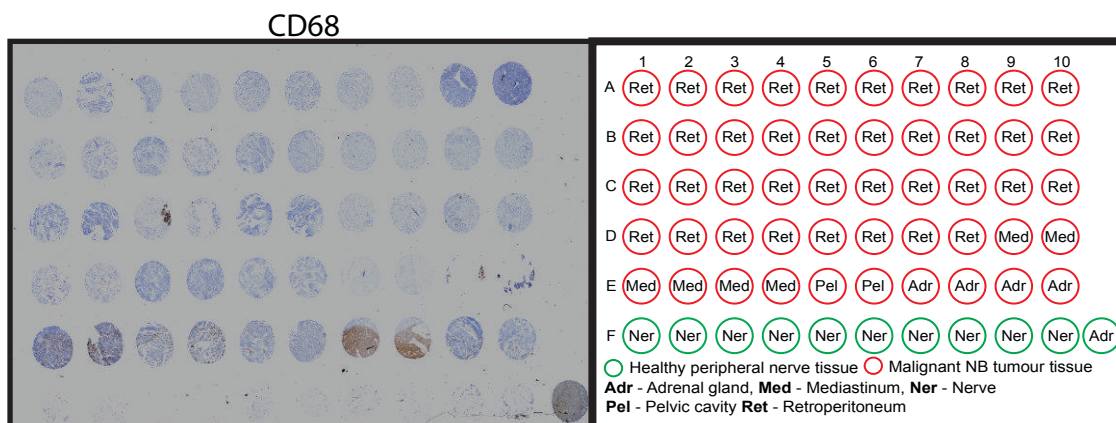
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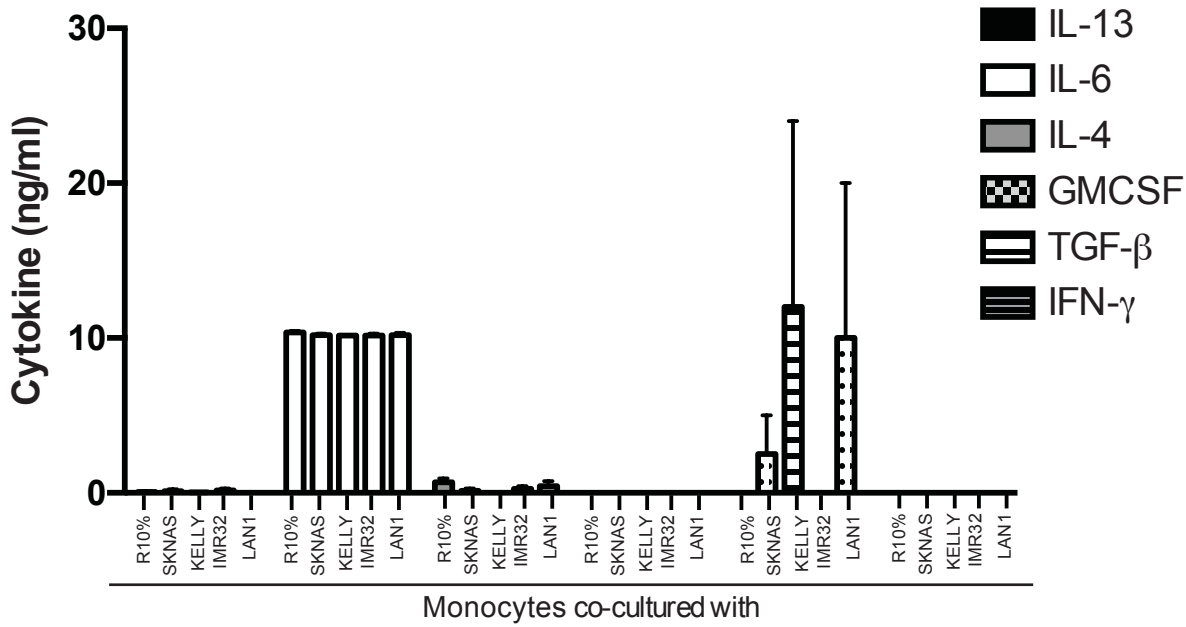
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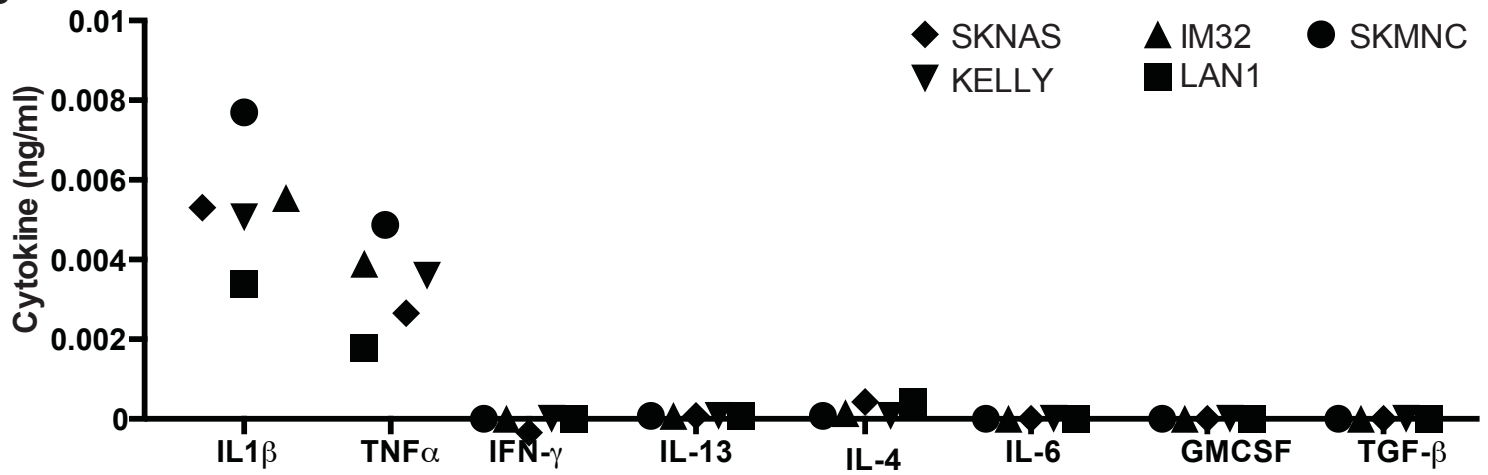
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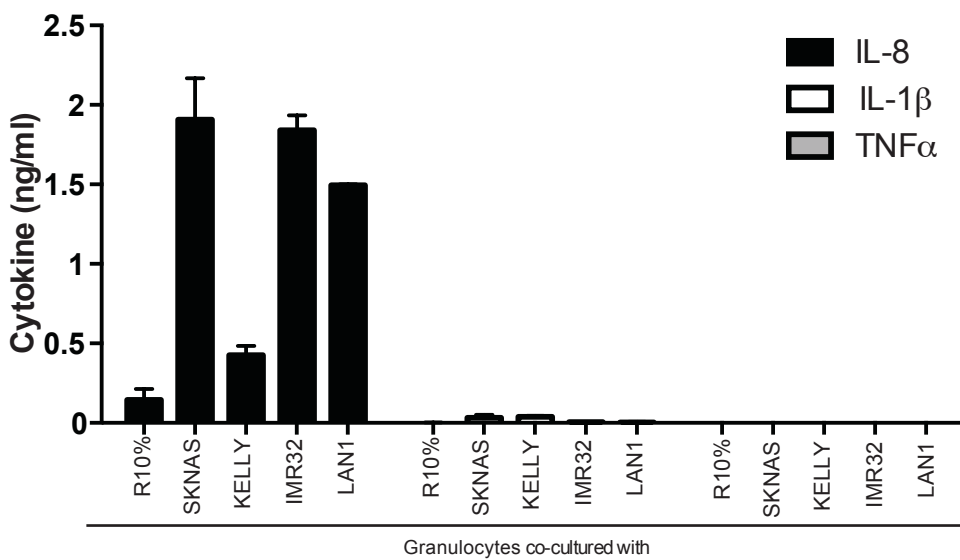
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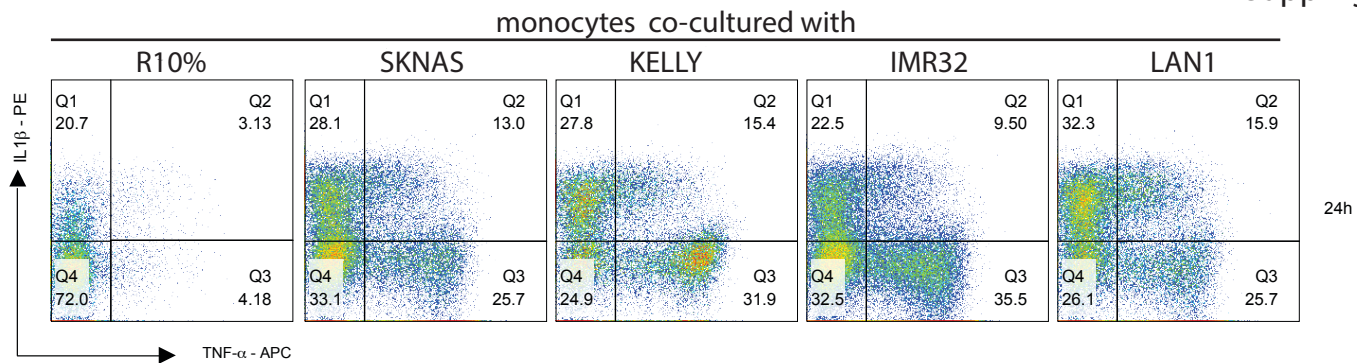
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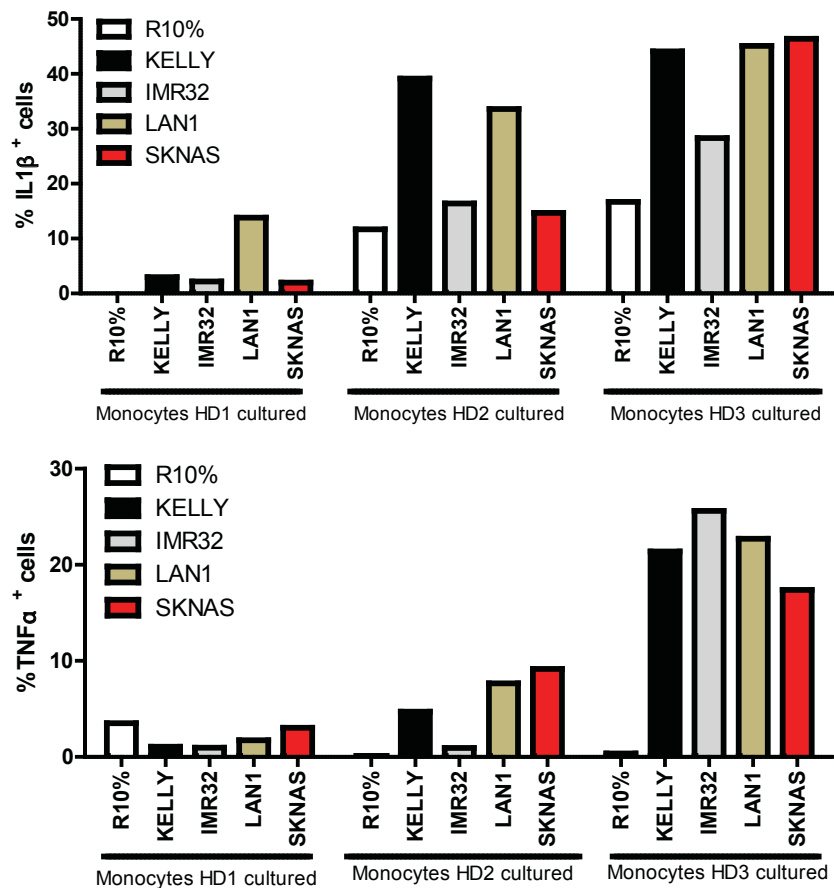
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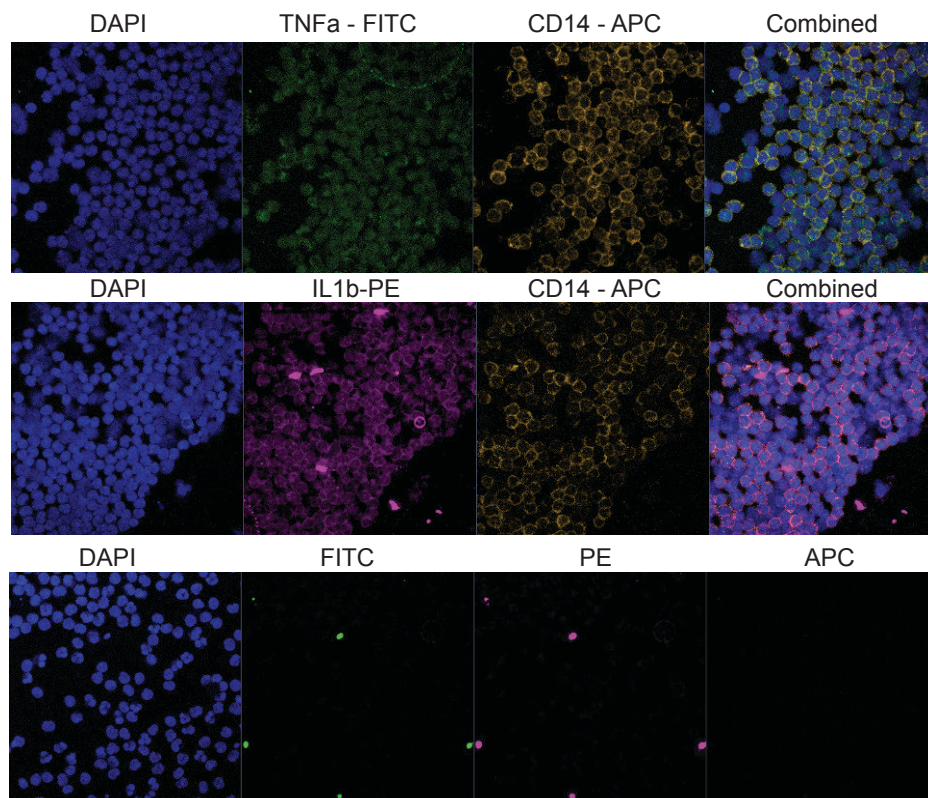
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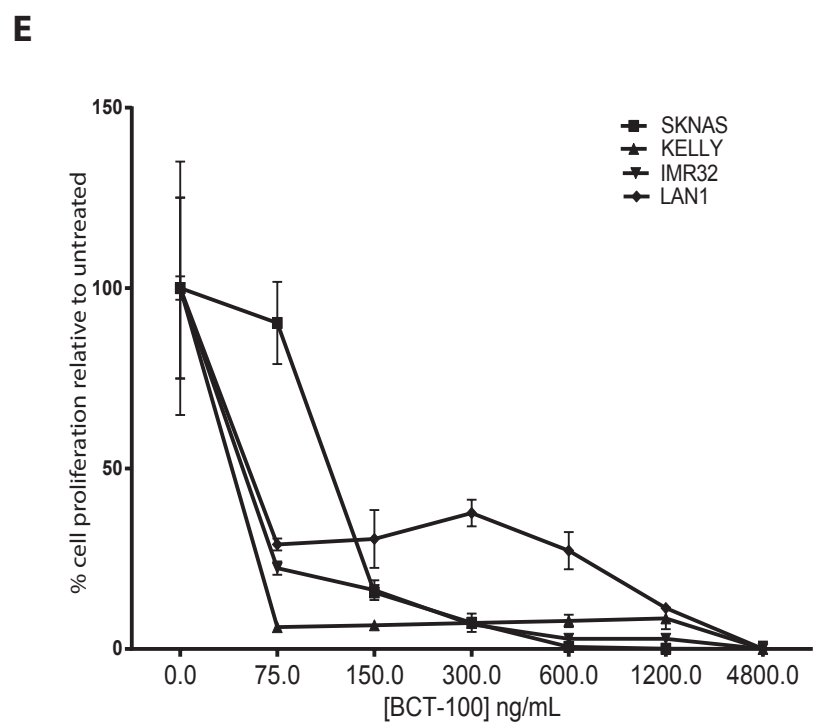
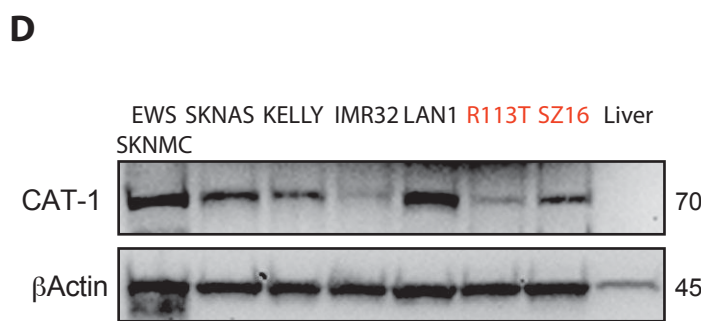
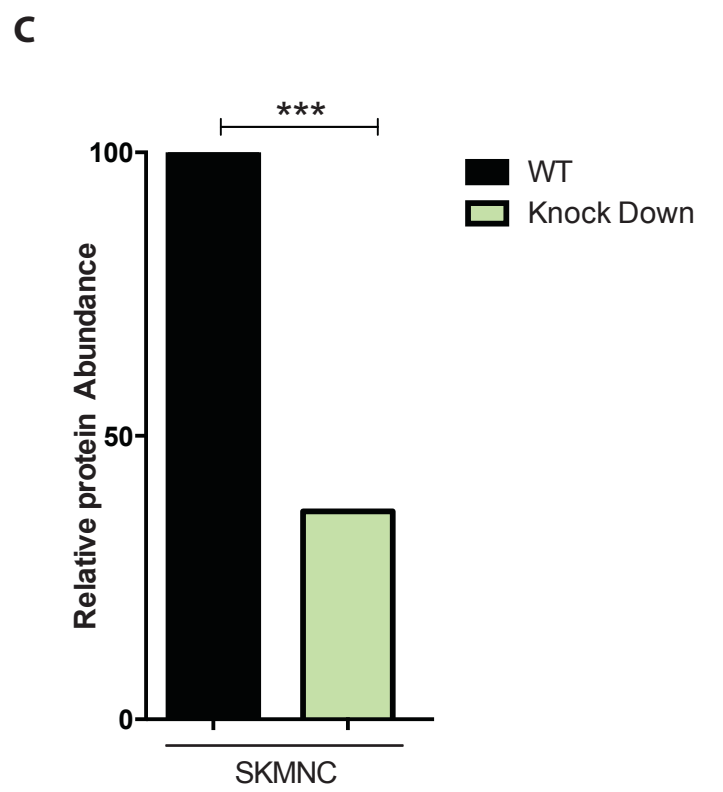
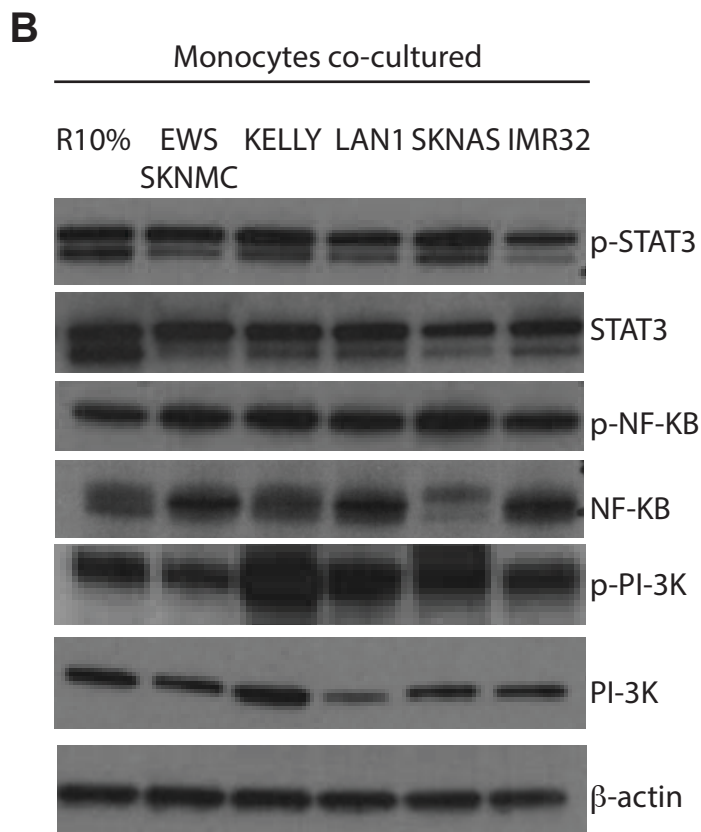
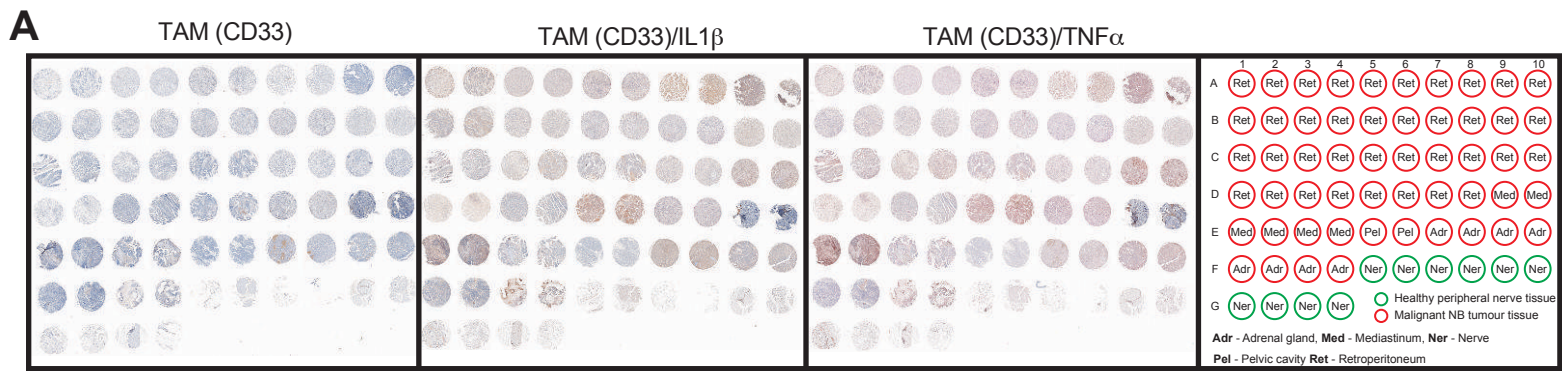


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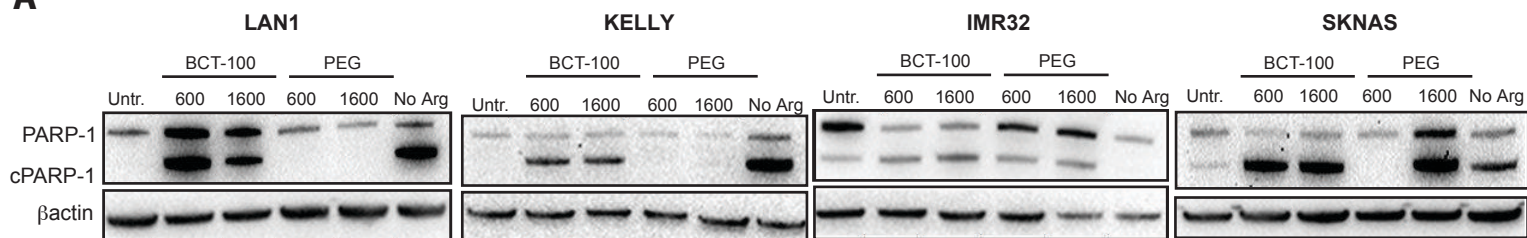


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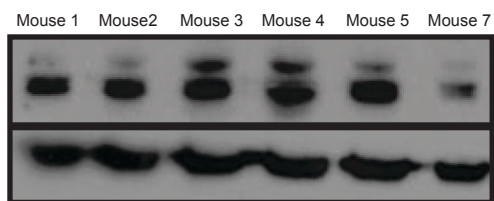




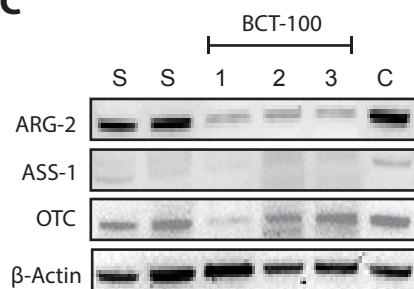
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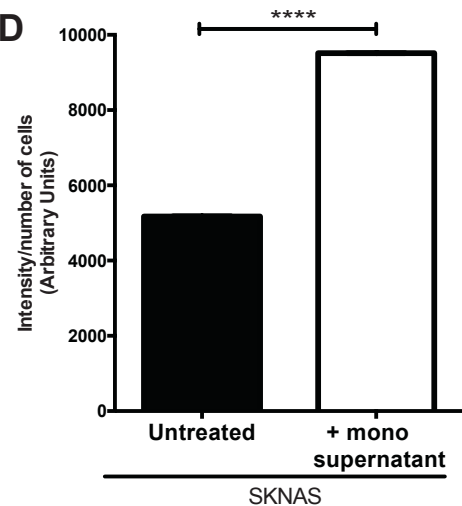
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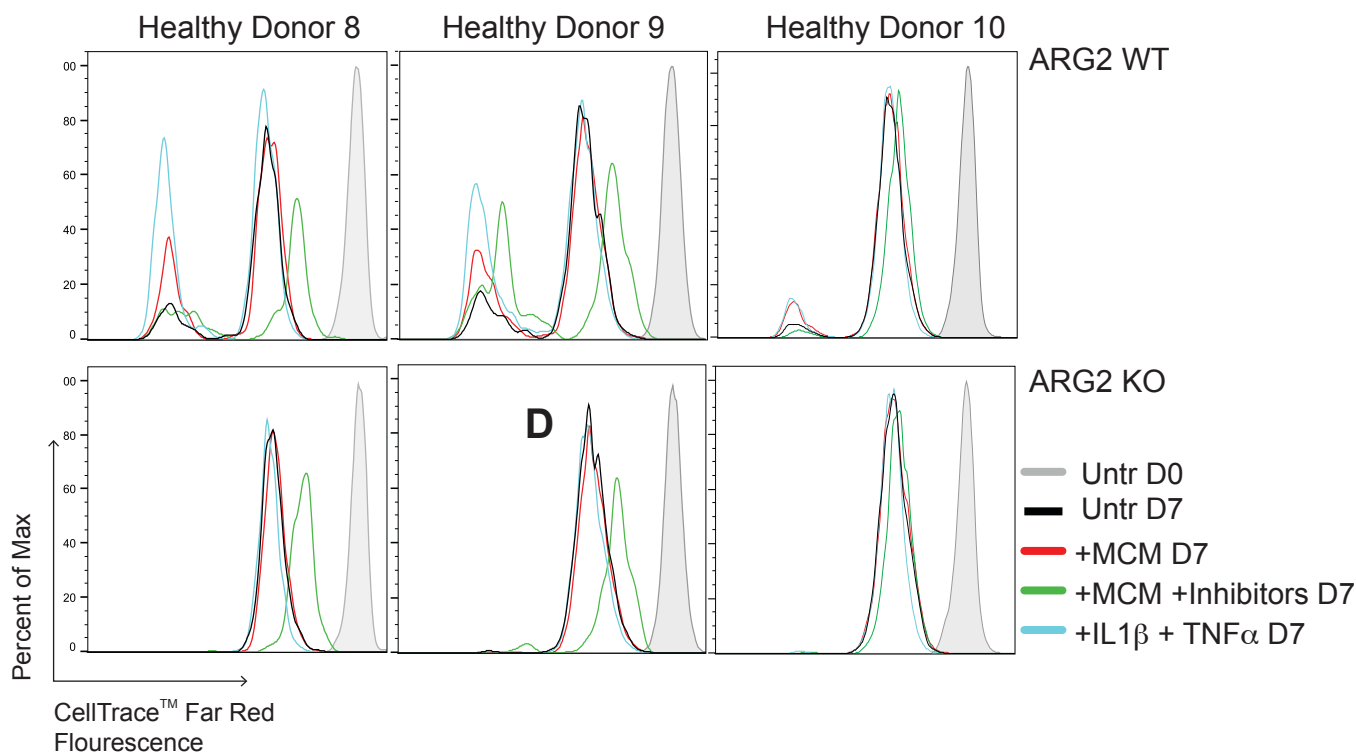
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