A novel role for an old target: CD45 for breast cancer immunotherapy

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ABSTRACT

Breast cancer subtypes have not shown significant response to current immunomodulatory therapies. Although most subtypes are treatable, triple negative breast cancer (TNBC), an aggressive highly metastatic cancer, comprising 10–20% of breast cancers, remains an unmet medical need. New strategies are needed in order to overcome flaws in the responsiveness to current TNBC therapies. Our aims were: first, to determine the efficacy of a novel immunomodulatory peptide, C24D, on TNBC and second, to elucidate the molecular mechanism by which C24D induces immune-modulating tumor killing. Using mass spectrometry analysis, we identified CD45 as the C24D binding receptor. In vitro and in vivo TNBC models were used to assess the efficacy of C24D in reversing TNBC-induced immunosuppression and in triggering immune-modulated tumor cell killing. The CD45 signal transduction pathway was evaluated by western blot and FACS analyses. We revealed that addition of PBMCs from healthy female donors to TNBC cells results in a cascade of suppressive CD45 intracellular signals. On binding to CD45’s extra-cellular domain on TNBC-suppressed leukocytes, the C24D peptide re-activates the Src family of tyrosine kinases, resulting in specific tumor immune response. In vitro, immune reactivation by C24D results in an increase of CD69+ and CD69+ NK cells, triggering specific killing of TNBC cells. In vivo, C24D induced CD8+ and activated CD56+ tumor infiltrated cells, resulting in tumor apoptosis. Our results should renew interest in molecules targeting CD45, such as the C24D peptide, as a novel strategy for TNBC immunotherapy.

Introduction

Overcoming tumor-induced immune suppression, immunotherapy has revolutionized the treatment of cancer, offering a more effective alternative for multiple types of tumors. Tumor escape mechanisms involve loss of antigenicity, immunosuppression and/or reduction of immunogenicity. Malignant cell escape mechanisms include MHC downregulation or upregulation of immunosuppressive checkpoint pathways, affecting the anti-tumor response of T cells. Monoclonal antibodies directed against these immune checkpoint axes, e.g., the programmed cell death protein 1 or programmed cell death ligand 1 (PD1/ PD-L1), in or without combination with cytotoxic T-lymphocyte-associated antigen 4 (CTLA-4), have been successfully used for the treatment of some cancers. Biologically, triple negative breast cancer (TNBC) is highly heterogeneous. TNBC has more tumor infiltrated lymphocytes, higher levels of PD-L1 expression on both tumor and immune cells and a greater number of nonsynonymous mutations. These characteristics often correlate with better responses to check point inhibitors and to tumor-specific neoantigens. However, multiple clinical trials have resulted in disappointing response rates and single-agent efficacy is still low.

Between 10% and 20% of breast cancers are TNBCs. Compared to other breast cancer sub-types, TNBC is more metastatic and has a higher rate of recurrence. TNBC tumors are negative for progesterone, estrogen and Her2, thereby explaining the lack of targeted therapies. Highly toxic chemotherapies are the most common clinical option. Ongoing efforts in translational research are needed to broaden current clinical applications and overcome unsolved issues.

In this study, we present the novel molecular mechanism of C24D, an immune modulating therapeutic peptide derived from the placentan immunomodulatory ferritin (PLIF), its effect on TNBC-dampened immune cells and the resulting immune-modulated TNBC cell killing. The C24D sequence consists of the last 24 amino acids of PLIF and is synthesized as a homodimeric peptide with a disulfide bond. We revealed that C24D binds to the CD45 receptor on T and NK cells. CD45 is a transmembrane protein tyrosine phosphatase receptor type C (PTPRC), with double opposing effects on T cell receptor (TCR) activity. On the one hand, CD45 plays an inhibitory function involving the dephosphorylation of the tyrosine 394 (Y394) in the lymphocyte-specific protein tyrosine kinase (Lck), preventing its activation. On the other hand, CD45 plays the role of activator when it dephosphorylates the tyrosine 505 (Y505), an inhibitory site at the C-terminal end of the non-receptor tyrosine-Src kinases. Activated Lck phosphorylates the immunoreceptor tyrosine-based activation motifs (ITAMs) of the T cell receptor (TCR)/CD3 complex. The phosphorylated ITAMs recruit the Zeta-chain-associated protein kinase
70 (ZAP-70), via its Src homology 2 (SH2) domains. Finally, for TCR activation, CD3-bound ZAP-70 is activated by both Lck and (trans-)autophosphorylation at the ZAP-70 tyrosine 493 (Y493).\textsuperscript{19} Downstream of protein tyrosine kinases is located the proto-oncogene VAV. Ligation of the activating receptor NKG2D on human NK cells results in phosphorylation of the signaling chain DAP10 by Src family kinases and the recruitment of VAV-1. VAV-1 is then phosphorylated by Src family kinases, as required for NK cell activation.\textsuperscript{20} 

In this study, we found that the Src family of tyrosine kinases in PBMCs of TNBC patients is inhibited. We also demonstrated that by binding to the CD45 receptor on the TNBC-suppressed leukocytes, C24D breaks the inhibition and triggers an intracellular cascade of signals which results in immune-cell activation and specific killing of TNBC cells.

**Material and methods**

**Human cell lines and culture conditions**

MDA-MB-231 and MDA-MB-468 (obtained from ATCC, Biological Industries, Kibbutz Beit Ha Emek, Israel) is a human triple-negative breast cancer cell line that is Her2-neu, estrogen receptor (ER)- and progesterone receptor (PR)-negative. MCF-10A (obtained from ATCC) are normal breast cells. Cells were grown in DMEM supplemented with 10% FCS, 1-glutamine (2 mM), Na-pyruvate (1 mM), penicillin (100 µg/ml), streptomycin sulfate (0.1 mg/ml) and nystatine (12.5 µg/ml) complete culture medium (Biological Industries). Cultures were maintained at 37°C in a humidified 5% CO₂ incubator.

A large stock of cells was prepared to maintain homogeneity and tumorigenicity of the cell line. Cells were not used beyond passage five and examined for mycoplasma (Mycoplasma detection kit, Biological Industries) at least once every six months.

C24D peptide synthesis: The C24D peptide is a 25 amino acid homodimer peptide with a di-sulphide bond at CG (CG-HHLLRPRRRRRKPSITFIPILRSP). An irrelevant scrambled peptide (CG-PPIPPLKRIRLTRPLHRPHHS) (S-C24D) was used as control. Both peptides were synthesized by Synpeptide Co., Ltd (Shanghai, China). The C24D peptide and the control peptide were also synthesized with the addition of Biotin or FITC C-terminal Ahx, 6-aminohexanoic acid residue to the peptide sequence. HPLC showed purity >97%.

**PBMC-tumor cell co-cultures and efficacy of peptide**

**PBMC isolation:**

PBMCs were isolated from blood of healthy female donors or breast cancer patients, obtained from the Blood Bank Mada Tel HaShomer or Davidoff Cancer Center, respectively. The protocol was approved by the Institutional Review Board at Rabin Medical Center, Israel (0667–14-RMC). Samples were isolated by Ficoll–Hypaque density gradient (d = 1.077 g/ml, Ficoll-Paque Plus, GE Healthcare, Upsalla, Sweden) by centrifugation at 650 × g for 30 minutes.

**Tumor cells:**

MDA-MB-231 and MDAM-MB-468 at 4x10⁵/2 ml, derived from an exponentially growing monolayer were incubated in complete medium overnight in 6 well plates. PBMC in complete RPMI medium, supplemented with 5% AB serum instead of FBS (Biological Industries), were added onto the tumor cells (2x10⁴/ml). The C24D peptide was added immediately at 0.1, 10, 30 and 60 µg/ml and incubated for various times at 37°C, 5% CO₂. S-C24D was used as negative control. A dose response analysis revealed that 10 µg/ml was the most effective dose for *in vitro* experiments, based on IFNγ secretion and tumor cell density (Supplementary Figure 1). MCF-10A normal breast cells were used as control, following the same protocol as described for tumor cells.

At the end of the experiments, lymphocytes were extracted from co-cultures, centrifuged and re-suspended in PBS for FACS analysis. Only samples containing more than 98% CD45+ cells after extraction were selected for the experiments.

The CD45+ cells were identified by FACS analysis with an anti-CD45-PB antibody (Supplementary Figure 2). Supernatant from co-cultures recovered from centrifugation of the lymphocytes was centrifuged at 1200 rpm for 10 minutes and stored at –80°C for cytokines determination.

**Cytokines:**

Human Interferon gamma (IFNγ), Tumor Necrosis Factor alpha (TNFa), Interleukin 2 (IL-2) and IL-1β were determined by ELISA Ready SET-Go (eBioscience, San Diego, CA, USA).

**Tumor cell density:**

For the determination of tumor cell density, co-cultures were washed once with PBS and cell density was documented by a light inverted microscope (Olympus IX73). For the determination of tumor cell killing, tumor cells were removed from plates and subjected to FACS analysis. Tumor apoptosis was determined in gated tumor cells using an AnnexinV/PI kit (MEBCYT® Apoptosis Kit by MBL, MA, USA). Previous to the addition of Annexin/PI, an anti-CD45-PB (Monoclonal Antibody IOTest Beckman Coulter, Marseille, France) was added to tubes for identification of leukocytes to be discarded in the FACS analysis.

**Tumor xenograft growth in nude mice**

The animal experimental procedures used in the present study were approved by the Animal Care and Use Committee of Tel Aviv University (TAU 06–01–20220), in accordance with their guidelines. In total, 30 BALB/c nude mice (female; 5–6 weeks of age; each weighing 20–25 g) were purchased from Envigo (Jerusalem, Israel). Tumor xenografts were generated by subcutaneously injecting 4 × 10⁶ MDA-MB-231 cells into the right nude-Balb/C mice flank. Tumor volumes were measured every other day using micrometer calipers and were calculated according to the following formula: tumor volume (mm³) = 0.5 x D x d², where d and D represent the shortest and the longest diameters, respectively.

Six days after tumor injection, when the xenograft grew to approximately 100 mm³, the mice were divided randomly into four groups: a negative control group, which was to be treated with PBS (n = 6), a second negative control group, to be treated with 60 µg/mouse S-C24D (scrambled C24D) in 200 µl PBS, and two C24D treated groups (C24D: 60 µg/mouse in 200 µl...
PBS and C24D: 300 µg/mouse in 200 µl PBS), n = 8 in each of the latter 3 groups.

Fresh human PBMCs from 2 different female donors were incubated with C24D or S-C24D or PBS (60 µg or 300/2 x 10^6 cells in 0.4 ml PBS) for 5 minutes before the first intravenous injection (IV). The PBMCs from the 2 donors were injected separately, in half of the mice in each group. The subsequent IV injections of C24D, S-C24D or PBS were administered twice a week for 2 weeks.

At the end of the experiment (13 days after the administration of the PBMCs), the mice were bled, for the determination of human IFNγ in serum and sacrificed, for the extraction of tumors. The tumors were processed for immunofluorescence.

**Immunofluorescent staining**

Tumor tissues underwent 4% formalin fixation overnight at 4°C and paraffin embedding. The specimens were then sliced into 5-µm-thick sections and deparaffinized in xylene and rehydrated in alcohol. After antigen retrieval at pH = 6, sections were incubated for 1 hour with primary antibodies: CD8 (Rabbit Monoclonal, 108 R-14, 1:50, Cell Marque, Rockling, CA, USA), CD45 (Mouse Monoclonal, 145 M-94, 1:50, Cell Marque), activated CD56 (Mouse Monoclonal, 156 M-84, 1:50, Cell Marque), Cleaved Caspase-3, (Rabbit Monoclonal, 9664S, 1:100, Cell Signaling, Danvers, MA, USA), HLA-A (Rabbit Monoclonal, Ab52922, 1:100, Abcam). Slides stained with an IgG irrelevant antibody followed by secondary antibodies served as negative controls. Human PBMCs in paraffin slides served as positive controls. The fluorescence detection was performed with Alexa Fluor 488-AffiniPure Donkey Anti-Mouse IgG (H + L) (715–545-151, 1:200); Cy3-AffiniPure Donkey Anti-Rabbit IgG (H + L) (711–165-152, 1:200); both from Jackson Immunoresearch Laboratories, Westgrove, Pennsylvania, USA). DAPI (4’,6-Diamidino-2-P henylinodole, Dilactate) (BLG-422801, 1:400, Biologend, San Diego, CA, USA) was used as counterstain. Fluorescence observation and image acquisition were performed using a Leica TCS SP5 confocal laser-scanning microscope (Leica Microsystems, Germany). Quantitative Image Analysis was preformed using FIJI (Image J2) image analysis software in 5 fields/slide in 5 different slides.

**FACS analysis of C24D binding to cells**

Fresh lymphocytes (0.5x10^6/50 µl PBS) were incubated with the following antibodies for multicolor staining: CD3-PC5.5, CD4-PC7, CD8-KO, CD56-PE, CD16-APC, NKG2D-APF750, CD45RA-APC, CD45RO-PC5.5 and CD14-KO (Beckman Coulter) and with 5 µg/50 µl C24D-FITC peptide for 40 minutes at room temperature. The cells were then washed twice (10 min., 1200 rpm, 4°C). Activated lymphocytes obtained from co-cultures with tumor cells were subjected to the same procedure. To ensure specificity, a Fluorescence Minus One (FMO) control was used to identify and gate cells in the context of data spread due to the presence of multiple fluorochromes in any given panel. In brief, for each sample, two tubes were used: one tube containing cells stained with the antibodies identifying the various PBMC subpopulations and a control peptide-FITC labeled and a second tube containing cells stained with the antibodies identifying the different PBMC subpopulations and with the C24D-FITC peptide. The percentage C24D/CD45 binding on the cell surface in each subpopulation was calculated after subtracting the fluorescence obtained for the control tube. The cells were washed with PBS and suspended in 0.5 ml PBS for FACS (Coulter Navios flow cytometer, Indianapolis, IN, USA). Data were analyzed with the Kaluza software (Beckman Coulter, IN, USA).

Anti-CD45-APC antibody was used for the double staining determination of the C24D+/CD45+ cells.

For the determination of the activated PBMC subpopulations, PBMCs from the co-cultures with tumor cells were counted and transferred to a FACS tube (0.5x10^6 cells/50 µl PBS) and incubated with the following antibodies: CD4-PC7, CD8-KO, CD56-PE, NKG2D-APF750, CD14-KO, CD69-PC5, CD57-FITC, CD137-EC (Beckman Coulter). Gated live cells were analyzed using Coulter Navios FACS, Kaluza software.

**Mass Spectrometry analysis for the identification of the specific peptide binding protein**

Fresh PBMCs (15x10^6) from human donors were centrifuged and transferred to a chilled microtub. Lysis buffer (150 mM NaCl, 1% triton X-100, 50 mM Tris HCl pH 8, with the addition of proteases inhibitors) was added to the cells for 20 minutes on ice. After centrifugation for 5 minutes at 1000 rpm, the supernatant was transferred to a new microtube for precipitation. Protein concentration was determined by BCA protein assay kit (Thermo Scientific, Rockford, IL, USA) following manufacturer instructions. Cell protein (50 µg/200 µl lysis buffer) from fresh lymphocytes was incubated with 15 µg biotinylated C24D peptide for 1 hour at 4°C at constant shaking. After incubation, the biotinylated peptide which bonded to specific proteins formed a complex. The complex was precipitated using streptavidin magnetic beads (µMACS streptavidin Kit, Miltenyi Biotec GmbH, Cat. Number 130–074-101). The complex was placed in a microcolumn in the magnetic field of a µMACS Separator. After rinsing the column, the target molecules which bonded to the biotinylated probe were eluted. The eluted sample was subjected to SDS PAGE in a 10% gel. The gels were stained with Imperial Protein Stain (Thermo Scientific, Cat. Number 24615) for mass spectrometry analysis.

Mass spectrometry analysis was done by The Smoler Proteomics Center, Technion, Haifa, Israel. Samples were reduced, modified and digested by trypsin, analyzed by LC-MS/MS on a Q exactive plus (Thermo). Data was analyzed by Discoverer Software, version 1.4, using the Sequest search engine, vs the human Uniprot database. The data was quantified using the same software. Protein identification was filtered for proteins identified with FDR <0.01 with at least 2 peptides. Proteins which were precipitated without biotinylated peptide served as controls. Proteins which had at least 4-fold more mass-to-charge ratio than the control were considered for further analysis.

Following mass spectrometry analysis, the eluted purified sample obtained from precipitation of the lysed PBMCs with biotinylated C24D was subjected to western blot analysis using and anti-CD45 antibody (Abcam, Cat. Number ab123522).
**Proximity ligation assay:**
The method was modified for Peptide–Target interaction previously described.21 In brief, a 96 well, high binding plate, was coated with His Tag antibody (Monoclonal IgG1, R&D Systems, USA) for 2 hours at room temperature. After washing with PBS-0.05% Tween, recombinant Human CD45 with a C-terminal 6 His Tag (R&D Systems, USA) was added to coated plates overnight at 4°C. Following 3 plate washes, Biotinylated C24D at various concentrations (0.05, 0.5, 1, 5 and 10 µg/ml) was added for 2 hours at room temperature. Streptavidin–HRP (R&D Systems, USA) was added to plates for 2 hours followed by TMB subtract for proximity Peptide/Target detection. HCl 1 M was used to stop the chemical reaction. Proximity ligation was detected by ELISA reader at 450 nm. A biotinylated scrambled version of C24D was used as negative control and a Biotinylated Anti CD45 antibody (Abcam) was used as positive control.

**Blocking of the CD45 receptor**
The anti-CD45 antibody 10 µg/ml (Abcam, Cat. Number ab123522) and/or the peptide CD45 inhibitor VI (Calbiochem, Burlington, Massachusetts, USA, Cat. Number 530197) were used for blocking the extracellular domain of CD45 in the PBMCs used in the relevant experiments. The blockers were added before addition of PBMCs to the tumor plates, as per manufacturer’s instructions. The efficacy of the blockers is presented in Supplementary Figure 3.

**CD45 signal transduction evaluation**
PBMC was added onto the tumor cells (2x10^6/ml) and 10 µg/ml of C24D was added immediately thereafter. Co-cultures were incubated for 5, 15, 30, 60 minutes and 24 hours at 37°C, 5% CO2, depending on each protein’s phosphorylation.

At the end of each incubation period, lymphocytes were extracted, and a sample of the extracted cells was submitted to FACS analysis to corroborate that the sample for western blot analysis contains more than 98% leukocytes (Supplementary figure 2). The extracted leukocytes were then centrifuged and re-suspended in 0.12 ml of lysis buffer (20 mM Tris-HCl pH 7.5, 150 mM NaCl, 1 mM NaF, 2 mM Na3VO4, 1% NP40, 10 mM b-glycerophosphate, 30% glycerol, 1 mM EDTA, 0.5% sodium-deoxycholate, 0.5% protease inhibitor cocktail), followed by one freeze-thaw cycle of 20 min. Cells were harvested and centrifuged (14,000 rpm, 15 min, 4°C). The supernatants were collected, and aliquots were separated on 10% SDS PAGE, followed by Western blotting with anti-phospho-Lck Y505 (0.5 µg/ml, ab4901, Abcam, Cambridge, UK), anti-phospho-Lck Y394 (0.25 µg/ml, ab201567, Abcam), anti-phospho-VAV-1 Y174 (0.23 µg/ml, ab76225, Abcam) and anti-phospho-ZAP70 Y493 (1 µg/ml, ab194800, Abcam). GAPDH (1 µg/ml, ab9485, Abcam) was added as a control for sample loading. After several washings, the secondary antibody, IRDye 800CW Goat anti-Rabbit or IRDye 680CW Goat anti-mouse (1 mg/ml, LI-COR, Nebraska, USA) was added for 1 h.

Quantification methods: The membrane was analyzed by Odyssey 2.1 (Infrared Imaging System) for specific band identification. Quantification of phosphorylation was done by Image J (NIH, USA). Percentage (%) of maximal phosphorylation of phosphorylated proteins were first normalized to the levels obtained with GAPDH respectively, and the activation values were normalized for each time point vs its control, incubated with a control peptide (e.g. C24D + lymphocytes vs. lymphocytes control). The values obtained were then expressed as % of maximal activation that was observed in each experiment, at each time point.

For the corroboration of the results obtained in the western blot analysis, FACS analysis was performed with the same antibodies, followed by incubation with a secondary antibody (anti-rabbit-PE, Jackson ImmunoResearch, cat: 711–116–152). Intracellular staining for the phosphorylation of the tyrosine 505 and 394 in Lck, the tyrosine 493 in ZAP-70 and the tyrosine 174 in VAV-1, was performed using a FIX & PERM Cell Permeabilization kit consisting of matched Fixation and Permeabilization Reagent (R&D Systems, Minneapolis, MN, USA).

**Statistical analysis**
To determine whether the data from test samples were significantly different from control samples, student’s t-tests were performed using GraphPad Software (GraphPad Software, La Jolla, CA). ANOVA, followed by Tukey-Kramer, was used for multiple comparisons. Significance was defined as p<0.05.

**Results**

**C24D induces re-activation of leukocytes, resulting in TNBC cell killing**
The immune-modulating tumor killing effect of C24D was previously demonstrated on MCF-7 (ER+) and T47D (HLA-A2) breast cancer cells, co-cultured with human PBMCs from healthy donors.16 In this study, we first set out to determine if C24D also induces effective immune-modulated tumor killing of TNBC cells (MDA-MB-231 and MDA-MD-468).

Observation by microscope revealed that the addition of 10 µg/ml C24D to co-cultures of human PBMCs from healthy donors with MDA-MB-231 resulted in massive tumor apoptosis (Figure 1a). Similar immune-modulated killing results were achieved with a second TNBC cell line: MDA-MD-468 (Supplementary Figure 4b). Treatment of tumors with C24D or with S-C24D (scrambled C24D), in the absence of PBMCs, both resulted in no effect (supplementary Figure 4a). In co-cultures with PBMC alone, we only noted a minor effect on tumor apoptosis.

To confirm the microscope observations, we performed FACS analysis using AnnexinV/PI. With PBMCs from 14 different healthy female donors, the percentage of AnnexinV/PI positive tumor cells (i.e., MDA-MB-231 cell killing) varied from 15% to 100%, with an average of 45.9 ± 30.6% (Figure 1b). Figure 1c illustrates an example of one such FACS analysis study, showing the results of experiment number 11.

To establish which of the leukocyte sub-populations is activated by C24D and involved in the tumor cell killing, we performed a series of FACS analyses using PBMCs from 11 healthy donors.
Figure 1. C24D induced re-activation of leukocytes, resulting in killing of TNBC cells: a. Representative pictures of tumor cell density in co-cultures of MDA-MB-231 cells with PBMCs from healthy female donors, documented using inverted microscope (Bars = 200 μm). b. Tumor cell killing by FACS analysis, using AnnexinV/PI (N = 14). c. Representative FACS dot blot (experiment #11) of AnnexinV/PI gated and stained tumor (CD45 negative) cells treated with 10 μg/ml C24D for 4–6 days. d. Percentage of CD4+/CD69+ and CD4+/CD57+ cells in co-cultures of PBMC with MDA-MB-231 cells, ± C24D, by FACS analysis. e. Percentage of CD8+/CD69+, CD8+/CD56+, and CD8+/CD56+/CD137+ cells among co-cultured CD4+ and CD8+ T/NK cells and CD69+ and CD56+ cells, in the co-cultures, by FACS analysis. f. Percentage of CD14+ and CD137+ in the co-cultures, by FACS analysis. g. Percentage of CD14+ and CD137+ in the co-cultures, by FACS analysis. h. Representative FACS analysis showing double staining of CD4, CD8 or CD56 gated cells with either CD69 or CD57, comparing effect of C24D versus control. i. Effect of C24D on IFNγ secretion (pg/ml) in supernatants of PBMCs alone or PBMCs co-cultured with MDA-MB-231 (n = 8). j. Effect of C24D on secretion of IL-2 (pg/ml) (n = 7). k. Effect of C24D on secretion of TNFα (pg/ml) (n = 7). l. Effect of C24D on secretion of IL-1β (pg/ml). Analysis of results presented as Mean ± SD.

Female donors. Addition of 10 μg/ml C24D to each PBMC/MDA-MB-231 co-culture resulted in a significant increase in the percentage of CD4+/CD69+ cells (from 2.68 ± 1.45% to 4.1 ± 1.58%, p < 0.046, Figure 1d), CD8+/CD69+ cells (from 22.6 ± 14.8% to 33.23%, p < 0.05, Figure 1e) and CD56+/CD69+ (from 19.9 ± 13.61% to 39.1 ± 16.0%, p < 0.033, Figure 1f). CD57 expression was not influenced by the addition of C24D to the co-cultures nor was the percentage of NKG2D+ and CD45RO+ cells. A representative FACS analysis of the activated cells sub-populations is shown in Figure 1h.

In contrast to T and NK activation resulting from addition of C24D, the percentage of activated monocytes (CD14+/CD137+) increased, but non-significantly, from 31.6 ± 9.4% to 45.2 ± 5.3%, p < 0.19 (Figure 1g). CD14 is a marker for circulating monocytes and CD137 (a member of the tumor necrosis factor receptor family) is expressed on
activated monocytes. Addition of C24D to the co-cultures had a non-significant effect on the secretion of IL-1β by monocytes (Figure 1i).

Addition of C24D to the MDA-MB-231/PBMC co-cultures for 4–6 days resulted in a significant increase in secretion of cytokines: 90% of IFN-γ (Figure 1i, p < 0.03), 92.2% of IL-2 (Figure 1j, p < 0.0007) and 98% of TNF-α (Figure 1k, p < 0.04). Similar results were observed in co-cultures of PBMCs with MDA-MB-468 (Supplementary Figure 5). No effect was noted when C24D was added to PBMCs alone (Figure 1(i, j, k, l)).

**Treatment with C24D inhibited tumor growth in vivo**

Using a human TNBC mouse model adapted from Zhang et al. (Figure 2a), we aimed to determine if the *in vitro* cytotoxic effect mediated by C24D would also be observed in a meaningful *in vivo* model. For this purpose, we measured tumor growth over time, IFN-γ in serum and tumor infiltrating human T and activated NK cells in tumor histological sections.

Results showed a significant reduction in tumor growth in mice treated with 60 or 300 µg C24D per mouse, compared to treatment with PBS or S-C24D as controls (p < 0.05, Figure 2b). In Figure 2c, we show the tumors extracted from 5 mice in each of the 4 groups, 13 days after treatment. At the end of the experiment, mice were bled for the determination of human IFN-γ in serum. A significant increase of 68% and 82% (p < 0.05) in human IFN-γ secretion was observed in the 60 and 300 µg groups, versus the group with S-C24D treatment (Figure 2d).

**Immunofluorescent analysis of tumor sections**

The presence of tumor infiltrating and human T and activated NK cells was revealed by an immunofluorescent analysis of tumor sections. A significant number of tumor infiltrating human leukocytes were found in the samples from mice treated with C24D, compared to the S-C24D group (Figure 2e, f, g). Compared to treatment with S-C24D, treatment with both 60 and 300 µg doses of C24D significantly induced the infiltration of human CD45+ cells (Figure 2e, green) in the human tumor (red) (p < 0.015 and p < 0.023 respectively).

The following tumor infiltrating human CD45+ sub-populations in mice treated with C24D were found in the immunohistology samples: activated CD56+ (stained with anti-human activated CD56 antibody in green, Figure 2f) and CD8+ cells (stained with an anti-human CD8 antibody in red, Figure 2g). Treatment with 60 or 300 µg C24D per mouse resulted in a significant increase in tumor-infiltrating NK (CD56+) cells (p < 0.011 and p < 0.042, respectively).

A similar pattern was observed for activated human CD8 T cells (p < 0.07 and p < 0.1, respectively). The non-significant results of human CD8 T cells might be explained by the low number of cells counted.

The presence of infiltrating human T and activated human NK cells supports the significant increase in tumor cell apoptosis (Figure 2h). The lower dose (60 µg) induced a more significant tumor cell apoptosis (p < 0.0013) than the higher dose (p < 0.049).

**Stability of C24D**

We performed two C24D stability studies: one in whole blood (ChemPartner Co, Ltd.) and a second, in plasma (in our laboratory). 10 µM (60 µg/ml) of C24D has a T1/2 of 165.03 minutes in human whole blood (Supplementary Figure 3c). In plasma, target saturation was evaluated by measuring C24D binding to the CD45 receptor, using two different methods: FACS analysis and ELISA. By FACS analysis, we observed that 1.5 µM (10 µg/ml) of C24D bound over 70% of the receptors of the studied PBMC sub-populations (Figure 3a). In the ELISA assay, 3 hours after adding 1.5 µM of C24D in the presence of human plasma, binding to CD45 decreased by 20%.

**CD45 revealed as the C24D binding receptor**

In an effort to elucidate the molecular mechanism which may explain the above immune-modulated tumor killing, we first proceeded to identify the receptor to which C24D binds. Using PBMCs from healthy female donors and C24D conjugated to Alexa-488 dye, by FACS analysis, we determined the leukocyte sub-populations to which C24D binds (Figure 3a). Surprisingly, we observed that all 9 of the tested T, NK and monocyte sub-populations bound C24D (Figure 3b; a representative dot blot FACS analysis).

Subsequently, we subjected the entire leukocyte population to mass spectrometry analysis. Using PBMCs from 8 different female donors, of which 2 were from breast cancer patients, we precipitated, per donor, the entire leukocyte population with biotinylated C24D. Each isolated complex was subjected to mass spectrometry analysis. The receptor we identified was: CD45 (PTPRC, X6R433 Uniprot), (Figure 3c). The ubiquitous nature of CD45 explained why all the 9 tested T, NK and monocyte sub-populations bound C24D.

To validate the mass spectrometry results, we performed a double-staining FACS analysis with an anti CD45 monoclonal antibody and the C24D-Alexa-488 conjugated peptide (Figure 3d). We observed that 100% of the PBMCs were stained by both antibody and C24D, thereby confirming that CD45 is the binding receptor. This result also suggests that the anti-CD45 antibody and C24D bind to different epitopes of the extra cellular domain of CD45.

Further validation of the mass spectrometry results were obtained by western blot analysis. The protein precipitated with biotinylated C24D, followed by streptavidin magnetic beads and eluted through a magnetic column, was subjected to gel electrophoresis. The membrane was stained with an anti CD45 antibody. Results revealed that the precipitated protein was CD45 (Figure 3e).

To demonstrate direct interaction between C24D and CD45 we performed a Proximity Ligation Assay using a biotinylated C24D peptide. Results showed a specific concentration dependent binding curve of C24D to an immobilized CD45 recombinant protein (Figure 3f).

To support the above results, we blocked the CD45 receptor on PBMCs with a monoclonal antibody and a commercially available blocking peptide and evaluated IFN-γ secretion.
Without blocking, a 2.4-fold increase in IFN-γ was found in the supernatants of C24D treated co-cultures of PBMCs with MDA-MB-231 cells, compared to S-C24D treated cells. In contrast, the results obtained on co-cultures on which the CD45 receptor was blocked, were similar to control (91 ± 35 pg/ml), S-C24D treated cells (108 ± 20 pg/ml) and in C24D treated cells (107 ± 27 pg/ml).

C24D restores the expression of CD45 receptors, previously reduced by exposure to tumor cells

We compared the percentage of C24D which bound to leukocyte sub-populations of PBMCs from healthy female donors versus breast cancer patients. We found that a significantly lower percentage of C24D bonded to the CD56 and NKG2D...
NK cells sub-populations in breast cancer patients ($p < 0.0007$ and $p < 0.035$ respectively) (Figure 3(a, b)).

We also compared C24D binding to PBMC from healthy donors, before versus after exposure to TNBC cells. Exposure of PBMCs to MDA-MB-231 tumor cells resulted in a significant decrease in the percentage of bonded C24D to the following sub-populations: CD3 ($p < 0.005$), CD4 ($p < 0.002$), CD8 ($p < 0.0001$), CD56 ($p < 0.0001$) and NKG2D ($p < 0.03$). Treatment with C24D reversed the decrease of C24D binding to CD8 and NKG2D ($p < 0.05$), and partially reversed the decrease of C24D binding to the CD3, CD4 and CD56 PBMC sub-populations. The percentage of C24D binding to CD16 was not affected by exposure to tumor cells (Figure 3h). A representative dot blot FACS analysis is presented in Figure 3i.

**C24D binding to CD45 induces the phosphorylation of Src protein kinases**

We then investigated the effect of the C24D/CD45 binding on the CD45 signaling pathway. The phosphorylation of proteins involved in the CD45 signaling pathway was determined by western blot analysis (Figure 4a) of co-cultures of MDA-MB-231 cells and PBMC from healthy female donors.

Interestingly, we found that addition of PBMCs to the MDA-MB-231 cells resulted in an increase in the phosphorylation of the tyrosine (Y) 505 in the Lck tyrosine kinase and a decrease in the phosphorylation of the Y394 in the Lck, causing inhibition of Lck. Additionally, we found a decrease in the phosphorylation of the Y493 in the ZAP70 protein kinase and in the proto-oncogene VAV-1, resulting in a deactivation of ZAP70 and VAV-1.

Addition of C24D to the above co-cultures reversed the inhibitory signaling observed in the presence of MDA-MB-231. In 13 separate experiments, we demonstrated that addition of C24D to PBMC/MDA-MB-231 co-cultures significantly reversed immune suppression by de-phosphorylation of Lck Y505 ($p < 0.027$) and phosphorylation ZAP70 Y493 ($p < 0.0063$) and VAV1 ($p < 0.0028$) (Figure 4b). Also, Lck Y394 increased, but not significant statistically ($n = 9$, $p < 0.27$). Blocking the extracellular domain of CD45 canceled the immune-reactivating effect.
effect of treatment with C24D to the PBMC/MDA-MB-231 co-cultures (Figure 4(c, d)).

To further confirm the above results, we performed FACS analysis of gated live cells (Figure 4(e, f)). PBMCs were identified using a CD45 antibody and the CD45+ cells were gated for further analysis of protein phosphorylation. FACS analysis results demonstrated that addition of C24D to the PBMC/MDA-MB-231 co-cultures resulted in
a reduction of 38.1 ± 6.2% versus control of the phosphorylation of Y505 and an increase in the phosphorylation of Y394 in Lck. Upon treatment with C24D, phosphorylation of the Y493 in ZAP70 increased by 50.13 ± 10.2% versus control and by 36.2 ± 4.0% in VAV-1 (Figure 4e). A representative dot blot FACS analysis is depicted in Figure 4f).

**C24D reverses CD45 inhibitory signals in PBMCs from TNBC patients**

After evaluating the effect of C24D on the CD45 signaling pathway of healthy-donor-PBMC/MDA-MB-231 co-cultures, we studied the effect of C24D on protein phosphorylation in fresh PBMCs from 5 TNBC patients. Phosphorylation of PBMC proteins was evaluated by western blot analysis (Figure 5(a, b)). Addition of C24D induced the de-phosphorylation of Lck Y505 (in average from 118.0 ± 26.3% to 79.2 ± 3.6%), the phosphorylation of the Y394 in Lck (from 87.1 ± 20.4% to 117.4 ± 36.2%), thereby activating Lck. In consequence, ZAP70 Y493 and VAV-1 were phosphorylated (from 92.9 ± 4 to 116.4 ± 10.4% and from 68.9 ± 30.0 to 88.4 ± 26.9%, respectively), activating ZAP70 and VAV-1 (Figure 5c). These results were similar to those from healthy-donor-PBMC/MDA-MB-231 co-cultures with C24D.

In contrast, no effect was observed on Lck, ZAP70 and VAV-1 activation when fresh PBMCs from healthy donors, in the absence of tumors, were incubated with C24D for 5 to 24 hours, suggesting that C24D has no effect on cells which are not immunosuppressed (Figure 5(d, e)). These results were reinforced by the lack of increase in INF-γ, TNF-α and IL-2 secretion after the addition of C24D to a co-culture of MCF-10A normal breast cells and PBMCs from healthy donors (Figure 5f).

The cytokine concentrations obtained in control cells vs C24D

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**Figure 5. C24D reversed CD45 suppressive signals in PBMCs from TNBC patients.** a. PBMCs from 5 metastatic breast cancer patients were incubated with the C24D peptide for 5 to 60 minutes. Cells were lysed and subjected to SDS PAGE. Individual western blot analysis of lysed cells blotted with antibodies to detect the protein phosphorylation. b. Statistical analysis, per TNBC patient, of western blots, depicting phosphorylation of proteins, normalized to control. c. Mean (%) ± SD of phosphorylation of the Lck, ZAP-70 and VAV-1 proteins in the 5 TNBC patients, normalized to control. d. Representative western blot analysis of protein phosphorylation in fresh PBMCs from a healthy female donor. C24D was added to the PBMCs for at least 24 hours. e. Percentage of protein phosphorylation of 5 separate experiments (Mean ± SE, no statistical difference in protein phosphorylation was observed). f. Effect of C24D on secretion of INF-γ, IL-2 and TNF-α in supernatants of PBMCs from healthy female donors, co-cultured with normal breast cell MCF-10A, ± C24D (n = 3). Results presented as percentage of cytokine secretion normalized to control. g. MCF-10A (breast cancer normal cells) killing by FACS analysis, using AnnexinV/PI (N = 3). Representative FACS dot blot (experiment #1) of AnnexinV/PI gated and stained breast normal cells (CD45 negative) treated with 10 μg/ml C24D for 4–6 days. Analysis of results presented as Mean ± SD.
C24D molecular mechanism

Figure 6a depicts the suggested CD45 molecular mechanism observed when human PBMCs are exposed to TNBC cells. Figure 6b depicts the reversing effect on tumor-immunosuppressed leukocytes resulting from binding of C24D to the CD45 receptor.

Discussion

Despite enormous efforts to find successful therapies, TNBC treatment remains an unmet medical need. The large number of clinical trials currently running on checkpoint inhibitors for the treatment of breast cancers\textsuperscript{23,24} reflects the great interest in (and need for) finding better therapies. The main challenge for developing effective TNBC immunotherapies is the elucidation of the flaws in the responsiveness of breast cancers to immunotherapy. New strategies are thus needed to overcome these hurdles.\textsuperscript{25,26}

In our study, we revealed a novel role (immunotherapy) for an old target (CD45). The molecular mechanism deployed by this new/old target is fundamentally different from that of currently reported immune-checkpoint strategies and may overcome many of the difficulties encountered so far in breast cancer immunotherapy. The CD45 receptor is a cell surface glycoprotein with tyrosine phosphatase activity. It is expressed on all nucleated hematopoietic cells, except for mature erythrocytes and platelets\textsuperscript{17,26,27}. CD45 has been identified as a rheostat of T and B cell activation, doing so by positively and negatively regulating Src family protein tyrosine kinases and thereby, the signaling cascade in leukocytes.\textsuperscript{28–30}

We observed that on exposure to TNBC cells, PBMCs from healthy female donors showed no significant immune response against the tumor cells. We revealed that the lack of response against the TNBC cells might be due to the increase in the phosphorylation of the tyrosine 505 and decrease in the phosphorylation of the tyrosine 394 in Lck, thus deactivating Lck (one of the members of the Src family protein tyrosine kinases). In consequence, the tyrosine 493 in ZAP70 was dephosphorylated, inhibiting TCR activation.\textsuperscript{31–33} Decrease in ZAP70 activation affected VAV-1 phosphorylation, resulting in NK deactivation (Figure 6a).\textsuperscript{34}

Conversely, treatment with C24D reverses the CD45 inhibitory signals in PBMCs which were exposed to tumors. Mass

\begin{figure}[h]
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\caption{Figure 6. Schematic diagram: Suggested model of how C24D reverses tumor-induced immune suppression through the CD45 signaling pathway. a. In PBMCs from healthy donors exposed to TNBC tumors, phosphorylation of the inhibitory tyrosine (y) 505 in Lck (a member of the Src tyrosine kinases) increased, phosphorylation of the tyrosine 394 in Lck decreased, de-activating the Lck protein in the peripheral blood leukocytes. In consequence, tyrosine 493 in ZAP70, and VAV-1 were de-phosphorylated, de-activating ZAP70 and VAV-1. As a result, TCR activity decreased, dampening immune response. b. Binding of the C24D peptide to the CD45 receptor on leukocytes which were previously exposed to TNBC cells, reverses the suppression by de-phosphorylation of the tyrosine 505 and phosphorylation of the tyrosine 394 in Lck. Consequently, the tyrosine 493 in ZAP70 and VAV-1 are phosphorylated, resulting in TCR activation, tumor cell killing and secretion of IFN-γ, IL-2 and TNF-α.}
\end{figure}
spectrometry results revealed that C24D binds to a short isoform of the human extra cellular domain of CD45, the receptor-type tyrosine-protein phosphatase C (PTPRC). The binding receptor was validated by western blot analysis using an anti-CD45 antibody to confirm CD45 as the protein precipitated from PBMCs with C24D. C24D/CD45 interaction was demonstrated by a Proximity Ligation Assay. Validation of CD45 as C24D binding receptor was also demonstrated when the effect of C24D on IFN-γ secretion was canceled by blocking the extracellular domain of CD45. Upon binding to the CD45 receptor of tamed PBMCs, C24D induces the activation of the CD45 signal transduction pathway.

It has been reported that loss of CD45 results in hyperphosphorylation of the Lck C-terminal tail (Y505), thereby reducing the amount of active Lck. The reduction in Lck impairs T cell development when TCR signaling is required. We observed a significant decrease in CD45 binding to almost all the tested sub-populations (CD3, CD4, CD8 T cells and CD56 and NKG2D NK cells) in PBMCs from healthy donors co-cultured with MDA-MB-231 cells. The significant decrease in the percentage of CD45 binding to PBMCs suggests a loss of CD45 (or of a specific CD45 isoform) after exposure to TNBC.

In PBMCs from breast cancer patients, we observed a lower percentage of CD45 binding only to NK cells (CD56, CD16 and NKG2D positive cells). The difference in patients compared to co-cultures, is possibly due to the direct contact between the cells in the co-cultures.

The percentage of CD45 binding to T and NK cells in PBMCs co-cultured with MDA-MB-231 was restored after treatment with C24D. Concurrently, C24D treatment induced Lck, ZAP70 and VAV-1 phosphorylation, resulting in reactivation of the CD45-signaling cascade (Figure 6b) and the triggering of massive tumor cell killing.

Increase in CD69 expression reflects an immunoreactive phenotype associated with enhanced cytotoxicity against various target cells. Increase of CD69 expression in NK cells in breast cancer patients after immunotherapy was associated with better survival and longer disease free survival compared to patients with lower receptor expression.

We found that addition of C24D to the MDA-MB-231/ PBMC co-cultures induced a significant increase in activated CD4+/CD69+, CD8+/CD69+ T cells and CD56+/CD69+ NK cells and a significant increase in type 1 cytokines which are critical for optimal anti-tumor response: IFN-γ, TNF-α and IL-2. In contrast, C24D had no significant effect on CD14+CD137L+ activated monocytes. Furthermore, IL-1β secretion did not change upon C24D treatment. Increase in circulating CD8+ or CD4+ cells positive for CD57 (a marker of terminal differentiation) has been associated with poor prognosis in several solid and hematological tumors. Additionally, an inverse correlation between circulating CD57+ and tumor-infiltrating NK cells was found in Her2+ breast cancer patients. Treatment with C24D to PBMC/MDA-MB-231 co-cultures did not result in an increase of CD57+ clones, enabling effective tumor cell killing.

We also evaluated the effect of treatment with C24D in an in vivo model using TNBC cells in nude mice transplanted with human PBMCs. Analysis of the xenografts from the C24D-treated mice showed that tumor growth was arrested with both the 60 and the 300 μg/mouse doses. However, better results were achieved with 60 μg/mouse. Also in the in vitro studies, higher than optimal doses resulted in a lower response. Such bell-shaped dose responses, resulting from immunological feedback, drug-receptor binding affinity, target saturation and drug stability have been reported. Also anti-PD1 antibodies display similar bell-shaped dose-response and receptor saturation characteristics. The combination of the relatively long stability of C24D and receptor saturation may explain the reduction of the immune response with amounts larger than the optimal dose.

In the serum of the treated mice, we found an increase in human INF-γ. An immuno-fluorescent analysis of the MDA-MB-231 xenografts demonstrated a significant increase in tumor cell apoptosis, explained by a significant increase in infiltrated human CD45+ cells, composed of human CD8 + T cells and activated human CD56+ NK cells. Clinical studies have found a connection between tumor-infiltrated NK and CD8+ cells and anti-cancer immune responses linked to breast cancer patients’ survival. Particularly with TNBC, a 10% increase in tumor-infiltrating lymphocytes resulted in improved prognosis. Moreover, a recent study correlated TNBC patients’ survival (up to 18-year follow-up) with high frequency of CD8+ tumor infiltrated cells.

Since CD45 is a ubiquitous receptor expressed on all leukocytes, we performed preliminary assays to test C24D’s specificity. We treated PBMCs exposed to normal breast cells (MCF-10A) with C24D and evaluated killing and cytokine secretion. In contrast to PBMCs exposed to TNBC cells, addition of C24D did not induce an immune-stimulating effect, suggesting that C24D only affects immune-suppressed cells.

CD45 has been called a signaling “gatekeeper”, filtering weak or false signaling events and allowing effective signaling. Considering the results presented herein, it is possible that breast tumors exploit CD45’s gatekeeper function, suppressing immune response, which is subsequently reversed by treatment with C24D.

Given the central role played by CD45 in the immune system, and given the positive results obtained with C24D treatment, we propose that CD45 can be a promising therapeutic target for the reactivation of immune responses in cancer. We hope this study rejuvenates the interest in delineating new roles in immunotherapy for the “old” CD45 target.

However, further studies are essential for the development of an effective immunomodulatory compound. Studies for the identification of the C24D/CD45 binding epitope and for a more comprehensive elucidation of the anti-tumor mechanism have been initiated and will undoubtedly shed light on this novel therapeutic strategy.

**List of abbreviations:**

- **TNBC**: triple negative breast cancer
- **PD1/PDL-1**: the programmed cell death protein 1 or programmed cell death ligand 1
- **CTLA-4**: cytotoxic T-lymphocyte-associated antigen 4
- **PTPRC**: protein tyrosine phosphatase receptor type C
Institutional available
Davidoff
RY
involved
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study-
Availability
Peripheral
approved
IL-2
Lck
Imanu
10.1007/s00262-019-
17(5):574–582.
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