

Identification of Immunogenic MHC Class II Human HER3 Peptides that Mediate Anti-HER3 CD4⁺ Th1 Responses and Potential Use as a Cancer Vaccine



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ABSTRACT

The HER3/ERBB3 receptor is an oncogenic receptor tyrosine kinase that forms heterodimers with EGFR family members and is overexpressed in numerous cancers. HER3 overexpression associates with reduced survival and acquired resistance to targeted therapies, making it a potential therapeutic target in multiple cancer types. Here, we report on immunogenic, promiscuous MHC class II-binding HER3 peptides, which can generate HER3-specific CD4⁺ Th1 antitumor immune responses. Using an overlapping peptide screening methodology, we identified nine MHC class II-binding HER3 epitopes that elicited specific Th1 immune response in both healthy donors and breast cancer patients. Most of these peptides were not identified by current binding algorithms. Homology

assessment of amino acid sequence BLAST showed >90% sequence similarity between human and murine HER3/ERBB3 peptide sequences. HER3 peptide-pulsed dendritic cell vaccination resulted in anti-HER3 CD4⁺ Th1 responses that prevented tumor development, significantly delayed tumor growth in prevention models, and caused regression in multiple therapeutic models of HER3-expressing murine tumors, including mammary carcinoma and melanoma. Tumors were robustly infiltrated with CD4⁺ T cells, suggesting their key role in tumor rejection. Our data demonstrate that class II HER3 promiscuous peptides are effective at inducing HER3-specific CD4⁺ Th1 responses and suggest their applicability in immunotherapies for human HER3-overexpressing tumors.

Introduction

Oncodrivens are proteins overexpressed in tumor cells that promote proliferation and growth, but counteract cellular senescence, contributing to tumor malignancy. Such oncoviral addiction in tumor cells makes these proteins a promising target for developing new immunotherapies relevant to a large number of patients. Human ERBB3

receptor tyrosine kinase 3 (HER3/ERBB3) is a member of the ERBB family of growth receptors that has several ligands, including heregulin and neuregulin. Although lacking intrinsic kinase activity, HER3 is a critical heterodimerization partner for other members of the family (EGFR and HER2) contributing to the growth, proliferation, and survival of tumor cells (1). HER3 is an established oncoviral in multiple cancers because HER3 overexpression has been detected in breast, melanoma, colorectal, prostate, lung, and ovarian cancers (2). The HER2-HER3 heterodimer is responsible for the most potent ligand-induced tyrosine phosphorylation and downstream signaling via the PI3K-AKT pathway, utilizing the six p85-binding motifs present in HER3 intracellular domain (3, 4). In HER2-overexpressing (HER2^{POS}) breast cancer, HER3 is identified as one of the most prominent inducers of therapy escape, leading to trastuzumab resistance and hyperactivation of PI3K-AKT-mediated signaling (3). Transcriptional/translational upregulation of HER3 is linked to resistance to MEK/RAF inhibitors in melanoma (5), castration-resistant prostate cancer (6), platinum-resistant ovarian cancer (7), and EGFR TKI-resistant non-small cell lung cancer (NSCLC; refs. 8, 9). In triple-negative breast cancer (TNBC), HER3 overexpression has been independently identified as a prognostic marker of poor survival (10, 11), and combined antagonism of EGFR and HER3 enhances responses to PI3K-AKT inhibitors in TNBC preclinical and clinical samples (12). Multiple monoclonal antibodies targeting HER3 are currently being tested at various stages of preclinical and clinical studies; however, therapeutic potential of a HER3-specific immunotherapy has not been comprehensively tested. Developing HER3-specific cellular immunotherapy can be a novel and efficient treatment strategy for multiple cancer types overexpressing HER3 in order to improve patient prognosis and survival.

Although current immunotherapies focus on cytotoxic CD8⁺ T-cell activity, “helper signals” from CD4⁺ T cells have been deemed

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essential for the proliferation, recruitment, and effector function of CD8⁺ T cells in tumors (13, 14). Help from CD4⁺ T cells is necessary for memory T-cell survival during recall expansion (15), and IL-2 secreted from CD4⁺ T cells in the tumor microenvironment has been shown to increase CD8⁺ T-cell proliferation and granzyme B production (16). Absence of CD4⁺ T-cell help affects survival and clonal expansion of CD8⁺ T cells, and defective recall responses by CD8⁺ T cells have been reported in CD4^{-/-} mice (17). In a head and neck cancer model, HER3-specific helper T-cell responses induce cytolytic CD4⁺ T-cell activity against tumor cells in an HLA-DR-restricted manner (18). These studies highlight the potential for developing CD4⁺ T cell-based immunotherapy for cancer, and as previously observed by our lab and others, oncogenes represent an excellent choice for developing such therapeutic strategies (19).

A previous study from our lab demonstrates gradual loss of HER3-specific CD4⁺ Th1 immune responses in the peripheral circulation of TNBC patients compared with healthy donors and other breast cancer subtypes, and this loss of immune response negatively associates with patient outcome (20). In the current study, we investigated whether promiscuous MHC class II HER3 peptides could be identified to develop an immunotherapy strategy for HER3-overexpressing cancers. Herein, we present a peptide screening methodology for identifying promiscuous class II epitopes that are capable of inducing tumor-specific CD4⁺ T-cell responses. We identified nine immunogenic class II epitopes and investigated the therapeutic efficacy of HER3 peptide-pulsed type I DC (HER3-DC1) treatment in multiple preclinical models of breast cancer and melanoma and in both preventive and therapeutic settings.

Materials and Methods

HER3 expression in cancer and correlation with survival

The Cancer Genome Atlas (TCGA) RNA sequencing (RNA-seq) expression data (FPKM) for *HER3* (*ERBB3*) in 12 types of cancers were downloaded from the Genomic Data Commons Data Portal (GDC; <https://portal.gdc.cancer.gov>; RRID:SCR_014514). Project IDs included in analysis were as follows: TCGA-BLCA, bladder urothelial carcinoma ($n = 408$); TCGA-BRCA, breast invasive carcinoma ($n = 1092$); TCGA-COAD, colon adenocarcinoma ($n = 456$); TCGA-HNSC, head and neck squamous cell carcinoma ($n = 501$); TCGA-LUAD, lung adenocarcinoma ($n = 515$); TCGA-LUSC, lung squamous cell carcinoma ($n = 501$); TCGA-OV, ovarian serous cystadenocarcinoma ($n = 376$); TCGA-PAAD, pancreatic adenocarcinoma ($n = 177$); TCGA-PRAD, prostate adenocarcinoma ($n = 496$); TCGA-SKCM, skin cutaneous melanoma ($n = 468$); and TCGA-STAD, stomach adenocarcinoma ($n = 380$). Only samples with available sequence reads were chosen for further analysis. *HER3* mRNA expression in primary and normal tissues was assessed and graphed as a boxplot using a customized R script (Supplementary File). For tumor samples, only data from primary tissues were used. An unpaired *t* test was used for differential analysis of the data set.

Survival plots were generated by the cBioPortal (<https://www.cbioportal.org>; RRID:SCR_014555) survival tool. Samples were sorted in the descending order, according to the median value of the *HER3* gene expression, and the list was divided into two groups in the middle of the list: High expression was defined as the top 50% of the list and low expression as the bottom 50% of the list.

Peptide library generation

The amino acid sequence for native human HER3 protein (1,342 amino acids, Uniprot Accession number P21860) was divided into two

peptide libraries: the extracellular domain (ECD, aa 20–643) and intracellular domain (ICD, aa 665–1,342). Libraries consisted of peptide fragments 15 amino acids in length, with a 10-amino acid overlap between adjacent sequences, generating 123 ECD and 134 ICD peptides (GenScript). BRAF-negative control peptide was purchased from GenScript. Peptides were reconstituted according to solubility instructions (GenScript) to 1 mg/mL concentrations. ICD peptides p37, p54, and p55 were unable to be synthesized. Identified immunogenic human HER3 peptides were used for testing in preclinical mouse models as well, due to significant sequence homology.

Human samples

Autologous monocyte fractions were obtained from leukapheresis products from the peripheral blood of normal donors and breast cancer patients. Normal donor samples ($n = 11$) were purchased through the Cell Therapies Core at H. Lee Moffitt Cancer Center and Research Institute, which performed the elutriation to generate monocyte- and lymphocyte-rich fractions. Breast cancer patient elutriation fractions were obtained under the Moffitt Cancer Center (MCC) 19000 protocol and collected during 2017–2021. Inclusion criteria included samples from newly diagnosed, operable breast cancer patients with: (i) incident invasive primary ductal breast cancer prior to surgery with Her2/neu overexpression via IHC; (ii) age over 18; and (iii) consent to participate. Patients with bilateral breast cancer were not excluded. Exclusion criteria included: (i) patients with no plan for definitive surgery; (ii) no detectable residual breast disease after diagnostic biopsy; and (iii) patients unable to receive neoadjuvant herceptin-based chemotherapy. The control group was composed of unaffected, seemingly healthy individuals with inclusion criteria: (i) no prior history of systemic cancer treatment (i.e., chemotherapy or hormone therapy); (ii) age over 18; (iii) consent to participate; and (iv) BIRADS 1–3 breast imaging within the past 12 months. Exclusion criteria included: (i) active diagnosis of autoimmune disease (i.e., rheumatoid arthritis, lupus, Sjogren syndrome) and (ii) systemic steroid use within the past 12 months.

Elutriation fractions were washed in PBS (cat. #SH30028LS, Fisher Scientific) and lysed with ACK RBC lysis buffer containing NH₄Cl (0.15M, cat. #A649-500; Fisher Scientific), KHCO₃ (10 mmol/L, cat. #P184-500, Fisher Scientific) and Na₂EDTA (0.5 mmol/L; cat. #E7889-100 mL, Sigma-Aldrich), with pH 7.2, when necessary to remove remaining RBCs. Processed fractions were stored in liquid nitrogen until further use.

Breast cancer patient ($n = 10$) and healthy control ($n = 6$) samples were used for *in vitro* validation of identified HER3 class II peptides. All clinical samples were obtained by informed written consent from the subjects, following an Institutional Review Board-approved protocol. All research involving human subjects was performed in accordance with the Declaration of Helsinki and was approved by the IRB at Advarra.

Human DC generation

Monocytes were differentiated into dendritic cells (DC) through the addition of recombinant human (rh)GM-CSF (50 ng/mL; cat. #215-GM-050, R&D Systems) and rhIL4 (10 ng/mL; cat. #204-IL-050, R&D Systems) and cultured in macrophage serum-free media (cat. #12065074; Thermo Fisher Scientific) for 24 hours at 37°C and 5% CO₂. DCs were then pulsed with HER3 ECD or ICD peptides (10 µg/mL), a negative control (BRAF class II p8), or left unpulsed. After 24 hours, DCs were matured and polarized to a type I phenotype (DC1) with the addition of rhIFN-γ (1,000 IU/mL; cat. #2851F-100, R&D Systems), followed by LPS (0.2 µg/mL; cat.

#L4391-1MG, Sigma-Aldrich) 6 hours prior to harvest. Mature DC1 cells were harvested after 72 hours of culture. Immature DCs (iDC) remained in rhGM-CSF and rhIL4 for 48 hours prior to harvesting. DCs were preserved in 10% DMSO (cat. #D5879-500 mL, VWR) and 90% human AB serum (cat. #S11550, Atlanta Biologicals) and stored in liquid nitrogen until use.

Culture medium

Complete media consisted of RPMI-1640 growth media (cat. #MT-10-040-CM, Corning) supplemented with 10% heat-inactivated fetal bovine serum (FBS; cat. #S11550 Atlanta Biologicals), 0.1 mmol/L nonessential amino acids (cat. #25-025-CI, Corning), 1 mmol/L sodium pyruvate (cat. #MT-25-000-CI, Corning), 100 mg/mL streptomycin, 100 U/mL penicillin (cat. #MT-30-002-CI, Fisher Scientific), 50 mg/mL gentamycin (cat. #15750-060, Gibco), 0.5 mg/mL amphotericin B (cat. #400-104, GeminiBio), and 0.05 mmol/L 2-mercaptoethanol (cat. #21985-023, Invitrogen).

Peptide screening

Mature HER3-DC1 were cocultured in complete media with autologous lymphocytes at a 1:10 ratio in a 12-well tissue culture plate and incubated at 37°C and 5% CO₂. The lymphocyte-rich apheresis fractions were processed as described above. After 24 hours, rhIL2 (5 IU/mL) was added to induce the proliferation of CD4⁺ T cells. Following 8 to 10 days of coculture, T cells were harvested and restimulated in a 96-well plate with iDCs (10⁵ T cells with 10⁴ iDCs for 10:1 ratio) pulsed with the corresponding HER3 class II peptide, a negative class II control (BRAF class II p8, 10 µg/mL), or unpulsed iDCs.

Culture supernatants were collected after 24 hours, and IFN-γ production was measured using a Human Quantikine IFN-γ ELISA kit (cat. #SIF50, R&D Systems) after 1:20 or 1:30 sample dilution. The screening process proceeded using a 10-peptide pool, 5-peptide pool, and individual peptide (peptide concentration 10 µg/mL throughout the screening sequence) sequential scheme (Supplementary Fig. S1A) for both ECD and ICD libraries, separately. Libraries were first screened in pools of 10 peptides and run on three independent donor samples. The 10-peptide pools that showed ≥1.5-fold increase in IFN-γ production in two of the three donors were further pursued in 5-peptide pools. Common positive 5-peptide pools were then tested as individual peptides and screened in additional donor samples to identify peptides that could reproducibly induce an increased CD4⁺ Th1 immune response. Due to the variability in donor sample size and the progressive nature of the peptide screening methodology, the sample number tested for each identified peptide was inconsistent; however, individual peptide candidates were screened across a minimum of five donor samples, with a threshold response rate of ≥1.5-fold to be considered immunogenic.

Whole HER3 ECD/ICD protein restimulation

Immunogenicity of the identified HER3 class II epitopes was confirmed through sensitization of CD4⁺ T cells with HER3-DC1 as described above, followed by restimulation with iDCs pulsed with the corresponding HER3 class II peptide, a negative peptide control (BRAF class II p8), native HER3 ECD (cat. #NBP2-52128-0.05 mg, Novus Biologicals) or ICD (cat. #10201-H20B1, SinoBiological) whole protein sequence, or a negative whole protein control (Hemocyanin-Keyhole Limpet Native protein; cat. #SRP6195, Sigma). The same number of T cells, HER3-DC1, and iDCs were used for this experiment and 10 µg/mL of whole protein and/or peptide was used throughout this experiment. Culture supernatants were collected after 24 hours,

and IFN-γ production was measured using the Human Quantikine IFN-γ ELISA kit after 1:20 or 1:30 sample dilution.

ELISPOT assays

To evaluate anti-HER3 CD4⁺ Th1 immune responses in breast cancer patients, IFN-γ production was compared between healthy controls and patient samples using the human IFN-γ ELISPOT kit (cat. #HIFNgp-1M/10, Cellular Technologies Limited). ELISPOT plates precoated with human IFN-γ capture antibody were incubated with nine HER3 class II peptides (4 µg/well), media only (untreated/negative control), or anti-human CD3 (Orthoclone OKT3, cat. #73337989, Johnson and Johnson, treated/positive control, 15 ng/mL). Cryopreserved PBMCs were plated (2 × 10⁵ cells/well) in CTL-Test medium supplemented with 1% L-glutamine (provided with the kit) and incubated at 37°C, 5% CO₂ for 48 hours. Per manufacturer's protocol, plates were washed, and detection antibody (anti-human IFN-γ Biotin; 100 mg/mL) was added to each well. After incubation at room temperature for 2 hours, 1:1,000-diluted streptavidin-AP was added and incubated for 30 minutes. Plates were washed again before substrate solution was added, and plates were incubated for 15 minutes to allow for color development. Plates were washed with tap water, dried overnight at room temperature, and spot-forming cells (SFC) were counted using an automated reader (Immunospot Cellular Technology Limited). Because inter-replicate variability in ELISPOT was low, an empiric method of determining antigen-specific responses was used as previously shown (21). In brief, positive response to an individual class II HER3 peptide was defined as (i) threshold minimum of 20 SFC/2 × 10⁵ cells in experimental wells after subtracting unstimulated background; and (ii) approximately 2-fold increase of antigen-specific SFCs over background. Three metrics of CD4⁺ Th1 response were defined for each group: (i) overall anti-HER3 responsivity (proportion of patients responding to ≥1 peptide), (ii) response repertoire (mean number of reactive peptides), and (iii) cumulative response across nine identified class II HER3 peptides (total SFC/1 × 10⁶ cells).

HLA typing of donor samples

Lymphocyte fractions of healthy donors (*n* = 11) used in peptide screening HER3 ECD and ICD libraries were sent for HLA-DRB1, HLA-DQB1, and HLA-DPB1 allele typing (American Red Cross). Alleles expressed by each donor were cross-referenced with the CD4⁺ Th1 response induced by each identified HER3 class II peptide. Each HER3 peptide tested against a donor expressing a particular HLA allele was indicated with a (+) if ≥1 instance of this allele demonstrated a peptide-specific Th1 immune response (IFN-γ production increased ≥1.5-fold compared to negative control) to the corresponding peptide. Because HER3 ECD and ICD libraries were screened using different donor samples, this resulted in two groupings.

NetMHCIIpan 4.0 prediction algorithm

Epitope prediction was done using NetMHCIIpan 4.0 (22) as implemented by www.iedb.org (23). A series of 15-mer peptides were serially generated (one amino acid difference) from the HER3 protein. Each peptide was then predicted for its binding affinity (nmol/L) with each HLA allele type by identifying a 9-mer binding core. Peptide candidates with affinity <500 nmol/L were labeled WB (weak binding) and <50 nmol/L as SB (strong binding).

MixMHCIIpred prediction algorithm

Epitope prediction was done using the MixMHCIIpred algorithm (24, 25). A series of random 12–25 amino acid (aa) peptides

were generated from the HER3 protein sequence by the algorithm. A percentile rank was provided for each of the nine HER3 peptides out of all 12–25aa random peptides, for each of the 38 HLA allele types. Best score for percentile rank was 0 and the worst score was 100. The best alleles with the highest binding affinity for each peptide were also identified.

Cell lines and reagents

The human TNBC cell lines MDA-MB-231 (ATCC HTB-26, RRID: CVCL_0062), MDA-MB-468 (ATCC HTB-132, RRID:CVCL_0419), HCC1143 (ATCC CRL-2321, RRID:CVCL_1245), Hs578T (ATCC HTB-126, RRID:CVCL_0332), BT-549 (ATCC HTB-122, RRID: CVCL_1092), SK-BR-3 (ATCC HTB-30, RRID: CVCL_0033) and mouse breast cancer cell lines 4T1 (ATCC CRL-2539, RRID: CVCL_0125) and EO771 (ATCC CRL-3461, RRID:CVCL_GR23) were obtained from the American Type Culture Collection. The TUBO murine breast cancer cell line (a kind gift from Dr. Wei Zen Wei, Wayne State University) was cloned from a spontaneous mammary tumor in BALB/c mice transgenic for the rat *HER2/neu* gene (BALB-HER2/neuT; ref. 26). The M05 cell line was a kind gift from Dr. Shari Pilon-Thomas (Moffitt Cancer Center), and cells were grown in media containing 0.8 mg/mL G418 (neomycin; cat. #30234CR, Corning; refs. 27, 28). M05 is derived from the B16 murine melanoma cell line, transfected to express ovalbumin, and transgene expression was maintained using G418 as the selection antibiotic in the culture media. Cells were grown at 37°C in a humidified 5% CO₂ incubator in complete media. All cell lines used in the study tested negative for *mycoplasma* using a mycoplasma kit (PlasmoTest, cat. #rep-pt1, InvivoGen). Cell lines at passages 2–5 were used for all experiments. For all *in vitro* experiments, recombinant human and mouse IFN- γ were purchased from R&D Biosystems (human: cat. #285IF-100; mouse: cat. #485-MI-100).

BLASTP analysis

Amino acid sequence homology between identified human HER3 peptides and murine ErbB3 sequence (UniProtKB; Q61526) was evaluated by using the BLASTP program (http://blast.ncbi.nlm.nih.gov/Blast.cgi?PROGRAM=blastp&PAGE_TYPE=BlastSearch&LINK_LOC=blasthome RRID:SCR_001010). The percentage of amino acid sequence homology has been reported.

Western blots

For protein expression, cells were seeded (4×10^5 cells/well for human TNBC cells, 2×10^5 cells/well for mouse TNBC cells) in six-well plates and cell lysates were prepared in RIPA buffer (cat. #20-188, EMD Millipore) containing protease inhibitor cocktail (10 μ L/mL; cat. #P8340-1ML, Sigma-Aldrich) and phosphatase inhibitor (100 μ L/mL; cat. #A32957, Pierce) by incubating for 20 minutes at 4°C. Cell lysates were centrifuged at $14,000 \times g$ for 20 minutes at 4°C, and total protein in the supernatant was collected and stored at –86°C until further use. Protein concentration was measured using Bradford protein assay (cat. #5000006, Bio-Rad). For Western blotting, 20 μ g protein was resolved on 4% to 12% SDS-PAGE (GenScript) and transferred onto PVDF membranes (cat. #IPVH00010, Millipore) using eBlot L1 wet transfer system (GenScript). Membranes were blocked with 5% BSA/TBS-T for 1 hour at room temperature, and then incubated with following primary antibodies overnight at 4°C: HER3 (cat. #12708S), phospho-Stat1 (Tyr701; 58D6; cat. #9167S, Cell Signaling Technology), cleaved caspase-3 (Asp175; cat. #9661S, Cell Signaling Technology), phospho-p44/42 MAPK (Erk1/2; Thr202/Tyr204; cat. N#9101S, Cell

Signaling Technology; RRID:AB_2315112), phospho-Akt (Ser473; cat. #9271S, Cell Signaling Technology; RRID:AB_329825; all at 1:1,000 dilution), and β -actin (C4; cat. #sc-47778, Santa Cruz Biotechnology; RRID:AB_2714189; 1:3,000 dilution). Membranes were probed with anti-rabbit IgG HRP-linked secondary antibody (cat. #7074S, Cell Signaling Technology, RRID:AB_2099233) or goat anti-mouse IgG (H + L)–HRP conjugated secondary antibody (cat. #172-1011, Bio-Rad, RRID:AB_11125936; 1:3,000 dilution) for 1 hour at room temperature. Protein expression was detected with ECL Western blotting substrate (cat. #32106, Pierce) using an AmershamTM Imager 600 image acquisition system. Analysis and quantification of Western blot images were performed using ImageJ software (<http://rsb.info.nih.gov/ij/>, RRID:SCR_003070).

Immunofluorescence

Cells were seeded (4×10^5 cells/well for human cells (MDA-MB-468, SK-BR-3), 2×10^5 cells/well for mouse cells (4T1, TUBO, and M05) in six-well plates, each well containing three 12-mm round glass coverslips (cat. #12-545-80, Fisher Scientific). After cells reached 70% to 80% confluence, cells were washed twice in PBS, fixed with 4% paraformaldehyde for 15 minutes at room temperature, and washed three times with PBS. Cells were permeabilized with 0.02% Triton X-100 (cat. #T8787—50 mL, Sigma; in PBS) for 10 minutes at room temperature, washed three times with PBS, and blocked with 5% BSA/PBS blocking buffer for 1 hour at room temperature. Cells were incubated with primary anti-HER3 (cat. #12708S, Cell Signaling Technology) in 3% BSA/PBS (1:500 dilution) overnight at 4°C. The next day, cells were washed in PBS three times and incubated in Alexa Fluor 594-conjugated goat-anti-rabbit secondary antibody (1:5,000 dilution; cat. #8889S, Cell Signaling Technology) and FITC-conjugated goat anti-mouse secondary antibody (1:5,000 dilution; cat. #115-095-003 Jackson ImmunoResearch Inc, RRID: AB_2338589) for 1 hour at room temperature. Cells were washed three times in PBS, and the coverslips were mounted onto sterile glass slides using VECTA-SHIELD Antifade Mounting Medium with DAPI (cat. #H-1200, Vector Laboratories). Slides were sealed with clear nail varnish and allowed to cure overnight at 4°C in the dark. Immunofluorescence images were obtained using Zeiss Apotome.2 fluorescence microscope (Carl Zeiss Inc.).

Mouse DC generation

Bone marrow was harvested from 6- to 8-week-old BALB/c (RRID: IMSR_ORNL:BALB/cRI) and C57BL/6 (RRID: IMSR_JAX:000664) mice (purchased from Charles River Laboratories), as described previously (29). Briefly, femurs and tibias were harvested from mice, bone marrow cells were flushed to obtain a single-cell suspension in PBS, and red blood cells were lysed using ACK lysis buffer. Cells (2×10^6 cells/mL) were then cultured in complete media containing recombinant human Flt3L (25 ng/mL; cat. #10778-670, VWR/Pepro-Tech) and recombinant mouse IL6 (30 ng/mL; cat. #406ml025, R&D Systems), and incubated for 6 days at 37°C and 5% CO₂. On day 6, cells were harvested, washed with RPMI-1640, and cultured with recombinant mouse GM-CSF (50 ng/mL; cat. #415-ML-050, R&D Systems) and recombinant mouse IL4 (10 ng/mL; cat. #404ml050, R&D Systems) overnight for DC differentiation. The cells were matured with DC1-polarizing cytokines: CPG/ODN1826, a TLR9 agonist (10 ng/mL; cat. #NC9685794, InVivoGen), and LPS, a TLR4 agonist (20 ng/mL). DCs were pulsed with the nine immunogenic class II human HER3 peptides (10 μ g/mL) 6 to 8 hours later and harvested the following morning prior to injection into mice.

Mouse models

Female BALB/c and C57BL/6 mice (6–8 weeks old) were housed at the Animal Research Facility of the H. Lee Moffitt Cancer Center and Research Institute. The study protocol was designed in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the NIH. The protocol was reviewed and approved by the Institutional Animal Care and Use Committee at the University of South Florida. Mice were observed daily and were euthanized by CO₂ inhalation at the end of the study or if a solitary subcutaneous tumor exceeded endpoint (250 mm² for 4T1 and 400 mm² for TUBO and M05), following the American Veterinary Medical Association Guidelines. All efforts were made to minimize suffering.

Preventive models

BALB/c and C57BL/6 mice ($n = 8$ – 10 /group) were vaccinated with either unpulsed mature DC1 cells or HER3-DC1 cells subcutaneously (1×10^6 cells/mouse) twice a week for a total of six doses. Two weeks after the last DC injection, naïve and vaccinated BALB/c mice received either 4T1 (50,000 cells/mouse) or TUBO (250,000 cells/mouse) tumor cells, and C57BL/6 mice received M05 (300,000 cells/mouse) tumor cells, administered subcutaneously on the opposite flank to the vaccination site. Tumor size was measured using caliper and recorded every 2 to 3 days. Mouse tumor area was determined by the formula length \times width (mm²). Each experiment was performed three times.

Therapeutic models

For therapeutic models, either 4T1 (50,000 cells/mouse) or TUBO (250,000 cells/mouse) were injected subcutaneously in BALB/c mice, and M05 cells (300,000 cells/mouse) were injected in C57BL/6 mice ($n = 8$ – 10 mice/group). After 7 to 10 days (7 days for 4T1 and TUBO, 10 days for M05) when tumors were palpable, mice received intratumoral injection of either unpulsed mature DC1 cells or HER3-DC1 (2×10^6 cells/mouse for 4T1 and 1×10^6 cells/mouse for TUBO and M05, in 50 μ L PBS) once (TUBO and M05) or twice (4T1) a week for a total of six doses. Control mice received PBS intratumoral injection. Tumor growth was measured with caliper and recorded every 2 to 3 days, and tumor area was determined by the formula length \times width (mm²). Each experiment was performed three times.

CD4⁺ T-cell depletion in therapeutic models

BALB/c mice were injected intraperitoneally with monoclonal CD4 depletion antibody [InVivoMAb, anti-mouse CD4 (GK1.5); cat. #BE0003-1, Bio X Cell; RRID:AB_1107636; 300 μ g/mouse] starting 3 days before subcutaneous TUBO tumor cell injection (250,000 cells/mouse) and was continued twice a week until endpoint. When tumors were palpable, mice were randomized into two groups. One group of mice continued receiving CD4 depletion antibody only, and the other group received once-weekly intratumoral HER3-DC1 (1×10^6 cells/mouse, in 50 μ L PBS) along with CD4 depletion antibody. Another group of TUBO tumor-bearing mice received intratumoral HER3-DC1 only, and the control group of mice received PBS intratumorally. Tumor growth was measured with caliper and recorded every 2 to 3 days, and tumor area was determined by the formula length \times width (mm²).

Flow cytometry

For functional analyses, mice ($n = 3$ /group) were sacrificed 2 weeks after the last HER3-DC1 injection, and tumors, spleen, and lymph nodes (tumor-draining and nondraining) were collected under sterile conditions for *in vitro* assays. Single-cell suspensions were prepared

from the tumor samples by enzymatic digestion with HBSS (Fisher Scientific; cat. #MT-21-022-CM), containing 1 mg/mL collagenase (cat. #C9891 and C-5138), 0.1 mg/mL DNase I (cat. #DN25), and 2.5 U/mL hyaluronidase (cat. #H-6254-1G; all purchased from Millipore Sigma), by constant stirring for 2 hours at room temperature. Tumor digests were strained through a 100- μ m cell strainer, and ACK lysis buffer was used to remove red blood cells from tumor digests. Resulting cell suspensions were further strained through a 70- μ m and a 30- μ m cell strainer to generate single-cell suspension. Splenocytes and lymph nodes were processed in PBS. Samples were macerated using the back of a syringe plunger in a 10-cm culture dish, followed by red blood cell lysis using ACK lysis buffer. After lysis, cell suspensions were strained through a 70- μ m and a 30- μ m cell strainer sequentially. Cell suspensions were centrifuged at 1,500 rpm for 5 minutes, and cells were resuspended in complete media and counted before plating.

To analyze distribution and phenotype of immune cell populations, 1×10^6 cells were incubated with Live/Dead Zombie near IR (cat. #423106, BioLegend) for 30 minutes at room temperature. After washing once with PBS, cells were stained with the following mouse antibodies for surface expression analysis: CD3-APC (cat. #553066, Clone 145-2C11, BD Biosciences), CD45-PE/Cy7 (cat. #103114, Clone 30-F11, BioLegend), or CD45-BUV395 (cat. #564279, Clone 30-F11, BD Biosciences), CD4-BUV805 (cat. #564922, Clone GK1.5, BioLegend), CD8-Pacific Blue (cat. #558106, Clone 53-6.7, BD Biosciences), CD44-FITC (cat. #553133, Clone IM7, BD Biosciences), CD62L-BUV395 (cat. #740218, Clone MEL-14, BD Biosciences). Cells were incubated in the antibody solution made in staining buffer, which was made with PBS (without calcium or magnesium, 100 mL/L; cat. #14200166, GIBCO), sodium azide (1 g/L; cat. #S2002, Sigma) and BSA (1% final W/V; cat. #BP1605-100, Fisher Scientific) in water, for 20 minutes on ice and protected from light, following the manufacturer's instructions. Flow cytometry acquisition was performed using an LSRII (BD Biosciences) cytometer, and FACS data analysis was performed with FlowJo software (FlowJo, RRID:SCR_008520).

Functional restimulation/coculture assays

To investigate antigen specificity following HER3-DC1 administration in mice, spleens and lymph nodes were harvested 2 weeks after the last vaccination, processed as stated above, and splenocytes from control and vaccinated/treated mice were plated in 48-well tissue culture plates at 2×10^6 cells/mL in 1% FBS-supplemented RPMI and rested for 24 hours at 37°C and 5% CO₂. After 24 hours, cells were either pulsed (2 μ g/mL) with individual HER3 class II immunogenic peptides, a negative class II control peptide, or left unpulsed. For lymph nodes, lymphocytes were plated in a 96-well plate with DCs, previously matured as described above, and pulsed with HER3 class II peptides, a negative class II control (OT-II), or left unpulsed (lymphocyte:DC ratio 10:1). Following 72 hours of incubation, culture supernatant was collected, and IFN- γ secretion was measured after 1:20 dilution of the supernatant, using mouse IFN- γ Quantikine ELISA Kit (cat. #PMIF00, R&D Systems). The same protocol was followed for all mouse models included in the study.

Intracellular staining for IFN- γ secretion

Tumors were collected ($n = 3$ – 5 /group) from control and HER3-DC1-treated TUBO tumor-bearing mice. Tumor-infiltrating lymphocytes (TIL) were isolated following the protocol described above and cocultured with mature DC1 cells pulsed with individual HER3 peptides (10:1 TIL:DC; i.e., 10^6 TIL: 10^5 DC in 1 mL total volume). Intracellular staining was performed using the BD Cytofix/Cytoperm Plus Fixation/Permeabilization Kit with BD GolgiPlug

Protein transport inhibitor containing Brefeldin A (cat. #555028, BD Biosciences). Briefly, 6 hours after TIL:DC coculture, GolgiPlug was added to inhibit intracellular protein transport (1 $\mu\text{L}/10^6$ cells) for 12 hours. Cells were harvested the next day, and surface staining with CD45-BUV395 (cat. #564279, Clone 30-F11, BD Biosciences), CD4-BUV805 (cat. #612900, Clone GK1.5, BD Biosciences), and CD8-Pacific Blue (cat. #558106, Clone 53-6.7, BD Biosciences) was performed as described above. Cells were fixed and permeabilized following the manufacturer's protocol and were stained for intracellular IFN- γ -PE (cat. #554412, Clone XMG1.2, BD Biosciences). Acquisition was performed using an LSRII (BD Biosciences) cytometer, and FACS data analysis was performed with FlowJo software (FlowJo).

Statistical analyses

To compare immune response generated by the peptides, ELISA data were analyzed by multiple *t* test, without correction for multiple comparisons, using GraphPad Prism software (RRID:SCR_002798). Each row was analyzed individually, without assuming consistent standard deviation. Data are represented as mean \pm SEM. For analyzing immune response across HLA alleles, statistical significance was determined using Mann-Whitney test (ns, $P > 0.05$), and data are represented as mean \pm SEM. To compare tumor growth between groups, data were analyzed using multiple *t* test without correction for multiple comparisons. Each row was analyzed individually, without assuming a consistent standard deviation, and data are represented as mean \pm SEM. A log-rank (Mantel-Cox) test was used to determine differences between the survival curves. Unpaired two-tailed *t* test was performed to analyze Western blot data. For all analyses, significance threshold was considered as *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$.

Results

HER3 is overexpressed in multiple cancers

HER3 is an established oncodriver that contributes to the growth, proliferation, and survival of cancer cells (30, 31). Analysis of *HER3* RNA-seq expression data in 12 cancers from the GDC demonstrated significantly increased *HER3* expression in bladder ($P = 0.0045$), breast ($P = 1.15\text{E}^{-12}$), lung adenocarcinoma ($P = 1.77\text{E}^{-19}$), prostate ($P = 6.63\text{E}^{-11}$), and stomach cancers ($P = 1.31\text{E}^{-5}$; **Fig. 1A**). Although not statistically significant, a similar trend of high *HER3* expression was also noted in pancreatic cancer ($P = 0.1843$). Conversely, *HER3* mRNA expression in tumor tissues was significantly lower than healthy tissues in HNC ($P = 7.27\text{E}^{-7}$) and lung squamous cell carcinoma ($P = 1.82\text{E}^{-7}$), indicating cancer type specificity. Due to lack of data on normal tissue *HER3* mRNA expression in TCGA database, comparison of ovarian serous cystadenocarcinoma and melanoma could not be performed; and no change in *HER3* expression was noted in colon adenocarcinoma ($P = 0.1259$) or esophageal cancer ($P = 0.758$). We used the cBioPortal survival tool to further investigate if elevated/suppressed *HER3* expression correlated with overall patient survival. Whereas in breast cancer high versus low *HER3* expression had no statistically significant correlation with overall survival ($P = 0.182$), high *HER3* expression showed a significantly negative correlation with overall survival in melanoma ($P = 0.0246$; **Fig. 1B–C**). Although *HER3* mRNA expression was not significantly different between normal pancreas and pancreatic adenocarcinoma, a significant negative correlation was observed between high *HER3* expression and overall survival ($P = 6.248\text{E}^{-3}$) in pancreatic cancer (Supplementary Fig. S1B). A positive correlation between high *HER3* expression and overall survival was indicated in head and

neck squamous cell carcinoma ($P = 3.913\text{E}^{-3}$; Supplementary Fig. S1C). We did not observe a statistically significant correlation between *HER3* expression and overall survival in any of the other cancers investigated (Supplementary Fig. S1D–S1K). These results suggested an impact of increased expression of *HER3* in specific malignancies.

Identification of HER3 class II peptide epitopes

To identify immunogenic class II epitopes from the *HER3* protein, human DCs were pulsed with overlapping 15-mer peptides from a *HER3* library, rapidly matured into a type I phenotype (DC1), and cocultured with autologous naïve CD4⁺ T cells (Supplementary Fig. S2). Following restimulation of peptide-primed CD4⁺ T cells with the matching class II peptide or negative control, supernatants were collected to measure peptide-specific Th1 responses by IFN- γ secretion. Peptide libraries were sequentially screened in pools of 10-peptides, 5-peptides, and individual peptides based on significant fold increases in IFN- γ compared with control (see Materials and Methods and Supplementary Fig. S1A). Both ECD and ICD *HER3* peptide libraries were screened separately on three healthy donor samples with two representative screenings reported for ECD (samples 1–2; **Fig. 1D** and **E**) and ICD libraries (sample 3–4; **Fig. 1F** and **G**) to identify immunogenic *HER3* peptides and confirm reproducibility of peptide-specific Th1 immune responses. Screening of *HER3* ECD 10-peptide pools revealed p11–20, p81–90, and p91–100 as inducing a significant increase in Th1 immune responses compared with the negative control (**Fig. 1D** and **E**). Further breakdown into 5-peptide pools showed p11–15, p81–85, and p91–95 had a comparable significant increase in IFN- γ . Lastly, the corresponding 15 individual peptides were screened, identifying peptides p12 ($P = 0.0281$), p81 ($P = 0.0041$), p84 ($P = 0.0023$), and p91 ($P = 0.0161$) in sample 1 and likewise, p12 ($P = 0.0052$), p81 ($P = 0.0021$), p84 ($P = 0.0009$), and p91 ($P = 0.0012$) in sample 2. Similarly, p31–40, p51–60, and p81–90 demonstrated significant increases in Th1 immune responses when screening *HER3* ICD 10-peptide pools (**Fig. 1F** and **G**). Of the three donor samples used in *HER3* ICD 10-peptide pool screening, the first donor sample demonstrated an overall lack in response to pool p41–50 for both the control- and peptide-stimulated T cells (Supplementary Fig. S3A). Prior to continued screening of *HER3* ICD 10-peptide pools, all peptides within the pool p41–50 were screened individually, with p41 inducing a significant increase ($P = 0.0158$) in IFN- γ (Supplementary Fig. S3B) and was included as an epitope candidate. To validate this discovery, individual peptides from two additional unresponsive pools were screened in the ECD (Supplementary Fig. S3C–S3D) and ICD (Supplementary Fig. S3E and S3F) libraries; no other positive peptides were identified. 5-Peptide pools were then screened, which revealed p36–40, p51–55, and p86–90 as common positive pools. Thirteen peptides were then screened individually, in addition to the five individual peptides within pool p56–60, which showed a significant increase in IFN- γ in sample 3. Overall, five individual *HER3* ICD peptides demonstrated a common significant increase in IFN- γ compared with the negative control: p38 ($P = 0.0055$), p52 ($P = 0.0119$), p86 ($P = 0.0020$), and p89 ($P = 0.0028$) in sample 3 and, similarly, p38 ($P = 0.0043$), p41 ($P = 0.0124$), p52 ($P = 0.0018$), p86 ($P = 0.0016$), and p89 ($P = 0.0012$) in sample 4. Taken together, four *HER3* ECD and five *HER3* ICD peptides were identified as potential class II epitopes through the sequential peptide screening of *HER3* ECD and ICD peptide libraries: *HER3*aa56–70, *HER3*aa401–415, *HER3*aa416–430, and *HER3*aa451–465, *HER3*aa850–864, *HER3*aa865–879, *HER3*aa920–934, *HER3*aa1090–1104, and *HER3*aa1105–1119 (Supplementary Table S1). Screening schema for the ECD and ICD

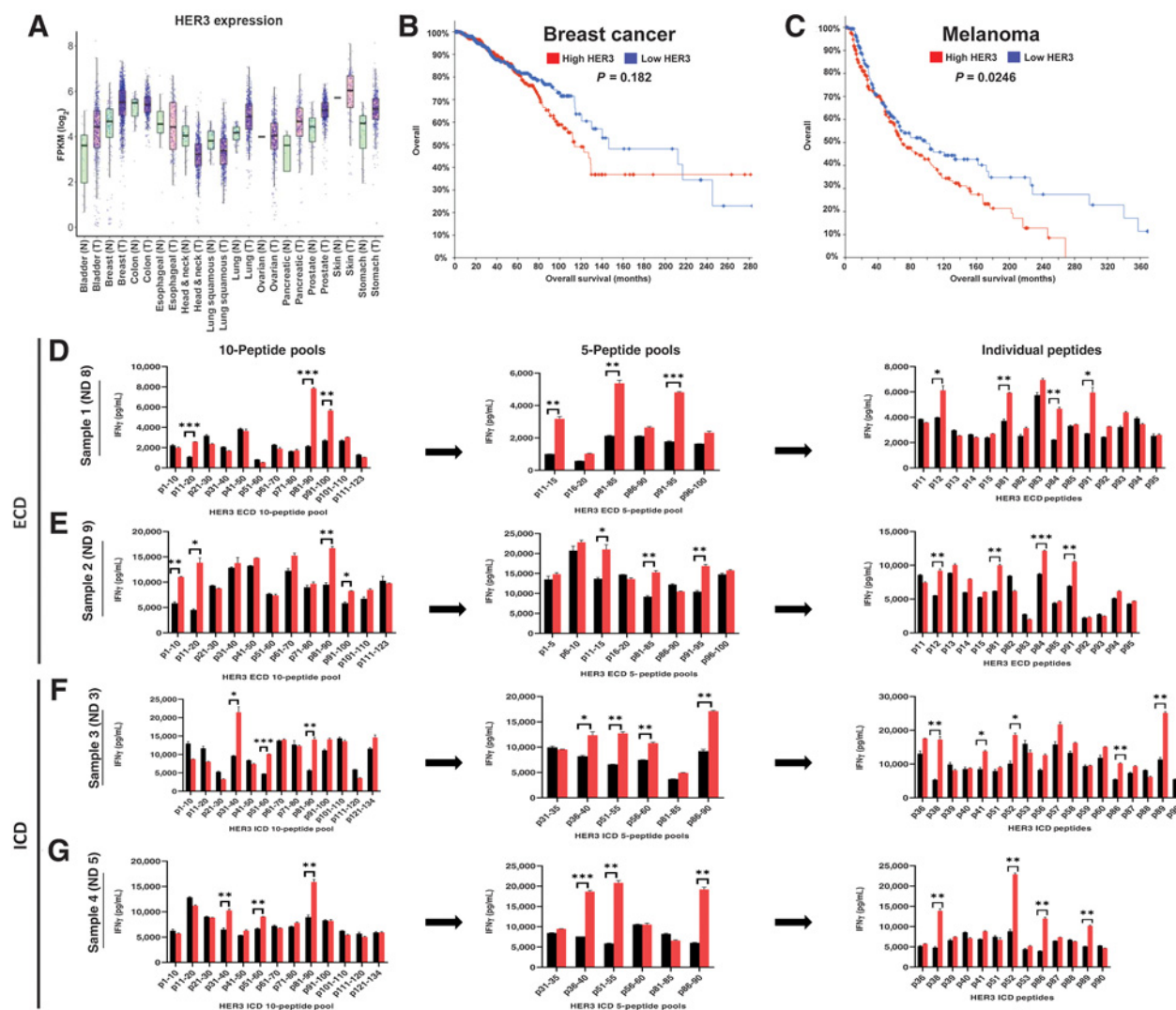


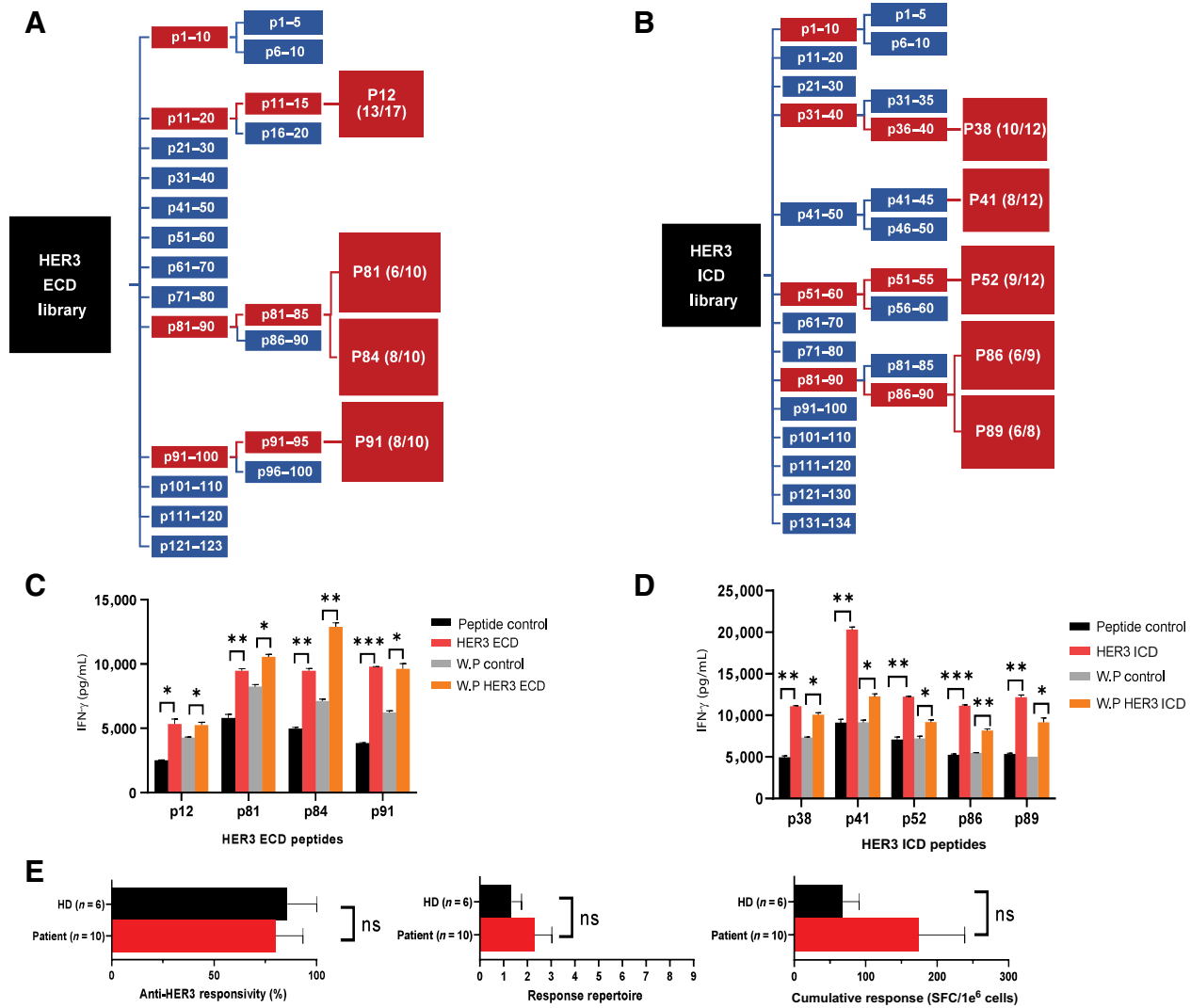
Figure 1. HER3 expression in cancer, and HER3 peptide screening of ECD and ICD class II peptide libraries. **A**, Expression of *HER3* mRNA in normal (N) and tumor (T) tissues obtained from RNA-seq data from the GDC across cancer types (see Materials and Methods). **B**, Correlation between percentage of HER3 expression and overall patient survival (in months) in breast cancer. Samples were sorted in the descending order of HER3 expression and put into two groups: high HER3 (red) and low HER3 (blue). **C**, Correlation of overall patient survival with high HER3 (red) versus low HER3 (blue) expression in melanoma. *P* value indicated in individual graphs. **D** and **E**, IFN- γ production at each screening step for sample 1 [normal donor (ND 8); **D**] and sample 2 (ND 9; **E**) when stimulated with ECD peptides. **F** and **G**, IFN- γ production at each screening step for sample 3 (ND 3; **F**) and sample 4 (ND 5; **G**) upon stimulation with ICD peptides. **D–G**, IFN- γ response to negative peptide control (black) compared with HER3 peptides (red) with an immunogenic response threshold of ≥ 1.5 -fold increase. Data represented as mean \pm SEM with statistical significance determined using a multiple *t* test without correction for multiple comparisons. Each row was analyzed individually, without assuming consistent SD. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001.

libraries at each step are shown in Fig. 2A–B. The nine HER3 class II peptides were recognized as candidate HER3 epitopes and screened across 6–13 additional samples (Supplementary Table S1) to confirm the reproducibility of the peptide-specific CD4⁺ Th1 immune responses (Supplementary Fig. S4A–S4F).

HER3 epitopes induce HER3-specific Th1 immune responses *in vitro*

To confirm the identified HER3 peptides were competent epitopes *in vitro*, HER3-peptide-primed CD4⁺ T cells were restimulated with the corresponding whole HER3 ECD or ICD protein.

HER3 ECD peptides demonstrated a significant increase in IFN- γ production when peptide-primed CD4⁺ T cells were restimulated with the matching class II peptide [(p12, *P* = 0.0181), (p81 = 0.0075), (p84, *P* = 0.0025), (p91, *P* = 0.00004)], and responses were comparable to that of restimulation with the full HER3 ECD protein domain [(p12, *P* = 0.0417), (p81, *P* = 0.0121), (p84, *P* = 0.0036), p91 (*P* = 0.0155); Fig. 2C]. HER3 ICD peptide-primed CD4⁺ Th1 cells restimulated with the matching HER3 class II peptides similarly showed a significant response compared with the peptide negative control [(p38 (*P* = 0.0012), (p41, *P* = 0.0021), (p52, *P* = 0.0035), (p86, *P* = 0.0008), (p89, *P* = 0.0021)], which was analogous


Figure 2.

HER3 class II peptides demonstrate anti-HER3 immune responses *in vitro*. **A** and **B**, ECD and ICD peptide libraries were screened sequentially in 10-peptide pools, 5-peptide pools, and individual peptides with an immunogenic response threshold of ≥ 1.5 -fold increase in IFN- γ production between peptide and control restimulated CD4⁺ T cells (see **Fig. 1**), ultimately identifying four ECD and five ICD HER3 class II peptides. Each schematic is representative of the combined responses across samples used in the peptide screening ($n = 3$), indicating reproducible significant immunogenic response compared with the class II control in ≥ 2 samples and number of donor responses in parentheses (red). **C** and **D**, Peptide-primed CD4⁺ T cells were restimulated with matching class II peptide (HER3 ECD or HER3 ICD), class II-negative control (peptide control), whole HER3 domain protein (WP HER3 ECD or WP HER3 ICD), or whole protein control (WP control). Data represented as mean \pm SEM with statistical significance determined using multiple t test without correction for multiple comparisons. Each row was analyzed individually, without assuming a consistent SD. *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. **E**, PBMCs from healthy donors (HD, black bar, $n = 6$) or breast cancer patients (Patient, red bar, $n = 10$) were individually stimulated with the nine HER3 class II peptides and analyzed via IFN- γ ELISPOT. Left, percentage of subjects responding to ≥ 1 HER3 peptide (anti-HER3 responsiveness). Middle, mean number of peptides inducing anti-HER3-specific immunity (response repertoire). Right, total IFN- γ spots (mean total SFC/1e⁶ cells) from stimulation with HER3 peptides (cumulative response). Data represented as mean \pm SEM with statistical significance determined using Mann-Whitney test. ns, not significant.

to that of peptide-primed CD4⁺ Th1 cells restimulated with the full HER3 ICD protein domain [(p38, $P = 0.0131$), (p41, $P = 0.0191$), (p52, $P = 0.0430$), (p86, $P = 0.0072$), (p89, $P = 0.0159$); **Fig. 2D**].

To determine if the identified HER3 peptides induced comparable anti-HER3 Th1 immune responses in healthy donor and breast cancer patient samples *in vitro*, PBMCs from healthy donor controls ($n = 6$) and breast cancer patients ($n = 10$) were sensitized to the nine HER3 peptides, and IFN- γ responses were evaluated by ELISPOT (**Fig. 2E**). There was no significant difference between healthy donor controls and breast cancer patients in anti-HER3 responsiveness ($P = 0.0820$),

response repertoire ($P = 0.4717$), or cumulative responses ($P = 0.3132$). Together, these results indicate CD4⁺ T cells specific to the nine HER3 class II peptide candidates could reproducibly elicit the desired HER3-specific Th1 immune response in different donor and breast cancer patient samples and were competent epitopes *in vitro*.

HER3 class II peptides represent promiscuous epitopes

To evaluate if the identified HER3 class II peptides were restricted to specific HLA alleles, donor samples used in both ECD and ICD peptide library screenings were analyzed for MHC class II HLA allele

expression (Supplementary Table S2). A total of 35 different alleles were expressed across HLA-DR (13 alleles), HLA-DP (11 alleles), and HLA-DQ (11 alleles), with minimal overlap in expression across donor samples (≤ 3 samples expressed the same HLA allele). Allele expression was cross-referenced with the corresponding HER3 ECD and HER3 ICD (Supplementary Table S3) peptide-specific Th1 immune responses. Together, these data confirmed high variability in HLA expression across samples and demonstrates the promiscuous binding of identified HER3 peptides across multiple MHC II HLA alleles, indicating the application of identified HER3 epitopes for a HER3-DC1 vaccine with capabilities for widespread application.

Class II predictive algorithm did not identify HER3 peptides as candidate epitopes

MHC II prediction algorithm, NetMHCIIpan 4.0, was used to predict class II peptides from the full HER3 protein sequence, and predicted results were compared with those obtained from the experimental peptide screening. NetMHCIIpan predicted approximately 458 class II peptide sequences (15-mer) with >700 different 9-mer binding cores exhibiting strong binding affinity (<50 nmol/L; **Table 1**). HER3 ECD p91 was the only peptide with predicted strong binding affinity with multiple HLA alleles. The remaining HER3 peptides were predicted to establish only weak binding interactions (50–500 nmol/L) with approximately 1–19 different HLA alleles. However, the MixMHCIIpred algorithm predicted a strong binding affinity for all nine HER3 peptides, with p91 showing the strongest binding affinity to most of the 38 tested HLA alleles (**Table 1**). Most of the peptides showed strong binding affinity toward a handful of HLA alleles, although each showed significant immunologic responses when tested experimentally. These results indicate the inability of the NetMHCIIpan prediction algorithm, as well as inconsistencies between algorithms, to identify the majority of HER3 class II peptides as candidate epitopes from the full HER3 protein and supports the indispensability of an empirical approach to epitope identification for tumor antigens.

Antigen-specific responses and delayed tumor growth by preventive vaccination

To investigate the antitumor immune response generated by the nine identified human HER3 peptides, we investigated both preventive and therapeutic efficacy of HER3 peptide-pulsed DC1 (HER3-DC1) in preclinical murine models. Because we observed high *HER3* expression in breast cancer and melanoma compared with normal tissues in our TCGA analysis, we chose murine mammary carcinoma models 4T1 and TUBO, which represent human TNBC and HER2^{pos} breast cancer, respectively, and the M05 murine melanoma model. Western blots showed HER3 protein expression in all three cell lines, with TUBO cells showing the highest and 4T1 showing the lowest HER3 expression (**Fig. 3A**). Immunofluorescence staining showed HER3 expression in 4T1 and TUBO cells, primarily confined to the cellular surface of both cell lines (**Fig. 3B**). In human TNBC and HER2^{pos} breast cancer cells, we observed HER3 expression, although the expression level varied considerably across cell lines (Supplementary Fig. S5A and S5B). To test the human peptides in murine models, we tested the amino acid sequence similarity between the identified human HER3 peptides and full-length murine *ErbB3* amino acid sequence using the BLASTP program. We observed 100% sequence homology between murine *ErbB3* and human HER3 peptides ECD p12, ECD p81, ECD p84, ICD p38, and ICD p86; 93% sequence homology with ICD p41, ICD p52, and ICD p89; and 87% sequence homology with ECD p91 (Supplementary Table S4).

Next, we evaluated the preventive efficacy of HER3-DC1 in BALB/c mice by vaccination with either unpulsed DC1 or HER3-DC1, followed by 4T1 or TUBO tumor challenge. To investigate immune response and antigen specificity of the response to HER3 peptides in murine models, splenocytes from vaccinated and naïve control mice were restimulated with the HER3 peptides for 72 hours *ex vivo*, followed by an IFN- γ ELISA using culture supernatants. We observed significantly higher IFN- γ secretion from HER3-DC1-vaccinated mice compared with control mice (**Fig. 3C**; Supplementary Fig. S5C). No significant difference in splenocyte IFN- γ secretion from unpulsed DC1 vaccinated mice compared with controls was observed, suggesting immune responses were specifically stimulated by HER3 peptides and was not a result of nonspecific immune stimulation by mature, nonprimed DC1 vaccination.

Similarly, IFN- γ in supernatants from cocultures of the lymph node-derived lymphocytes with DC1 pulsed with cognate HER3 peptides showed significantly increased IFN- γ secretion from HER3-DC1-vaccinated BALB/c mice compared with naïve controls (**Fig. 3D**; Supplementary Fig. S5D). These data suggest that the development of systemic immune responses in HER3-DC1-vaccinated mice is HER3 peptide-specific and not induced by unpulsed mature DC1.

Two weeks after the last vaccination, HER3-DC1-vaccinated or unvaccinated control mice were challenged with either 4T1 or TUBO tumor to compare preventive efficacy of HER3-DC1 vaccine. HER3-DC1-vaccinated mice challenged with 4T1 showed significantly delayed tumor growth ($P = 0.007$) and extended survival, whereas there was no significant difference in tumor growth and survival between control and unpulsed DC1-vaccinated mice ($P = 0.9782$; **Fig. 3E**; Supplementary Fig. S5E). Tumor growth in HER3-DC1-vaccinated mice was also significantly delayed compared with the unpulsed DC1 mice ($P = 0.0086$), suggesting the role of anti-HER3 Th1 response in preventing tumor growth over unpulsed DC1 vaccine.

In the TUBO model, we observed significantly delayed tumor growth in HER3-DC1-vaccinated mice compared with the unvaccinated controls ($P = 0.0191$; **Fig. 3F**). Despite clear preventive benefit and systemic immune response after HER3-DC1 vaccination, the NetMHCIIpan4.0 algorithm did not predict binding affinity of any of the nine HER3 peptides (Supplementary Table S5), further highlighting the limitation of false-negative predictions by algorithms. These data suggest HER3-DC1 vaccination stimulates HER3 antigen-specific immune responses and offers preventive benefit to delay tumor growth in both TNBC and HER2^{pos} breast cancer models.

Intratumoral HER3 DCs delay tumor growth and enhance immune infiltration

Next, we evaluated the therapeutic efficacy of HER3-DC1 in the aggressive murine mammary carcinoma model 4T1 (32), which mimics human TNBC. Intratumoral HER3-DC1 administration significantly delayed tumor growth compared with controls ($P = 0.0019$) and unpulsed DC1-treated mice ($P = 0.0183$), indicating therapeutic benefit of HER3-DC1 (**Fig. 4A and B**), and no significant difference was noted in tumor growth between control and unpulsed DC1-treated groups. We also observed a significant increase in survival rate in the HER3-DC1-treated group compared with control mice ($P = 0.0327$; **Fig. 4C**). Compared with the intratumoral route of delivery, subcutaneous HER3-DC administration did not show therapeutic benefit in the 4T1 model (Supplementary Fig. S5F and S5G).

We then investigated changes in the intratumoral immune landscape after intratumoral delivery of HER3-DC1, as well as in the tumor-draining lymph nodes, by flow cytometry. The flow gating

Table 1. Prediction of binding affinity from algorithms for identified HER3 peptides.

NetMHCIIpan 4.0								
p12: CEVVMGNLEIVLTGH			p91: AGRYISANRQLCYH			p38: VADFGVADLLPPDDK		
Allele	Core	Affinity (nmol/L)	Allele	Core	Affinity (nmol/L)	Allele	Core	Affinity (nmol/L)
DRB1_0101	MGNLEIVLT	314.72	DRB1_0101	IYISANRQL	25.22	DPA10202-DPB10301	FGVADLLPP	485.34
DRB1_0102	MGNLEIVLT	77.8	DRB1_0102	IYISANRQL	8.69	DQA10505-DQB10202	ADFGVADLL	296.18
DRB1_0103	VVMGNLEIV	252.16	DRB1_0103	IYISANRQL	60.96	DQA10505-DQB10301	ADFGVADLL	297.42
DRB1_1201	MGNLEIVLT	205.35	DRB1_0301	ISANRQLCY	82.01	p41: QLLYSEAKTIKWKMA		
DRB1_1301	VVMGNLEIV	395.9	DRB1_0401	IYISANRQL	249.1	Allele	Core	Affinity (nmol/L)
DPA10103-DPB10301	GNLEIVLTG	287.41	DRB1_0404	IYISANRQL	223.89	DRB1_0101	YSEAKTIK	84.71
DPA10103-DPB10601	VMGNLEIVL	406.22	DRB1_0405	IYISANRQL	224.29	DRB1_0102	YSEAKTIK	53.61
DPA10103-DPB11701	VMGNLEIVL	405.89	DRB1_0408	IYISANRQL	351.53	DRB1_0103	YSEAKTIK	265.34
DPA10201-DPB10301	GNLEIVLTG	265.48	DRB1_0701	IYISANRQL	36.7	DRB1_0401	YSEAKTIK	291.56
DPA10201-DPB10601	GNLEIVLTG	347.25	DRB1_0801	YISANRQLC	61.9	DRB1_0405	YSEAKTIK	499.3
DPA10201-DPB11701	VMGNLEIVL	351.93	DRB1_1001	IYISANRQL	159.52	DRB1_0408	YSEAKTIK	481.44
DPA10202-DPB10201	VMGNLEIVL	466.96	DRB1_1101	YISANRQLC	51.11	DRB1_0701	YSEAKTIK	164.51
DPA10202-DPB10301	GNLEIVLTG	163.68	DRB1_1104	IYISANRQL	13.08	DRB1_0801	YSEAKTIK	241.93
DPA10202-DPB10601	GNLEIVLTG	231.07	DRB1_1201	IYISANRQL	6.48	DRB1_1001	YSEAKTIK	414.37
DPA10202-DPB11701	GNLEIVLTG	241.87	DRB1_1301	IYISANRQL	39.63	DRB1_1101	YSEAKTIK	266.23
DPA10202-DPB12001	GNLEIVLTG	342.06	DRB1_1303	IYISANRQL	18.55	DRB1_1104	LYSEAKTI	108.79
DQA10505-DQB10301	VVMGNLEIV	477.04	DRB1_1501	IYISANRQL	87.9	DRB1_1201	LYSEAKTI	134.48
p81: SWPPHMHNFVFSNL			DRB1_1601	YISANRQLC	266.47	DRB1_1303	LYSEAKTI	113.19
Allele	Core	Affinity (nmol/L)	DRB3_0202	IYISANRQL	69.67	DRB3_0202	YSEAKTIK	272.05
DRB1_0101	HMHNFVFS	273.57	DRB4_0101	IYISANRQL	233.16	DRB5_0101	YSEAKTIK	47.08
DRB1_0102	MHNFVFSN	146.74	DRB4_0103	IYISANRQL	233.16	DRB5_0202	YSEAKTIK	64.72
DRB1_0405	MHNFVFSN	425.91	DRB5_0101	IYISANRQL	44.32	DPA10103-DPB10301	SEAKTIKW	333.38
DRB1_0701	MHNFVFSN	452.1	DRB5_0202	IYISANRQL	14.96	DPA10201-DPB10301	SEAKTIKW	346.25
DRB1_1104	MHNFVFSN	394.75	DPA10103-DPB10201	ISANRQLCY	455.66	DPA10202-DPB10301	LYSEAKTI	345.3
DRB1_1201	MHNFVFSN	168.11	DPA10103-DPB10301	ISANRQLCY	80.12	DQA10505-DQB10301	YSEAKTIK	487.56
DRB1_1303	HMHNFVFS	301.54	DPA10103-DPB10601	ISANRQLCY	197.88	p52: VPDLLEKGERLAQPQ		
DRB1_1501	MHNFVFSN	498.15	DPA10103-DPB11101	ISANRQLCY	452.08	Allele	Core	Affinity (nmol/L)
DRB5_0202	HMHNFVFS	384.57	DPA10103-DPB11701	ISANRQLCY	239.62	DRB1_0102	DLLEKGERL	233.55
DPA10103-DPB10201	HNFSVFSNL	375.28	DPA10103-DPB12001	ISANRQLCY	153.46	DRB5_0202	DLLEKGERL	444.92
DPA10103-DPB10301	HNFSVFSNL	200.28	DPA10201-DPB10201	ISANRQLCY	450.12	p86: GCLASESSEGHVTGS		
DPA10103-DPB10601	HNFSVFSNL	335.46	DPA10201-DPB10301	ISANRQLCY	88.26	Allele	Core	Affinity (nmol/L)
DPA10103-DPB11701	HNFSVFSNL	421.32	DPA10201-DPB10601	ISANRQLCY	187.3	DQA10505-DQB10301	SESSEGHVT	229.07
DPA10103-DPB12001	HNFSVFSNL	374.2	DPA10201-DPB11101	ISANRQLCY	423.49	p89: EAELQEKVSMCRSRS		
DPA10201-DPB10201	HNFSVFSNL	451.48	DPA10201-DPB11701	ISANRQLCY	220.75	Allele	Core	Affinity (nmol/L)
DPA10201-DPB10301	HNFSVFSNL	255.55	DPA10201-DPB12001	ISANRQLCY	155.54	DRB1_0102	LQEKVSMCR	235.64
DPA10201-DPB10601	HNFSVFSNL	425.74	DPA10202-DPB10201	ISANRQLCY	428.27	DRB1_0801	KVSMCRSRS	250.44
DPA10201-DPB12001	HNFSVFSNL	462.6	DPA10202-DPB10301	ISANRQLCY	78.91	DRB1_1101	KVSMCRSRS	330.87
DPA10202-DPB10301	HNFSVFSNL	318.24	DPA10202-DPB10601	ISANRQLCY	162.84	DRB1_1104	KVSMCRSRS	106.56
P84: TTIGGRSLYNRGFSL			DPA10202-DPB11101	ISANRQLCY	330.55	DRB1_1201	KVSMCRSRS	214.85
Allele	Core	Affinity (nmol/L)	DPA10202-DPB11701	ISANRQLCY	212.1	DRB1_1303	LQEKVSMCR	339.59
DRB1_0101	SLYNRGFSL	300.1	DPA10202-DPB12001	ISANRQLCY	139.22	DRB5_0101	LQEKVSMCR	249.76
DRB1_0102	SLYNRGFSL	182.32	DQA10505-DQB10301	AGRYISAN	317.66	DRB5_0202	LQEKVSMCR	276.61
DRB1_0701	SLYNRGFSL	477.72	NOTE: 15-mer sequences with selected 9-mer binding core for predicted HLA class II allelic binding affinity.					
DRB1_1104	IGGRSLYNR	265.23	Affinity (nmol/L) <50 nm: strong binding.					
DRB1_1201	SLYNRGFSL	229.23	Affinity (nmol/L) 50–500 nm: weak binding.					
DRB1_1303	SLYNRGFSL	215.23						
DRB5_0202	SLYNRGFSL	207.95						
DQA10505-DQB10301	TIGGRSLYN	309.62						
MixMHCIIpred								
Peptide	BestAllele		%Rank_best (12–25aa)					
p12 CEVVMGNLEIVLTGH	DRB3_02_02		11					
p81 SWPPHMHNFVFSNL	DRB1_10_01		5.28					
p84 TTIGGRSLYNRGFSL	DRB1_12_01		3.4					
p91 AGRYISANRQLCYH	DPA1_01_03 DPB1_03_01		0.163					
p38 VADFGVADLLPPDDK	DPA1_01_03 DPA1_02_01 DPB1_11_01		1.64					
p41 QLLYSEAKTIKWKMA	DRB1_03_01		4.27					
p52 VPDLLEKGERLAQPQ	DRB1_08_01		1.21					
p86 GCLASESSEGHVTGS	DRB1_04_04		11.3					
p89 EAELQEKVSMCRSRS	DPA1_01_03 DPB1_105_01 DPB1_126_01		2.03					

Top, Class II-binding affinity predictions (nmol/L) from NetMHCIIpan 4.0 algorithm for nine identified HER3 peptides. Bottom, Class II-binding affinity predictions from MixMHCIIpred algorithm for nine identified HER3 peptides. The “BestAllele” column shows specific HLA allele(s) that had the best binding prediction. The “%Rank_best (12–25aa)” column shows the percentile rank among all randomly generated peptide sizes 12–25 amino acid; best score = 0, worst score = 100.

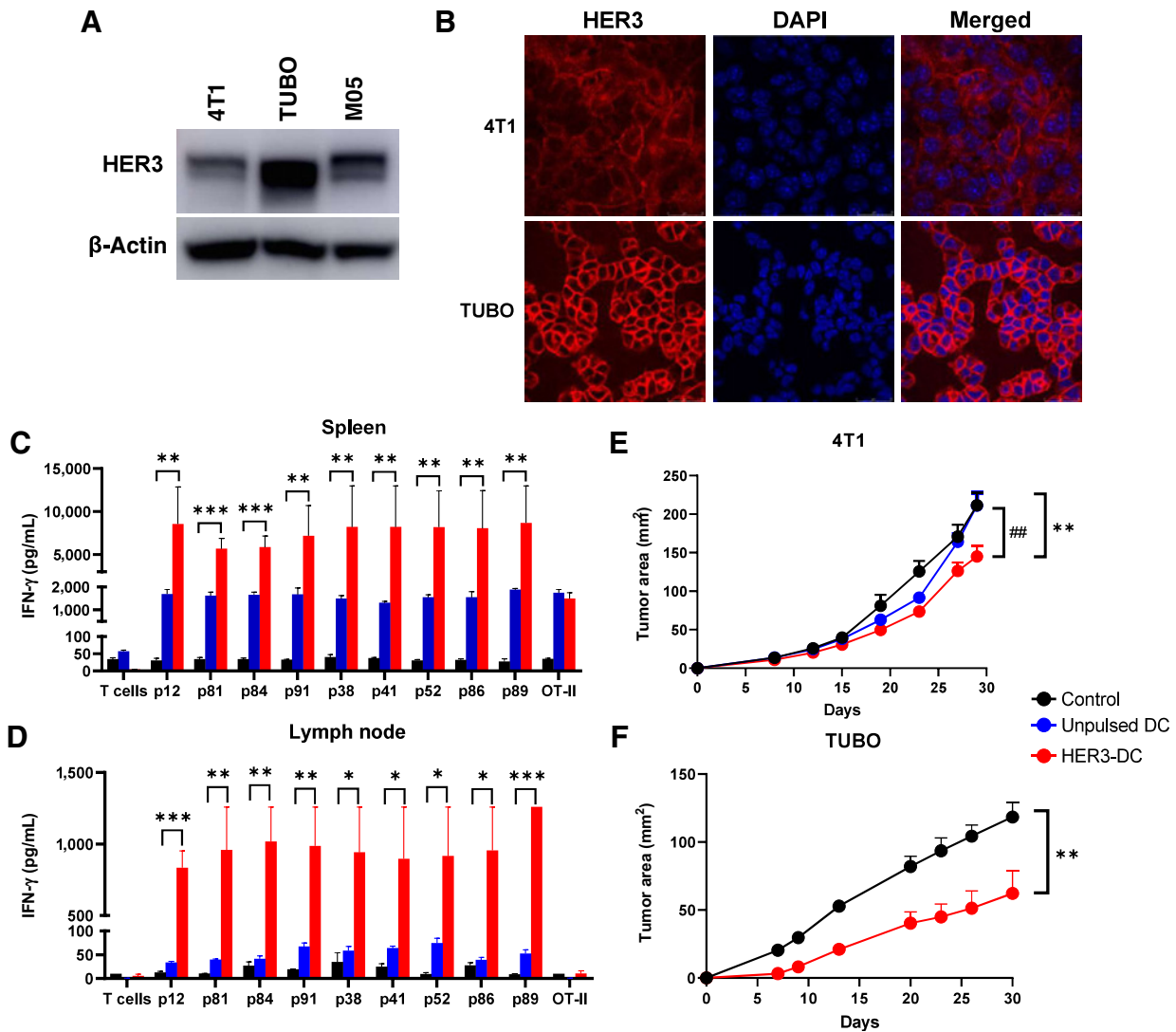
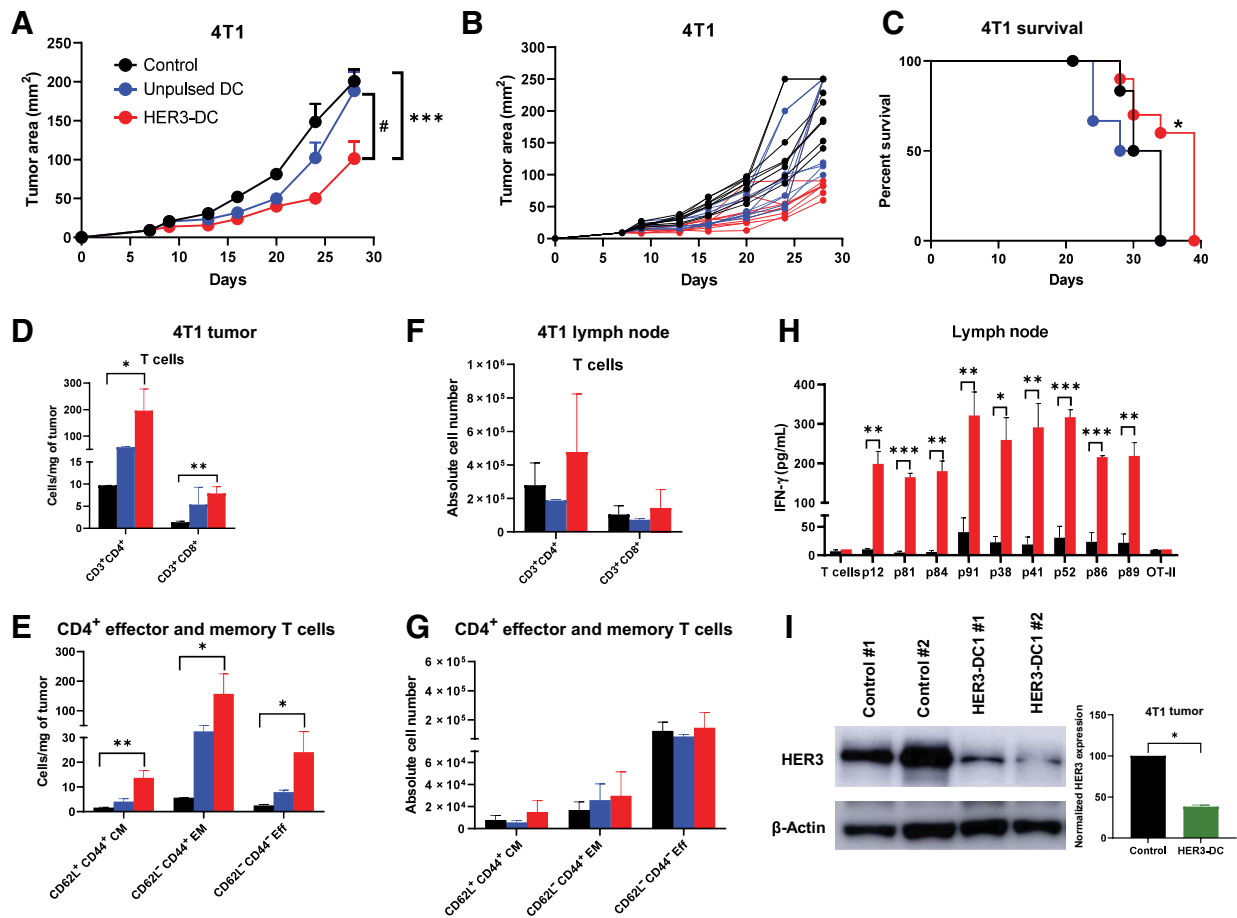


Figure 3. HER3-DC1 vaccination elicits peptide-specific immune response and delays tumor growth. **A**, Immunoblotting of murine tumor cell lines 4T1, TUBO, and M05 to detect HER3. β -Actin: loading control. **B**, Immunofluorescence for HER3 (red) and nucleus (DAPI, blue) in 4T1 and TUBO murine mammary tumor cells (image magnification: 1,200 \times). **C** and **D**, Individual HER3 peptide-specific immune responses in spleens (**C**) and lymph node-derived immune cells (**D**) from control (black), unpulsed mature DC1 (blue), and HER3-DC1 (red) vaccinated BALB/c mice ($n = 3$). Spleens were processed, and splenocytes were restimulated with the HER3 peptides for 72 hours to detect IFN- γ by ELISA. Lymph node-derived lymphocytes were cocultured with DC1 pulsed with individual HER3 peptides for 72 hours to detect IFN- γ by ELISA. **E**, Tumor growth after 4T1 tumor challenge in control (black), unpulsed mature DC1 (blue), and HER3-DC1 (red) vaccinated mice ($n = 7-10$ mice/group). Mice were challenged 2 weeks after the last vaccination and were monitored until endpoint. *, control versus HER3-DC1; #, unpulsed DC1 versus HER3-DC1. **F**, TUBO tumor growth in control (black) and HER3-DC1 (red) vaccinated mice ($n = 7-10$ mice/group). Mice were challenged 2 weeks after the last vaccination. Data represented as mean \pm SEM with statistical significance determined using multiple t test without correction for multiple comparisons. Each row analyzed individually, without assuming a consistent SD. *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$; ##, $P \leq 0.01$.

strategy to identify CD4⁺ and CD8⁺ TILs and their subtypes is shown in Supplementary Fig. S6. Flow analysis revealed significantly higher CD4⁺ ($P = 0.03$) and CD8⁺ ($P = 0.0042$) T-cell infiltration per milligram of tumor in HER3-DC1-treated mice compared with control tumors (Fig. 4D; Supplementary Fig. S7A). We observed a significant increase in the CD62L⁻CD44⁺ effector memory (EM, $P = 0.0304$), CD62L⁺CD44⁺ central memory (CM, $P = 0.0057$), and CD62L⁻CD44⁻ effector ($P = 0.0412$) T-cell populations in HER3-DC1-treated tumors compared with controls (Fig. 4E; Supplementary Fig. S7B). We then compared immune phenotypes of cells

isolated from the draining lymph nodes of the HER3-DC1 versus control mice and observed no statistically significant increases in total CD4⁺ ($P = 0.6492$) and CD8⁺ ($P = 0.7699$) populations in the HER3-DC1 group compared with controls (Fig. 4F), and no statistically significant increases in CD4⁺ CM, EM, and effector T-cell abundance in the HER3-DC1 group compared with the controls was seen (Fig. 4G).

To evaluate systemic immune responses after intratumoral HER3-DC1 administration, lymph nodes harvested from control and treated mice were cocultured with DC1 cells pulsed with individual HER3


Figure 4.

Intratumoral HER3-DC1 administration elicits peptide-specific immune responses and delays tumor growth. **A**, Tumor growth in the 4T1 murine mammary carcinoma model. BALB/c mice bearing subcutaneous 4T1 tumors received either intratumoral PBS (black), unpulsed mature DC1 (blue), or HER3 peptide-pulsed DC1 (red; $n = 10$ mice/group), starting on day 7 when tumors were palpable. Tumor growth was monitored until endpoint and was compared between control and HER3-DC1, as well as between unpulsed DC1 and HER3-DC1. *, control versus HER3-DC1; #, unpulsed DC1 versus HER3-DC1. **B**, Individual tumor growth for each mouse from control (black)-, unpulsed DC1 (blue)-, and HER3-DC1 (red)-treated groups. **C**, Percent survival in the 4T1 mouse model. Control, black; unpulsed DC1, blue; HER3-DC1, red. **D**, Intratumoral CD3⁺CD4⁺ and CD3⁺CD8⁺ T-cell infiltration per milligram of tumor in control (black)-, unpulsed DC1 (blue)-, and HER3-DC1 (red)-treated mice. Absolute number of immune cells was compared between control and HER3-DC1 groups. **E**, Frequency of CD62L⁺CD44⁺ central memory (CM), CD62L⁺CD44⁺ effector memory (EM), and CD62L⁻CD44⁻ effector (Eff) T-cell populations within intratumoral CD4⁺ cells between control- (black) and HER3-DC1-treated (red) tumors. The unpulsed DC1 (blue) group was not included in any statistical analyses. **F**, Absolute number of CD3⁺CD4⁺ and CD3⁺CD8⁺ T cells in lymph nodes of control (black)-, intratumoral unpulsed DC1 (blue)-, and HER3-DC1 (red)-treated mice. Cell numbers were compared in control versus HER3-DC1 groups. Data shown are the representative from three independent experiments. **G**, Absolute numbers of CD4⁺ CM, EM, and Eff T-cell populations in lymph nodes of control (black), unpulsed DC1 (blue), and HER3-DC1 (red) mice. Data shown are the representative from three independent experiments. **H**, Lymphocytes from the lymph nodes of control and treated mice were cocultured with DC1 pulsed with individual HER3 or OT-II (negative control) peptides. Culture supernatants were collected after 72 hours, and IFN- γ was measured by ELISA (control: black bar; HER3-DC1: red bar). **I**, Total protein isolated from *in vivo* tumor samples was analyzed by Western blotting to compare HER3 protein expression after intratumoral HER3-DC1 (green) administration with respect to the control (black). β -Actin: loading control. Data represented as mean \pm SEM with statistical significance determined using multiple t test without correction for multiple comparisons. Each row was analyzed individually, without assuming a consistent SD. A log-rank (Mantel-Cox) test was used to determine differences between the survival curves. Unpaired two-tailed t test was performed to analyze Western blot data. *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$; #, $P \leq 0.01$.

peptides or OT-II (as negative control) for 72 hours. ELISA with culture supernatants revealed significantly higher IFN- γ secretion in the HER3-DC1 group, compared with the control, for all nine peptides (Fig. 4H). Lastly, we investigated HER3 protein expression from *in vivo* tumor samples by immunoblotting to determine if intratumoral DC administration altered receptor protein expression in tumor cells. Total HER3 protein expression was significantly reduced in HER3-DC1-treated tumors compared with control tumors ($P = 0.0369$; Fig. 4I), suggesting molecular cross-talk between HER3-

DC1-induced immune responses and oncodriver signaling can affect receptor protein expression in 4T1 tumor cells.

HER3-DC1 delays tumor growth and enhances immune infiltration in the TUBO model

HER3 is the most potent dimerization partner of HER2 that facilitates downstream signaling, contributing to the growth and proliferation of tumor cells. Hyperactivation of HER3 has also been identified as one of the primary mechanisms of therapeutic resistance

to HER2-targeted therapies in HER2^{POS} breast cancer (33, 34). Therefore, we investigated the therapeutic efficacy of HER3-DC1 in a HER2-resistant TUBO murine mammary tumor model (35). We observed significantly delayed tumor growth in mice that received intratumoral HER3-DC1 compared to control- ($P = 0.0028$) and unpulsed DC1-treated mice ($P = 0.0185$). Tumor regression occurred in ~30% of HER3-DC1-treated mice, and growth was significantly delayed in the remaining mice, compared with the controls (Fig. 5A; Supplementary

Fig. S8A). This resulted in a significant improvement in survival of HER3-DC1-treated mice compared with the control (median survival 64 days vs. 46 days, $P = 0.0002$) and unpulsed DC1-treated groups (median survival 64 days vs. 49 days, $P = 0.0015$; Fig. 5B).

To evaluate immune infiltration into the tumor microenvironment after HER3-DC1 intratumoral delivery, we collected tumors from control- and HER3-DC1-treated mice and analyzed immune cell abundance by flow cytometry. Compared with control tumors, we

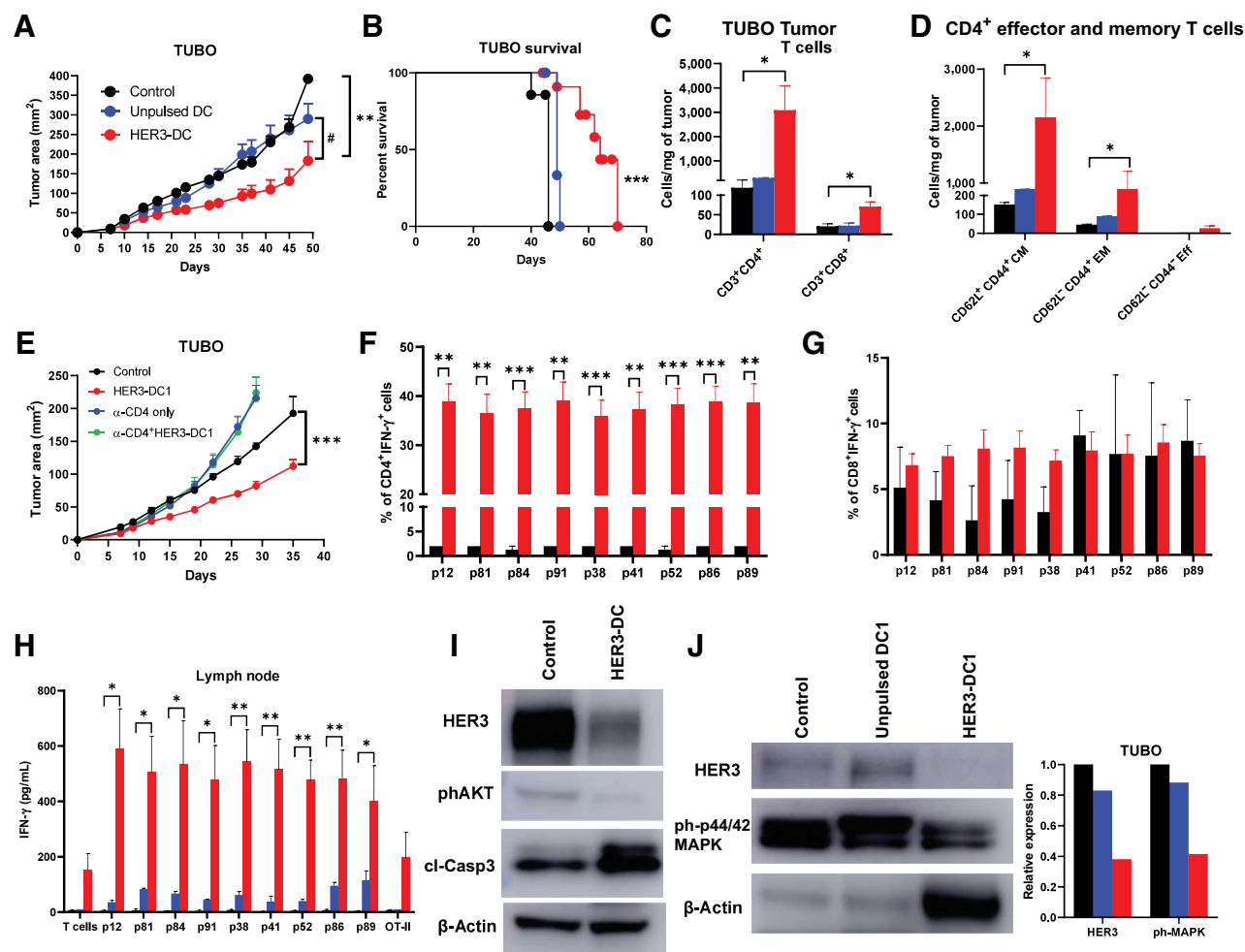


Figure 5.

Intratumoral HER3-DC1 delays tumor growth and enhances immune infiltration in a HER2^{POS} TUBO therapeutic model in a CD4-dependent manner. **A**, Tumor growth in the TUBO murine mammary carcinoma model. BALB/c mice were injected with TUBO tumor cells, and on day 7, mice received either PBS control (black), unpulsed mature DC1 (blue), or HER3-DC1 (red) intratumorally once weekly for six doses ($n = 10$ mice/group). Tumor growth was monitored until endpoint and compared in control versus HER3-DC1 (+) and unpulsed DC1 versus HER3-DC1 (#) mice. **B**, Percent survival in TUBO mouse model. Control: black; unpulsed DC1: blue; HER3-DC1: red. **C**, CD3⁺CD4⁺ and CD3⁺CD8⁺ T cells per milligram of tumors from mice (**A**) after intratumoral DC injection was compared between control (black) and HER3-DC1 (red) groups. No statistical analyses were performed for the unpulsed DC1 (blue) mice ($n = 3$ /group). **D**, Abundance of CD4⁺ central memory (CD62L⁺CD44⁺ CM), effector memory (CD62L⁻CD44⁺ EM), and effector (CD62L⁻CD44⁻ Eff) T cells in control (black) versus HER3-DC1 mice (red) per milligram of tumor tissue. Data shown are representative from three independent experiments. **E**, Tumor growth of TUBO tumors after CD4 depletion. BALB/c mice were injected with anti-CD4 antibodies 3 days before subcutaneous TUBO tumor injection. When tumors were palpable, mice received either PBS control (black), intratumoral HER3-DC1 once weekly (red) for six doses, CD4 depletion antibody alone (blue; continued twice weekly until endpoint), or HER3-DC1 (green) with CD4 depletion. Tumor growth was monitored until endpoint. **F** and **G**, Percentage of CD4⁺IFN- γ ⁺ (**F**) and CD8⁺IFN- γ ⁺ (**G**) TILs in the tumors from control (black) versus HER3-DC1 (red) mice from **E**. **H**, Coculture of the lymph node immune cells with HER3 peptide-pulsed DC1 for 72 hours to detect IFN- γ via ELISA. Control: black bar; unpulsed DC1: blue bar; HER3-DC1: red bar. **I**, Western blot for HER3, phosphorylated AKT (pAkt), and cleaved caspase-3 (clCasp-3) with total protein isolated from control- and HER3-DC1-treated TUBO tumors. β -Actin: loading control. **J**, Western blot for HER3 and phosphorylated p44/42 MAPK (ph-p44/42 MAPK) from control-, unpulsed DC1-, and HER3-DC1-treated TUBO tumors. β -Actin: loading control. Data represented as mean \pm SEM with statistical significance determined using multiple t test without correction for multiple comparisons. Each row was analyzed individually, without assuming a consistent SD. A log-rank (Mantel-Cox) test was used to determine differences between the survival curves. *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$; #, $P \leq 0.01$.

observed significantly higher infiltration of CD4⁺ ($P = 0.0449$) and CD8⁺ ($P = 0.0201$) T cells in tumors from HER3-DC1-treated mice (Fig. 5C; Supplementary Fig. S8B). Within the CD4⁺ population, we noted significantly amplified abundance of CD62L⁺CD44⁺ CM ($P = 0.0446$) and CD62L⁻CD44⁺ EM ($P = 0.0458$), but not CD62L⁻CD44⁻ effector T cells (Fig. 5D). We also observed significantly increased CD8⁺ T cells, specifically CD62L⁺CD44⁺ CM ($P = 0.0131$), and CD62L⁻CD44⁺ EM T cells ($P = 0.033$) in tumors from HER3-DC1-treated mice compared with the controls (Supplementary Fig. S8C). When CD4⁺ T cells were selectively depleted, tumor growth was comparably faster than the control group, and the antitumor effect of HER3-DC1 was completely abolished (Fig. 5E), highlighting the necessity of CD4⁺ T cells for functional activity of the class II peptide-pulsed HER3-DC1. When TILs from these mice were cocultured with DC1 pulsed with individual HER3 peptides, a significantly higher percentage of CD4⁺IFN- γ ⁺ cells in HER3-DC1-treated TILs were noted in response to each of the HER3 peptides compared with the untreated controls. These data indicate HER3-specific IFN- γ secretion by CD4⁺ T cells following HER3-DC1 administration (Fig. 5F; Supplementary Fig. S8D). On the contrary, no statistically significant difference was noted in the percentage of CD8⁺IFN- γ ⁺ cells between groups (Fig. 5G). In HER3-DC1-treated lymph nodes, we observed minimal, but not statistically significant, increase in the number of CD4⁺ and CD8⁺ T cells (Supplementary Fig. S8E). Together, these data suggest intratumoral HER3-DC1 can stimulate CD4⁺ immune responses, which, in turn, can activate CD8⁺ T cells.

Coculture of the lymph node immune cells with HER3 peptide-pulsed DC1 for 72 hours confirmed antigen specificity, with significantly augmented IFN- γ secretion in the HER3-DC1-treated group compared with controls (Fig. 5H). Similarly, elevated IFN- γ secretion was noted when HER3-DC1 splenocytes were restimulated with HER3 peptides, compared with the controls (Supplementary Fig. S8F).

Next, we examined whether HER3-DC1 intratumoral administration alters the molecular identity of TUBO tumor cells. Western blots on tumor cell lysates revealed downregulation of HER3 protein in HER3-DC1-treated tumors compared with controls. We also observed diminished phosphorylated AKT in HER3-DC1 tumors, along with an upregulation of apoptosis marker, cleaved caspase-3, compared to the control tumors (Fig. 5I). Downregulation of phosphorylated p44/42 MAPK in HER3-DC1 tumors was also noted, whereas no difference in HER3 and phospho-p44/42MAPK expression was observed in tumors from unpulsed DC1-treated mice compared to the controls (Fig. 5J). These data suggest HER3-DC1-induced Th1 immune responses can modulate molecular signaling to interfere with onco-driver expression and activation and induce tumor cell apoptosis.

HER3-DC1 vaccination prevents tumor development in the M05 model

Because TCGA and cBioPortal analyses indicated high HER3 expression in skin cutaneous melanoma compared with normal tissue and a negative correlation between high HER3 expression and overall patient survival, we tested the preventive and therapeutic efficacy of HER3-DC1 in an M05 murine melanoma model, derived from the original B16 melanoma cell line transfected with the ovalbumin gene (36). Immunoblotting of M05 cell lysates showed moderate HER3 protein expression (Fig. 3B), whereas substantial surface expression of HER3 was detected by immunofluorescence (Fig. 6A). Therefore, we tested the preventive efficacy of HER3-

DC1 in C57BL/6 mice. Compared with controls, splenocytes from HER3-DC1-vaccinated mice showed significantly enhanced peptide-specific immune responses when restimulated with HER3 peptides (Fig. 6B). Similarly, when immune cells isolated from the lymph nodes of vaccinated mice were cocultured with HER3 peptide-pulsed DC1 *ex vivo*, we observed significantly higher IFN- γ secretion compared with naïve controls (Fig. 6C). When naïve and vaccinated mice were challenged with M05 tumor 2 weeks after the last vaccination, tumor growth was significantly delayed, and survival was improved in vaccinated mice compared with naïve controls ($P = 0.0004$; Fig. 6D; Supplementary Fig. S8G). These data suggest HER3-DC1 vaccination induces peptide-specific, systemic immune responses in C57BL/6 mice.

Intratumoral HER3-DC1 diminishes tumor growth in the M05 melanoma model

Next, we investigated the therapeutic efficacy of intratumoral HER3-DC1 treatment in a subcutaneous M05 murine melanoma model. An OT-II-pulsed DC1 group was included as a positive control, since the M05 cell line expresses ovalbumin. Compared with controls and unpulsed DC1-treated mice, HER3-DC1 treatment significantly delayed tumor growth (Fig. 6E; control vs. HER3-DC1, $P \leq 0.000001$; unpulsed DC1 vs. HER3-DC1, $P < 0.0001$). HER3-DC1 treatment was comparable with OT-II-DC1 treatment in terms of tumor growth and tumor regression. We observed tumor regression in 20% of HER3-DC1-treated mice, whereas tumor growth was significantly slower in 40% of the remaining mice, with a steady growth in the rest. This resulted in significantly improved survival in HER3-DC1-treated mice compared with controls ($P \leq 0.0001$, median survival control = 25 days, HER3-DC1 = undefined; Fig. 6F). We next assessed how intratumoral HER3-DC1 treatment modulated the tumor immune landscape. Like the murine mammary carcinoma models, we observed significantly increased CD4⁺ T-cell infiltration in HER3-DC1 mice compared with the controls ($P = 0.0339$). A significantly enhanced CD8⁺ T-cell infiltration was also noted in the HER3-DC1-treated group, which could contribute to the antitumor efficacy of the intratumoral HER3-DC1 observed ($P = 0.0079$; Fig. 6G). For CD4⁺ T-cell subsets, EM ($P = 0.0208$) and CM ($P = 0.0199$) populations were significantly enhanced in the HER3-DC1 group, with no differences in effector T-cell populations (Fig. 6H).

We investigated the systemic immune responses mediated by intratumoral HER3-DC1 treatment. Lymph node immune cells from control and treated mice were assessed for lymphoid markers. As previously seen in the breast cancer models, we observed significantly enhanced CD4⁺ T cells in HER3-DC1-treated mice ($P = 0.0085$), which was comparable to the positive control OT-II-DC1 group; however, no significant difference was noted in the frequency of CD8⁺ T cells (Fig. 6I). Within the CD4⁺ population, we noted significantly increased EM ($P = 0.0058$), CM ($P = 0.0052$), and effector ($P = 0.0016$) T cells (Fig. 6J). We evaluated IFN- γ secretion by immune cells residing in the lymph node of treated and control mice. As previously seen in the breast cancer models, we noted significantly enhanced IFN- γ secretion in the HER3-DC1 group compared with the control, when lymph node immune cells were cocultured with HER3 peptide-pulsed DCs (Fig. 6K). In the OT-II-DC1-treated group, a significant positive response was noted only when cocultured with the OT-II peptide-pulsed DC1 ($P = 0.00567$), but not the HER3 peptides, indicating the antigen specificity of the response. We also detected significantly higher IFN- γ secretion from splenocytes isolated from the

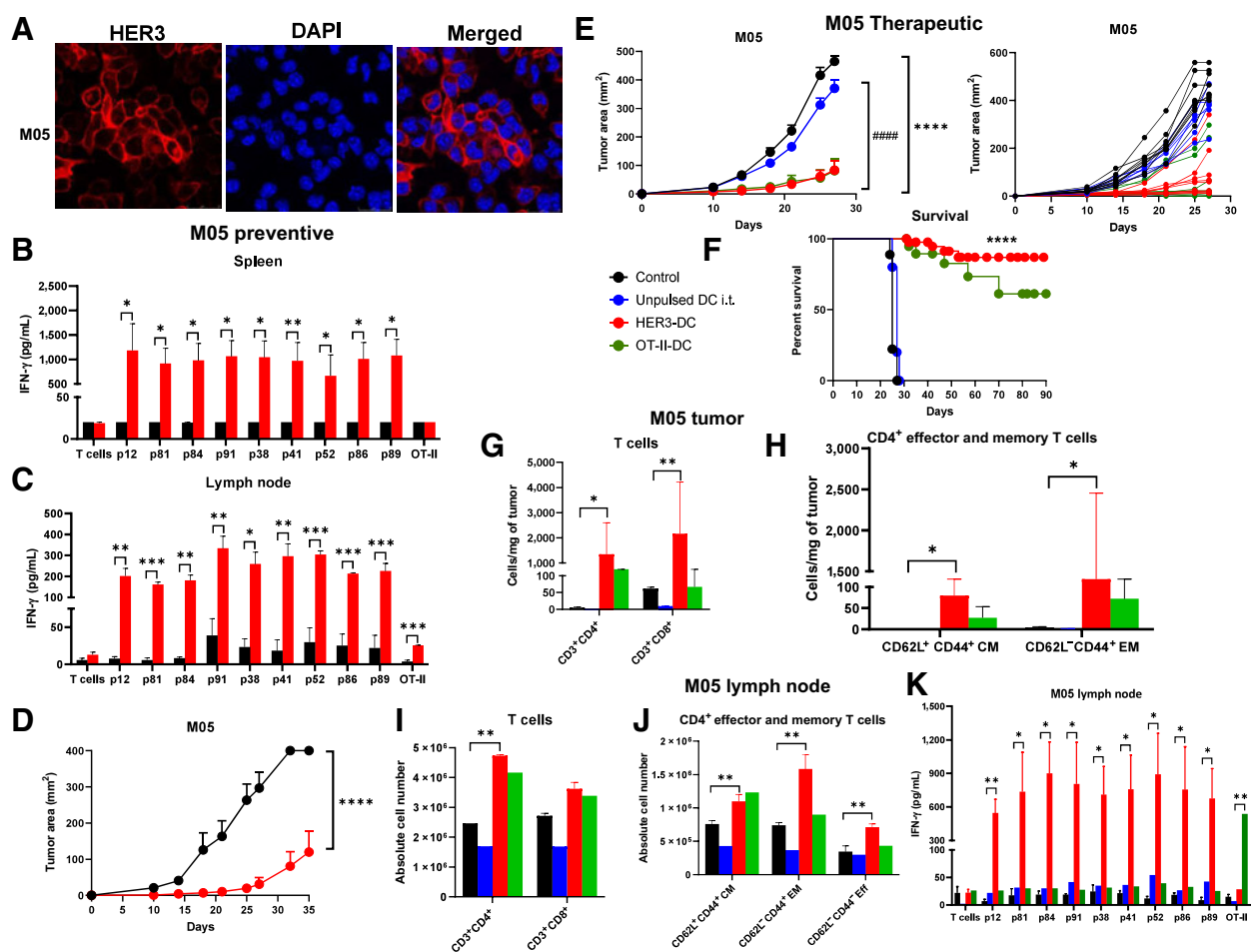


Figure 6. Vaccination prevents tumor development in a preventive HER3⁺ melanoma model and diminishes tumor growth and improves survival in a therapeutic HER3⁺ melanoma model. **A**, Immunofluorescence for HER3 (red) surface expression in M05 cells. DAPI (blue): nucleus (image magnification: 1,200×). **B**, HER3 peptide-specific immune responses ($n = 3-5$ /group). Splenocytes from control (black) and vaccinated (red) mice were restimulated with individual HER3 peptides for 72 hours to detect IFN- γ by ELISA. **C**, Lymph node lymphocytes from control (black) and HER3-DC1 (red) vaccinated mice were cocultured with individual HER3 peptide-pulsed DCs to detect antigen-specific immune response in HER3-DC1 vaccinated mice by ELISA. **D**, Preventive model: Two weeks after the last vaccine, C57BL/6 mice ($n = 10$ mice/group) were challenged with M05 tumor cells. Tumor growth was monitored until endpoint in control (black) versus HER3-DC1 vaccinated (red) mice. **E**, Therapeutic setting: C57BL/6 mice were injected subcutaneously with the M05 murine melanoma cells in the left flank, and upon palpable tumor formation on day 10, mice were randomized into four groups ($n = 10$ mice/group). Tumor growth was monitored in mice receiving PBS (black), unpulsed DC1 (blue), HER3-DC1 (red), and OT-II peptide-pulsed DC1 (green) intratumorally once weekly for 6 weeks. Tumor growth was compared between control and HER3-DC1 (+) and unpulsed DC1 vs. HER3-DC1 (#) groups. Individual tumor growth curve for each mouse shown on the right. **F**, Percent survival in M05 mouse model for control (black)-, unpulsed DC1 (blue)-, HER3-DC1 (red)-, and OT-II-DC1 (green)-treated mice. **G**, Intratumoral infiltration of CD3⁺CD4⁺ and CD3⁺CD8⁺ T cells was compared between control (black) and HER3-DC1 (red) mice from **E**. **H**, CD4⁺ central memory (CD62L⁺CD44⁺ CM) and effector memory (CD62L⁻CD44⁺ EM) T-cell infiltration per milligram of tumors in control (black) versus HER3-DC1 (red) mice from **E**. **I**, Absolute number of CD3⁺CD4⁺ and CD3⁺CD8⁺ T cells in the lymph nodes of control- versus HER3-DC1-treated mice from **E**. **J**, Abundance of CD4⁺ CM, EM, and effector (Eff) T cells in the lymph nodes of control versus treated mice. For **G-J**, Unpulsed DC1 (blue) and OT-II-DC1 (green) groups were not included in the statistical analyses. **K**, Lymph node lymphocytes from control and treated mice (**E**) were cocultured with individual HER3 peptide-pulsed DC1 for 72 hours to detect IFN- γ by ELISA. Responses were compared between control and HER3-DC1 groups for HER3 peptides, and control versus OT-II-DC1 groups to OT-II peptide-pulsed DCs. Data shown are the representative from three independent experiments and are represented as mean \pm SEM with statistical significance determined using multiple t test without correction for multiple comparisons. Each row was analyzed individually, without assuming a consistent SD. A log-rank (Mantel-Cox) test was used to determine differences between the survival curves. *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$; ****, $P \leq 0.0001$; #####, $P \leq 0.0001$.

HER3-DC1-treated M05 mice compared with controls (Supplementary Fig. S8H). Together, these data suggest HER3-DC1 induces potent antitumor immune response in a HER3⁺ murine melanoma model, resulting in tumor regression/delayed growth and improved survival by enhancing intratumoral and systemic CD4⁺ T cells and secretion of IFN- γ .

Discussion

HER3/ERBB3, a member of the ERBB family of growth receptors, has gained momentum as a therapeutic target, owing to its multifaceted role in tumor development, growth, and therapy resistance (1). In TCGA, HER3 expression was found to predict worse survival in

melanoma patients. Interestingly, in response to BRAF/MEK inhibitor treatment, melanoma cells are shown to adapt and escape therapy by upregulating HER3 expression via the FOXP3 transcription factor (5). DC1 pulsed with promiscuous MHC class II HER3 peptides demonstrated a dramatic therapeutic effect in a murine melanoma model that expressed increased HER3, suggesting the possibility of clinically developing a HER3 immunotherapy in melanoma patients expressing HER3, especially those treated with BRAF/MEK inhibitors to possibly prevent escape through HER3.

We did not find a significant impact of high HER3 expression on breast cancer survival, which could be attributed to the small sample size. However, high HER3 expression has been previously identified as a prognostic marker of poor overall and disease-free survival in TNBC (10) and has been shown to contribute to therapy resistance (37). Upregulated HER3 expression and activation have also been documented as one of the primary compensatory mechanisms allowing therapy escape in HER2^{POS} breast cancer via restimulation of the PI3K/AKT pathway (38). We utilized a HER2^{POS} mammary tumor model resistant to HER2-targeted therapies, which was sensitive to HER3-mediated CD4⁺ Th1 immune response. This raises the possibility that HER3 peptides can be used to overcome resistant HER2 populations.

CD4⁺ T cells are critical players in regulating antitumor immune responses, and Th1 cells secreting IFN- γ contribute to elimination of tumors through multiple mechanisms, including antibody class switching, CD8⁺ T-cell help, modulation of innate effectors, such as macrophages and NK cells, and maturation of DCs (39). This study showed increased production of IFN- γ by CD4⁺ Th1 cells in both human CD4⁺ T cells *ex vivo* and in murine preclinical models of mammary carcinoma and melanoma. CD4⁺ T cells accumulated in the tumors of mice treated with HER3 peptide-pulsed DC1, suggesting a contribution of the CD4⁺ response in tumor regression. We validated the significance of CD4⁺ T cells in stimulating antitumor immune responses by class II peptide-pulsed DC1 treatment. HER3-DC1 intratumoral administration significantly enhanced CD4⁺ T-cell infiltration, and restimulation with HER3 peptide-pulsed DC resulted in increased intracellular IFN- γ secretion by those CD4⁺ TILs. This observation highlights the CD4-dependent mechanism of action of the HER3-DC1.

In the 4T1 and TUBO models, HER3-DC1 intratumoral injection and subsequent Th1 immune responses result in delayed tumor growth and downregulation of HER3 expression in tumor cells, which indicates possible correlation at the mechanistic level between HER3 expression and the antitumor effect. We observed significantly decreased total HER3 protein in the HER3-DC1 tumors obtained from both 4T1 and TUBO models, suggesting that intratumoral delivery of HER3-DC1 may induce cellular cross-talk downstream and inhibit HER3 protein expression, which contributes to tumor growth and proliferation. For TUBO tumors, we observed HER3-DC1 tumors had downregulation of phospho-AKT and phospho-p44/42MAPK, the primary signaling effectors downstream of HER3, suggesting HER3-DC1 may interfere with activation and function of HER3 in tumors. Also, prominent upregulation of the apoptosis marker cleaved caspase-3 indicated intratumoral cell death due to HER3-DC1. The Th1 cytokine IFN- γ can downregulate expression of HER family of receptor proteins *in vitro* (35) and, hence, HER3-DC1 treatment may reduce HER3 expression by CD4⁺ Th1 stimulation.

The peptide library screening method described here identified promiscuous class II epitopes that were not clearly predicted by current binding algorithms. Only ECD p91 was predicted to have high binding

affinity, whereas none of the remaining eight peptides were indicated to be high-affinity binders. Previous reports have shown high rates of false negatives for immunogenic peptide epitopes that are intermediate- or low-affinity MHC binders (40–43). The MixMHCIIpred algorithm predicted strong binding affinity of all nine peptides identified empirically. However, it should be considered that binding affinity alone is not the only predictor of immunologic response generated by immunogenic peptides, and hence, the large number of random peptides generated by the algorithm that showed a strong binding affinity will still need to be tested individually and experimentally for immune responses *in vitro*, to avoid false-positive epitopes. Absence of defined cutoff percentile rank also leads to subjective interpretation of strong versus weak binding. Our identified peptides showed promiscuous binding across multiple MHC II HLA alleles, suggesting broad application across patients, whereas MixMHCIIpred algorithm showed variation of binding affinity of the peptides across HLA alleles. Lastly, the stark difference of predictive accuracy between the two algorithms used in this study further highlights the efficiency of our empirical approach to accurately identify immunologically responsive peptides. MHC I prediction algorithms have shown success in identifying immunogenic class I epitopes that can elicit potent CD8⁺ T-cell responses *in vivo*; however, the identification of MHC class II epitopes that activate CD4⁺ T cells has been less successful (24, 41–46). This discrepancy can be attributed to the highly polymorphic nature of MHC II molecules, which results in different binding patterns across the diverse HLA alleles (43–45, 47–49). Unlike MHC I, MHC II molecules consist of an open binding groove, allowing for longer peptides of 9–25 amino acid length to bind at different locations across the groove's open surface (43, 45, 46, 48). Our peptide screening attempted to circumvent several limitations in MHC II prediction algorithms to identify immunogenic class II epitopes for use in therapeutic development. First, a library of 15-mer peptides was generated for the protein of interest, with a 10-amino acid overlap between adjacent sequences. This design aimed to account for differences in the binding pattern across the MHC II open binding groove. Second, screening the full peptide library sequentially in pools of 10 peptides, 5 peptides, and individual peptides rapidly highlighted regions within the protein that may harbor immunogenic peptide sequences. Screening the full library essentially surveys all regions of the protein for epitope candidates, minimizing the probability of false negatives. Third, peptide candidates were screened across numerous donor samples to ensure the peptide-specific immune responses were reproducible and demonstrated promiscuous binding for widespread therapeutic application. Lastly, a key benefit to this methodology is that it takes an entirely experimental approach to epitope identification by recapitulating the *in vivo* processes of antigen processing and presentation, T-cell priming, and immune activation to isolate peptides capable of acting as competent epitopes *in vitro* and *in vivo*. Because this overlapping peptide library can be created and rapidly tested, it offers a novel avenue to MHC II epitope discovery against other known oncodrivers. We have, thus, begun to interrogate such oncodrivers for CD4⁺ Th1 peptide epitopes. Our study may also open the avenue for potential use of the DC vaccine platform to generate oncodriver-specific CD4⁺ T cells for novel adoptive T-cell therapies. In summary, using an overlapping 15-mer peptide library derived from the HER3 oncodriver, we identified promiscuous MHC II epitopes capable of driving an anti-HER3 CD4⁺ Th1 response that can have therapeutic impact in breast cancer and melanoma, and can be developed for cancer therapy in these diseases.

Authors' Disclosures

K.N. Kodumudi reports a patent for PCT/US2020/050689 pending to Moffitt Cancer Center. K.S.M. Smalley reports personal fees from Elsevier outside the submitted work. P.A. Forsyth reports personal fees from AbbVie Inc., Bayer, Bristol Myers Squibb, Boehringer Ingelheim, Inovio, NCI-NIH-NCRI, Novocure, Novellus, Physical Sciences Oncology Network, Tocagen, Ziopharm, and grants from CDMRP, Department of Defense, Moffitt Cancer Center of Excellence, NIH/NCI, Pfizer, and State of FL Bankhead Coley outside the submitted work. H. Han reports personal fees from Lilly and grants from Department of Defense outside the submitted work. H. Soliman reports personal fees from PUMA, Novartis, Eisai, Seattle Genetics, and AstraZeneca outside the submitted work. B.J. Czerniecki reports grants from CDMRP DOD during the conduct of the study; other support from ImmunoRestoration outside the submitted work; and a patent for identification of immunogenic MHC class II peptides for immune-based therapy (patent# 10,829,538) issued to the University of Pennsylvania. No disclosures were reported by the other authors.

Authors' Contributions

A. Basu: Conceptualization, formal analysis, validation, investigation, visualization, methodology, writing—original draft, project administration, writing—review and editing. **G.K. Albert:** Conceptualization, formal analysis, validation, investigation, visualization, methodology, writing—original draft, project administration, writing—review and editing. **S. Awshah:** Validation, investigation, methodology, writing—review and editing. **J. Datta:** Conceptualization, formal analysis, validation, investigation, methodology, writing—review and editing. **K.N. Kodumudi:** Conceptualization, formal analysis, supervision, validation, investigation, visualization, methodology, project administration, writing—review and editing. **C. Gallen:** Investigation, methodology, writing—review and editing. **A. Beyer:** Investigation. **K.S.M. Smalley:** Writing—review and editing. **P.C. Rodriguez:** Writing—review and

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