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(54) **IMMUNE CELLS EXPRESSING A
HYPOXIA-DEPENDENT CHIMERIC
ANTIGEN RECEPTOR TARGETED TO EGFR**

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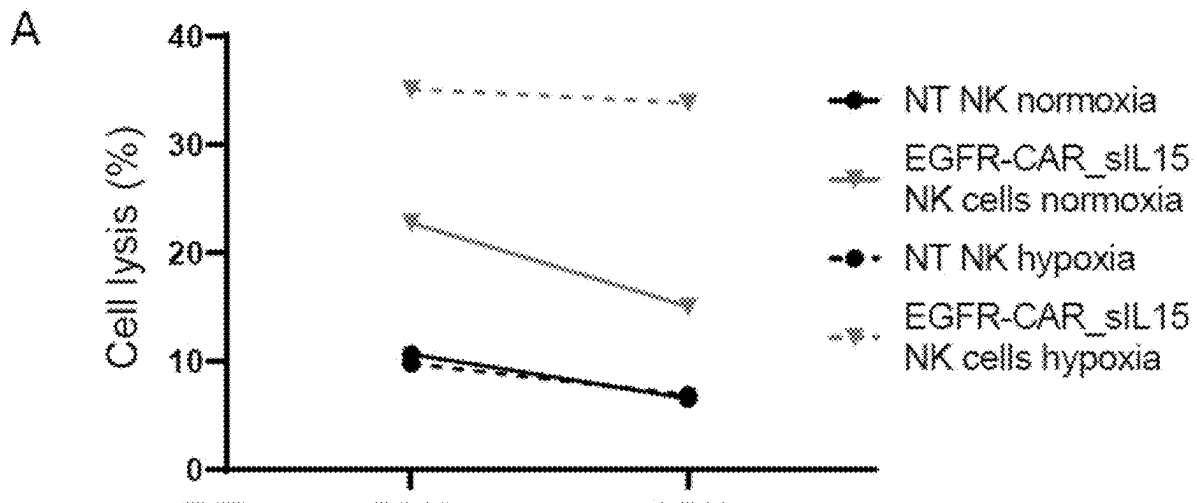
2319/95 (2013.01); *C12N 2510/00* (2013.01)

(57)

ABSTRACT

Described are immune cells expressing a chimeric antigen receptor targeted to EGFR and having an oxygen-dependent degradation domain. The chimeric antigen receptor is co-expressed in immune cells, e.g., NK cells with a soluble form of human IL-15. The inclusion of an oxygen-dependent degradation domain substantially restricts the activity of the immune cells expressing the chimeric antigen receptor to the more hypoxic environment of solid tumor tissue. The NK cells expressing the chimeric antigen receptor are useful for treating breast cancer that has low or no HER2 expression.

Specification includes a Sequence Listing.



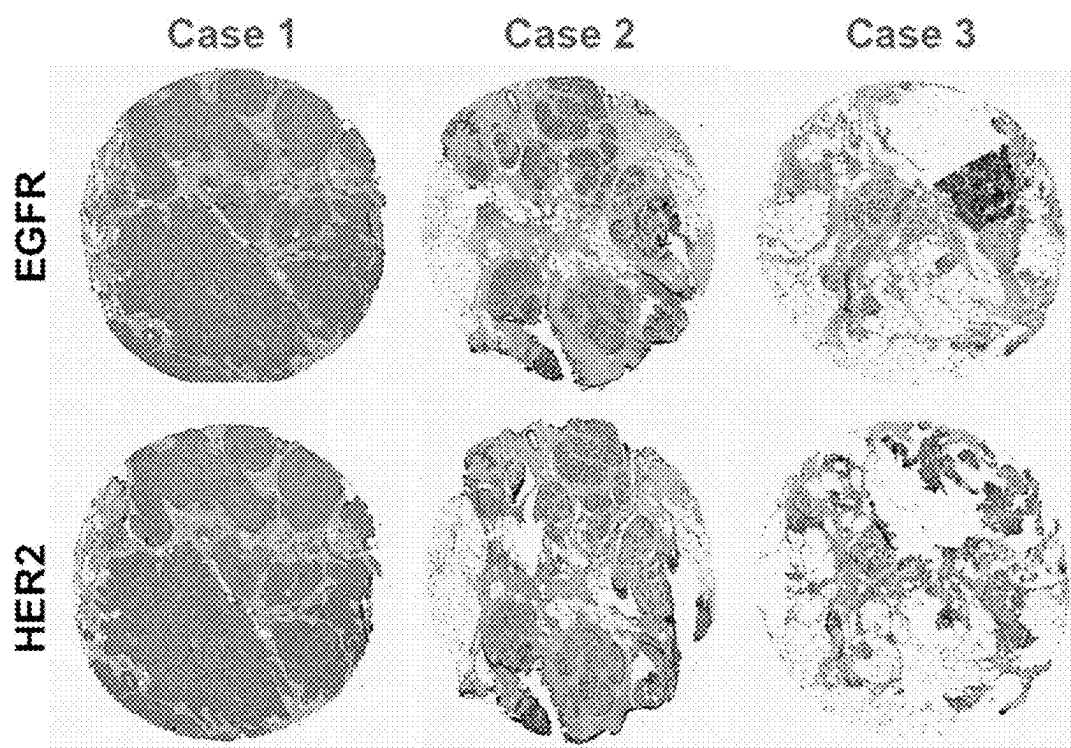


FIG 1



FIG 2

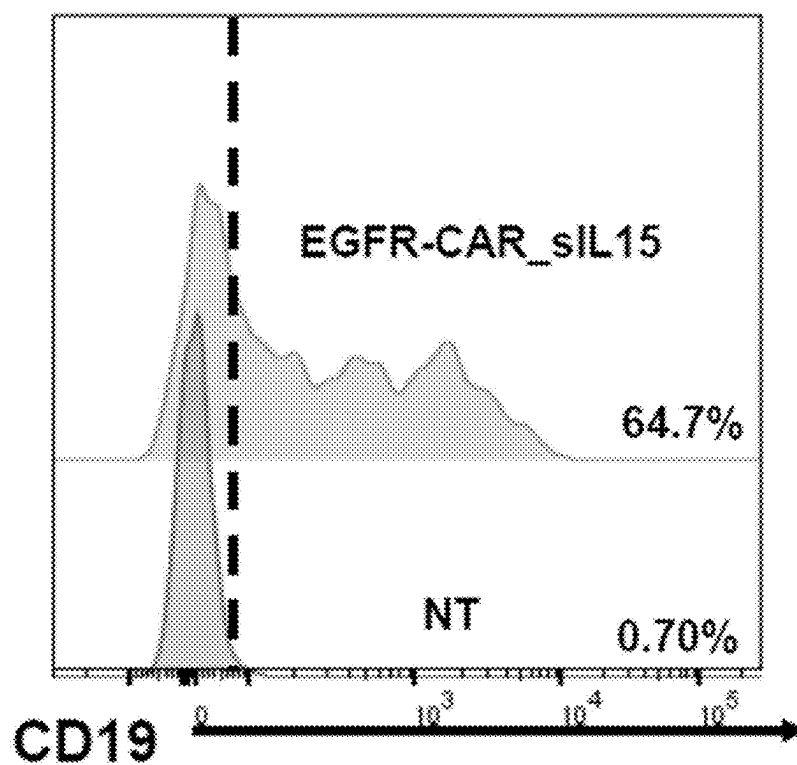


FIG 3

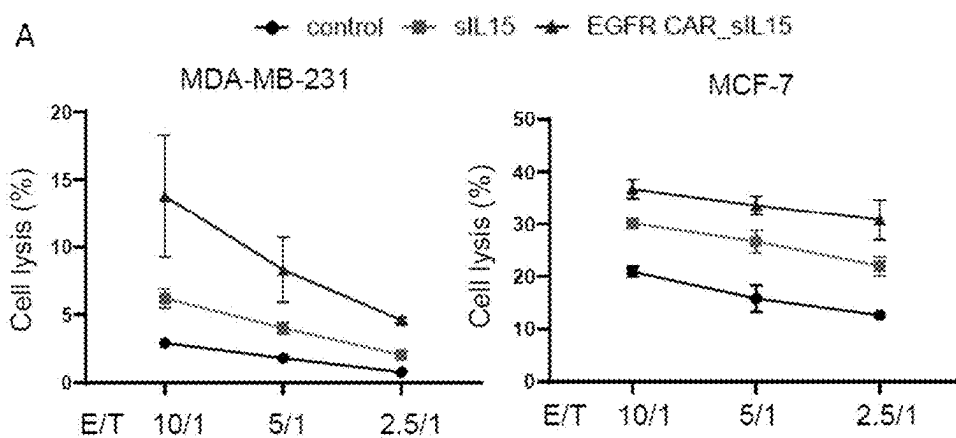


FIG 4A

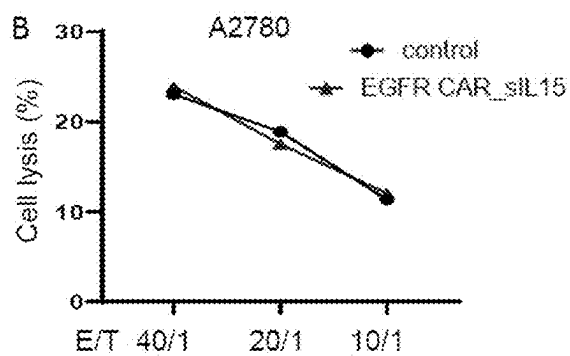


FIG 4B

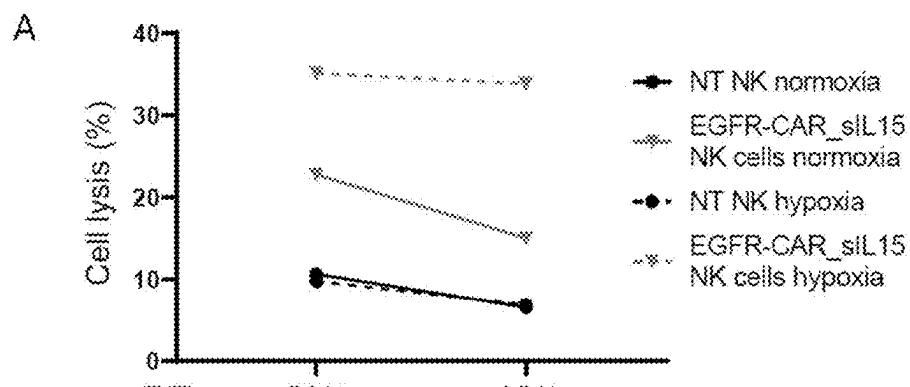


FIG 5A

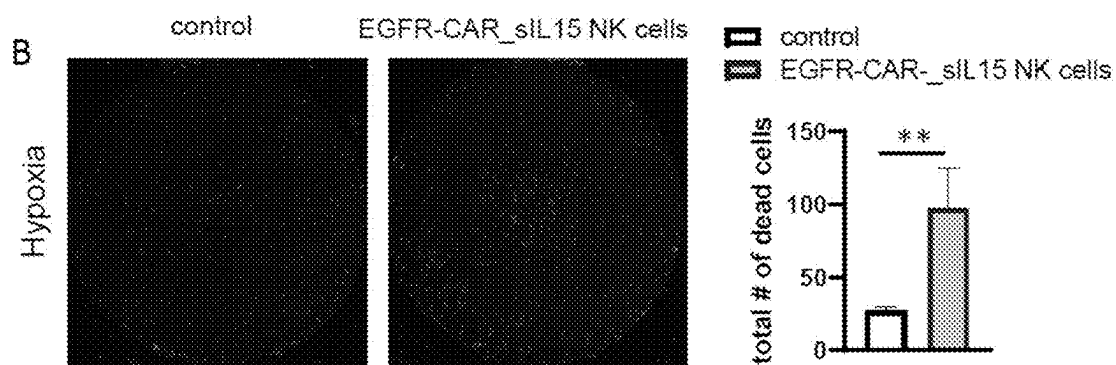


FIG 5B

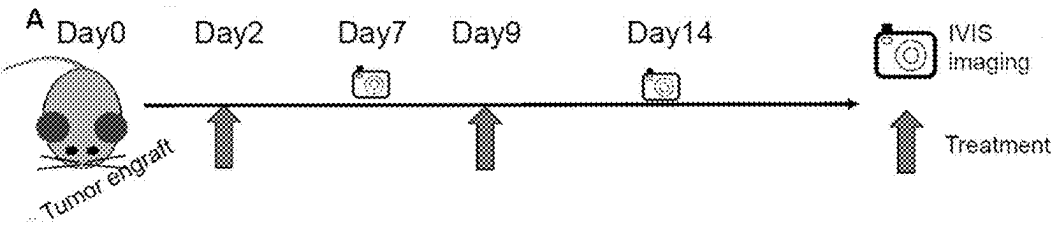


FIG 6A

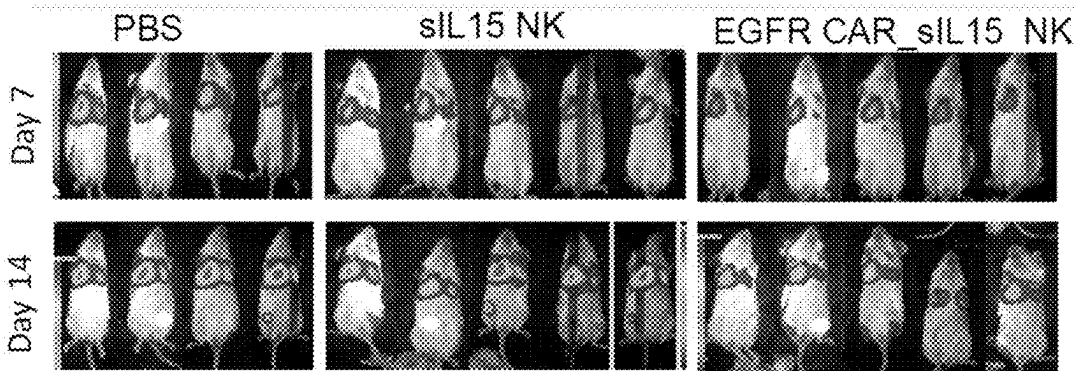


FIG 6B

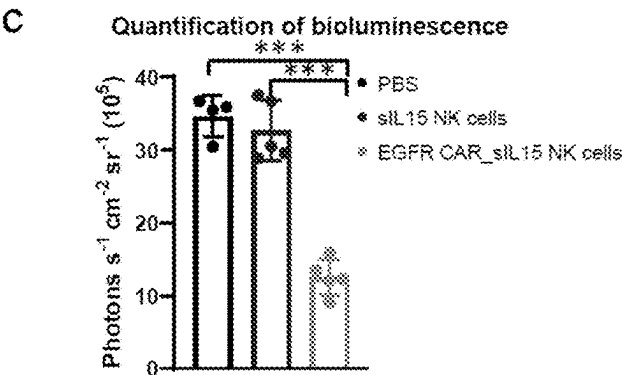


FIG 6C

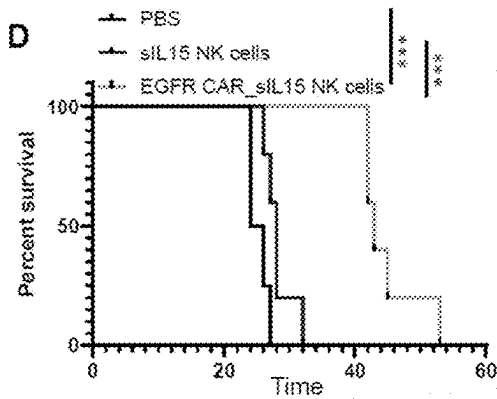


FIG 6D

IMMUNE CELLS EXPRESSING A HYPOXIA-DEPENDENT CHIMERIC ANTIGEN RECEPTOR TARGETED TO EGFR

CLAIM OF PRIORITY

[0001] This application claims the benefit of U.S. Provisional Application Ser. No. 63/603,759, filed on Nov. 29, 2023. The entire contents of the foregoing are incorporated herein by reference.

SEQUENCE LISTING

[0002] This application contains a Sequence Listing which has been submitted electronically in XML format and is hereby incorporated by reference in its entirety. Said XML copy, created on Mar. 3, 2025, is named 40056-0089001_SL.xml and is 56,711 bytes in size.

TECHNICAL FIELD

[0003] This disclosure relates to treating cancer using immune cells expressing a chimeric antigen receptor (CAR) targeted to epidermal growth factor receptor (EGFR) having an intracellular oxygen-dependent degradation domain (ODD). The CAR is co-expressed with human IL-15, e.g., a soluble form of human IL-15.

BACKGROUND

[0004] Breast cancer is the most common cancer and the second leading cause of cancer death in women in North America, with an estimated 268,600 new cases of advanced breast cancer annually. Approximately 155,000 women are living with metastatic breast cancer in the United States with a median overall survival of 3.5 years. About 60-65% of breast cancers are estrogen receptor (ER) positive/human epidermal growth factor 2 (HER2) negative. More effective and less toxic therapies replacing or combined with conventional chemotherapy represent an area of unmet need in patients with HER2 negative metastatic breast cancer who have progressed through first- and second-line therapies.

[0005] HER2-low breast cancer is defined by HER2 immunohistochemistry 1+ or 2+ with negative fluorescence in situ hybridization. The clinical development of novel anti-HER2 agents has the potential to improve the treatment armamentarium for a subgroup of HER2-low breast cancer patients. Patients with HER2-low breast cancer are not currently considered candidates for HER2-targeted therapy. The HER2-low population accounts for up to 45-55% of all breast cancers. However, the anti-HER2 agents currently approved for the treatment of HER2-positive breast cancer have little or no activity in HER2-low tumors. Currently, there are limited treatment options for patients with HER2-low breast cancer, including estrogen receptor positive (ER+) breast cancer and triple-negative breast cancer (TNBC), which lacks expression of estrogen receptor (ER), progesterone receptor (PR), and HER2. This represents an area of unmet need in patients who progressed after initial targeted therapy.

SUMMARY

[0006] Described herein are studies showing that, in some cases, EGFR is expressed or highly expressed on primary breast cancer tumor cells even when HER2 expression is at low levels. Also described are studies showing that EGFR is

expressed on human MDA-MB-231, MDA-MB-468, and MCF-7 BC cell lines, as previously reported¹. To target the elevated expression of EGFR that we observed certain breast cancer tumor cells, we developed an EGFR-targeted CAR (EGFR-CAR) that includes an anti-EGFR single-chain variable fragment (scFv) targeting both wild-type EGFR and EGFRvIII. The CAR includes an anti-EGFR scFv, a spacer domain, a transmembrane domain, a CD28 co-stimulatory domain, and CD38. Given that IL-15 activates and improves the survival and activation of CD8 T cells and NK cells, we thus incorporated a soluble form of IL-15 (sIL-15) into the EGFR-CAR construct. We fused codon-optimized sIL-15 to EGFR-CAR using a Thossea assigna virus 2A-like ribosomal skip sequence (T2A). We linked a truncated, non-signaling form of CD19 (tCD19) to sIL-15 via porcine teschovirus-1 2A-like self-cleaving peptide (P2A). The tCD19 can serve as a marker to monitor CAR expression cells and can be used as a suicide gene switch with an anti-CD19 antibody. To reduce the activity of the CAR in non-cancer tissue (e.g., outside of the hypoxic tumor microenvironment), the EGFR-CAR is expressed as a fusion protein with the oxygen-dependent degradation domain (ODD) of HIF-1 α . Solid tumor tissue is often hypoxic as a result of the high proliferation and high metabolic demand of tumor cells. Thus, tumor tissue can be below 5%, 4% or even below 2% O₂ (normal tissue is generally 5-15% O₂). The ODD of HIF1 α directs degradation of HIF1 α under hypoxic conditions. Under these conditions, prolines in the ODD are hydroxylated allowing the protein to be recognized by von Hippel-Lindau tumor suppressor, part of an E3 ubiquitin ligase complex that ubiquitinates HIF1 α and targets it for proteasomal degradation.

[0007] When an ODD is included in a CAR, it directs degradation outside of tumor tissue where the O₂ level is higher. Overall the EGFR-CAR with an ODD and sIL-15 is referred to as EGFR-CAR_sIL15 (FIG. 2). Importantly, the EGFR-CAR_sIL15 is expressed in NK cells, which have allogeneic, "off-the-shelf" (not patient-specific) potential. NK cells can be manufactured from different sources such as peripheral blood cells, umbilical cord blood-derived NK cells, hematopoietic stem cells (e.g., CD34+ cells), induced pluripotent stem cells, etc.

[0008] The ODD can comprise or consist of the sequence: APAAGDTIISLDFG-SNDTETDDQQLLEEVPLYNDVMLPSPNEKLQIN-LAMSPPLT AETPKPLRSSADPAL-NQEVALKLEPNPESLELSFTMPQIQDQTPSPSDGSTRQSSPE PNSPSEYCFYVDSMDMVNEFKLELVEKLF AE-DTEAKNPFTQDQTDLDLEMLAPYIP MDDDDFQLRSFDQLSPLESSSASPESASPQSTVTVFQ (SEQ ID NO: 10). It can be followed by a ribosomal skip sequence, e.g.: AEGRGSLLTCGDVEENPGP (SEQ ID NO: 13).

[0009] The soluble IL-15 can comprise or consist of the sequence: GIHVFI LGCF SAGLPKTEANWVNVISDLK-KIEDLIQSMHIDATLYTESDVHPSCKV TAMKCFLLQLVISLESGDASIHDTVENLILANNSLSSNG-NVTESGCKECELEE KNIKEFLQSFVHVHVMFINTS (SEQ ID NO:11), which can be preceded by a signal sequence that directs secretion from a mammalian cell, e.g.: MYRMQLLSIALSLALVINS (SEQ ID NO: 63) and it can be followed by a ribosomal skip sequence, e.g., GSGATNFSLLKQAGDVEENPGP (SEQ ID NO: 14), which in turn can be followed by a truncated CD19 receptor

that acts both as a marker and a suicide switch. This truncated CD19 receptor can comprise or consist of the sequence:

(SEQ ID NO: 47)
 MPPPRLLFFLLFLTPMEVRPEEPLVVKVEEGDNAVLQCLKGTS
 DTG
 PTQQLTWSRESPLKPKLKLGLPLGIHMRPLAIWLFIFNVSSQ
 Q
 MGGFYLCQPGPPSEKAWQPGWTVNVEGSGELFRWNVSDLGGLGCG
 LKNRSSEGPSSPSGKLMSPKLYVWAKDRPEIWEGEPPCLPPRDSL
 NQSLSQDLTMAPGSTLWLSCGVPPDSVSRGPLSWTHVHPKGP
 KSL
 LSLELKDDRPARDMVMETGLLLPRATAQDAGKYCHRGNTMSF
 HLEITARPVLWHLLRTGGWKVSAVTLAYLIFCLCSLVGILHLQR
 ALVLRKR.

[0010] The mature EGFR-CAR, with an ODD domain, can comprise or consist of the amino acid sequence:

(SEQ ID NO: 57)
 DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRP
 GQSPKLLIYKVSDFRSGVPDRFSGSGGTDFTLKISRVEAEDLGI
 YYCFQGSHIPPTFGGGTKLEIKRAAGGGSGGGSGGGSGVQLQ
 QSGSEMARPGASVKLPCKASGDTFTSYWMHWKQRHGHGPEWIGN
 IYPGSGGTNYAEKPKNKVTLTVDRSSRTVYMHLSRLTSEDSAVVY
 CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
 VVGGVLACYSLLVTVAFIIFWVRSKRSLHSDYMNMTPRRPGPT
 RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNQLYNELNLG
 RREEYDVLDKRRGRDPGEMGGKPRRKNPQEGLYNELQDKMAEAYS
 EIGMKGERRRGKGGHDGLYQGLSTATKDTYDALHMQALPPRAPAAG
 DTIISLDFGSNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
 SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
 DQTPSPSDGSTRQSSPEPNPSEYCFYVSDMVNEFKLELVEKLF
 AEDTEAKNPFSTQDTDLDEMLAPYIPMDDDFQLRSFDQLSPLES
 SSASPESASPQSTVTVFQ.

[0011] When followed by a ribosomal skip sequence and sIL-15, the amino acid sequence of EGFR-CAR_sIL15 without the CAR signal sequence, but with the sIL-15 signal sequence, comprises or consists of:

(SEQ ID NO: 58)
 DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRP
 GQSPKLLIYKVSDFRSGVPDRFSGSGGTDFTLKISRVEAEDLGI
 YYCFQGSHIPPTFGGGTKLEIKRAAGGGSGGGSGGGSGVQLQ
 QSGSEMARPGASVKLPCKASGDTFTSYWMHWKQRHGHGPEWIGN
 IYPGSGGTNYAEKPKNKVTLTVDRSSRTVYMHLSRLTSEDSAVVY
 CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPKFWVLV

-continued
 VVGGVLACYSLLVTVAFIIFWVRSKRSLHSDYMNMTPRRPGPT
 RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNQLYNELNLG
 RREEYDVLDKRRGRDPGEMGGKPRRKNPQEGLYNELQDKMAEAYS
 EIGMKGERRRGKGGHDGLYQGLSTATKDTYDALHMQALPPRAPAAG
 DTIISLDFGSNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
 SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
 DQTPSPSDGSTRQSSPEPNPSEYCFYVSDMVNEFKLELVEKLF
 AEDTEAKNPFSTQDTDLDEMLAPYIPMDDDFQLRSFDQLSPLES
 SSASPESASPQSTVTVFQAEGRGSLTTCGDVEENPGPMYRMQLLS
 CIALSLALVTNSGIHVFILGCFSGAGLPKTEANWVNVISDLKKIED
 LIQSMHIDATLYTESDVHPSCVKVTAMKCFLLELQVISLES
 GDASI
 HDTVENLIILANNSLSSNGNVTESGCKECELEEKNIKEFLQSFV
 HIVQMFINTS.

[0012] Thus, the mature proteins expressed on the surface of an immune cell (an NK cell, a T cell, an NKT cell, a macrophage, and gamma delta T cell) include:

(SEQ ID NO: 57)
 DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRP
 GQSPKLLIYKVSDFRSGVPDRFSGSGGTDFTLKISRVEAEDLGI
 YYCFQGSHIPPTFGGGTKLEIKRAAGGGSGGGSGGGSGVQLQ
 QSGSEMARPGASVKLPCKASGDTFTSYWMHWKQRHGHGPEWIGN
 IYPGSGGTNYAEKPKNKVTLTVDRSSRTVYMHLSRLTSEDSAVVY
 CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
 VVGGVLACYSLLVTVAFIIFWVRSKRSLHSDYMNMTPRRPGPT
 RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNQLYNELNLG
 RREEYDVLDKRRGRDPGEMGGKPRRKNPQEGLYNELQDKMAEAYS
 EIGMKGERRRGKGGHDGLYQGLSTATKDTYDALHMQALPPRAPAAG
 DTIISLDFGSNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
 SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
 DQTPSPSDGSTRQSSPEPNPSEYCFYVSDMVNEFKLELVEKLF
 AEDTEAKNPFSTQDTDLDEMLAPYIPMDDDFQLRSFDQLSPLES
 SSASPESASPQSTVTVFQ
 and

(SEQ ID NO: 11)
 GIHVFILGCFSGAGLPKTEANWVNVISDLKKIEDLIQSMHIDATLY
 TESDVHPSCVKVTAMKCFLLELQVISLES
 GDASIHDTVENLIILAN
 NSLSSNGNVTESGCKECELEEKNIKEFLQSFVHIVQMFINTS

[0013] In some cases, a mature, truncated CD19 receptor is also expressed by the cells. Described herein is a nucleic acid molecule encoding a fusion protein comprising a chimeric antigen receptor and an oxygen-dependent degradation domain (ODD), wherein the chimeric antigen receptor comprises: (i) an scFv that binds human EGFR; (ii) a spacer

domain; (iii) a transmembrane domain; (iv) a costimulatory domain; and (v) a CD3 ζ signaling domain.

[0014] In some embodiments: the scFv comprises: a light chain CDR1 comprising RSSQNIVHNNGITYLE (SEQ ID NO: 4), a light chain CDR2 comprising KVSDRFS (SEQ ID NO: 5), a light chain CDR3 comprising FQGSHIPPT (SEQ ID NO: 6), a heavy chain CDR1 comprising SYWMH (SEQ ID NO: 7), a heavy chain CDR2 comprising NIYPGSGGTNYAEKFKN (SEQ ID NO: 8), and a heavy chain CDR3 comprising SGGPYFFDY (SEQ ID NO: 9); the ODD is a human HIF1 α ODD; the ODD comprises or consists of

APAAGDTIISLDFG-SNDTETDDQQLLEEVPLYNDVMLPSPNEKLQIN-LAMSPPLT AETPKPLRSSADPAL-NQEVALKLEPNPESLELSFTMPQIQDQTPSPSDGSTRQSSPE PNPSPSEYCFYVDSMDVNEFKLELVEKLEFAE-DTEAKNPFSTQDITDLEMLAPYIP

MDDDFQLRSFDQLSPLESSASPESASPQSTVTVFQ (SEQ ID NO: 10); the scFv comprises a VL domain comprising: DILMTQSPSLSPVSLGDQASIS-CRSSQNIVHNNGITYLEWYLQRPQGSPKLLIYKVSDRFGVPDRFSGSGSGTDFTLKISRVEAE-DLGIYYCFQGSHIPPTFGGGTKLEIKR AA (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions (in some embodiments, the amino acid substitutions are not in the CDRs) followed by a VH domain comprising or consisting of QVQLQQSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHG-PEWIGNIY PGSGGTNY-AEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYC

TRSGGPYFFD YWGQGTTLVSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions (in some embodiments, the amino acid substitutions are not in the CDRs); the scFv comprises a VL domain comprising or consisting of an amino acid sequence at least 95%, 98% or 99% identical to: DILMTQSPSLSPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRPQGSPKLLIYKVSDRFGVPDRFSGSGSGTDFTLKISRVEAE-DLGIYYCFQGSHIPPTFGGGTKLEIKR AA (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions (in some embodiments, the amino acid substitutions are not in the CDRs) followed by a VH domain comprising or consisting of an amino acid sequence at least 95%, 98% or 99% identical to QVQLQQSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHG-PEWIGNIY PGSGGTNY-AEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYC

TRSGGPYFFD YWGQGTTLVSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions (in some embodiments, the amino acid substitutions are not in the CDRs); the spacer region comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: H and 24-34; the chimeric antigen receptor comprises: the spacer comprises or consists of a sequence selected from the group consisting of: SEQ ID NOs: H and 24-34; the transmembrane domain comprises or consists of a sequence selected from the group consisting of SEQ ID NOs: 15-23; the costimulatory domain comprises or consists of a sequence from the group consisting of SEQ ID NOs: 36-40, and the CD3 signaling domain comprises or consists of SEQ ID NO: 35; the nucleic acid molecule further encodes human IL-15 or a soluble variant of human IL-15; the nucleic acid molecule further encodes human IL-15 or a soluble variant of human IL-15 that is co-expressed with the fusion protein;

the soluble variant of IL-15 comprises or consists of: GIHV-FILGCFSAAGLPKTEANWVNVISDLKKIEDLIQSMHI-DATLYTESDVHPSCKV TAMKCFLLELQVISLESG-DASIHDTVENLIILANNSLSSNGNVTESGCKECEEELEE KNIKEFLQSFVHIVQMFINTS (SEQ ID NO: 11) the nucleic acid molecule further includes a nucleotide sequence encoding signal sequence preceding the nucleotide sequence encoding SEQ ID NO: 11.

[0015] In some embodiments: the fusion protein comprises or consists of the amino acid sequence:

(SEQ ID NO: 57)
DILMTQSPSLSPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRP
QGSPKLLIYKVSDRFGVPDRFSGSGSGTDFTLKISRVEAEDLGI
YYCFQGSHIPPTFGGGTKLEIKRAAGGGGSGGGGSGGGGSGVQLQ
QSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHGPEWIGN
IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYY
CTRSGGPYFFDYWGQGTTLVSSPKSCDKTHTCPPCPDPKFVWL
VVGGLVACYSLLVTVAFIIFWVRSKRSRLHSDYMMNTPRRPGPT
RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQGGQNQLYNELNLG
RREEYDVLDKRRGRDPEMGGKPRRNQPEGLYNELQDKDMAEAYS
EIGMKGERRRGKGGHGLYQGLSTATKDTYDALHMQALPPRPAAG
DTIISLDFGSDNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSPSEYCFYVDSMDVNEFKLELVEKLE
AEDTEAKNPFSTQDITDLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVFQ

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions. In some embodiments, the amino acid substitutions are not in the targeting domain (e.g., the scFv, the VL domain and VH domain, and/or the CDRs).

[0016] In some embodiments: the nucleic acid molecule encodes an amino acid sequence comprising or consisting of the amino acid sequence:

(SEQ ID NO: 57)
DILMTQSPSLSPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRP
QGSPKLLIYKVSDRFGVPDRFSGSGSGTDFTLKISRVEAEDLGI
YYCFQGSHIPPTFGGGTKLEIKRAAGGGGSGGGGSGGGGSGVQLQ
QSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHGPEWIGN
IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYY
CTRSGGPYFFDYWGQGTTLVSSPKSCDKTHTCPPCPDPKFVWL
VVGGLVACYSLLVTVAFIIFWVRSKRSRLHSDYMMNTPRRPGPT
RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQGGQNQLYNELNLG
RREEYDVLDKRRGRDPEMGGKPRRNQPEGLYNELQDKDMAEAYS
EIGMKGERRRGKGGHGLYQGLSTATKDTYDALHMQALPPRPAAG
DTIISLDFGSDNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM

-continued

SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF
AEDTEAKNPFSTQDSDLLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVTFQ

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions; or the amino acid sequence:

(SEQ ID NO: 58)
 DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRP
 GQSPKLLIYKVSDFRSGVDPDRFSGSGSGTDFTLKISRVEAEDLGI
 YYCFQGSHIPPTFGGGTKLEIKRAAGGGGSGGGGSGGGGSGVQLQ
 QSGSEMARPGASVKLPCKASGDTFTSYWMHWKQRHGHGPEWIGN
 IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYY
 CTRSGGPYFFDYWGQTTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
 VVGVLACYSLLVTVAFIIFWVRSKRRLHSDYMNMTPRRPGPT
 RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNLYNELNLG
 RREEYDVLDKRRGRDPEMGGKPRRKNPQEGLYNELQDKDMAEAYS
 EIGMKGERRRGKGDGLYQGLSTATKDTYDALHMQALPPRPAAG
 DTIISLDFGSNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF
AEDTEAKNPFSTQDSDLLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVTFQAEGRGSLTTCGDVEENPGPMYRMQLLS
CIALSLALVTNSGIHFVILGCFSGAGLPKTEANWVNVISDLKKIED
 LIQSMHIDATLYTESDVHPSCKVTAMKCFLLLELQVISLESGLDASI
 HDTVENLIILANNSLSSNGNVTESGCKECELEEKNIKEFLQSFV
 HIVQMFINTS

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions. In some embodiments, the amino acid substitutions are not in the targeting domain (e.g., the scFv, the VL domain and VH domain, and/or the CDRs).

[0017] Also disclosed is an immune cell (e.g., a T cell, an NK cell, an NKT cell, a macrophage, or a gamma delta T cell) harboring a nucleic acid described herein. In some embodiments: the immune cell expresses human IL-15 or soluble variant thereof; the soluble variant of IL-15 comprises or consists of: GIHVILGCFSGAGLPKTEANWVNVISDLKKIEDLIQSMHIDATLYTESDVHPSCKVTAMKCFLLLELQVISLESGLDASIHDTVEN-LIILANNSLSSNGNVTESGCKECELEE KNIKE-FLQSFVHIVQMFINTS (SEQ ID NO:11); and the immune cell is an NK cell.

[0018] Also described is a method for treating breast cancer comprising administering a composition comprising an immune cell described herein.

[0019] Also described is fusion protein comprising a chimeric antigen receptor and an oxygen-dependent degradation domain (ODD), wherein the chimeric antigen receptor

comprises: (i) an scFv that binds human EGFR; (ii) a spacer domain; (iii) a transmembrane domain; (iv) a costimulatory domain; and (v) a CD3 ζ signaling domain.

[0020] In various embodiment of the fusion protein: the scFv comprises: a light chain CDR1 comprising RSSQNIVHNNGITYLE (SEQ ID NO: 4), a light chain CDR2 comprising KVSDRFS (SEQ ID NO: 5), a light chain CDR3 comprising FQGSHIPPT (SEQ ID NO: 6), a heavy chain CDR1 comprising SYWMH (SEQ ID NO: 7), a heavy chain CDR2 comprising NIYPGSGGTNYAEKFKN (SEQ ID NO: 8), and a heavy chain CDR3 comprising SGGPYFFDY (SEQ ID NO: 9); the ODD is a human HIF1 α ODD; the ODD comprises or consists of APAAGDTIIS-LDFGSNDTETDDQQLLEEVPLYNDVMLPSP-NEKLQINLAMSPPLPT AETPKPLRSSADPAL-NQEVALKLEPNPESLELSFTMPQIQDQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF AEDTEAKNPFSTQDSDLLEMLAPYIPMDDDFQLRSFDQLSPLESSASPESASPQSTVTVTFQ (SEQ ID NO: 10); the scFv comprises a VL domain comprising: DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRPQGSPKLLIYKVSDFRSGVDPDRFSGSGTDFTLKISRVEAEDLGIYYCFQGSHIPPTFGGGTKLEIKRAAGGGGSGGGGSGGGGSGVQLQ (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions followed by a VH domain comprising or consisting of QVQLQQSGSEMARPGASVKLPCK-ASGDTFTSYWMHWKQRHGHGPEWIGNIYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYCTRSGGPYFFDY YWGQTTTLTVSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions; and the fusion protein comprises or consists of the amino acid sequence:

(SEQ ID NO: 57)
 DILMTQSPLSLPVSLGDQASISCRSSQNIHNNGITYLEWYLQRP
 GQSPKLLIYKVSDFRSGVDPDRFSGSGTDFTLKISRVEAEDLGI
 YYCFQGSHIPPTFGGGTKLEIKRAAGGGGSGGGGSGGGGSGVQLQ
 QSGSEMARPGASVKLPCKASGDTFTSYWMHWKQRHGHGPEWIGN
 IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYY
 CTRSGGPYFFDYWGQTTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
 VVGVLACYSLLVTVAFIIFWVRSKRRLHSDYMNMTPRRPGPT
 RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNLYNELNLG
 RREEYDVLDKRRGRDPEMGGKPRRKNPQEGLYNELQDKDMAEAYS
 EIGMKGERRRGKGDGLYQGLSTATKDTYDALHMQALPPRPAAG
 DTIISLDFGSNDTETDDQQLLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF
AEDTEAKNPFSTQDSDLLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVTFQ

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions. In some embodiments, the amino acid substitutions are not in the targeting domain (e.g., the scFv, the VL domain and VH domain, and/or the CDRs).

I. Chimeric Antigen Receptor

[0021] A chimeric antigen receptor (CAR) refers to an artificial immune cell receptor that is engineered to recognize and bind to a surface antigen. A NK cell that expresses a CAR polypeptide is referred to as a CAR NK cell. CARs have the ability to redirect immune cell specificity and reactivity toward a selected target in a non-MHC-restricted manner. The non-MHC-restricted antigen recognition gives CAR NK cells the ability to recognize an antigen independent of antigen processing, thereby bypassing a major mechanism of tumor escape. A CAR can also be expressed by other immune effector cells, including, but not limited to, T cells ("CAR T cells") and direct T cell killing to cells expressing the target of the CAR.

[0022] There are various generations of CARs, each of which contains different components. First generation CARs join an antibody-derived scFv to the CD3 intracellular signaling domain of the T cell receptor through a spacer region (also called a hinge domain) and a transmembrane domain. Second generation CARs incorporate an additional co-stimulatory domain (e.g., CD28, 4-1BB, or ICOS) to supply a co-stimulatory signal. Third generation CARs contain two co-stimulatory domains (e.g., a combination of CD27, CD28, 4-1BB, ICOS, or OX40) fused with the TCR CD3 ζ chain. Any generation of CAR is within the scope of the present disclosure.

[0023] The CAR described herein are fusion proteins comprising an extracellular domain that recognizes a tumor cell antigen (e.g., a single chain fragment (scFv) of an antibody or other antibody fragment), a spacer, a transmembrane domain, at least one co-stimulatory domain and an intracellular domain comprising a signaling domain of the T cell receptor (TCR) complex (e.g., CD3 ζ). A CAR is often fused to a signal peptide at the N-terminus for surface expression.

(a) Target Binding scFv

[0024] An EGFR CAR can include: (i) an scFv that binds EGFR; (ii) a spacer domain; (iii) a transmembrane domain; (iv) a costimulatory domain; and (v) a CD32 signaling domain. In various embodiments: the spacer region comprises 5-300 amino acids. The EGFR scFv can comprise: nucle The EGFR scFv can comprise: (a) a light chain variable domain that is at least 90%, 95% or 98% identical to: (SEQ ID NO: 2); and (b) a heavy chain variable domain that is at least 90%, 95% or 98% identical to: (SEQ ID NO:3). The light chain variable domain can precede the heavy chain variable domain and they can be joined by a linker that includes 5-20 amino acids, preferably G and S. For example, the EGFR scFv can comprise or consist of: DILMTQSPLSLPVSLGDAQISCRSSQNIVHNNGI-TYLEWYLQRPQGSPKLLIYKV SDRFSGVPDRFSGSGSGTDFTLKISRVEAE-DLGIYYCFQGSHIPPTFGGGTKLEIKR AAGGGSGGGSGGGGSQVQLQQSGSEMARPGASVKLPCKASGDTFTSYWMH VVKQRHGHGPEWIGNIYPGSGGTNY-AEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYC-TRSGGPYFFD YWGQGTTLTVSS (SEQ ID NO:1). Thus, the scFv can include a VL domain DILMTQSPLSLPVSLGDAQISCRSSQNIVHNNGI-TYLEWYLQRPQGSPKLLIYKV SDRFSGVPDRFSGSGSGTDFTLKISRVEAE-DLGIYYCFQGSHIPPTFGGGTKLEIKR AA (SEQ ID NO: 2) followed by a VH domain comprising or consisting of

QVQLQQSGSEMARPGASVKLPCKASGDTFTSYWMH VVKQRHGHGPEWIGNIY PGSGGTNY-AEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYC-TRSGGPYFFD YWGQGTTLTVSS (SEQ ID NO: 3). A linker having the sequence: GGGGSGGGGSGGGGS (SEQ ID NO: 60) or GGGGSGGGGS (SEQ ID NO: 61) can be located between the VH and VL domains.

[0025] The spacer region can comprise or consist of an amino acid sequence selected from the group consisting of SEQ ID NOs: H and 24-34; the transmembrane domain can comprise or consist of: PKSCDKTHTCPPCPDPK (SEQ ID NO: 59); a CD4 transmembrane domain, a CD8 transmembrane domain, a CD28 transmembrane domain (either CD28 or CD28 (M)), and a CD32 transmembrane domain; the costimulatory domain selected from the group consisting of: SEQ ID NOs: 36-40, a CD28 costimulatory domain, CD28gg costimulatory domain, a 41-BB costimulatory domain, an OX40 costimulatory domain, and a 2B4 costimulatory domain.

[0026] The chimeric antigen receptor can comprise or consist of an amino acid sequence that is 95%, 96%, 97%, 98%, 99% or 100% identical to any of the following sequences, or has 1, 2, 3, 4 or 5 single amino acid substitutions or deletions (preferable deletions are amino terminal or carboxy terminal) compared to, SEQ ID NO: A or B. To calculate percent identity, gapped BLAST can be utilized as described in Altschul et al., Nucleic Acids Res. 25 (17): 3389-3402, 1997. When utilizing BLAST and Gapped BLAST programs, the default parameters of the respective programs (e.g., XBLAST and NBLAST) are used.

[0027] The amino acid sequence of the CAR can be preceded by a signal sequence, for example: MDWIWRIL-FLVGAATGAHS (SEQ ID NO: 12) or MLLLVTSLLL-CELPHPAFLIP (SEQ ID NO: 62)

[0028] Also disclosed are nucleic acid molecules encoding a chimeric antigen receptor, wherein the chimeric antigen receptor comprises: an scFv described herein (e.g., SEQ ID NO: 1); a spacer comprising a sequence selected from the group consisting of: PKSCDKTHTCPPCPDPK (SEQ ID NO: 59) and SEQ ID NOs: 24-34; a transmembrane domain comprising a sequence selected from the group consisting of SEQ ID NOs: 15-23; a costimulatory domain comprising a sequence selected from the group consisting of SEQ ID NOs: 36-40, and a CD3 ζ signaling domain comprising SEQ ID NO: 35 (or any of SEQ ID NOs: 50-56).

[0029] Also disclosed are immune cells harboring any nucleic acid molecule described herein.

[0030] Also disclosed are methods of treating breast cancer (e.g., a breast cancer patient with a HER2-low breast cancer, including: 1) estrogen receptor positive (ER+), HER-2 low breast cancer; and 2) HER2-low, triple-negative breast cancer) the method comprising administering a therapeutically effective amount of immune cells, e.g., NK cells, expressing an EGFR-CAR_sIL15. An EGFR-CAR_sIL15 NK cell or EGFR-CAR_sIL15 T cell can also be used to treat HER2 expressing breast cancer.

(b) Transmembrane Domain

[0031] A CAR disclosed herein can contain a transmembrane domain, which can be a hydrophobic alpha helix that spans the membrane. As used herein, a transmembrane domain refers to any protein structure that is thermodynamically stable in a cell membrane, preferably a eukaryotic cell membrane.

[0032] The transmembrane domain of a CAR as provided herein can be a CD28 transmembrane domain having the sequence: FWLVVVGGVLACYSLLVTVAIFWV (SEQ ID NO: 16). Other transmembrane domains can be used including those shown below in Table 1.

TABLE 1

Examples of Transmembrane Domains				
Name	Accession	Length	Sequence	
CD3z	JO4132.1	21 aa	LCYLLDGILFIYGVILTALPL (SEQ ID NO: 15)	
CD28	NM_006139	27 aa	FWVLVVVGGVLACYSLLVTVAFIIFWV (SEQ ID NO: 16)	
CD28 (M)	NM_006139	28 aa	MFWVLVVVGGVLACYSLLVTVAFIIFWV (SEQ ID NO: 17)	
CD4	M35160	22 aa	MALIVLGGVAGLLFLFIGLGIF (SEQ ID NO: 18)	
CD8tm	NM_001768	21 aa	IYIWAPLAGTCGVLLSLVIT (SEQ ID NO: 19)	
CD8tm2	NM_001768	23 aa	IYIWAPLAGTCGVLLSLVITLY (SEQ ID NO: 20)	
CD8tm3	NM_001768	24 aa	IYIWAPLAGTCGVLLSLVITLYC (SEQ ID NO: 21)	
4-1BB	NM_001561	27 aa	IISFFLALTSTALLFLFF LTLRFSSV (SEQ ID NO: 22)	
NKG2D	NM_007360	21 aa	PFFFCFIAVAMGIRFIIMVA (SEQ ID NO: 23)	

(c) Spacer Region

[0033] A CAR or described herein can include a spacer region located between the EGFR targeting domain (i.e., an EGFR targeted scFv or variant thereof) and the transmembrane domain. Without being bound by theory, the spacer region can function to provide flexibility to the CAR, or domains thereof, or to prevent steric hindrance of the CAR,

or domains thereof. A variety of different spacers can be used. Some of them include at least portion of a human Fc region, for example a hinge portion of a human Fc region or a CH3 domain, or variants thereof. For example, a suitable spacer can comprise or consist of the sequence: PKSCDKTHTCPPCPDPK (SEQ ID NO: 59). Table 2 below provides 10 various spacers that can be used in the CARs or polypeptides described herein.

TABLE 2

Examples of Spacers		
Name	Length	Sequence
a3	3 aa	AAA
linker	10 aa	GGGSSGGGSG (SEQ ID NO: 24)
IgG4 hinge (S→P) (S228P)	12 aa	ESKYGPPCPPCP (SEQ ID NO: 25)
IgG4 hinge	12 aa	ESKYGPPCPSCP (SEQ ID NO: 26)
IgG4 hinge (S228P) + linker Also called HL	22 aa	ESKYGPPCPPCPGGSSGGGSG (SEQ ID NO: 27)
CD28 hinge	39 aa	IEVMYPPPYLDNEKSNGTIIHVKGKHLCPSPLPFGPSKP (SEQ ID NO: 28)
CD8 hinge-48aa	48 aa	AKPTTTPAPRPPTPAPTIASQPLSLRPEACRP AAGGAVHTRGLDFACD (SEQ ID NO: 29)
CD8 hinge-45aa	45 aa	TTTPAPRPPTPAPTIASQPLSLRPEACRPAAG GAVHTRGLDFACD (SEQ ID NO: 30)
IgG4 (HL-CH3) Also called: IgG4 (HL-ACH2) (includes S228P in hinge)	129 aa	ESKYGPPCPPCPGGSSGGGSGGQPREPQV YTLPPSQEEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTTPVLDSDGSFFLYSRL TVDKSRWQEGNVFSCSVMEALHNHYTQKS LSLSLGK (SEQ ID NO: 31)

TABLE 2-continued

Examples of Spacers		
Name	Length	Sequence
IgG4 (L235E, N297Q)	229 aa	ESKYGPPCPSCPAPEFEGGSPSVFLFPPKPKD TLMISRTPEVTCVVVDVSQEDPEVQFNWYVD GVEVHNAKTKPREEQFQSTYRVVSVLTVLHQ DWLNGKEYKCKVSNKGLPSSIEKTIKAKGQP REPQVYTLPPSQEEMTKNQVSLTCLVKGFYP SDIAVEWESNGQPENNYKTTPVLDSDGSFF LYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 32)
IgG4 (S228P, L235E, N297Q)	229 aa	ESKYGPPCPPCPAPEFEGGSPSVFLFPPKPKD TLMISRTPEVTCVVVDVSQEDPEVQFNWYVD GVEVHNAKTKPREEQFQSTYRVVSVLTVLHQ DWLNGKEYKCKVSNKGLPSSIEKTIKAKGQP REPQVYTLPPSQEEMTKNQVSLTCLVKGFYP SDIAVEWESNGQPENNYKTTPVLDSDGSFF LYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 33)
IgG4 (CH3) Also called IgG4 (ACH2)	107 aa	GQPREPQVYTLPPSQEEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPVLDSDG SFPLYSLRTVDKSRWQEGNVFSCSVMEALH NHYTQKSLSLSLGK (SEQ ID NO: 34)

[0034] Some spacer regions include all or part of an immunoglobulin (e.g., IgG1, IgG2, IgG3, IgG4) hinge region, i.e., the sequence that falls between the CH1 and CH2 domains of an immunoglobulin, e.g., an IgG4 Fc hinge or a CD8 hinge. Some spacer regions include an immunoglobulin CH3 domain (called CH3 or ΔCH2) or both a CH3 domain and a CH2 domain. The immunoglobulin derived sequences can include one or more amino acid modifications, for example, 1, 2, 3, 4 or 5 substitutions, e.g., substitutions that reduce off-target binding.

[0035] The hinge/linker region can also comprise an IgG4 hinge region having the sequence ESKYGPPCPSCP (SEQ ID NO: 26) or ESKYGPPCPPCP (SEQ ID NO: 25). The hinge/linker region can also comprise the sequence ESKYGPPCPPCP (SEQ ID NO: 25) followed by the linker sequence GGGSSGGGSG (SEQ ID NO: 24) followed by IgG4 CH3 sequence: GQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNHNYTQKSLSLSLGK (SEQ ID NO: 34). Thus, the entire linker/spacer region can comprise the sequence: ESKYGPPCPPCPGGSSGGGSGGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNHNYTQKSLSLSLGK (SEQ ID NO: 31). In some cases, the spacer has 1, 2, 3, 4, or 5 single amino acid changes (e.g., conservative changes) compared

to SEQ ID NO: 31. In some cases, the IgG4 Fc hinge/linker region that is mutated at two positions (L235E; N297Q) in a manner that reduces binding by Fc receptors (FcRs).

(d) Intracellular Signaling Domains

[0036] A CAR constructs described herein contains one or more intracellular signaling domains (e.g., CD3ζ, and optionally one or more co-stimulatory domains), which are the functional end of the receptor. Following antigen recognition, receptors cluster and a signal is transmitted to the cell.

[0037] CD3ζ is the cytoplasmic signaling domain of the T cell receptor complex. CD3ζ contains three immunoreceptor tyrosine-based activation motifs (ITAMs), which transmit an activation signal to the cell after the cell is engaged with a cognate antigen. In some cases, CD3ζ provides a primary cell activation signal but not a fully competent activation signal, which requires a co-stimulatory signal.

[0038] Accordingly, in some examples, the CAR polypeptides disclosed herein may further comprise one or more co-stimulatory signaling domains in addition to CD3ζ. For example, the co-stimulatory domain CD28 and/or 4-1BB can be used to transmit a proliferative/survival signal together with the primary signaling mediated by CD3ζ. The co-stimulatory domain(s) are located between the transmembrane domain and the CD3ζ signaling domain. Table 3 includes examples of suitable co-stimulatory domains together with the sequence of the CD3ζ signaling domain.

TABLE 3

CD3ζ Domain and Examples of Co-stimulatory Domains			
Name	Accession	Length	Sequence
CD3ζ	J04132.1	113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPFEMGGKPRRKNPQEGLYNELQKDK MAEAYSEIGMKGERRRGKHGDL YQGL STATK DTYDALHMQALPPR (SEQ ID NO: 35) ITAMs 1-3 underlined

TABLE 3-continued

CD3 ζ Domain and Examples of Co-stimulatory Domains			
Name	Accession	Length	Sequence
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLFNQLQKDK MAEAFSEIGMGERRRGKGGHDLFQGLSTATK DTFDALHMQALPPR (SEQ ID NO: 50)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLFNQLQKDK MAEAFSEIGMGERRRGKGGHDLFQGLSTATK DTYDALHMQALPPR (SEQ ID NO: 51)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLYNELQKDK MAEAYSEIGMGERRRGKGGHDLFQGLSTATK DTFDALHMQALPPR (SEQ ID NO: 52)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLYNELQKDK MAEAFSEIGMGERRRGKGGHDLFQGLSTATK DTFDALHMQALPPR (SEQ ID NO: 53)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLFNQLQKDK MAEAYSEIGMGERRRGKGGHDLFQGLSTATK DTFDALHMQALPPR (SEQ ID NO: 54)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLFNQLQKDK MAEAFSEIGMGERRRGKGGHDLFQGLSTATK DTFDALHMQALPPR (SEQ ID NO: 55)
CD3 ζ variant		113 aa	RVKFSRSADAPAYQQGQNQLYNELNLGRREEYD VLDKRRGRDPEMGGKPRRKNPQEGLFNQLQKDK MAEAFSEIGMGERRRGKGGHDLFQGLSTATK DTYDALHMQALPPR (SEQ ID NO: 56)
CD28	NM_006139	42 aa	RSKRSLHSDYMNMTPRRPGPTRKHYQPYAPP RDFAAYRS (SEQ ID NO: 36)
CD28g g*	NM_006139	42 aa	RSKRSGGHSYMNMTPRRPGPTRKHYQPYAP PRDFAAYRS (SEQ ID NO: 37)
41BB	NM_001561	42 aa	KRGRKKLLYIFKQPFMRPVQTQEEEDGSCRFPP EEEEGGCEL (SEQ ID NO: 38)
OX40	NM_003327	42 aa	ALYLLRRDQRLPPDAHKPPGGGSFRTPIQEEQA DAHSTLAKI (SEQ ID NO: 39)
2B4	NM_016382	120 aa	WRRKRKEKQSETSPKEFLTIYEDVKDLKTRRNHE QEQTFFGGGSTIYSMIQSSAPTSQEPAYTLYS LIQPSRKSGSRKRNHSPSFNSTIYEVIGKSQPKA QNPARLSRKELENFVYVS (SEQ ID NO: 40)

[0039] In some examples, the CD3 ζ signaling domain comprises an amino acid sequence that is at least 90%, at least 95%, at least 98% identical to SEQ ID NO: 35. In such instances, the CD3 ζ signaling domain has 1, 2, 3, 4, or 5 amino acid changes (preferably conservative substitutions) compared to SEQ ID NO: 35. In other examples, the CD3 ζ signaling domain is SEQ ID NO: 35 or a CD3 ζ variant (e.g., any of SEQ ID NOs: 50-56).

[0040] In various embodiments: the co-stimulatory domain is selected from the group consisting of: a co-stimulatory domain depicted in Table 3 or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications, a CD28 or CD28gg co-stimulatory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications, a 4-1BB co-stimulatory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications and an OX40 co-stimula-

tory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications. In certain embodiments, a 4-1BB co-stimulatory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications is present in the CAR polypeptides described herein.

[0041] In some embodiments, there are two co-stimulatory domains, for example, a CD28 co-stimulatory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications (e.g., substitutions) and a 4-1BB co-stimulatory domain or a variant thereof having 1-5 (e.g., 1 or 2) amino acid modifications (e.g., substitutions). In various embodiments the 1-5 (e.g., 1 or 2) amino acid modification are substitutions. In various embodiments, the co-stimulatory domain is amino terminal to the CD3 ζ signaling domain and a short linker consisting of 2-10, e.g., 3 amino acids (e.g., GGG) is can be positioned between the co-stimulatory domain and the CD3 ζ signaling domain.

II. Preparation of CAR NK Cells

[0042] In some cases, the CAR can be produced using a vector in which the CAR open reading frame is followed by a T2A ribosome skip sequence and a truncated EGFR (EGFRt), which lacks the cytoplasmic signaling tail, or a truncated CD19R (also called CD19t). In this arrangement, co-expression of EGFRt or CD19t provides an inert, non-immunogenic surface marker that allows for accurate measurement of gene modified cells, and enables positive selection of gene-modified cells, as well as efficient cell tracking of the therapeutic immune cells in vivo following adoptive transfer. Efficiently controlling proliferation to avoid cytokine storm and off-target toxicity is an important hurdle for the success of immune cell immunotherapy. The EGFRt or the CD19t incorporated in the CAR lentiviral vector can act as suicide gene to ablate the CAR+ NK cells in cases of treatment-related toxicity.

[0043] The CAR, including the ODD, can be followed by a ribosomal skip sequence (e.g., GSGATNFSLLKQAGDVEENPGP (SEQ ID NO: 14) or LEGGGEGRGSLTTCGDVEENPGPR; SEQ ID NO: 45) and a truncated CD19 receptor (also called CD19t) having a sequence that is at least 90%, at least 95%, at least 98% identical to or identical to:

(SEQ ID NO: 64)

MPPPRLLFFLLFLTPMEVRPEEPLVVKVEEGDNAVLQCLKGTSDG
 PTQQLTWSRESPLKPKLKLGLPGLGIHMRPLAIWLFIFNVSQQ
 MGGFYLCQPGPPSEKAWQPGWTVNVEGSGELFRWNVSDLGGLGCG
 LKNRSSEGPSSPSGKLMSPKLYVWAKDRPEIWEGEPPCPVPRDSL
 NQLSQDLTMAPGSTLWLSCGVPPDSVSRGPLSWTHVHPKPKSL
 LSLELKDDRPARDMMVMETGLLLPRATAQDAGKYCHRGNTMSF
 HLEITARPVLWHWLLRTGGWKVSAVTLAYLIFCLCSLVGILHLQR
 ALVLRKR.

[0044] Any CAR described herein can be produced by any means known in the art, though preferably it is produced using recombinant DNA techniques. Nucleic acids encoding the several regions of the chimeric receptor can be prepared and assembled into a complete coding sequence by standard techniques of molecular cloning known in the art (genomic library screening, overlapping PCR, primer-assisted ligation, site-directed mutagenesis, etc.) as is convenient. The resulting coding region can be inserted into an expression vector and used to transform a suitable expression host cell line. A suitable host cell line includes, for example, an NK cell. An expression vector encoding a CAR or polypeptide described herein can be a viral vector. Suitable viral vectors, including lentiviral vectors, are known in the art and can be used in any of the methods described herein. In some aspects, any of the transduced immune cells described herein can be autologous or allogenic. For example, suitable cell populations can include allogenic NK cells, autologous NK cells, allogenic NK cells and autogenic NK cells.

[0045] The CAR can be transiently expressed in an NK cell population by an mRNA encoding the CAR. The mRNA can be introduced into immune cells by electroporation (Wiesinger et al. 2019 *Cancers* (Basel) 11:1198).

[0046] Disclosed herein, amongst other things, are methods of making any CAR NK II described herein by introducing into a NK cell a vector comprising a nucleotide sequence encoding the CAR. Disclosed herein, amongst other things, are methods of making a population of NK cells comprising a nucleic acid encoding any CAR or polypeptide described herein.

IV. Treatment of Cancer

[0047] Aspects of the present disclosure provide methods for treating a subject having breast cancer by administering immune cells expressing a CAR described here.

(a) Subjects

[0048] A subject to be treated by the methods described can be a human patient having a breast cancer with low or no HER2 expression, e.g., breast cancer that is HER2 immunohistochemistry (IHC) 1+ or 2+ with negative fluorescence in situ hybridization (FISH).

(b) Treatment Regimens

[0049] Aspects of the present disclosure provide methods of treating breast cancer comprising administering a lymphodepletion treatment (e.g., cyclophosphamide) in combination with CAR immune cells, which can be administered locally or systemically. Generally, the methods include administering a therapeutically effective amount of a population of CAR NK cells as described herein, to a subject who is in need of, or who has been determined to be in need of, such treatment. The population of EGFR CAR_{sIL}-15 immune cells in all compositions and methods disclosed herein can be autologous or allogenic.

[0050] Any subject suitable for the treatment methods described herein can receive a lymphodepleting therapy to reduce or deplete the endogenous lymphocytes of the subject. Lymphodepletion refers to the destruction of endogenous lymphocytes, which is commonly used prior to immunotransplantation and immunotherapy. Lymphodepletion can be achieved by administering a lymphodepleting agent and/or irradiation (e.g., stereotactic radiation). A lymphodepleting agent can be any molecule capable of reducing, depleting, or eliminating endogenous lymphocytes when administered to a subject. In some examples, the lymphodepleting agents are cytotoxic agents that specifically kill lymphocytes. Non-limiting examples of lymphodepleting agents include cyclophosphamide, fludarabine, gemcitabine, methotrexate, doxorubicin, and etoposide phosphate. In some cases the lymphodepletion treatment is non-myeloablative.

[0051] Methods described herein can include a conditioning regimen comprising a single lymphodepleting agent (e.g., cyclophosphamide) or multiple lymphodepleting agents (e.g., cyclophosphamide and fludarabine). The subject to be treated by the methods described herein can receive one or more doses of the one or more lymphodepleting agents for a period suitable for reducing or depleting the endogenous lymphocytes of the subject (e.g., 1-5 days).

[0052] The subject can then be administered any of the CAR immune cells described herein after administration of the lymphodepleting therapy as described herein. For example, the one or more lymphodepleting agents can be administered to the subject 1-5 days (e.g., 1, 2, 3, 4, or 5 days) prior to administering the CAR immune cells.

[0053] Methods described herein can include redosing the subject with CAR immune cells. In some examples, the subject is administered a lymphodepleting treatment prior to redosing of the anti-CAR immune cells. Each dose of the CAR immune cells can be the same or the doses can be ascending or descending.

[0054] Methods described herein can include redosing the subject with CAR immune cells. In some examples, the subject is administered 3-6 doses of the CAR immune cells, each of which is administered 1-15 days after the prior dose. Each dose of CAR immune cells can be the same or the doses can be ascending or descending.

(c) Administration

[0055] An effective amount of a therapy (e.g., EGFR-CAR_sIL15 NK cells) can be administered to a subject (e.g., a human) in need of the treatment via any suitable route (e.g., administered locally or systemically to a subject). Suitable modes of administration include injection, infusion, instillation, or ingestion. Injection includes, without limitation, intravenous, intramuscular, intra-arterial, intrathecal, intraventricular, intradermal, intraperitoneal, and subcutaneous injection and infusion.

[0056] An effective amount, or therapeutically effective amount, refers to the amount of each active agent required to confer therapeutic effect on the subject, either alone or in combination with one or more other active agents. Effective amounts vary, as recognized by those skilled in the art, depending on the particular condition being treated, the severity of the condition, the individual patient parameters including age, physical condition, size, gender and weight, the duration of treatment, the nature of concurrent therapy, if any, the specific route of administration and like factors within the knowledge and expertise of the health practitioner. The amelioration of one symptom associated with the condition, cancer, or disease is enough to confer therapeutic effect on the subject. These factors are well known to those of ordinary skill in the art and can be addressed with no more than routine experimentation. It is generally preferred that a maximum dose of the individual components or combinations thereof be used, that is, the highest safe dose according to sound medical judgment. It will be understood by those of ordinary skill in the art, however, that a patient may insist upon a lower dose or tolerable dose for medical reasons, psychological reasons or for virtually any other reasons.

[0057] Disclosed herein, amongst other things, methods of administering to a subject in need thereof (e.g., a subject having breast cancer), a therapeutic amount of any disclosed cell population comprising a nucleic acid encoding an EGFR-CAR_sIL15 described herein. Disclosed herein, amongst other things, methods of administering to a subject in need thereof (e.g., a subject having breast cancer), a therapeutic amount of any disclosed cell population expressing any an EGFR-CAR_sIL15 described herein.

[0058] The materials, methods, and examples are illustrative only and not intended to be limiting. All publications, patent applications, patents, sequences, database entries, and other references mentioned herein are incorporated by reference in their entirety for any and all purposes.

[0059] Other features and advantages of the described compositions and methods will be apparent from the following detailed description and figures, and from the claims.

DESCRIPTION OF DRAWINGS

[0060] FIG. 1 depicts expression of EGFR and HER2 in breast cancer. EGFR and HER2 immunohistochemistry of tumor tissue from patients with primary breast cancer using an anti-human EGFR antibody. Representative data of 3 out of 24 patients are presented.

[0061] FIG. 2 depicts a schematic of the structure of an EGFR-CAR_sIL15 CAR. SP, signal peptide; TM, trans-membrane domain; ODD, oxygen dependent degradation domain of HIF1 α ; T2A, ribosomal skip sequence; Codon Opm I15, Codon-optimized soluble IL-15; P2A, porcine teschovirus 2A ribosomal skip sequence; tCD19, truncated CD19 receptor.

[0062] FIG. 3 depicts the results of an analysis of expression of EGFR CAR_sIL15 and control vector transduction of UCB-derived NK cells as assessed by tCD9 expression.

[0063] FIGS. 4A-4B depict the result of an in vitro functional evaluation of EGFR-CAR_sIL15 NK cells. (A-B) Cytotoxicity against MDA-MB-231 and MCF-7 (EGFR+) cells (A) and A2780 (EGFR-) (B) cells were tested after co-culture of EGFR_sIL15-CAR NK cells, NK cells expressing IL-15 (sIL15NK), or unmodified NK cells at different E:T ratios analyzed using ^{51}Cr assay.

[0064] FIGS. 5A-5B depict the results of an in vitro functional evaluation of EGFR-CAR_sIL15 NK cells under hypoxia. (A) Cytotoxicity against MDA-MB-231 (EGFR+) cells was tested after co-culture of EGFR_sIL15-CAR NK cells or unmodified NK cells under hypoxia or normoxia condition at different effector (E):target (T) ratios analyzed using ^{51}Cr assay. (B) The cytotoxicity was analyzed by microscope under hypoxia conditions at E:T=2:1. PI was used to stain dead cells (bright red).

[0065] FIGS. 6A-6D depict the results of a study showing that EGFR-CAR_sIL15 NK cells improve tumor rejection and prolong survival of mice bearing breast tumor that metastasized to lung. (A) Schematic of metastatic breast cancer model (5×10^5 cells). (B) Time-lapse luciferase imaging of breast cancer tumor mice with indicated treatments (4×10^6 CAR NK cells/infusion). (C) Quantification of bioluminescence images. (D) Survival analysis of the breast tumor model that metastasized to the lung.

DETAILED DESCRIPTION

Examples

[0066] The invention is further described in the following examples, which do not limit the scope of the invention described in the claims.

Example 1: Expansion and Transduction of Natural Killer Cells

[0067] We used an efficient retrovirus-based transduction system to optimize the manufacturing of CAR NK cells. Briefly, we expanded UCB NK cells on a K562 feeder cell layer expressing membrane-bound interleukin-21 (IL-21) and 4-1BB ligand (4-1BBL). The expanded NK cells were infected with a replication-defective retroviral vector expressing EGFR-CAR_sIL15 prior to further expansion. This platform improved transduction efficiency and survival of NK cells¹¹. Using our system, we achieved ~60-70% transduction of human NK cells, which is comparable to the transduction of T cells with the same retrovirus expressing a CAR (data not shown). As we incorporated sIL-15 into our

vector to enhance survival of CAR NK cells in vivo, for the control, we also generated a vector expressing sIL-15 and tEGFR without the EGFR-CAR (FIG. 3).

Example 2: Assessment of Cytotoxicity

[0068] We tested EGFR-CAR_sIL15 NK cells for cytotoxicity against the two MDA-MB-231 and MCF-7 BC cell lines, both of which are EGFR⁺ and HER2⁻. To do this, we co-cultured EGFR-CAR_sIL15 NK cells, NK cells only expressing sIL15 (sIL15 NK) and unmodified control NK cells, with tumor cells at different effector:target (E:T) ratios. We analyzed cytotoxicity using ⁵¹Cr assay. EGFR-CAR_sIL15 NK cells killed BC cells substantially better than sIL15 NK and unmodified control NK cells (FIG. 4A). We also measured the cytotoxicity against A2780 cells, which are EGFR⁻. There was no cytotoxicity difference between unmodified control NK cells and EGFR-CAR_sIL15 NK cells targeting EGFR-cells (FIG. 4B).

Example 3: Assessment of Activity in Hypoxia and Normoxia

[0069] Hypoxia is a characteristic feature shared by most advanced solid tumors, which promotes a heterogeneous landscape of oxygen levels and metabolic activities. It has been reported that hypoxia is a physical cue for licensing CAR-T cell or immune cell activation, specifically in the tumor microenvironment. Therefore, we tested EGFR-CAR_sIL15 NK cells for cytotoxicity against the EGFR MDA-MB-231 BC cell line under hypoxia conditions. Due to the ODD in the construct, EGFR-CAR_sIL15 NK cells

killed BC cells substantially better under the hypoxia condition than that under the normoxia condition. In contrast, there is no difference for nontransduced NK cells with a very low level of cytotoxicity at both the normoxia and hypoxia conditions. The data indicated that NK cells engineered with the EGFR-CAR_sIL15 vector fused with ODD could be safe and effective in targeting solid tumors (FIGS. 5A-5B).

[0070] To test the efficacy of EGFR-CAR_sIL15 NK cells in vivo, we first injected (iv) 5×10⁵ MDA-MB-231 cells expressing luc per mouse to establish a metastatic BC mouse model (FIG. 6A). We manufactured frozen, off-the-shelf EGFR-CAR_sIL15 NK cells and sIL15 control NK cells from a same UCB unit. Two days after tumor implantation, we randomized mice into three groups that received three cycles of PBS, off-the-shelf frozen EGFR-CAR_sIL15 NK cells, or control sIL15 NK cells given intravenously. We monitored tumor progression using luc-based imaging starting 7 days post-tumor implantation. EGFR-CAR_sIL15 NK cells were significantly more effective than the other two groups at improving the rejection of breast tumors and survival (FIGS. 6B-6D).

OTHER EMBODIMENTS

[0071] It is to be understood that while the invention has been described in conjunction with the detailed description thereof, the foregoing description is intended to illustrate and not limit the scope of the invention, which is defined by the scope of the appended claims. Other aspects, advantages, and modifications are within the scope of the following claims.

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GGGGSGGGGS QVQLQQSGSE MARPGASVKL PCKASGDTFT SYWMHWVKQR HGHGPEWIGN 180
IYPGSGGTNY AEKFKNKVTL TVDRSSRTVY MHL SRLTSED SAVYYCTRS GPFYFDYWGQ 240
GTTTLTVSS          248

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	mol_type = protein	
	organism = synthetic construct	
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SYWMH		5
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	mol_type = protein	
	organism = synthetic construct	
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SEQ ID NO: 9	moltype = AA length = 9	
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	mol_type = protein	
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SGGPYFFDY		9
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	mol_type = protein	
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APAAGDTIIS LDFGSNDTET DDQQLLEVPL YNDVMLPSPN EKLQINILAM SPLPTAETPK		60
PLRSSADPAL NQEVALKLEP NPESLELSFT MPQIQDQTPS PSDGSTRQSS PEPNSPSEYC		120
FVVDSDMVNE FKLELVEKLF AEDTEAKNPF STQDTDLDE MLAPYIPMDD DFQLRSFDQL		180
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CFLLELQVIS LESGDASIHD TVENLIILAN NSLSSNGNVT ESGCKECEEL EEKNIKEFLQ		120
SFVHIVQMFI NTS		133
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	mol_type = protein	
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SEQUENCE: 14		
GSGATNF SLL KQAGDVEENP GP		22
SEQ ID NO: 15	moltype = AA length = 21	
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source	1..21	
	mol_type = protein	
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SEQUENCE: 15		
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SEQ ID NO: 16	moltype = AA length = 27	
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	mol_type = protein	
	organism = synthetic construct	
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SEQ ID NO: 17	moltype = AA length = 28	
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SEQUENCE: 17		
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	mol_type = protein	
	organism = synthetic construct	
SEQUENCE: 18		
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SEQ ID NO: 19	moltype = AA length = 21	
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source	1..21	
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SEQUENCE: 19		
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SEQ ID NO: 20	moltype = AA length = 23	
FEATURE	Location/Qualifiers	
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SEQUENCE: 20		
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SEQ ID NO: 21	moltype = AA length = 24	
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source	1..24	
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source	1..27	
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	organism = synthetic construct	
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source	1..10	
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SEQ ID NO: 25	moltype = AA length = 12	
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FEATURE	Location/Qualifiers	
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SEQUENCE: 26		
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SEQ ID NO: 27	moltype = AA length = 22	
FEATURE	Location/Qualifiers	
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SEQUENCE: 27		
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SEQUENCE: 28		
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source	1..48	
	mol_type = protein	
	organism = synthetic construct	
SEQUENCE: 29		
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SEQ ID NO: 30	moltype = AA length = 45	
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source	1..45	
	mol_type = protein	
	organism = synthetic construct	
SEQUENCE: 30		
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SEQ ID NO: 31	moltype = AA length = 129	
FEATURE	Location/Qualifiers	
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	mol_type = protein	
	organism = synthetic construct	
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VEWESNGQPE NNYKTPPV L DSDGSFFLYS RLTVDKSRWQ EGNVFSCSV M HEALHNHYTQ		120
KSLSLSLGK		129
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FEATURE	Location/Qualifiers	

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DSDGSFFLYS RLTVDKSRWQ EGNVFSCSV MHEALHNHYTQ KSLSLSLGK 229

SEQ ID NO: 33 moltype = AA length = 229
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AKGQPREPQV YTLPPSQEEM TKNQVSLTCL VKGFYPSDIA VEWESNGQPE NNYKTTTPVL 180
DSDGSFFLYS RLTVDKSRWQ EGNVFSCSV MHEALHNHYTQ KSLSLSLGK 229

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DGSFFLYSRL TVDKSRWQEG NVFSCSV MHEALHNHYTQ KSLSLSLGK 107

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SEQ ID NO: 37 moltype = AA length = 41
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SEQUENCE: 37
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SEQ ID NO: 38 moltype = AA length = 42
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mol_type = protein
organism = synthetic construct

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SEQ ID NO: 39 moltype = AA length = 42
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mol_type = protein
organism = synthetic construct

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LFRWNVSDLG	GLGCGLKNRS	SEGPSSPSGK	LMSPKLYVWA	KDRPEIWEGE	PPCLPPRDSL	180
NQSLSQDLTM	APGSTLWLSC	GVPPDSVSRG	PLSWTHVHPK	GPKSLLSLEL	KDRPARDMW	240
VMETGLLLPR	ATAQDAGKY	CHRGNLTMSF	HLEITARPV	WHWLLRTGGW	KVSAVTLAYL	300
IFCLCSLVGI	LHLQRALVLR	RKR			323	
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	mol_type = protein					
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SEQUENCE: 50						
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SEQUENCE: 52						

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ELQKDKMAEA	FSEIGMKGER	RRGKGHDGLF	QGLSTATKDT	FDALHMQALP	PR	112

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ELQKDKMAEA	FSEIGMKGER	RRGKGHDGLY	QGLSTATKDT	FDALHMQALP	PR	112

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SGVPDRFSGS	GSGETDFTLKI	SRVEAEDLGI	YYCFQGSHP	PTFGGGTKLE	IKRAAGGGGS	120
GGGSGGGGS	QVQLQSGSE	MARPGASVKL	PCKASGDTFT	SYWMHWVKQR	HGHGPEWIGN	180
IYPGSGGTNY	AEKFKNKVTL	TVDRSSRTVY	MHLSRLTSED	SAVYYCTRS	GPYFFDYWGQ	240
GTTLTVSSPK	SCDKTHTCPP	CPDPKFWVLV	VVGGVLACYS	LLVTVAFIIF	WVRSKRSRL	300
HSDYMMNTPR	RPGPTRKHYQ	PYAPPRDFAA	YRSRVKFSRS	ADAPAYQQGQ	NQLYNELNLG	360
RREEYDVLDK	RRGRDPEMGG	KPRRKNPQEG	LYNELQKDKM	AEAYSEIGMK	GERRRGKGHD	420
GLYQGLSTAT	KDTYDALHMQ	ALPPRAPAAG	DTIISLDLFGS	NDTETDDQQL	EEVPLYNDVM	480
LPSPNEKLQN	INLAMSPLPT	AETPKPLRSS	ADPALNQEVA	LKLEPNPESL	ELSFTMPQIQ	540
DQTPSPSDGS	TRQSSPEPNS	PSEYCFYVDS	DMVNEFKLEL	VEKLFAEDTE	AKNPFSTQDT	600
DLGLEMLAPY	IPMDDDFQLR	SFDQLSPLES	SSASPESASP	QSTVTVFQ		648

SEQ ID NO: 58 moltype = AA length = 820
FEATURE Location/Qualifiers
source 1..820
 mol_type = protein
 organism = synthetic construct

SEQUENCE: 58

DILMTQSPLS	LPVSLGDQAS	ISCRSSQNIV	HNNGITYLEW	YLQRPQGSPK	LLIYKVSDRF	60
SGVPDRFSGS	GSGETDFTLKI	SRVEAEDLGI	YYCFQGSHP	PTFGGGTKLE	IKRAAGGGGS	120
GGGSGGGGS	QVQLQSGSE	MARPGASVKL	PCKASGDTFT	SYWMHWVKQR	HGHGPEWIGN	180
IYPGSGGTNY	AEKFKNKVTL	TVDRSSRTVY	MHLSRLTSED	SAVYYCTRS	GPYFFDYWGQ	240
GTTLTVSSPK	SCDKTHTCPP	CPDPKFWVLV	VVGGVLACYS	LLVTVAFIIF	WVRSKRSRL	300
HSDYMMNTPR	RPGPTRKHYQ	PYAPPRDFAA	YRSRVKFSRS	ADAPAYQQGQ	NQLYNELNLG	360
RREEYDVLDK	RRGRDPEMGG	KPRRKNPQEG	LYNELQKDKM	AEAYSEIGMK	GERRRGKGHD	420
GLYQGLSTAT	KDTYDALHMQ	ALPPRAPAAG	DTIISLDLFGS	NDTETDDQQL	EEVPLYNDVM	480
LPSPNEKLQN	INLAMSPLPT	AETPKPLRSS	ADPALNQEVA	LKLEPNPESL	ELSFTMPQIQ	540
DQTPSPSDGS	TRQSSPEPNS	PSEYCFYVDS	DMVNEFKLEL	VEKLFAEDTE	AKNPFSTQDT	600
DLGLEMLAPY	IPMDDDFQLR	SFDQLSPLES	SSASPESASP	QSTVTVFQAE	GRGSLTTCGD	660
VEENPGPMYR	MQLLSIALS	LALVTNSGIH	VFILGCFASG	LPKTEANWVN	VISDLKKIED	720
LIQSMHIDAT	LYTESDVHPS	CKVTAMKCFL	LELQVISLES	GDAISIHDTVE	NLIILANNSL	780

-continued

SSNGNVTESG CKECEELEEK NIKEFLQSFV HIVQMFINTS				820	
SEQ ID NO: 59	moltype = AA length = 17				
FEATURE	Location/Qualifiers				
source	1..17				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 59					
PKSCDKTHTC PPCDPK				17	
SEQ ID NO: 60	moltype = AA length = 15				
FEATURE	Location/Qualifiers				
source	1..15				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 60					
GGGSGGGGS GGGGS				15	
SEQ ID NO: 61	moltype = AA length = 10				
FEATURE	Location/Qualifiers				
source	1..10				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 61					
GGGSGGGGS				10	
SEQ ID NO: 62	moltype = AA length = 22				
FEATURE	Location/Qualifiers				
source	1..22				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 62					
MLLVTSLLL CELPHPAFLL IP				22	
SEQ ID NO: 63	moltype = AA length = 20				
FEATURE	Location/Qualifiers				
source	1..20				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 63					
MYRMQLLSCI ALSLALVTNS				20	
SEQ ID NO: 64	moltype = AA length = 323				
FEATURE	Location/Qualifiers				
source	1..323				
	mol_type = protein				
	organism = synthetic construct				
SEQUENCE: 64					
MPPPRLLFFL LFLTPMEVRP	EEPLVVKVEE	GDNVAVLQCLK	GTSDGPTQQL	TWSRESPLKP	60
FLKLSLGLPG LGIHMRLPLAI	WLFIFNVSSQ	MGGFYLCQPG	PPSEKAWQPG	WTVNVEGSGE	120
LFRWNVSDLG GLGCGLKNRS	SEGPPSPSGK	LMSPKLYVWA	KDRPEIWEGE	PPCVPPRDSL	180
NQSLSQDLTM APGSTLWLSC	GVPPDSVSRG	PLSWTHVHPK	GPKSLLSLEL	KDDRPARDMW	240
VMETGLLLPR ATAQDAGKYY	CHRGNTMSF	HLEITARPVL	WHWLLRTGGW	KVSAVTLAYL	300
IFCLCSLVGI LHLORALVLR	RKR				323

1. A nucleic acid molecule encoding a fusion protein comprising a chimeric antigen receptor and an oxygen-dependent degradation domain (ODD), wherein the chimeric antigen receptor comprises:

- (i) an scFv that binds human EGFR;
- (ii) a spacer domain;
- (iii) a transmembrane domain;
- (iv) a costimulatory domain; and
- (v) a CD3 ζ signaling domain;

wherein the scFv comprises: a light chain CDR1 comprising RSSQNIVHNNGITYLE (SEQ ID NO: 4), a light chain CDR2 comprising KVSDRFs (SEQ ID NO: 5), a light chain CDR3 comprising FQGSHIPPT (SEQ ID NO: 6), a heavy chain CDR1 comprising SYWMH (SEQ ID NO: 7), a heavy chain CDR2 comprising

NIYPGSGGTNYAEKFKN (SEQ ID NO: 8), and a heavy chain CDR3 comprising SGGPYFFDY (SEQ ID NO: 9).

2. (canceled)

3. The nucleic acid molecule of claim 1, wherein the ODD is a human HIF1 α ODD.

4. The nucleic acid molecule of claim 3, wherein the ODD comprises or consists of

(SEQ ID NO: 10)
 APAAGDTIISLDFGSDNTETDDQQLVEEVPLYNDVMLPSPNEKLQN
 INLAMSPPLTAETPKPLRSSADPALNQEVALKLEPNPSELELSFT
 MPQIQDQTPSPSDGSTRQSSPEPNSPSEYCFYVDSMDVNEFKLEL

-continued

VEKLFAEDTEAKNPFSTQDLDLEMLAPYIPMDDDFQLRSFDQL
SPLESSASPESASPQSTVTVFQ.

5. The nucleic acid molecule of claim 1, wherein the scFv comprises a VL domain comprising: DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGI-TYLEWYLQRPQGSPKLLIYKVSDFRS GVPDRFSGSGSGTDFTLKISRVEAEDLGIYYCFQG-SHIPPTFGGGTKLEIKRAA (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions followed by a VH domain comprising or consisting of QVQLQQSGSEMARPGASVKLPCKASGDTFT-SYWMHWVKQRHGHGPEWIGNIYPGSGG TNY-AEKFKNKVTLTVDRSSRTVYMHLRLTSEDSAVYYC-TRSGGPYFFDYWGQGTTLT VSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions; optionally wherein the amino acid substitutions are not in the CDRs.

6. The nucleic acid molecule of claim 1, where the scFv comprises a VL domain comprising or consisting of an amino acid sequence at least 95%, 98% or 99% identical to: DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGI-TYLEWYLQRPQGSPKLLIYKVSDFRS GVPDRFSGSGSGTDFTLKISRVEAEDLGIYYCFQG-SHIPPTFGGGTKLEIKRAA (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions followed by a VH domain comprising or consisting of an amino acid sequence at least 95%, 98% or 99% identical to QVQLQQSGSEMARPGASVKLPCKASGDTFT-SYWMHWVKQRHGHGPEWIGNIYPGSGG TNY-AEKFKNKVTLTVDRSSRTVYMHLRLTSEDSAVYYC-TRSGGPYFFDYWGQGTTLT VSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions; optionally wherein the amino acid substitutions are not in the CDRs.

7. (canceled)

8. The nucleic acid molecule of claim 1, wherein the chimeric antigen receptor comprises: the spacer comprises or consists of a sequence selected from the group consisting of: SEQ ID NOs: 59 and 24-34; the transmembrane domain comprises or consists of a sequence selected from the group consisting of SEQ ID NOs: 15-23; the costimulatory domain comprises or consists of a sequence from the group consisting of SEQ ID NOs: 36-40, and the CD3 ζ signaling domain comprises or consists of SEQ ID NO: 35.

9. The nucleic acid molecule of claim 1, wherein the nucleic acid molecule further encodes human IL-15 or a soluble variant of human IL-15.

10. The nucleic acid molecule of claim 1, wherein the nucleic acid molecule further encodes human IL-15 or a soluble variant of human IL-15 that is co-expressed with the fusion protein.

11. The nucleic acid molecule of claim 9, wherein the soluble variant of IL-15 comprises or consists of: GHV-FILGCFSAGLPKTEANWVNVISDLKKIEDLIQSMHI-DATLYTESDVHPSCKVTAMK CFLLELQVISLESGLDASIHDTVENLILANNSLSSNGNVTESGCKECELEEKNI-KEFLQSF VHIVQMFINTS (SEQ ID NO:11).

12.-13. (canceled)

14. The nucleic acid molecule of claim 1 encoding an amino acid sequence comprising or consisting of the amino acid sequence:

(SEQ ID NO: 57 [[A]])

DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRP
GQSPKLLIYKVSDFRS GVPDRFSGSGSGTDFTLKISRVEAEDLGI
YYCFQGSHIPPTFGGGTKLEIKRAAGGGSGGGSGGGSGVQVQLQ
QSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHGPEWIGN
IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLRLTSEDSAVYY
CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
VVGGLACYSLLVTVAFIIFWVRSKRSLRLHSDYMNMTPRRPGPT
RKHYQPYAPPRDFAAYRSRVKFSRADAPAYQQGQNLYNELNLG
RREYDVLDRGRDPGEMGGKPRRKNPQEGLYNELQDKMAEAYS
EIGMKGERRRGKGDGLYQGLSTATKDTYDALHMQALPPRAPAAG
DTIISLDFGSNDTETDDQGLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF
AEDTEAKNPFSTQDLDLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVFQ

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions or the amino acid sequence:

(SEQ ID NO: 58[B])

DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRP
GQSPKLLIYKVSDFRS GVPDRFSGSGSGTDFTLKISRVEAEDLGI
YYCFQGSHIPPTFGGGTKLEIKRAAGGGSGGGSGGGSGVQVQLQ
QSGSEMARPGASVKLPCKASGDTFTSYWMHWVKQRHGHGPEWIGN
IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLRLTSEDSAVYY
CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPKFWVLV
VVGGLACYSLLVTVAFIIFWVRSKRSLRLHSDYMNMTPRRPGPT
RKHYQPYAPPRDFAAYRSRVKFSRADAPAYQQGQNLYNELNLG
RREYDVLDRGRDPGEMGGKPRRKNPQEGLYNELQDKMAEAYS
EIGMKGERRRGKGDGLYQGLSTATKDTYDALHMQALPPRAPAAG
DTIISLDFGSNDTETDDQGLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVDSMDVNEFKLELVEKLF
AEDTEAKNPFSTQDLDLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVFQAEGRGSLTTCGDVEENPGPMYRMQLLS
CIALSLALVTNSGIHVFIILGCFSAAGLPKTEANWVNVISDLKKIED
LIQSMHIDATLYTESDVHPSCKVTAMKCFLELQVISLESGLDASI
HDTVENLILANNSLSSNGNVTESGCKECELEEKNIKEFLQSFV
HIVQMFINTS

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions.

15. An immune cell harboring a nucleic acid of claim 1.

16. The immune cell of claim 15, wherein the immune cell is selected from the group consisting of: a T cell, an NK cell, an NKT cell, a macrophage, and a gamma delta T cell.

17. The immune cell of claim 15, wherein the immune cell expresses human IL-15 or soluble variant thereof.

18. The immune cell of claim 17, wherein the soluble variant of IL-15 comprises or consists of:

(SEQ ID NO: 11)
GIHVFILGCFSGAGLPKTEANWVNVISDLKKIEDLIQSMHIDATLY
TESDVHPSCVKVTAMKCFLLLELQVISLESGDASIHDTVENLIILAN
NSLSSNGNVTESGCKECELEEKNKEFLQSFVHVHIVQMFINTS.

19. The immune cell of claim 18, wherein the immune cell is an NK cell.

20. A method for treating breast cancer comprising administering a composition comprising an immune cell of claim 1.

21. A fusion protein comprising a chimeric antigen receptor and an oxygen-dependent degradation domain (ODD), wherein the chimeric antigen receptor comprises:

- (i) an scFv that binds human EGFR;
- (ii) a spacer domain;
- (iii) a transmembrane domain;
- (iv) a costimulatory domain; and
- (v) a CD3 ζ signaling domain;

wherein the scFv comprises: a light chain CDR1 comprising RSSQNIVHNNGITYLE (SEQ ID NO: 4), a light chain CDR2 comprising KVSDFRS (SEQ ID NO: 5), a light chain CDR3 comprising FQGSHIPPT (SEQ ID NO: 6), a heavy chain CDR1 comprising SYWMH (SEQ ID NO: 7), a heavy chain CDR2 comprising NIYPGSGGTNYAEKFKN (SEQ ID NO: 8), and a heavy chain CDR3 comprising SGGPYFFDY (SEQ ID NO: 9).

22.-23. (canceled)

24. The fusion protein of claim 21, wherein the ODD comprises or consists of APAAGDTIISLDFG-SNDTETDDQQLEEVPLYNDVMLPSPNEKLQIN-LAMSPLPTAETPK PLRSSADPAL-NQEVALKLEPNPESLELSFTMPQIQDQTPSPSDGSTRQSSPEPNPSEYCFY VSDMVNEFKLELVEKLF AE-DTEAKNPFSTQDLDLEMLAPY-IPMDDDFQLRSFDQLS PLESSASPESASPQSTVTVFQ (SEQ ID NO: 10).

25. The fusion protein of claim 21, wherein the scFv comprises a VL domain comprising: DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRPQGSPKLLIYKVSDRFS GVPDRFSGSGSGTDFTLKISRVEAEDLGIYYCFQG-SHIPPTFGGGTKLEIKRAA (SEQ ID NO: 2) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions followed by a VH domain comprising or consisting of QVQLQQSGSEMARPASVKLPCKASGDTFT-SYWMHWVKQRHGHGPEWIGNIYPGSGG TNY-AEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYYC-TRSGGPYFFDYWGQGTTLT VSS (SEQ ID NO: 3) with no more than 10, 5, 4, 3, 2 or 1 single amino acid substitutions; optionally wherein the amino acid substitutions are not in the CDRs.

26. The fusion protein of claim 21, wherein the fusion protein comprises or consists of the amino acid sequence:

(SEQ ID NO: 57 [A])
DILMTQSPLSLPVSLGDQASISCRSSQNIVHNNGITYLEWYLQRP
QGSPKLLIYKVSDRFSGVPDRFSGSGSGTDFTLKISRVEAEDLGI
YYCFQGSHPPTFGGGTKLEIKRAAGGGGSGGGGSGGSGVQLQ
QSGSEMARPASVKLPCKASGDTFTSYWMHWVKQRHGHGPEWIGN
IYPGSGGTNYAEKFKNKVTLTVDRSSRTVYMHLSRLTSEDSAVYY
CTRSGGPYFFDYWGQGTTLTVSSPKSCDKTHTCPPCPDPEKFWVLV
VVGGLACYSLLVTVAFIIFWVRSKRSRLHSDYMNMTPRRPGPT
RKHYQPYAPPRDFAAYRSRVKFSRSADAPAYQQGQNLYNELNLG
RREEYDVLDKRRGRDPEMGGKPRRKNPQEGLYNELQKDKMAEAYS
EIGMKGERRRGKGDGLYQGLSTATKDTYDALHMQALPPRAPAAG
DTIISLDFGSDTETDDQQLEEVPLYNDVMLPSPNEKLQINLAM
SPLPTAETPKPLRSSADPALNQEVALKLEPNPESLELSFTMPQIQ
DQTPSPSDGSTRQSSPEPNPSEYCFYVSDMVNEFKLELVEKLF
AEDTEAKNPFSTQDLDLEMLAPYIPMDDDFQLRSFDQLSPLES
SSASPESASPQSTVTVFQ

with no more than 10, 5, 4, 3, 2, 1 or no single amino acid substitutions.

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