The Natural Cytotoxicity Receptors in Health and Disease

Alexander David Barrow*1, Claudia Jane Martin1 and Marco Colonna2*

1 Department of Microbiology and Immunology, Peter Doherty Institute for Infection and Immunity, University of Melbourne, Melbourne, VIC, Australia, 2 Department of Pathology and Immunology, Washington University School of Medicine, St. Louis, MO, United States

The Natural Cytotoxicity Receptors (NCRs), NKp46, NKp44, and NKp30, were some of the first human activating Natural Killer (NK) cell receptors involved in the non-MHC-restricted recognition of tumor cells to be cloned over 20 years ago. Since this time many host- and pathogen-encoded ligands have been proposed to bind the NCRs and regulate the cytotoxic and cytokine-secreting functions of tissue NK cells. This diverse set of NCR ligands can manifest on the surface of tumor or virus-infected cells or can be secreted extracellularly, suggesting a remarkable NCR polyfunctionality that regulates the activity of NK cells in different tissue compartments during steady state or inflammation. Moreover, the NCRs can also be expressed by other innate and adaptive immune cell subsets under certain tissue conditions potentially conferring NK recognition programs to these cells. Here we review NCR biology in health and disease with particular reference to how this important class of receptors regulates the functions of tissue NK cells as well as confer NK cell recognition patterns to other innate and adaptive lymphocyte subsets. Finally, we highlight how NCR biology is being harnessed for novel therapeutic interventions particularly for enhanced tumor surveillance.

Keywords: receptors, natural killer cell, immunoregulation, disease association, tissue homeostasis

INTRODUCTION

Natural Killer (NK) cells constitute a population of large, granular lymphocytes that are located in the blood, lymphoid organs, such as the thymus and spleen, and non-lymphoid organs, such as the liver and uterus, as well as tissues, such as skin (1–4). NK cells were originally identified based on their ability to lyse certain tumor and virally infected cells (5, 6). In contrast, normal healthy cells that express sufficient levels of MHC class I molecules are spared from NK cell attack (7).

Subsequently, the process whereby the effector functions of developing NK cells are adapted to the levels of MHC class I molecules expressed by host cells, termed NK cell “education”, was described (8, 9). It was during this exciting era of discovery that the functional activity of NK cells was shown to be exquisitely controlled by inhibitory receptors specific for MHC class I molecules, such as the Ly49 receptors in mice (10), and the Killer immunoglobulin-like receptors (KIR) in humans (11–14). Fully educated NK cells efficiently lyse target cells lacking MHC class I molecules, implying the existence of a set of activating NK cell receptors for non-MHC class I ligands expressed on target cells that are either not present or expressed at much lower levels on healthy cells, although formal molecular evidence for this was lacking during this period.

In contrast to T and B lymphocytes, NK cells lack the expression of rearranging cell-surface antigen receptors and so it remained unclear how NK cells might be aroused by surface ligands expressed by tumor or virus-infected cells. Even though NK cells lack expression of the T cell
receptor (TCR), they still retain expression of the ζ chain from the CD3 signaling complex. These data suggested that NK cells express cell-surface receptors that might share similar downstream signaling mechanisms to the TCR but are regulated by different ligands. For example, whereas T cells recognize short antigenic peptides in the context of MHC class I, NK cells can be inhibited by the expression of MHC class I molecules. Thus, NK cells express activating receptors that may recognize ligands expressed by tumor and virus-infected cells in a non-MHC-restricted fashion. Whilst the function and downstream signaling events of the low affinity Fc receptor, CD16, that mediates antibody-dependent cellular cytotoxicity (ADCC) were well-defined at this time (15–17), the identity of other activating receptors and their cognate ligands involved in the natural cytotoxicity of NK-susceptible targets cells were largely unknown, which stimulated further research into the receptors that trigger NK cell cytotoxicity.

The Natural cytotoxicity receptors (NCRs) were originally identified by the Moretta group almost 20 years ago in a series of elegant redirected lysis experiments using human NK cells (18–20). The NCR family are comprised of three type I transmembrane (TM) receptors, termed NKp46, NKp44, and NKp30, which are encoded by the genes, NCR1, NCR2, and NCR3, respectively. Even though the NCRs were discovered based on their ability to induce NK cell cytotoxicity of monoclonal antibody (mAb)-coated tumor cell targets, the blocking of individual NCR activity using soluble mAbs had only a mild effect on NK cell cytotoxicity and different tumor cells varied in their susceptibility. Combinations of soluble mAbs to the NCRs were found to have a much stronger blocking effect for selected tumor cell-lines indicating that the NCRs can cooperate with each other to mediate NK cell cytotoxicity of certain tumor cell-types (18, 20, 21). These results suggest that the coordinated expression of NCR ligands by different tumor cell-types as well as the level of NCR expression by different NK cell clones governs NK cell cytotoxicity, which is counterbalanced by the level of MHC class I expressed for a given tumor cell type.

Recent success in identifying the various ligands for the NCRs is now beginning to illuminate the biological roles that this important class of receptors plays in NK cell surveillance of malignant or pathogen-infected cells and in different tissue microenvironments. One striking aspect that has become apparent is that each NCR may interact with several different pathogen- and host-encoded molecules that can either be expressed on cell-surfaces, secreted or shed extracellularly, or incorporated into the extracellular matrix (ECM). Moreover, cytokines expressed in the local tissue microenvironment can influence which particular NCR isofrom is expressed. These different NCR isoforms have now been shown to deliver either activating or inhibitory signaling functions depending on their interaction with ligand. Here we review NCR biology in the different tissue NK cell populations as well as other innate and adaptive immune subsets, their functional interactions with a diverse set of cellular- and pathogen-encoded ligands, and the potential for these interactions to be harnessed for tumor immunotherapy.

**NKp46**

Molecular cloning of the cDNA for NKp46 (also known as natural cytotoxicity receptor 1, NCR1) revealed an open reading frame (ORF) encoding a 46 kDa type I TM protein belonging to the immunoglobulin (Ig) superfamily characterized by two extracellular C2-type Ig-like domains followed by a stalk region (Figure 1) (21). The two Ig-like domains of NKp46 are arranged in a V-shaped conformation positioned at an angle of 85° to each other similarly to the D1D2 domains of KIRs and Leukocyte Ig-like receptors [LILR, also known as ILTs (22)] that share a distant ancestral evolutionary relationship with NKp46 (23, 24). The cytoplasmic domain of NKp46 lacks an Immunoreceptor Tyrosine-based Activation Motif (ITAM), instead the TM domain contains a positively charged arginine (R) residue in their hydrophobic TM domains that can form a salt bridge with a corresponding aspartate (D) residue in the TM domains of the ITAM adaptors; CD3, FcR, or DAP12, respectively. The cytoplasmic domains of the NCRs do not encode any inherent signaling capacity with the exception of NKp44 that contains a putative ITIM sequence (red) in its cytoplasmic tail and thus maintains potential for inhibitory signaling.
ortholog has also been cloned and maps to mouse chromosome 7, the syntenic region of human chromosome 19 (21).

The expression of NKp46 on NK cells is conserved across all mammalian species (26). In humans, NKp46 is expressed by all CD56dimCD16+ and CD56brightCD16− human NK cells irrespective of their activation status (19). Cross-linking with anti-NKp46 mAb results in calcium release and the secretion of IFN-γ and TNF-α by NK cells and blocking NKp46 signaling with specific mAbs can result in reduced NK cell cytotoxicity of certain tumor cell-lines, although the most potent blocking activity is observed in combination with mAbs to other NCRs (19, 21). Subsequent studies have now shown that NKp46 is also expressed by innate lymphoid cells (ILCs) of group 1 (ILC1) and a subset of group 3 ILCs (NCR+ ILC3) (27, 28), γδ T cells (29, 30), a population of oligoclonally expanded intraepithelial (IEL) cytotoxic T lymphocytes (CTL) (31) and a population of IL-15-dependent innate-like IEL lacking surface TCR expression (32) in celiac disease patients, and umbilical cord blood (UCB) T cells cultured in IL-15 (33). NKp46 is also expressed by malignant NK, NKT, and T cell lymphomas (32, 34–36) (Table 1).

**NKp44**

The functional activity of NK cells against tumor cells deficient in the expression of MHC class I molecules is greatly enhanced by culture in IL-2, suggesting that NK cells upregulate activating receptors for additional non-MHC ligands. Whereas, NKp30 and NKp46 are constitutively expressed by resting NK cells obtained from peripheral blood, the expression of NKp44, also known as natural cytotoxicity receptor 2 (NCR2), is upregulated on NK cells stimulated by IL-2, IL-15 or IL-1β, particularly on the CD56bright subset (20, 37–39). NKp44 is a 44 kDa protein comprised of a single extracellular V-type Ig (IgV) domain followed by a long stalk region and a hydrophobic TM domain containing a charged lysine residue that mediates association with the ITAM adaptor, DAP12 (also known as KARAP and TYROBP) (Figure 1) (40, 41). Structural studies have shown that the IgV domain of NKp44 forms a saddle-shaped dimer with a positively charged groove on one side of the protein (42).

The gene for NKp44 (NCR2) is encoded in the human TREM receptor locus at 6p21.1, which is centromeric of the MHC (20, 43) and is also found in several primates species but not in mice (44). Three major transcripts are expressed from NCR2 (NKp44-1, −2, and −3) that have been investigated in detail. Whereas, NKp44-2 and NKp44-3 are predicted to encode proteins with short cytoplasmic domains with no inherent signaling capacity, NKp44-1 is predicted to encode a protein with a long cytoplasmic tail containing the amino acid sequence “ILYHTV” that conforms to the sequence of an Immunoreceptor Tyrosine-based Inhibition Motif (ITIM). The NKp44-1 isoform thus has the potential for inhibitory as well as activating signaling since this isoform also retains the capacity to associated with DAP12. A molecular characterization of the NKp44-1 ITIM showed that it has the potential to be phosphorylated in the NK92 NK cell-line but could not recruit the phosphotyrosine phosphatases, SHP-1 or SHP-2, or the 5′-inositol phosphatase, SHIP, in order to mediate cellular inhibition (45). Thus, it was concluded that the activating function of NKp44-1 is not influenced by the presence of the cytoplasmic ITIM. Nevertheless, NKp44-1 has the potential for dual signaling functions through the cytoplasmic ITIM and association with the ITAM adaptor, DAP12.

Like the other NCRs, NKp44 has also been shown to be expressed by ILC1 (46, 47), ILC3 (48, 49), γδ T cells (30, 50), oligoclonally expanded IEL CTL in celiac disease patients (31), and UCB T cells cultured in IL-15 (33), in addition to plasmacytoid dendritic cells where NKp44 may be involved in the regulation of type I interferon secretion (51) (Table 1).

**NKp30**

NKp30, also known as natural cytotoxicity receptor 3 (NCR3), was identified as a 30 kDa protein that, similarly to NKp46, is expressed on all mature resting and activated NK cells (18). Molecular cloning of the NKp30 cDNA revealed an open reading frame predicted to encode one extracellular IgV domain and a hydrophobic TM domain with a charged arginine residue capable of associating with the ITAM adaptors, CD3ζ and/or FcRγ (Figure 1) (18, 88). The crystal structure of NKp30 reveals some structural similarity to CTLA-4 and PD-1 and NKp30 is thus considered a member of the CD28 family of receptors (83, 89). Both NKp30 and NKp46 have reduced surface expression on adaptive memory NK cells most likely due to the downregulated expression of the FcRγ signaling chain required for the surface expression of these receptors (90, 91).

In humans, NCR3 is encoded in the class III region of the Major Histocompatibility Complex (MHC) but was found to be a pseudogene in 13 strains of mice with the exception of *Mus caroli* (92). Six alternatively spliced transcripts are transcribed from the NKp30 gene, termed NKp30a-f. Three NKp30 isoforms, NKp30a, NKp30b, and NKp30c, which differ in their cytoplasmic tails due to alternative splicing in exon 4, have been studied in detail. Whilst NKp30a and NKp30b evoke NK cell activation, the NKp30c isoform was shown to elicit secretion of the immunosuppressive cytokine, IL-10, from NK cells (93). NKp30 has also been shown to be expressed by γδ T cells (30), CD8+ T cells (94), and UCB T cells cultured in IL-15 (33).

**NCRs AND THEIR LIGANDS IN CANCER**

The NCRs were first characterized based on their ability to evoke the cytotoxic and cytokine-secreting functions of NK cells toward tumor cell-lines in *vitro*. Several studies have now provided evidence that the NCRs are involved in tumor surveillance *in vivo*. For example, genetic deficiency of NKp46 in mice results in the impaired clearance of subcutaneous T lymphoma (95) and melanoma (96) tumors and melanoma lung metastases (97, 98). Moreover, transgenic overexpression of NKp46 resulted in the enhanced clearance of melanoma lung metastases (99). Intriguingly, enhanced NKp46 signaling elicited IFN-γ secretion and increased tumor deposition of fibronectin, which altered the solid tumor architecture and resulted in the decreased formation of melanoma metastases (100). Importantly, the T lymphoma and melanoma cell-lines used in these studies all expressed cell-surface ligands for NKp46. Moreover, in human patients with melanoma, normal melanocytes had negligible expression of NKp46 ligands in comparison to malignant melanocytes deep
TABLE 1 | Expression of Natural cytotoxicity receptors and their ligands.

| Receptor | Gene | Cellular expression | Ligands | Ligand-induced signal | References |
|----------|------|---------------------|---------|-----------------------|-----------|
| NKp46    | NCR1 | NK cells, ILC1, ILC3, Small intestine TCRαβ + CD8α + IEL, Small intestine TCRαβ−/−/CD8α− IEL, Expanded peripheral blood γδ T cells (Vα1+), NK, NKT and T lymphomas, Cord blood T cells cultured in IL-15 | Heparan sulfate (HS) glycosaminoglycans (GAGs) | Activation | (52) |
|          |      |                     | HA (hemagglutinin) of influenza virus | Activation | (53) |
|          |      |                     | HA of human vaccinia virus | Activation | (54) |
|          |      |                     | HN of avian Newcastle disease virus, Stendai virus and human parainfluenza virus | Activation | (53, 55–57) |
|          |      |                     | (DBL)-1 α domain of Plasmodium falciparum erythrocyte membrane protein (PfEMP1) | Activation | (58) |
|          |      |                     | Vimentin expressed on cells infected with Mycobacterium tuberculosis | Activation | (59, 60) |
|          |      |                     | Unidentified ligand expressed by Fusobacterium nucleatum | Activation | (61) |
|          |      |                     | C. glabrata Epa proteins Complement Factor P (properdin) | Activation | (62) |
|          |      |                     | HS GAGs | Activation | (63) |
|          |      |                     | Syndecan-4 (in cis) | Inhibition | (64) |
|          |      |                     | HA of influenza virus | Activation | (65) |
|          |      |                     | HN of avian Newcastle disease virus, Stendai virus and human parainfluenza virus | Activation | (66) |
|          |      |                     | PDGF-DD | Inhibition | (67) |
|          |      |                     | Nidogen-1 | Activation | (68) |
|          |      |                     | PCNA | Activation | (69, 70) |
|          |      |                     | NKp44L expressed on tumor cells, bystander CD4+ T cell during HIV infection, or cartilage-derived chondrocytes Domain III envelope protein from West Nile and Dengue viruses Unknown ligand(s) on Mycobacterium tuberculosis, M. bovis, Nocardia farcinica and Pseudomonas aeruginosa | Activation | (71–73) |
|          |      |                     | Unidentified ligand expressed by pancreatic β-cells | Activation | (74) |
|          |      |                     | C. glabrata Epa proteins Complement Factor P (properdin) | Activation | (75, 76) |
| NKp44    | NCR2 | NK cells, ILC1, ILC3, Plasmacytoid dendritic cells, Small intestine TCRαβ + CD8α + IEL, Expanded peripheral blood γδ T cells (Vα1+), Cord blood T cells cultured in IL-15 | HS GAGs | Activation | (62, 65) |
|          |      |                     | Syndecan-4 (in cis) | Inhibition | (66) |
|          |      |                     | HA of influenza virus | Activation | (67) |
|          |      |                     | HN of avian Newcastle disease virus, Stendai virus and human parainfluenza virus | Activation | (68) |
|          |      |                     | PDGF-DD | Inhibition | (69) |
|          |      |                     | Nidogen-1 | Activation | (70, 71) |
|          |      |                     | PCNA | Activation | (72) |
|          |      |                     | NKp44L expressed on tumor cells, bystander CD4+ T cell during HIV infection, or cartilage-derived chondrocytes Domain III envelope protein from West Nile and Dengue viruses Unknown ligand(s) on Mycobacterium tuberculosis, M. bovis, Nocardia farcinica and Pseudomonas aeruginosa | Activation | (73) |
|          |      |                     | Unidentified ligand expressed by pancreatic β-cells | Activation | (74) |
|          |      |                     | C. glabrata Epa proteins Complement Factor P (properdin) | Activation | (75, 76) |
| NKp30    | NCR3 | NK cells, CD8α+ T cells, Expanded peripheral blood γδ T cells (Vα1+), ILC2 | HS GAGs | Activation | (62, 77, 78) |
|          |      |                     | HA of human vaccinia virus | Inhibition | (63) |
|          |      |                     | pp65, major tegument protein of human cytomegalovirus | Inhibition | (64) |
|          |      |                     | (DBL)-1 α domain of Plasmodium falciparum erythrocyte membrane protein (PfEMP1) | Activation | (65) |
|          |      |                     | BAT3/BAG6 | Activation | (66) |
|          |      |                     | B7-H6 | Activation | (67) |
|          |      |                     | Galectin-3 | Activation | (68) |
|          |      |                     | β-1,3 glucan | Activation | (69) |

within melanoma lesions that stained strongly for NKp46-Fc showing that NKp46 ligands can be upregulated on malignantly transformed cells but are not expressed by healthy cells (101). Several cellular ligands for the NCRs have now been proposed (Table 1) and we discuss these here in the context of anti-tumor responses (Figure 2).

**Heparan Sulfate Glycosaminoglycans**

Heparan sulfate (HS) glycosaminoglycans (GAGs) are found on cell surfaces and within the ECM and consist of long, branched, anionic polysaccharides that are incorporated into proteins to form HS proteoglycans (HSPGs), such as the syndecans and glypican, that form an integral and dynamic component of normal tissue architecture (102, 103). HSPGs also play vital roles in tumor progression allowing cancer cells to proliferate, elude immunosurveillance, invade neighboring tissues, and metastasize to distal tissue sites from the primary tumor. The negative charge of HSPGs can also provide docking sites for the basic domains of various secreted factors, such as chemokines and growth factors like fibroblast growth factor (104).

All three NCRs have been reported to bind to different HS sequences. HS GAGs are heterogenous in structure with a preference for highly sulfated HS structures and with each NCR possessing a distinct HS binding specificity. NKp30 and NKp46 binding to HS is similar although the binding varies significantly. In contrast, NKp44 displays a very different binding pattern to NKp30 and NKp46 (52, 77). A cis interaction of NKp44 with the cell-associated HSPG, syndecan-4, has been reported. The interaction between syndecan-4 and NKp44 was shown to activate (cis) and inhibit (trans) NKp44 signaling (82–85).
FIGURE 2 | Expression of the NCRs and their Ligands. Schematic representation of the expression of the NCRs on NK cells and the NCR ligands by tumors (green) or pathogens or cells infected with pathogens (light blue). Nkp30 and Nkp46 are expressed by resting (pink) as well as activated (orange) NK cells, whereas Nkp44 is only expressed by NK cells activated with IL-2 or IL-15 (arrow). PCNA, BAT3 and Nkp44L are nuclear proteins but the pathways whereby they become expressed on the surface of tumor cells remain to be identified. Nkp44L is a splice variant of the Mixed-lineage leukemia 5 (MLL5) protein encoding a C-terminal modification reported to mediate cell-surface expression of Nkp44L on tumor cells. Similarly, the pathway whereby vimentin becomes expressed on the surface of M. tuberculosis infected cells also remains to be described.

dampen Nkp44 activity. Treatment of NK cells with soluble HS was proposed to disrupt the cis association with syndecan-4 and potentiate Nkp44 signaling (65).

It is an attractive hypothesis to speculate that NK cells could utilize the NCRs to sense changes in HS GAGs in the tumor microenvironment or possibly even during infection to activate NK cell cytotoxicity and IFN-γ secretion, particularly since several cancers exhibit aberrant regulation of key HS biosynthetic enzymes, such as 3-O- and 6-O-Sulfotransferases, and catabolic enzymes, such as heparanase and the HS endosulfatases, SULF1 and SULF2 (105). Conversely, pathogens or tumor cells could modify HS GAGs to evade NK cell surveillance. Indeed, the overexpression of telomere repeats binding factor 2 (TRF2), a key factor in telomere protection, can result in the upregulation of heparan sulfate-glucosamine 3-O-sulfotransferase 4 (HS3ST4) in cancer cells by binding to the interstitial telomeric repeat located within the HS3ST4 intron (106). Silencing of HS3ST4 expression using short hairpin RNAs in cancer cells resulted in increased tumor infiltration of activated NK cells. Thus, aberrant sulfation of cell-surface HS GAGs can modify the capacity for tumor surveillance by NK cells. However, the precise role of the NCRs and HS GAGs in NK cell tumor surveillance remains unclear since Nkp30-dependent NK cell cytotoxicity was unaffected by GAG-deficiency or heparanase treatment of tumor cell targets (78). It is possible that HS GAGs may also serve as “co-receptors” facilitating interactions with other NCR ligands, such as growth factors like PDGF-DD (67).

B7-H6
K562 cells are highly susceptible to lysis by human NK cells, which is mediated by NKP30-mediated recognition of a tumor cell-surface ligand. A proteomics approach designed to trap ligands bound to an NKP30 Fc-fusion protein by chemical cross-linking resulted in the co-immunoprecipitation of B7-H6 from K562 cell lysates (84). B7-H6 is type I TM protein possessing two extracellular Ig-like domains and is a member of the B7 family of costimulatory molecules. Like other members of the B7 family, the extracellular domain of B7-H6 is composed of a membrane distal IgV domain and a membrane proximal IgC domain (83). NKP30 was found to use the front and back β-sheets of its IgV domain to engage the side and face of the corresponding β-sandwich of B7-H6, whereas CTLA-4 and PD-1...
use only the front β-sheet of their Ig-like domains to contact their ligands. B7-H6 contacts NKp30 through the complementarity-determining region (CDR)-like loops of its IgV domain, thus resembling the binding interaction of antibodies with antigen, which is not observed for CTLA-4 binding to B7.1/B7.2 or PD-1 binding to PD-L1/PD-L2 (83).

B7-H6 expression was found to be negligible on normal cells but was highly expressed by a wide range of tumor cells, showing that cellular transformation serves as a mode of immunosurveillance in the innate immune system (84). In support of this, the expression of B7-H6 is upregulated by the proto-oncogene Myc (107) and is associated with greater overall survival of patients with oral squamous carcinoma (108). In contrast, some tumors are thought to escape NKp30 recognition by metalloprotease-mediated shedding of B7-H6 from the cell-surface (109). The soluble form of B7-H6 is detected in the serum of patients with hepatocellular carcinoma (110), metastatic gastrointestinal tumors (GIST) (111), neuroblastoma (112, 113), and peritoneal fluid from ovarian cancer patients (114), and is associated with impaired NKp30 expression and NK cell dysfunction as well as poor overall patient survival.

Differential expression of transcripts encoding the different NKp30 isoforms have also been associated with cancer prognosis. The preferential expression of the immunosuppressive NKp30c isoform over the activating NKp30a and NKp30b isoforms was associated with the reduced overall survival of GIST patients treated with imatinib mesylate (93, 111). The NKp30c isoform was reported to trigger secretion of the immunosuppressive cytokine IL-10 from NK cells and was associated with NK cell dysfunction characterized by defective NK cell degranulation and secretion of IFN-γ and TNF-α. Interestingly, polymorphism in the 3' untranslated region of NCR3 can result in the preferential expression of the immunosuppressive NKp30c isoform by NK cells and predicted the clinical outcome for GIST patients independently from KIT mutation (93). In pediatric neuroblastoma, expression of the immunostimulatory (NKp30a/b) versus immunosuppressive (NKp30c) isoforms is also associated with a higher risk of relapse and elevated serum levels of soluble B7-H6 that inhibit NK function were associated with the bone marrow metastasis and chemotherapy of neuroblastoma cells (112, 113). Expression of B7-H6 is also detected on monocytes and neutrophils treated with TLR ligands and pro-inflammatory cytokines. A soluble form of B7-H6 was also detected under these conditions and in sepsis patients suggesting B7-H6 may also regulate NK cell activity via NKp30 isoforms during inflammation (115).

Nuclear Proteins

NKp44 has been reported to bind to several intracellular ligands that have been proposed to be aberrantly expressed on the cell-surface of tumor or virus-infected cells. An NKp44 ligand was reported to be expressed on the surface of tumor or CD4+ T cell from HIV patients (71) that was dependent on expression of HIV gp41. Moreover, a peptide sequence derived from HIV gp41 was shown to upregulate the expression of this NKp44 ligand on CD4+ T cells (71). A yeast two-hybrid screen using a cDNA library from the Jurkat T cell-line was employed to find interacting partners for NKp44 (72) and a cDNA encoding a splice variant of the mixed-lineage leukemia protein-5 (MLL5) was subsequently cloned. MLL5 is normally a nuclear antigen but the cDNA sequence revealed a new exon resulting in a transcript encoding an MLL5 variant protein with an alternative C-terminal amino-acid sequence. This new protein termed NKp44L was reported to be expressed on the cell-surface and in the cytoplasm of various tumor cell-lines but not in normal tissues (72). However, the mechanism driving NKp44L cell-surface expression remains obscure. NKp44L expression has also been reported on human articular chondrocytes (73).

NKp44 was also reported to interact with Proliferating cell nuclear antigen (PCNA). PCNA is found in the nucleus and as a co-factor for DNA polymerase 6 that helps leading strand synthesis during DNA replication (116). PCNA was identified as an NKp44 ligand following a screen of a yeast surface display library using an NKp44-Fc fusion protein (69). Cell-surface expression of PCNA was reported following transfection into HeLa cells, which reduced NK cell cytotoxicity and IFN-γ secretion. PCNA was found to localize at the plasma membrane where it was shown to form an immunological synapse with NKp44 to mediate NK cell inhibition. The inhibition of NK cell signaling was reported to occur via the NKp44-1 splice variant that potentially encodes a cytoplasmic ITIM (69). The interaction of PCNA with NKp44 was proposed to form a unique interaction capable of transducing an inhibitory rather than an activating signal (69, 117). PCNA has also been shown to associate with Human Leukocyte Antigen (HLA) class I molecules on the cell-surface of tumor cells to form an inhibitory ligand for NKp44 and suppression of NK cell cytotoxicity (70). The authors suggested that NCR ligands could act as Damage-Associated Molecular Patterns (DAMPs) through association with HLA class I molecules (118). Interestingly, preferential expression of the transcript encoding the ITIM-bearing NKp44-1 isoform is associated with poor survival in acute myeloid leukemia and was proposed to be a novel target for checkpoint blockade on NK cells (119).

HLA-B-associated transcript 3 (BAT3), also known as Bcl2-associated anthogene 6 (BAG6), is another intracellular protein that was identified as an NKp30 ligand using a yeast two-hybrid screening approach (80). BAT3 is a multifunctional molecular chaperone that contributes to several cell processes including apoptosis, gene regulation, protein synthesis, protein quality control and protein degradation. For example, BAT3 can function by preventing the aggregation of misfolded and hydrophobic-patch containing proteins ensuring their correct delivery to the endoplasmic reticulum or the sorting of proteins that have mislocalized to the cytoplasm for proteosomal degradation (120). BAT3 also accumulates in the nucleus where it is involved in regulating apoptosis following DNA damage (121). BAT3 was proposed to be released from tumor cells to engage NKp30 and trigger NK cell activation (80, 81). Exosomal release of BAT3 was also reported to promote NK:DC cross-talk (82). In contrast, the release of a soluble form of BAT3 found in the plasma of CLL patients was found to suppress NK cell activation through competition for the exosomal form of BAT3 and other tumor ligands (122, 123).
Platelet-Derived Growth Factor-DD

Growth factors (GFs) are important for guiding various cellular and developmental processes and GF pathways are often dysregulated in cancer. The platelet-derived growth factor (PDGF) family is comprised of four polypeptides that can assemble into at least five dimeric isoforms, PDGF-AA, PDGF-BB, PDGF-AB, PDGF-CC and PDGF-DD, and which engage with the receptor tyrosine kinases, PDGF receptor (PDGFR)-α and PDGFR-β (124). Cancer cells frequently express PDGFs which can trigger autocrine and paracrine PDGFR signaling to promote tumor growth, proliferation, stromal recruitment, angiogenesis, epithelial to mesenchymal cell transition, and metastasis (125).

PDGF-DD was identified as a ligand for NKp44 by screening a library of secreted proteins with NKp44 GFP reporter cells (67). PDGF-DD induced NK cell degranulation and the secretion of IFN-γ and TNF-α from NK cells and ILC1 as well as TNF-α secretion from ILC3 in agreement with a previous study (67, 126). IFN-γ and TNF-α secreted by NK cells stimulated with PDGF-DD could induce cell cycle arrest in melanoma, ovarian, and breast tumor cell lines. Thus, innate immune cells have the capacity to sense the expression of GFs by tumor or tumor-associated stroma cells and potentially even infected or developing tissues to evoke NK cell responses. Remarkably, the upregulation of PDGF-DD-induced NK cell cytokines and chemokines and the downregulation of tumor cell cycle genes correlated with NCR2 expression and was associated with greater survival in a cohort of glioblastoma patients. Moreover, transgenic expression of NKp44 in mouse NK cells resulted in greater control of tumors expressing PDGF-DD showing that whilst PDGF-DD can support tumor growth it also exposes tumor cells to NK cell immunosurveillance via NKp44. The ability of NK cells and ILCs to engage in GF surveillance via NKp44 is a new immunological paradigm that remains to be fully explored (67). Interestingly, a polymorphism in PDGFDD is associated with serum IFN-γ levels in humans (127), suggesting the PDGF-DD/NKp44 interaction may play wider biological roles beyond cancer (Table 2).

Nidogen-1

The ECM protein, Nidogen-1 (NID1, also known as Entactin) was identified as an NKp44 ligand, in another screen to identify soluble ligands that might regulate NCR activity and NK cell function (68). NID1 is an essential component of the basement membrane (BM) where it plays a role in BM assembly and stabilization, as well as adhesion between cells and the ECM (128). Soluble NID1 can be detected in the serum from patients with ovarian and lung cancer that may be derived from proteolytic degradation by cathepsin-S (CatS) (129, 130). Thus, it is conceivable that the release of NID1 into extracellular fluids may regulate the activity of activated NK cells in the blood or at specific tissue sites. Indeed, CatS-degradation and release of NID1 was proposed to reflect the loss of BM integrity, which is frequently associated with invasive tumors. Soluble NID1 was able to inhibit cytokine secretion mediated by PDGF-DD or cross-linking with anti-NKp44 mAbs. Consequently, the release of soluble NID1 was proposed to be a novel immunosuppressive mechanism exploited by tumor cells to evade NK cell surveillance (68).

Galectin-3

Galectin-3 is a β-galactoside-binding lectin that has been reported to be a critical immune regulator in the tumor microenvironment (131). Galectin-3 can be expressed in the cytoplasm, nucleus, cell-surface, or extracellularly depending on the cell type and proliferative status. For example, extracellular galectin-3 can facilitate metastasis by promoting cell adhesion, invasiveness and immune evasion. The genetic manipulation of galectin-3 expression levels in tumor cells showed that downregulation of galectin-3 lead to tumor growth inhibition, whereas the upregulation of galectin-3 led to enhanced tumor growth in human cervical and breast cancer models. Moreover, soluble galectin-3 released from tumor cells was found to bind specifically to NKp30 thereby inhibiting NKp30-mediated cytotoxicity. Thus, it was proposed that the secretion of galectin-3 by tumors represents a novel pathway to escape NKp30-mediated NK cell immunosurveillance (85).

NCR-MEDIATED CONTROL OF TUMORICIDAL PATHWAYS

The NCRs have been proposed to bind to many cellular ligands which are implicated in NK cell surveillance of tumor cells. Many of these interactions have been shown to evoke the cytotoxic and cytokine-secreting functions of NK cells. However, it is also possible that the NCRs may regulate other anti-tumor pathways. For melanoma metastases, NK cells were attributed to play a major role (99). However, ILC1 are phenotypically closely related to conventional NK cells as both these cell-types express T-bet and secrete IFN-γ (132, 133). ILC1 are characterized by higher TRAIL expression than NK cells due to tissue-imprinting by TGF-β (134). TRAIL (TNFSF10) can bind to TRAIL receptors (TRAIL-Rs), such as TRAIL-R1 in humans and TRAIL-R2 in humans and mice, which carry death domains that can induce caspase-8-mediated apoptosis of TRAIL-R+ tumor cells. TRAIL expression can be induced on NK cells activated by various cytokines and interferons (135–137) and several reports have described NK cell-mediated clearance of TRAIL-sensitive transplantable tumors (135, 138), chemical-induced fibrosarcoma (139), liver metastases (135, 139, 140), and hematological malignancies (141). NKp46 was shown to regulate TRAIL surface expression resulting in the diminished cytotoxicity of TRAIL-R+ target cells by NKp46-deficient NK cells and ILC1 (142, 143). The NKp46-mediated pathway that regulates surface TRAIL expression remains to be fully characterized but appears to involve a post-transcriptional mechanism (142, 143). Whether NKp46 can regulate surface TRAIL expression when expressed by other immune cell subsets, such as γδ T cells and CD8+ T cells that have been expanded with IL-15, remains to be determined. These data show that NKp46 can endow NK cells or ILC1 with the ability to lyse TRAIL-sensitive tumors as well as contribute to
| NCR gene | Gene location | Disease association/reported trait | Reference (PubMed ID) | NCR Ligand gene | Gene location | NCR ligand disease association | Reference (PubMed ID) |
|----------|---------------|------------------------------------|----------------------|-----------------|---------------|-------------------------------|----------------------|
| NCR1     | 19q13.42      | Blood protein levels               | 29875488 30072576    | KMT2E (parental gene for NKp44L) | 7q22.3 | Intelligence (MTAG) | 29326435            |
|          |               |                                    |                      |                 |               | Educational attainment (MTAG) | 30038396            |
|          |               |                                    |                      |                 |               | Educational attainment (years of education) | 30038396            |
|          |               |                                    |                      |                 |               | Cognitive performance (MTAG) | 30038396            |
|          |               |                                    |                      |                 |               | No data available |                      |
|          |               |                                    |                      |                 |               | Post bronchodilator FEV1/FVC ratio | 26834245            |
|          |               |                                    |                      |                 |               | No data available |                      |
| NCR2     | 6p21.1        | Alzheimer's disease biomarkers (p-tau measurement) | 23362540 | PDGFD | 11q22.3 | Interferon gamma levels | 27989323            |
|          |               | Total ventricular volume (brain measurement) | 21116278 |          |           | Corticosteroid-induced adrenal suppression (peak cortisol < 350 nmol/L) | 29551627            |
|          |               | Telomere length                     | 29151059 |          |           | Corticosteroid-induced adrenal suppression (peak cortisol < 500 nmol/L) | 21347282            |
|          |               |                                    |                      |                 |               | LDL cholesterol |                      |
|          |               |                                    |                      |                 |               | Blood protein levels |                      |
|          |               |                                    |                      |                 |               | 3-hydroxypropylmercapturic acid levels in smokers | 26053186            |
|          |               | SDC4 (Syndecan-4)                   | 20q13.12 |          |           | Bacterial meningitis | 28928442            |
|          |               | NID1                                | 1q42.3 |          |           | Psoriasis | 20953189            |
|          |               |                                    |                      |                 |               | Velopharyngeal dysfunction | 29885589            |
|          |               |                                    |                      |                 |               | Lung function (FEV1/FVC) | 30595370            |
|          |               |                                    |                      |                 |               | Urate levels in overweight individuals | 25811787            |
|          |               |                                    |                      |                 |               | Cutaneous nevi | 21478494            |
|          |               |                                    |                      |                 |               | Blood protein levels | 30072576            |
|          |               |                                    |                      |                 |               | Waist-hip ratio | 30595370            |
|          |               |                                    |                      |                 |               | Facial morphology (factor 20) | 28441456            |
| NCR3     | 6p21.33       | C3 and C4 complement levels         | 23028341 | BAG6 | 6p21.33 | Small cell lung carcinoma | 28604730            |
|          |               |                                    |                      |                 |               | Diastolic blood pressure | 28739976            |
|          |               |                                    |                      |                 |               | Systolic blood pressure | 21909115            |
|          |               |                                    |                      |                 |               | Feeling worry | 27618447            |
|          |               |                                    |                      |                 |               | Hypertension | 29500382            |
|          |               |                                    |                      |                 |               | Neuroticism measurement | 21909115            |
|          |               |                                    |                      |                 |               | Psoriatic arthritis | 30552173            |
|          |               |                                    |                      |                 |               | Urinary tract infection frequency | 28928442            |
|          |               |                                    |                      |                 |               | Allergic disease (asthma, hay fever or eczema) | 29083406            |
|          |               |                                    |                      |                 |               | IgG glycosylation | 23382691            |
|          |               |                                    |                      |                 |               | Menopause (age at onset) | 29773799            |
|          |               |                                    |                      |                 |               | Laryngeal squamous cell carcinoma | 25194280            |
|          |               |                                    |                      |                 |               | Crohn's disease | 22936669            |
|          |               |                                    |                      |                 |               | Pulmonary function | 223924391           |
|          |               |                                    |                      |                 |               | No data available |                      |
|          |               |                                    |                      |                 |               | No data available |                      |
|          |               |                                    |                      |                 |               | No data available |                      |
TRAIL-mediated immunoregulation of infectious diseases, tissue inflammation (144–146), and potentially autoimmunity. It will also be interesting to see which NKP46 ligands can induce surface TRAIL expression in NK cells and ILC1 and the functional consequence for TRAIL-sensitive tumors in vivo.

Control of Malignancy by Unconventional NCR+ Lymphocyte Subsets

Until recently NCR expression was considered to be restricted to NK cells. The NCRs have now been shown to be expressed by CD8+ T cell populations and γδ T cells expanded with IL-15, in addition to ILC1 and subsets of ILC3. Moreover, antibody-mediated stimulation of NKP44 and NKP46 on "NK-like" IL-15-expanded IEL CTL elicits the secretion of IFN-γ showing that these NCRs can function independently of TCR stimulation in T lymphocytes (31). Moreover, a population of NKP30+ CD8+ T cells with anti-tumor potential was shown to be induced by IL-15 (94). These NKP30+ CD8+ T cells exhibited high "NK-like" anti-tumor activity and NKP30 synergized with TCR signaling to control tumor growth in a preclinical xenograft mouse model.

Murine γδ T cells do not express NKP46, however, all three NCRs are expressed on human γδ T cells following continuous stimulation with TCR agonists or mitogens in the presence of IL-2 or IL-15 (30). Interestingly, NCR induction was mostly restricted to the Vδ1+ T cell subset and not Vδ2+ T cells. Like NK cells, the NCR+ Vδ1+ T cells were highly cytolytic against primary leukemia cells and tumor cell-lines in redirected cytotoxicity assays and NCR triggering also enhanced the expression of IFN-γ. Based on these observations, IL-15 expanded populations of "NK-like" CD8+ or Vδ1+ T cells that express NCRs have tremendous potential to be used clinically for adoptive cancer immunotherapy (147). The acquisition of NCRs by Vδ1+ T cells and CD8+ T cells is thought to require strong TCR activation, which is consistent with the oligoclonal expansion of gut IEL CTL in celiac disease that express a highly restricted TCR repertoire (31).

Although unconventional NCR+ lymphocyte subsets have been implicated in tumor surveillance they may also promote immunosuppression and facilitate tumor immune evasion. For example, a unique CD3−CD56+ ILC population was recently described that inhibits tumor-infiltrating lymphocytes (TILs) from high-grade serous ovarian carcinomas (148). The CD3−CD56+ ILC population was associated with a reduced TIL expansion and altered TIL cytokine production. This novel population of CD3−CD56+ ILCs exhibited low cytotoxic potential, secreted IL-22, and exhibited a transcriptional profile overlapping that of NK cells and other ILCs. NKP46 was highly expressed by the regulatory CD3−CD56+ ILC population and NKP46 signaling promoted the ability to suppress TIL expansion in co-culture (148).

Group 2 ILCs (ILC2s) secrete large amounts of type 2 cytokines, such as IL-5, IL-9, and IL-13, which promote alternative macrophage activation, eosinophilia, and goblet cell hyperplasia to limit parasite infection (149, 150). Increased levels of hyperactivated ILC2s expressing NKP30 and Chemokine (C-C motif) receptor-homologous molecule expressed on Th2 cells (CRTH2) are found in patients with peripheral acute promyelocytic leukemia (APL). APL blasts were found to express high levels of surface B7-H6 and prostaglandin D2 (PGD2) that bind to NKP30 and CRTH2 (151) on ILC2s, respectively, to drive potent IL-13 secretion and activation of IL-13+ myeloid-derived suppressor cells (152). Disruption of this tumor immunosuppressive axis by specifically blocking PGD2, IL-13, and NKP30 signaling partially normalized ILC2 and MDSC levels resulting in enhanced survival in leukemic mice (152).

NCRs AND THEIR LIGANDS IN INFECTIOUS DISEASE

The central role of the NCRs in resistance to infectious disease is exemplified by the susceptibility of NKP46-deficient mice to infection with different microorganisms. For example,
human metapneumovirus (HMPV) causes acute respiratory tract infections in infants and children worldwide that can be fatal in immunosuppressed hosts. HMPV-infected cells express an unidentified ligand for NKp46 and NKp46-deficient mice are more susceptible to HMPV infection (153). The NCRs have also been shown to contribute to humoral immune responses and the generation of protective pathogen-specific antibodies. For example, deficiency in NKp46 resulted in the impaired maturation, function, and migration of NK cells to regional lymph nodes of mice infected with murine cytomegalovirus (MCMV). This was accompanied by a reduction in CD4+ T cell activation and follicular helper T cell generation leading to diminished B cell maturation in germinal centers and reduced titers of MCMV-specific antibodies (154).

Given the central role of NCRs in resistance to infectious disease, it is unsurprising that pathogens have evolved mechanisms to avoid NCR recognition. Herpesviruses are DNA viruses that have evolved various strategies of immune evasion as exemplified by their ability to establish latent infection with limited transcription of viral genes. Cells infected with human cytomegalovirus (HCMV) and human herpesvirus 6B (HHV6B) can express the NKp30 ligand, B7-H6. Both HMCV and HHV6B have evolved strategies to downregulate B7-H6 and ligands for other activating NK cell receptors, such as NKG2D, from the cell-surface, thus impairing NK cell recognition of virus-infected cells (155, 156). In the case of HCMV infection, HCMV expressed two gene products US18 and US20, that interfere with B7-H6 surface expression by promoting the lysosomal degradation of B7-H6 (155).

Many of the genes for the NCRs and their ligands are associated with infectious diseases highlighting the importance of the NCRs for pathogen recognition (Table 2). For example, NCR2 is associated with chronic periodontitis in pregnant women and NCR3 is associated with susceptibility to cold sores and shingles (Table 2). Here we document various NCR ligands that have been reported to play a role in pathogen surveillance by NK cells and other NCR+ immune cell subsets (Figure 2).

**Hemagglutinins and Hemagglutinin Neuraminidases**

Even though the NCRs were first identified based on the ability to lyse certain tumor cell-lines, some of the first NCR ligands identified were viral proteins. Hemagglutinins (HA) from influenza (53, 66) and hemagglutinin-neuraminidases (HN) from parainfluenza, Sendai, and Newcastle disease viruses (53, 55) expressed on infected cells were shown to bind to NKp46 and NKp44. HAs from poxviruses, such as vaccinia and ectromelia, were also identified as ligands for NKp30 and NKp46 (54). However, the NCRs do not bind to HA from measles virus, showing that whilst the NCRs are capable of recognizing a broad range of viral HA and HNs there is some selectivity, which is predominantly dependent on α-2,3- and α-2,6-sialylated O-glycans on NCRs (53, 55, 56).

The interaction of viral-encoded HA and HNs on the surface of infected cells results in NK cell activation via NKp46 and NKp44 (53, 55, 56) and binding of the HN of Newcastle disease virus to NKp46 results in activation of downstream signaling molecules in the ITAM pathway e.g., Syk and surface upregulation of TRAIL on murine NK cells (157). In support of these observations, mice deficient in NKp46 expression are more susceptible to infection with influenza A virus (158, 159). In contrast, the interaction of NKp30 with HA expressed on the cell-surface or shed from vaccinia-infected cells counteracted activation by NKp46 resulting in the inhibition of NK cell lysis (54), suggesting a novel pathway of viral immune escape via NKp30. Moreover, exposure of NK cells to influenza virions or soluble HA can result in lysosomal degradation of the CD3ζ signaling chain and a reduction of NK cell cytotoxicity mediated by NKp46 and NKp30 (160). In contrast, the cleavage of sialic acid residues from NKp46 by the influenza virus neuraminidase (NA) was proposed as a mechanism of immune evasion by disrupting NKp46-binding to the viral HA and inhibitors specific for the influenza NA enhanced NKp46 recognition and virus clearance (161).

**Human Cytomegalovirus Tegument Protein pp65**

Intracellular staining of human cytomegalovirus (HCMV) infected fibroblasts revealed an unidentified protein that interacted with NKp30-Fc fusion protein. Purified virions from HCMV-infected fibroblasts also bound to NKp30-Fc but not to control Fc-fusion proteins. These data suggested the unidentified viral protein that interacted with NKp30 may be incorporated into HCMV virus particles. The NKp30 Fc fusion was used to immunoprecipitate proteins from HCMV-infected fibroblasts and the HCMV-encoded tegument protein pp65 was subsequently identified as a ligand for NKp30 using mass spectrometry (79). HCMV pp65 was shown to bind directly to NKp30 and induce the dissociation of the signaling adaptor CD3ζ. HCMV-infected fibroblasts were found to be less susceptible to NK cell cytotoxicity compared to fibroblasts infected with a pp65-deficient HCMV or in the presence of blocking antibodies to NKp30. Thus, pp65 furnishes HCMV with a novel immune escape pathway by disrupting the surveillance of virus-infected cells by NKp30.
Flavivirus Proteins

NKp44 Fc-fusion proteins have also been shown to interact with purified envelope protein from the flaviviruses, dengue virus (DV), and West Nile Virus (WNV) and WNV virus-like particles. NKp44 was shown to specifically bind to domain III of the envelope protein from WNV, which also bound to NK cells expressing high levels of NKp44 and the binding of infectious WNV and WNV-infected cells stimulated NK cell degranulation and IFN-γ secretion in an NKp44-dependent manner (74). Unlike the interaction with viral hemagglutinins, NKp44 binding to domain III of the DV and WNV envelope proteins was independent of glycan-linked sialic acid residues on NKp44.

Bacterial NCR Ligands

In contrast to viral ligands, the NCRs have also been shown to bind directly to molecules encoded by bacteria and parasites in addition to host molecules exposed on the surface of cells infected with bacteria. For example, an Fc-fusion protein of NKp44 (NKp44-Fc), but not NKp30 or NKp46, was shown to bind to Mycobacterium bovis (M. bovis), M. tuberculosis, Nocardia farcinica, and Pseudomonas aeruginosa (75, 76). NKp44 was specifically shown to bind to cell wall components, such as arabinogalactan-peptidoglycan (mAGP) as well as to mycolic acids and arabinogalactan and derivatives, from M. tuberculosis. Whilst a direct role in NK cell cytotoxicity could not be demonstrated, anti-NKp44 antibodies inhibited mAGP and BCG-induced CD69 expression on IL-2-activated NK cells, suggesting NKp44 recognition of bacterial wall components may sustain NK cell activation during infection with M. tuberculosis (76). In contrast to NKp44, NKp46 was reported to play a dominant role in the lysis of mononuclear phagocytes infected with M. tuberculosis (59). NKp46 was subsequently shown to recognize vimentin expressed on the surface of cells infected with M. tuberculosis (60). Antibodies to vimentin inhibited NK cell lysis of monocytic cells infected with M. tuberculosis, whereas transfection of vimentin enhanced NK cell lysis of Chinese hamster ovary cells (60).

Fusobacterium nucleatum and Porphyromonas gingivalis are involved in the pathogenesis of periodontitis. The severity of periodontal disease is determined by the host’s immune response. Alveolar bone loss occurred in wild-type mice but was minimal in NKp46-deficient mice in an oral infection model using Fusobacterium nucleatum or Porphyromonas gingivalis (61). Expression of an NKp46 ligand by F. nucleatum was demonstrated by the binding of a NKp46-Fc and the activation of NKp46 reporter cells as well as the secretion of TNF-α from wild-type but not NKp46-deficient NK cells (61). Interestingly, NCR2, but not NCR1, is associated with chronic periodontitis in humans (Table 2) (162), which could reflect cross-talk in NCR signaling (163).

Parasite NCR Ligands

Parasite infection has also been shown to induce the expression of NCR ligands. Fc fusion proteins of NKp30 and NKp46 were shown to specifically bind to the Duffy binding-like (DBL)-1 α domain of Plasmodium falciparum erythrocyte membrane protein-1 (PfEMP-1), which is expressed on the cell-surface of parasitized erythrocytes (38). NK cells lysed erythrocytes infected with P. falciparum, which was blocked by antibodies to NKp30 and NKp46 with the greatest inhibition observed when both antibodies were used in combination.

Fungal NCR Ligands

NKp46 and NKp30 have been shown to interact with fungal ligands. Both mouse and human NKp46 were shown to bind to the emerging fungal pathogen, Candida glabrata but not to C. albicans, C. parapsilosis, C. kruzei, or Cryptococcus gattii (63). Fc fusion proteins from human and mouse NKp46 were specifically shown to bind to the C. glabrata adhesins Epa1, Epa6, and Epa7. The Epa proteins are lectins that bind to glycans to facilitate the C. glabrata infection of mammalian cells and mutation of O-linked glycosylation of Threonine 225 of mouse and human NKp46 abolished binding to C. glabrata Epa proteins (63). Moreover, the clearance of systemic C. glabrata infection was impaired in NKp46-deficient mice.

NKp30 mediates the recognition of Cryptococcus neoformans and C. albicans resulting in the formation NK cell-fungal conjugates and direct fungal killing (164). NKp30 was specifically shown to bind to the fungal cell wall component β-1,3 glucan to stimulate the polarization of cytotoxic granules in NK cell-fungal conjugates, which enhanced the release of perforin required for fungal cytotoxicity (86). Rather than blocking the interaction between NK cells and fungi, soluble β-1,3 glucan enhanced fungal killing (86). Fungal infections are the leading cause of death in AIDS patients and NK cells from HIV-infected patients had reduced expression of NKp30. IL-12 and β-1,3 glucan restored NK cell expression of NKp30 and fungal killing by NK cells (86, 164). The activation of src family kinases by NKp30 synergized with β1 integrin signaling and the activation of integrin-linked kinase (ILK) and Rac1, which converged into a central PI3 kinase signaling pathway that was required for fungal killing by NK cells (87).

Complement Factor P (properdin)

The complement system is an evolutionary ancient component of the innate immune response. Complement factor P (CFP, also known as properdin) is a plasma glycoprotein that binds to microbial surfaces and is the only known positive regulator of the alternative pathway of complement (165). CFP stabilizes the C3- and C5-convertase enzyme complexes that ultimately leads to the formation of the membrane attack complex and target cell lysis. NKp46 was shown to bind to CFP. Patients deficient in CFP are more susceptible to lethal Neisseria meningitidis due to an inability to activate the alternative pathway of complement (166). CFP was shown to bind to a recombinant NKp46 Fc-fusion protein and to activate NKp46 reporter cells but could not induce classical NK cell activation characterized by degranulation and IFN-γ secretion (64). Instead CFP induced the upregulation of genes encoding the leucine zipper protein FosB and the chemokine CXCL1 (also known as lymphotactin). Thus, CFP binding to NKp46 was proposed to activate a non-canonical pathway of gene expression in NK cells. In agreement with patients deficient in CFP, NKp46 and ILC1 were required for mice to survive infection with N. meningitidis. Moreover, the beneficial
effects of CFP supplementation for *N. meningitidis* infection were dependent on NKp46 and ILC1 (64).

**Unconventional Roles for NCRs in the Control of Infection**

NKp30 is also expressed on γδ T cells (30) bearing TCR Vδ1 but not TCR Vδ2. Circulating γδ T cells are normally devoid of NCR expression. However, all three NCRs were induced on γδ T cells following TCR engagement in conjunction with cytokines, such as IL-2 and IL-15 (30). Moreover, mAb cross-linking of NKp30 on NCR+ Vδ1 T cells, and to a lesser extent NKp44 and NKp46, induced the chemokines CCL1, CCL2 and CCL3, which suppressed HIV infection by binding the HIV co-receptor, CCR5 (167).

**NCRs AND THEIR LIGANDS IN DEVELOPMENT**

In steady state, human uterine NK (uNK) cells maintain endometrial arteries (169). However, during pregnancy, uterine NK cells maintain key developmental processes, such as trophoblast invasion, remodeling uterine vasculature, and the promotion of fetal growth (169). The tissue origins of uNK remain obscure and are thought to constitute a heterogeneous population derived from local endometrial NK cells as well as conventional NK cells recruited from the circulation (169, 170). Uterine NK cells possess limited cytotoxicity and actively secrete cytokines, chemokines, GFs, and angiogenic factors required for the correct remodeling of maternal spiral arteries and likely function by promoting the recruitment of invading trophoblast cells and promoting angiogenesis during pregnancy (171).

The NCRs have been reported to be expressed by uNK (171–173) and the ligands for NKp30 and NKp44 are also expressed by trophoblast cells (171). Consequently, trophoblast cells have the potential to directly stimulate uNK functions via the NCRs and ligation of the NCRs with specific antibodies resulted in the secretion of cytokines, chemokines, GFs, and angiogenic factors by uNK (171–173). Thus, NCR expression endows uNK with the capacity to orchestrate key developmental processes at the fetal-maternal interface following interactions with trophoblast cells or other NCR ligands expressed in decidua. Uterine NK and peripheral blood NK (pNK) operate in different tissue microenvironments and also differ in NCR expression. For example, resting pNK do not express NKp44 unlike uNK. The decidual stroma expresses the cytokines IL-15, IL-18, and TGF-β, that are required for successful pregnancy. Uterine NK were found to express distinct NKp30 and NKp44 splice variants compared to pNK, characterized by predominant expression of the inhibitory NKp30c and NKp44-1 isoforms and lower amounts of the activating NKp30α/b and NKp44-2/3 isoforms (174). A combination of IL-15, IL-18, and TGF-β shifted the pNK NCR splice variant expression profile toward that of uNK suggesting the decidual tissue microenvironment can confer a decidual-like uNK phenotype on pNK (174).

**NCRs AND THEIR LIGANDS IN ALLERGY**

Allergic disease is caused by inappropriate immune responses to harmless antigens driven by type 2 cytokines. Very few papers have documented a role for NK cells or the NCRs in allergic disease. However, recent studies have suggested that NK cells and the NCRs may be important in allergy due to the immunoregulatory activity of NK cells on other immune cell subsets (175) and NK cells have also been shown to drive allergic responses independently of T and B cells (176, 177). NKp46-deficient mice were shown to have reduced skin and airway hypersensitivity (178). In a delayed type IV hypersensitivity model of airway inflammation, the number of IFN-γ producing NK cells and the proliferation of OVA-specific T cells was reduced following intranasal OVA challenge in NKp46-deficient mice, suggesting that differences in the stimulation of T cells resulting in an altered cytokine profile upon challenge with ovalbumin (OVA) (178). These results suggested that NKp46 signaling enhances allergic responses. In contrast, a separate study using a delayed type I hypersensitivity model, showed that NKp46 signaling suppressed the development of airway eosinophilia following intranasal OVA challenge (179). It will be interesting to see which NKp46-expressing cell types, such as NK cells, ILC1 or ILC3, might be responsible for either suppressing (179) or augmenting (178) airway inflammation in models of type I and type IV hypersensitivity, respectively.

NCR expression by other lymphocyte subsets may also regulate allergic response. For example, whilst ILC2s secrete type 2 cytokines to limit parasite infection they are also important contributors to allergic inflammation (149, 150). Activated human ILC2s can express NKp30 and stimulation of ILC2s with B7-H6 induced the rapid production of type 2 cytokines, such as IL-13, which was blocked by anti-NKp30 blocking mAb and galectin-3 (180). Higher expression of B7-H6 is observed in skin lesion biopsies from patients with atopic dermatitis and stimulation of keratinocytes with proinflammatory as well as type 2 cytokines to limit parasite infection they are also important contributors to allergic inflammation (149, 150). Activated human ILC2s can express NKp30 and stimulation of ILC2s with B7-H6 expression leading to enhanced IL2 production of type 2 cytokines. Thus, the NKp30-B7-H6 interaction is a novel cell contact-dependent interaction that can mediate IL2 activation and is a potential target for the development of novel therapeutics for atopic dermatitis and possibly other atopic diseases (180).

**NCRs AND THEIR LIGANDS IN AUTOIMMUNITY**

NK cell have also been implicated in the development of autoimmune diseases, such as type 1 diabetes (181–184). NKp46-deficient mice were reported to be resistant to the development diabetes induced by streptozotocin and a soluble NKp46-Fc protein blocked the development of diabetes in mice (62). NKp46 was suggested to play a dominant role in NK cell cytotoxicity of human pancreatic β-cells through the recognition of an as yet unidentified NKp46 ligand expressed in the β-cell insulin granules (97). Anti-NKp46 mAbs that can induce the downregulation and lysosomal degradation of mouse and human
NKp46 have now been developed, which may be effective in the treatment of diabetes (185, 186).

Primary Sjögren’s syndrome (pSS) is a chronic autoimmune disease characterized by lymphocytic exocrinopathy that can affect up to 0.1 to 0.6% of the general population. An excess accumulation of NK cells in minor salivary glands of pSS patients is correlated with the severity of exocrinopathy. B7-H6, the ligand for NKp30, is expressed in minor salivary glands and a higher cell-surface expression of NKp30 on circulating NK cells is observed in pSS patients compared to healthy controls (187). A case control study of genetic polymorphisms revealed that a single nucleotide polymorphism (SNP) rs11575837 (G > A) in the promoter region of NCR3 was associated with reduced transcription and protection from pSS. These findings suggested that NK cells may promote NKp30-dependent inflammation in salivary glands and blockade of the NKp30-B7-H6 interaction may represent a novel clinical target for pSS (187).

**NCR-BASED CANCER IMMUNOTHERAPIES**

In addition to the development of therapeutic antibodies to the NCRs for potential use in the treatment of autoimmune diseases, such as diabetes (see above), several reports are now beginning to provide evidence that the NCRs and their ligands can be successfully targeted for cancer immunotherapy. Moreover, these systems have shown to be effective in various pre-clinical mouse tumor models highlighting the tremendous potential of NCR-based strategies for tumor immunotherapy.

**Virotherapy**

Reovirus is a double-stranded RNA virus that infects much of the population during childhood causing mild or subclinical infections. Interestingly, reovirus efficiently infects tumor cells and is currently being tested in human clinical trials as a potential cancer virotherapy although the mechanism whereby reovirus infected tumor cells are recognized and eliminated by the immune system is not well-understood. The reovirus σ1 protein forms an elongate trimer that serves as a viral attachment protein (188). Human and mouse NKp46 were shown to bind to reovirus σ1 protein in a sialic acid-dependent manner that led to NK cell activation in vitro (189) (Figure 2). NK cells and NKp46 were shown to be essential for the clearance of reovirus infection from the lungs of infected mice and for the success of reovirus-based tumor therapy (189).

**Bi-Specific T Cell Engagers (BiTE)**

The NKp30/B7-H6 interaction has been targeted using various therapeutic strategies. One approach has been to target T cells to B7-H6 expressing tumors using a BiTE that incorporated a single-chain Fv (ScFv) to B7-H6 fused to an anti-CD20 ScFv. The B7-H6-specific BiTE promoted T cell cytotoxicity and IFN-γ secretion against B7-H6+ tumor cells, which enhanced the survival of lymphoma-bearing mice and decreased the tumor burden of melanoma- and ovarian cancer-bearing mice (190). Similar conceptual approaches, have involved fusing the B7-H6 ectodomain (with the aim of engaging NKp30 on NK cells) to either an anti-Human Epidermal Growth factor 2 (HER2) scFv to enhance NK cell killing of HER2+ breast cancers (191) or an anti-CD20 scFv to target lymphomas (192, 193). Such reagents were reported to be effective at nM concentrations and to enhance ADCC mediated by therapeutic antibodies, such as trastuzumab, cetuximab, and rituximab (191–193).

**NCR-Based Chimeric Antigen Receptors (CARs)**

Yet another therapeutic strategy has been to engineer chimeric antigen receptor (CAR) constructs that incorporate either the NKp30 ectodomain (NKp30-CAR) or an anti-B7-H6 scFv (B7H6-specific CAR) with TM domains fused to the CD3ζ or and/or CD28 cytoplasmic signaling domains for cell-surface expression in T or NK cells intended for use in adoptive cancer immunotherapy (194–196). NKp30-CAR and B7-H6-specific CAR T cells elicited robust cellular cytotoxicity and IFN-γ secretion when co-cultured with B7-H6+ tumor cells. Dendritic cells (DCs) also enhanced IFN-γ secretion from NKp30-CAR expressing T cells, whereas B7-H6-specific CAR T cells exhibited little self-reactivity to DCs and monocytes expressing B7-H6 ligands. The adoptive transfer of T cells expressing NKp30-CAR or B7-H6-specific CAR T cells enhanced the survival of mice bearing RMA lymphoma cells transduced with B7-H6. Moreover, mice that remained tumor-free, were resistant to subsequent rechallenge with B7-H6-deficient parental RMA tumor cells (194, 195). NKp44- and NKp46-derived CARs have also been developed and were shown to elicit clinical efficacy for various solid tumors that express ligands for these receptors (197, 198).

**DISCUSSION**

The NCRs were originally identified as activating receptors expressed on NK cells. However, it is now apparent the NCRs are expressed by ILCs in addition to adaptive Vδ1+ and CD8+ T cells. The NCRs have been shown to synergize with TCR signaling as well as function independently on T cells. Thus, like NK cells, Vδ1+ and CD8+ T cells expressing NCRs hold potential for adoptive cancer immunotherapy (147, 199). It will be interesting to see how NCR+ Vδ1 T cells and CD8+ T cell functions are regulated by the range of NCR ligands now reported in the literature.

The NCRs can interact with soluble as well as membrane bound ligands. Such polyfunctionality is not uncommon in the immune system (25). For example, the activating immunoreceptor OSCAR signals via FcγR and possesses two Ig-like domains similar to NKp46 and genes for both proteins are encoded in the Leukocyte Receptor Complex (LRC), suggesting a common evolutionary origin (200). OSCAR interacts with at least three different types of ECM collagens to costimulate osteoclastogenesis (201–203) and also binds to the secreted collagen-like lectin, Surfactant Protein D (SP-D) (204). The LRC-encoded immunoreceptors, LAIR-1 and LAIR-2, also bind to multiple ECM collagens as well as SP-D (205–207). Moreover, NKG2D, a member of the C-type lectin family of activating NK cell receptors, is highly promiscuous and binds to a number of ligands.
NCRs all bind the same classes of ligands, such as viral HAs and cellular HS GAGs? Structural studies are clearly required to resolve many of these outstanding questions and may reveal fascinating insights into the polyfunctionality of the NCRs as well as intriguing comparisons for other immunoreceptor families. Such a structurally diverse set of soluble and surface bound NCR ligands might also be expected to induce different strengths and duration of activating or inhibitory NCR signaling that may regulate NK cell activity in different tissues and inflammatory settings. Recent data have raised the importance of secreted ligands in the “alternative activation” of NK cells and the NCRs no exception (64, 67, 210). For example, PDGF-DD has been reported to bind to NKp44 to induce the secretion of IFN-γ and TNF-α and chemokines, such as CCL1, CCL3, CCL4, XCL1 and XCL2 (67), whereas CFP binding to NKp46 only elicits the expression of XCL1 (64). Firstly, these data challenge the view that stimulation with soluble or shed ligands invariably leads to the desensitization of activating receptor pathways and NK cell inhibition. Secondly, the range of different response to soluble ligands suggests the NCR interaction with each ligand may be tailored to the tissue microenvironment for a specific function or to avoid immunopathology. For example, CFP binding to NKp46, is critical for resistance to N. meningitidis infection, which causes septicaemia and meningitis (64). It is conceivable that the reduced NK cell activation induced by the CFP/NKp46 interaction may be desirable in the meninges given the proximity of the inflammatory response to the brain. Further research into the molecular basis and cell biology of NCR signaling for surface-bound and soluble ligands is warranted to fully understand how soluble ligands, such as PDGF-DD and CFP, as well as shed ligands, such as MULTI, (210), evoke appropriate NK cell activation in different organs and tissues and within the tumor microenvironment (212). Such approaches will inform methods to enhance NK cell targeting and to stimulate or inhibit their functions in vivo and will lead to the development of novel clinical interventions (212, 213). For example, CARs have already been engineered for soluble growth factors, such as TGF-β, that function in T cells and promote anti-tumor responses in vivo (67, 214, 215).

Many of the proposed NCR ligands are normally localized to the nucleus or cytoplasmic proteins, such as BAT3, NKp44L, and PCNA. However, the cellular pathways responsible for the cell-surface localization of these ligands and their selective upregulation in malignant vs. non-malignant cells still remain to be characterized in molecular detail. These questions are critical for understanding how the NCR ligands regulate the activity of NK cells and other NCR+ immune cell subsets in different tissues and pathological processes. Interestingly, the NCRs and their ligands are associated with a diverse range of diseases and biological traits (Table 2). It might be expected that many of these disease associations are independent of the NCRs and result from changes to the primary biological function of the NCR ligand in question. However, it is also feasible that the NCRs and their ligands could play a pathogenetic role for some of these aforementioned diseases. Further research into the role of the NCRs and their ligands that are genetically associated with disease should be encouraged and will provide novel insights into the role of tissue NK cells for diseases that may have an as yet unrecognized immunopathological component.

AUTHOR CONTRIBUTIONS

AB, CM, and MC authored and edited the manuscript. CM contributed figures and tables.

ACKNOWLEDGMENTS

CM was the recipient of a Susan Keyes-Pearce Microbiology and Immunology laboratory experience award.

REFERENCES

1. Aw Yeang HX, Piersma SJ, Lin Y, Yang L, Malkova ON, Miner C, et al. Cutting edge: human CD49e- NK cells are tissue resident in the liver. J Immunol. (2017) 198:1417–22. doi: 10.4049/jimmunol.1601818
2. Crinier A, Milpied P, Escalier B, Piperoglou C, Galluso J, Balsamo A, et al. High-dimensional single-cell analysis identifies organ-specific signatures and conserved NK cell subsets in humans and mice. Immunity. (2018) 49:971–86 e5. doi: 10.1016/j.immuni.2018.09.009
3. Sojka DK, Plougastel-Douglas B, Yang L, Pak-Wittel MA, Artyomov MN, Ivanova Y, et al. Tissue-resident natural killer (NK) cells are cell lineages distinct from thymic and conventional splenic NK cells. eLife. (2014) 3:e01659. doi: 10.7554/eLife.01659
4. Sharkey AM, Xiong S, Kennedy PR, Gardner L, Farrell LE, Chazara O, et al. Tissue-specific education of decidual NK cells. J Immunol. (2014) 193:3026–32. doi: 10.4049/jimmunol.1501229
5. Kiessling R, Klein E, Wiggell H. “Natural” killer cells in the mouse. I. Cytotoxic cells with specificity for mouse Moloney leukemia cells. Specificity and distribution according to genotype. Eur J Immunol. (1973) 5:112–7. doi: 10.1002/eji.183050208
6. Herberman RB, Nunn ME, Lavrin DH. Natural cytotoxic reactivity of mouse lymphoid cells against syngeneic acid allogeneic tumors. I. Distribution of reactivity and specificity. Int J Cancer. (1975) 16:216–29.
7. Ljunggren HG, Kärre K. In search of the “missing self”: MHC molecules and NK cell recognition. Immunol Today. (1990) 11:237–44.
8. Bix M, Liao NS, Zijlstra M, Loring J, Jaenisch R, Raulet D. Rejection of class I MHC-deficient haemopoietic cells by irradiated MHC-matched mice. Nature. (1991) 349:329–31. doi: 10.1038/349329a0
9. Liao NS, Bix M, Zijlstra M, Jaenisch R, Raulet D. MHC class I deficiency: susceptibility to natural killer (NK) cells and impaired NK activity. Science. (1991) 253:199–202.
10. Karlhofer FM, Ribaudo RK, Yokoyama WM. MHC class I alloantigen specificity of Ly-49+ IL-2-activated natural killer cells. Nature. (1992) 358:66–70. doi: 10.1038/358066a0
11. Colonna M, Samaridis J. Cloning of immunoglobulin-superfamily members associated with HLA-C and HLA-B recognition by human natural killer cells. Science. (1995) 268:405–8.

12. Wagtmann N, Biassoni R, Cantoni C, Verdiann S, Malnati MS, Vitale M, et al. Molecular clones of the p58 NK cell receptor reveal immunoglobulin-related molecules with diversity in both the extra- and intracellular domains. Immunity. (1995) 2:239–49.

13. Colonna M, Brooks EG, Falco M, Ferrara GB, Strominger JL. Generation of allospecific natural killer cells by stimulation across a polymorphism of HLA-C. Science. (1993) 260:1121–4.

14. Malnati MS, Peruzzi M, Parker KC, Biddison WE, Ciccone E, Moretta A, et al. Identification and molecular characterization of NKp30, a novel triggering receptor involved in natural cytotoxicity mediated by human natural killer cells. J Exp Med. (1999) 190:1505–16.

15. Sivori S, Vitale M, Morelli L, Sanseverino L, Augugliaro R, Bottino C, et al. p46, a novel natural killer cell–specific surface molecule that mediates cell activation. J Exp Med. (1997) 186:1129–36.

16. Vivier E, Ackery M, Mechtlich H, Levine H, Rudd CE, et al. Phenotypic and functional plasticity of murine intestinal lymphocytes develop in the intestine and transform into lymphomas in celiac disease. Immunity. (2016) 45:610–25. doi: 10.1016/j.immuni.2016.07.018

17. Vivier E, Ackerly M, Rochet N, Anderson P. Structure and function of the CD16:zeta/gamma complex expressed on human natural-killer cells. Int J Cancer Suppl. Int J Cancer Suppl. (1992) 7:11–4.

18. Vivier E, da Silva AJ, Ackery M, Levine H, Rudd CE, et al. Association of a 70-kDa tyrosine phosphoprotein with the CD16: zeta: gamma complex expressed in human natural killer cells. Eur J Immunol. (1993) 23:1872–6. doi: 10.1002/eji.1830230821

19. F支出 D, Parolini S, Pessino A, Sivori S, Augugliaro R, Morelli L, et al. NKp44, a novel triggering surface molecule specifically expressed by activated natural killer cells, is involved in non-major histocompatibility complex-restricted tumor cell lysis. J Exp Med. (1998) 187:2055–72.

20. Pessino A, Sivori S, Bottino C, Malaspina A, Morelli L, Moretta L, et al. Molecular cloning of NKp46: a novel member of the immunoglobulin superfamily involved in triggering of natural cytotoxicity. J Exp Med. (1998) 188:953–60.

21. Dietrich J, Nakajima H, Colonna M. Human inhibitory and activating Ig-like receptors which modulate the function of myoidoid cells. Microbes Infect. (2000) 2:323–9. doi: 10.1016/S1286-4579(00)00294-X

22. Foster CE, Colonna M, Sun PD. Crystal structure of the human natural killer (NK) cell activating receptor NKp46 reveals structural relationship to other leukocyte receptor complex immunoreceptors. J Biol Chem. (2003) 278:46081–6. doi: 10.1074/jbc.M304912000

23. Pessino A, Sivori S, Bottino C, Malaspina A, Morelli L, Moretta L, et al. NKp44, a triggering receptor involved in tumor cell lysis by activated human natural killer cells, is a novel member of the immunoglobulin superfamily. J Exp Med. (1999) 189:787–96.

24. Shemesh A, Kugel A, Steinier N, Yezermsky M, Tirosh D, Edri A, et al. NKp44 and NKp30 splice variant profiles in decidua and tumor tissues: a comparative viewpoint. Oncotarget. (2017) 6:70912–23. doi: 10.18632/oncotarget.12292

25. Lanier LL, Coro CL, Wu J, Leong C, Phillips JH. Immunoreceptor DPAPD2 bearing a tyrosine-based activation motif is involved in activating NK cells. Nature. (1998) 391:703–7. doi: 10.1038/35642

26. Tomassello O, Olesc K, Vell E, Geourgeon C, Bléry M, Moqrich A, et al. Gene structure, expression pattern, and biological activity of mouse killer cell activating receptor–associated protein (KARAP)/DAP-12. J Biol Chem. (1999) 274:34115–9.

27. Mattioli L, Pesant M, Tontoriento PF, Molgopi M, Marcenaro E, Lugli E, et al. Priming of human resting NK cells by autologous M1 macrophages via the engagement of IL-1β, IFN-β, and IL-15 pathways. J Immunol. (2015) 195:2818–28. doi: 10.4049/jimmunol.1500325

28. Cantoni C, Bottino C, Vitale M, Pessino A, Augugliaro R, Malaspina A, et al. NKp44, a triggering receptor involved in tumor cell lysis by activated human natural killer cells, is a novel member of the immunoglobulin superfamily. J Exp Med. (2013) 267:117–24.

29. Dietrich J, Nakajima H, Colonna M. Human inhibitory and activating Ig-like receptors which modulate the function of myoidoid cells. Microbes Infect. (2000) 2:323–9. doi: 10.1016/S1286-4579(00)00294-X

30. Bonomi M, Velia F, Barrow AD, Forbes S, Beck S, Trowsdale J. The human TR EM receptor involved in natural cytotoxicity mediated by human natural killer cells. J Exp Med. (1998) 187:2055–72.

31. Foster CE, Colonna M, Sun PD. Crystal structure of the human natural killer (NK) cell activating receptor NKp46 reveals structural relationship to other leukocyte receptor complex immunoreceptors. J Biol Chem. (2003) 278:46081–6. doi: 10.1074/jbc.M304912000

32. Ponassi M, Cantoni C, Biassoni R, Conte R, Spallarossa A, Pesce A, et al. Structure of the human NK cell triggering receptor NKp46 ectodomain. Biochem Biophys Res Commun. (2003) 309:317–23. doi: 10.1016/j.bbrc.2003.08.007

33. Trowsdale J, Jones DC, Barrow AD, Traherne JA. Surveillance of cell and tissue perturbation by receptors in the LRC. Immunol. Rev. (2017) 267:116–37. doi: 10.1111/imr.12314

34. Cantoni C, Garzoni M, Biassoni R, Conte R, Spallarossa A, Moretta A, et al. The three-dimensional structure of the human NK cell receptor NKp44, a triggering partner in natural cytotoxicity. Struct. Eng. (1993) 113:75–34.

35. Allocock RJN, Barrow AD, Forbes S, Beck S, Trowsdale J. The human TREM gene cluster at 6p21.1 encodes both activating and inhibitory single IgV domain receptors and includes NKp44. Eur J Immunol. (2003) 33:567–77. doi: 10.1002/immu.200310033

36. De Maria A, Ugelotti E, Rutjens E, Mazza S, Radic L, Faravelli A, et al. NKp44 expression, phylogenesis and function in non-human primate NK cells. Int Immunol. (2009) 21:245–55. doi: 10.1093/intimm/dxn144

37. Campbell KS, Yusa S, Kikuchi-Maki A, Carino TL. NKp44 triggers NK cell activation through DAP12 association that is not influenced by a putative cytoplasmic inhibitory sequence. J Immunol. (2004) 172:899–906.

38. Cortez VS, Ulland TK, Cervantes-Barragan L, Bando JK, Robinson ML, Wang Q, et al. SMAD4 impedes the conversion of NK cells into ILC1-like cells by curtailing non-canonical TGF-β signaling. Nature. (2017) 548:140–7. doi: 10.1038/nn.4327

39. Correia DV, Fogli M, Hudspeth K, da Silva MG, Mavilio D, Silva Santos DM, et al. An innate source of natural killer and T-cell neoplasia. Am J Clin Pathol. (2011) 131:969–76. doi: 10.1093/ajcp/wgr005

40. Zeng C, Yusa S, Kikuchi-Maki A, Carino TL, Bando JK, Robinson ML, et al. SMAD4 impedes the conversion of NK cells into ILC1-like cells by curtailing non-canonical TGF-β signaling. Nature. (2017) 548:140–7. doi: 10.1038/nn.4327
IL-22 for mucosal immunity. *Nature.* (2009) 457:22–5. doi: 10.1038/nature07537

49. Cella M, Otero K, Colonna M. Expansion of human NK-22 cells with IL-7, IL-2, and IL-1beta reveals intrinsic plasticity. *Proc Natl Acad Sci USA.* (2010) 107:9161–6. doi: 10.1073/pnas.1005641107

50. von Lilienfeld-Toal M, Nattemann J, Feldmann G, Sievers E, Frank S, Strehl J, et al. Activated gammadelta T cells express the natural cytotoxicity receptor natural killer p 44 and show cytotoxic activity against myeloma cells. *Clin Exp Immunol.* (2006) 144:528–33. doi: 10.1111/j.1365-2249.2006.03078.x

51. Fuchs A, Cella M, Kondo T, Colonna M. Paradoxic inhibition of human natural interferon-producing cells by the activating receptor NKp44. *Blood.* (2005) 106:2076–82. doi: 10.1182/blood-2004-12-4802

52. Hecht M-L, Rosental B, Horlacher T, Hershkovitz O, De Paz JL, Noti C, et al. Regulation of natural cytotoxicity receptors by heparan sulfate/heparin sequences. *J Proteome Res.* (2009) 8:712–20. doi: 10.1021/pr080747c

53. Hershkovitz O, Jarahian M, Zilka A, Bar-Ilan A, Landau G, Jivov V, et al. Recognition of haemagglutinins on virus-infected cells by NKp46 activates lysis by human NK cells. *Nature.* (2001) 409:1055–60. doi: 10.1038/s769110

54. Jarahian M, Fiedler M, Cohnen A, Djandji D, Hämerling GJ, Gati C, et al. Modulation of NKp30- and NKp46-mediated natural killer cell responses by poxviral hemagglutinin. *PLoS Pathog.* (2011) 7:e1002195. doi: 10.1371/journal.ppat.1002195

55. Jarahian M, Watzi C, Fournier P, Arnold A, Djandji D, Zahedi S, et al. Activation of natural killer cells by Newcastle disease virus hemagglutinin-neuraminidase. *J Virol.* (2009) 83:8108–21. doi: 10.1128/JVI.00211-09

56. Arnon TI, Achdout H, Lieberman N, Gazit R, Gonen-Gross T, Katz G, et al. The mechanisms controlling the recognition of tumor- and virus-infected cells by NKp46. *Blood.* (2004) 103:664–72. doi: 10.1182/blood-2003-05-1716

57. McQuaid S, Loughran S, Power P, Maguire P, Walls D, Cusi MG, et al. Identification of a cellular ligand for the natural cytotoxicity receptor NKp44. *Blood.* (2005) 106:2076–82. doi: 10.1182/blood-2004-12-4802

58. Mavoungou E, Held J, Mewono L, Kremsner PG. Duffy binding-like domain is involved in the NKp46-mediated recognition of Plasmodium falciparum-parasitized erythrocytes by natural killer cells. *J Infect Dis.* (2007) 195:1521–31. doi: 10.1086/515579

59. Vankayalapati R, Garg A, Porgador A, Griffith DE, Klucar P, Safi H, et al. Role of NK cell-activating receptors and their ligands in the lysis of mononuclear phagocytes infected with an intracellular bacterium. *J Immunol.* (2005) 175:4611–7. doi: 10.4049/jimmunol.175.7.4611

60. Garg A, Barnes PF, Porgador A, Roy S, Wu S, Nanda JS, et al. Vimentin expressed on *Mycobacterium tuberculosis*-infected human monocytes is involved in binding to the NKp46 receptor. *J Immunol.* (2006) 177:6192–8. doi: 10.4049/jimmunol.177.6.6192

61. Chaushu S, Wilensky A, Gur C, Shapira L, Elboim M, Halftek G, et al. Direct recognition of Fusobacterium nucleatum by the human natural killer-activating receptor NKp44. *Blood.* (2013) 121:534–48 e19. doi: 10.1182/blood-2013-03-489054

62. Rosental B, Gaggero S, Bruschi M, Petretto A, Parodi M, Zotto GD, Lavarello C, et al. Human leukocyte antigen-B-associated transcript-3 positive dendritic cells release HLA-B-associated transcript-3 positive vesicles with antitumor activity. *Oncoimmunology.* (2016) 5:e1219827. doi: 10.1080/2162402X.2016.1219827

63. Hershkovitz O, Barrow AD, Edeling MA, Trifonov V, Luo J, Goyal P, Bohl B, et al. Natural killer cells control tumor growth by sensing a growth factor. *Cell.* (2018) 172:534–48 e19. doi: 10.1016/j.cell.2017.11.037

64. Gaggero S, Bruschi M, Petretto A, Parodi M, Zotto GD, Lavarello C, et al. Id agon-1 is a novel extracellular ligand for the NKp44 activating receptor. *Oncoimmunology.* (2018) 7:e1470730. doi: 10.2126420X.2018.1470730

65. Rosental B, Brusilovsky M, Hadad U, Oz D, Appel MY, Afergan E, et al. Proliferating cell nuclear antigen is a novel inhibitory ligand for the natural cytotoxicity receptor NKp44. *J Immunol.* (2017) 188:5693–702. doi: 10.4049/jimmunol.1102267

66. Hershkovitz O, Rosental B, Rosenberg LA, Navarro-Sanchez ME, Jivov S, Zilka A, et al. NKp44 receptor mediates interaction of the envelope glycoproteins from the West Nile and dengue viruses with NK cells. *J Immunol.* (2009) 183:2610–21. doi: 10.4049/jimmunol.0802806

67. Eison S, Batoni G, Cownoupas C, Stringaro A, Brancaziano FL, Colone M, et al. Direct binding of human NK cell natural cytotoxicity receptor NKp44 to the surfaces of mycobacteria and other bacteria. *Infect Immun.* (2008) 76:1719–27. doi: 10.1128/IAI.00870-07

68. Eison S, Cownoupas C, Aulicino A, Brancaziano FL, Maisetta G, Bottai D, et al. Interaction of *Mycobacterium tuberculosis* cell wall components with the human natural killer cell receptors NKp44 and Toll-like receptor 2. *Scand J Immunol.* (2013) 77:460–9. doi: 10.1111/sji.12052

69. Hershkovitz O, Jarahian M, Zilka A, Bar-Ilan A, Landau G, Jivov S, et al. Altered glycosylation of recombinant NKp30 hampers binding to heparan sulfate: a lesson for the use of recombinant immunoreceptors as an immunological tool. *Glycobiology.* (2008) 18:28–41. doi: 10.1093/glycob/cwn125

70. Warren HS, Jones AL, Bettadapura J, Parish CR. Evidence that the cellular ligand for the human NK cell activation receptor NKp30 is not a heparan sulfate glycosaminoglycan. *J Immunol.* (2005) 175:207–12.

71. Arnon TI, Achdout H, Levi O, Markel G, Saleh N, Katz G, et al. Inhibition of the NKp30 activating receptor by pp65 of human cytomegalovirus. *Nat Immunol.* (2005) 6:515–23. doi: 10.1038/ni1190

72. Pogge von Strandmann E, Simhadri JR, von Tresekwok S, Sasse S, Reiners KS, Hansen HP, et al. Human leukocyte antigen B-associated transcript 3 is released from tumor cells and engages the NKp30 receptor on natural killer cells. *Immunity.* (2007) 27:965–74. doi: 10.1016/j.immuni.2007.10.010

73. Daffler-Plenker J, Reiners KS, van den Boorn JG, Hansen HP, Putschli B, Barnert S, et al. RIG-I activation induces the release of extracellular vesicles with antitumor activity. *Oncoimmunology.* (2016) 5:e1219827. doi: 10.1080/2162402X.2016.1219827

74. Simhadri JR, Reiners KS, Hansen HP, Topolar D, Simhadri VL, Nohroudi K, et al. Dendritic cells release HLA-B-associated transcript-3 positive exosomes to regulate natural killer function. *PLoS ONE.* (2015) 3:e3377. doi: 10.1371/journal.pone.003377

75. Long Y, Wang Q, Marziutza RA. Structure of the human activating natural cytotoxicity receptor NKp30 bound to its tumor cell ligand B7-H6. *J Exp Med.* (2011) 208:703–14. doi: 10.1084/jem.20102548

76. Brandt CS, Baratin M, Yi EC, Kennedy J, Gao Z, Fox B, et al. The B7 family member B7-H6 is a tumor cell ligand for the activating natural
85. Wang W, Guo H, Geng J, Zheng X, Wei H, Sun R, et al. Tumor-released Galectin-3, a soluble inhibitory ligand of human NKP30, plays an important role in tumor escape from NK cell attack. *J Biol Chem.* (2014) 289:33311–9. doi: 10.1074/jbc.M114.546157

86. Li SS, Ogimoto H, Mansour MK, Xiang RF, Szabo L, Munro F, et al. Identification of the fungal ligand triggering cytotoxic PRR-mediated NK cell killing of *Cryptococcus* and *Candida*. *Nat Commun.* (2018) 9:751. doi: 10.1038/s41467-018-03014-4

87. Xiang RF, Li S, Ogimoto H, Stack D, Mody CH. β sulfatases in the modification of heparan sulfate proteoglycans within the tumor microenvironment and opportunities for novel cancer therapeutics. *Front Oncol.* (2014) 4:195. doi: 10.3389/fonc.2014.00195

88. Barrow et al. NCRs in Health and Disease

90. Schlums H, Cichocki F, Tesi B, Theorell J, Beziat V, Holmes TD, et al. NCRs in the mouse strain: NK cells are essential for protection against pulmonary B16 melanoma cell lines. *J Clin Invest.* (2012) 128:1661–72. doi: 10.1172/JCI62402

91. Joyce MG, Tran P, Zhuravleva MA, Jaw J, Colonna M, Sun PD. Crystal structure of human natural cytotoxicity receptor NKP30 and identification of its ligand binding site. *Proc Natl Acad Sci USA.* (2011) 108:6232–8. doi: 10.1073/pnas.1016221010

92. Schlums H, Cichocki F, Tesi B, Theorell J, Beziat V, Holmes TD, et al. Cytoxan-mediated ovarian cancer cells with altered signaling and effector function. *Immunity.* (2015) 42:443–56. doi: 10.1016/j.immuni.2015.02.008

93. Lee J, Zhang T, Hwang I, Kim A, Nitschke L, Kim M, et al. Epigenetic modification and antibody-dependent expansion of memory-like NK cells in human cytomegavirus-infected individuals. *Immunity.* (2015) 42:431–42. doi: 10.1016/j.immuni.2015.02.013

94. Hollyoake M, Campbell RD, Aguado B. NKP30 (NCR3) is a pseudogene in 12 inbred and wild mouse strains, but an expressed gene in Mus caroli. *Mol Biol Evol.* (2005) 22:1661–72. doi: 10.1093/molbev/msi162

95. Delahaye NF, Rusakiewicz S, Martins I, Ménard C, Roux S, Lyonnnet L, et al. Alternatively spliced NKP30 isoforms affect the prognosis of gastrointestinal stromal tumors. *Nat Med.* (2011) 17:790–7. doi: 10.1038/nm.2366

96. Correia MP, Stojanovic A, Bauer K, Juraeva D, Tykocinski L-O, Lorenz H-M, et al. Distinct human circulating NKP30+FcεRI+CD4+ T cell population exhibiting high natural killer-like antitumor potential. *Proc Natl Acad Sci USA.* (2018) 115:5898–90. doi: 10.1073/pnas.1710541115

97. Halfteck GG, Elboim M, Gur C, Achdout H, Ghadially H, Mandelboim O. Enhanced in vivo growth of lymphoma tumors in the absence of the NK-activating receptor NKP46/NCR1. *J Immunol.* (2009) 182:2221–30. doi: 10.4049/jimmunol.0801878

98. Lakshmikanth T, Burke S, Ali TH, Kimpfler S, Ursini F, Ruggeri L, et al. NCRs and DNAM-1 mediate NK cell recognition and lysis of human and mouse melanoma cell lines in vitro and in vivo. *J Clin Invest.* (2009) 119:1251–63. doi: 10.1172/JCI36022

99. Glasner A, Ghadially H, Gur C, Stanietisky N, Tsukerman P, Enk J, et al. Recognition and prevention of tumor metastasis by the NK receptor NKP46/NCR1. *J Immunol.* (2012) 188:2509–15. doi: 10.4049/jimmunol.1102461

100. Merzoug LB, Marie S, Satoh-Takayama N, Lesjean S, Albanesi M, Luche H, et al. Conditional ablation of NKp46+ cells with altered signaling and effector function. *Front Immunol.* (2017) 8:369. doi: 10.3389/fimmu.2017.00369

101. Horton NC, Mathew PA. NKp44 and natural cytotoxicity receptors as damage-associated molecular pattern recognition receptors. *Front Immunol.* (2015) 6:31. doi: 10.3389/fimmu.2015.00031

102. Reiners KS, Topolar D, Henke A, Simhadri VR, Kessler J, Sauer M, et al. Soluble ligands for NK cell receptors promote evasion of chronic
lymphocytic leukemia cells from NK cell anti-tumor activity. Blood. (2013) 121:3658–65. doi: 10.1182/blood-2013-01-476606

123. Binici J, Hartmann J, Herrmann J, Schreiber C, Beyer S, Güler G, et al. A soluble fragment of the tumor antigen BCL2-associated athanogene 6 (BAG-6) is essential and sufficient for inhibition of NKp30 receptor-dependent cytotoxicity of natural killer cells. J Biol Chem. (2013) 288:34295–302. doi: 10.1074/jbc.M113.483602

124. Fredriksson L, Li H, Eriksson U. The PDGF family: four gene products form five dimeric isoforms. Cytokine Growth Factor Rev. (2004) 15:197–204. doi: 10.1016/j.cytogfr.2004.03.007

125. Andrae J, Gallini R, Betscholtz C. Role of platelet-derived growth factors in physiology and medicine. Genes Dev. (2008) 22:1276–312. doi: 10.1101/gad.1653708

126. Glatter T, Killig M, Meisig J, Ommert I, Luetke-Eversloh M, Babic M, et al. ROXβ+ innate lymphoid cells acquire a proinflammatory program upon engagement of the activating receptor NKp44. Immunity. (2013) 38:1223–35. doi: 10.1016/j.immuni.2013.05.013

127. Ahola-Olli AV, Würtz P, Havulinna AS, Aalto K, Pitkänen N, Lehtimäki T, et al. Genome-wide association study identifies 27 loci influencing concentrations of circulating cytokines and growth factors. Am J Hum Genet. (2017) 100:40–50. doi: 10.1016/j.ajhg.2016.11.007

128. Ho MSP, Böse K, Mokkapati S, Nischt R, Smyth N. Nidogens-extracellular matrix linker molecules. Micros Res Tech. (2008) 71:387–95. doi: 10.1016/j.mretc.2007.05.007

129. Li L, Zhang Y, Li N, Feng L, Yao H, Zhang R, et al. Nidogen-1: a candidate biomarker for ovarian serous cancer. Ipn J Clin Oncol. (2015) 45:176–82. doi: 10.1093/jjco/hyu187

130. Willemsen N, Bager CL, Leeming DJ, Bay-Jensen A-C, Karsdal MA, et al. Transforming growth factor-β extracellular matrix linker molecules. Clin Cancer Res. (2017) 23:368–75. doi: 10.1158/1078-0432.CCR-16-0538

131. Farhad M, Rolig AS, Redmond WL. The role of Galectin-3 in apoptosis-inducing ligand-deficient mice. J Exp Med. (2011) 204:1409–14. doi: 10.1084/jem.20101287

132. Barrow AD, Colonna M. Innate lymphoid cell sensing of tissue vitality. Curr Opin Immunol. (2018) 56:82–93. doi: 10.1016/j.coi.2018.10.004

133. Cortez VS, Cervantes-Barragan L, Robinette ML, Bando JK, Wang Y, Geiger TL, et al. Transforming growth factor-β signaling guides the differentiation of innate lymphoid cells in salivary glands. Immunity. (2016) 44:1127–39. doi: 10.1016/j.immuni.2016.03.007

134. Smyth MJ, Cretney E, Takeda K, Wiltrout RH, Sedger LM, Kayagaki N, et al. Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) contributes to interferon γ-dependent natural killer cell protection from tumor metastasis. J Exp Med. (2001) 193:661–70. doi: 10.1084/jem.193.6.661

135. Stegmann KA, Björkström NK, Veber H, Ciesek S, Riese P, Wiegand J, et al. Interferon-alpha-induced TRAIL on natural killer cells is associated with control of hepatitis C virus infection. Gastroenterology. (2010) 138:1885–97. doi: 10.1053/j.gastro.2010.01.051

136. Kayagaki N, Yamaguchi N, Nakayama M, Takeda K, Akiba H, Tsutsui H, et al. Expression and function of TNF-related apoptosis-inducing ligand on murine activated NK cells. J Immunol. (1999) 163:1906–13.

137. Takeda K, Smyth MJ, Cretney E, Hayakawa Y, Yamaguchi N, Yagita H, et al. Involvement of tumor necrosis factor-related apoptosis-inducing ligand in NK cell-mediated and IFN-γamma-dependent suppression of subcutaneous tumor growth. Cell Immunol. (2001) 214:194–200. doi: 10.1006/cimm.2001.1896

138. Cretney E, Takeda K, Yagita H, Glaccum M, Peschon JJ, Smyth MJ. Increased susceptibility to tumor initiation and metastasis in TNF-related apoptosis-inducing ligand-deficient mice. J Immunol. (2002) 168:1356–61.

139. Seki N, Hayakawa Y, Brooks AD, Wine J, Wiltrout RH, Yagita H, et al. Tumor necrosis factor-related apoptosis-inducing ligand-mediated apoptosis is an important endogenous mechanism for resistance to liver metastases in murine renal cancer. Cancer Res. (2003) 63:207–13.
cogenic mice with consequences in mouse susceptibility to infection. J Immunol. (2018) 200:1982–7. doi: 10.4049/jimmunol.1701676

160. Mao H, Tu W, Liu Y, Qin G, Zheng J, Chan P-L, et al. Inhibition of human natural killer cell activity by influenza viros and hemagglutinin. J Virol. (2010) 84:11488–57. doi: 10.1128/JVI.02340-09

161. Bar-On Y, Seidel E, Tsukerman P, Mandelboim M, Mandelboim O. Influenza virus uses its neuraminidase protein to evade the recognition of two activating NK cell receptors. J Infect Dis. (2014) 210:410–8. doi: 10.1093/infdis/jiu094

162. Divaris K, Monda KL, North KE, Olshan AF, Reynolds LM, Hsueh K, et al. Selective cross-talk among natural cytotoxicity receptors in human natural killer cells. Eur J Immunol. (2003) 33:1235–41. doi: 10.1002/eji.2003323896

163. Augugliaro R, Parolini S, Castriconi R, Marcenaro E, Cantoni C, Nanni M, et al. Selective inhibition of NKp44 and NKp30 receptors in human NK cells. J Immunol. (2009) 182:1230–7. doi: 10.4049/jimmunol.2009.06.123

164. Li SS, Kyei SK, Timms-Cann M, Oghomo H, Jones GI, Shi M, et al. The NK receptor Nkp30 mediates direct fungal recognition and killing and is diminished in NK cells from HIV-infected patients. Cell Host Microbe. (2013) 14:387–97. doi: 10.1016/j.chom.2013.09.007

165. Kemper C, Atkinson JP, Hourcade DE. Properdin: emerging roles of a pattern-recognition molecule. Annu Rev Immunol. (2010) 28:131–55. doi: 10.1146/annurev-immunol-030409-101250

166. Sprong T, Roos D, Weemaes C, Neeleman C, Geesing CLM, Mollnes TE, et al. Deficient alternative complement pathway activation due to factor D deficiency by 2 novel mutations in the complement factor D gene in a family with meningococcal infections. Blood. (2006) 107:4865–70. doi: 10.1182/blood-2005-07-2820

167. Hudspheth K, Fogli M, Correia DV, Mikulak J, Roberto A, Della Bella S, Divaris K, Monda KL, North KE, Olshan AF, Reynolds LM, Hsueh K, et al. Inhibition of human natural killer cells regulate endometrial bleeding in women and are suppressed by the progesterone receptor modulator asoprisnil. J Clin Invest. (2014) 124:1872–9. doi: 10.1172/JCI68107

168. Filippovic I, Chiossone L, Vacca P, Hamilton RS, Inggegnere T, Doisne J-M, et al. Molecular definition of group 1 innate lymphoid cells in the mouse uterus. Nat Commun. (2018) 9:4492. doi: 10.1038/s41467-018-06918-3

169. Hanna J, Goldman-Wohl D, Hamani Y, Avraham I, Greenfield C, Nathan- Yaron S, et al. Decidual NK cells regulate endometrial bleeding in women and are suppressed by the progesterone receptor modulator asoprisnil. J Immunol. (2013) 191:2226–35. doi: 10.4049/jimmunol.1300958

170. Moffett A, Colucci F. Uterine NK cells act as regulators of the maternal-fetal interface. J Clin Invest. (2014) 124:1872–9. doi: 10.1172/JCI68107

171. Poirot L, Benoist C, Mathis D. Natural killer cells distinguish innocuous and destructive forms of pancreatic islet autoimmunity. Proc Natl Acad Sci USA. (2004) 101:8102–7. doi: 10.1073/pnas.0402065101

172. Alba A, Planas R, Clemente X, Carrillo J, Ampudia R, Puertas M-C, et al. Natural killer cells are required for accelerated type 1 diabetes driven by interferon-beta. Clin Exp Immunol. (2008) 151:467–75. doi: 10.1111/j.1600-0528.2007.03580.x

173. Traherne JA, Jiang W, Valdes AM, Hollenbach JA, Jaryamane J, Lana JE, et al. KIR haplotypes are associated with late-onset type 1 diabetes in European-American families. Genes Immun. (2016) 17:8–12. doi: 10.1038/genes.2015.44

174. Weng Y, Yuan W, Guo H, Jiang Y. High frequency of activated NKp46(+) natural killer cells in patients with new diagnosed of latent autoimmune diabetes in adults. Autoimmunity. (2015) 48:267–73. doi: 10.3109/08916934.2014.990629

175. Yossel R, Gur C, Shemesh A, Guttman O, Hadad U, Nedevtski S, et al. Targeting natural killer cell reactivity by employing antibody to NKp46: implications for type 1 diabetes. PLoS ONE. (2015) 10:e0118936. doi: 10.1371/journal.pone.0118936

176. Berhani O, Glasner A, Kahan S, Duev-Cohen A, Yamin R, Horwitz E, et al. Human anti-NKp46 antibody for studies of NKp46-dependent NK cell function and its applications for type 1 diabetes and cancer research. Eur J Immunol. (2018) 48:1002–7. doi: 10.1002/eji.201847611

177. Rusakiewicz W, Nocturne G, Lazure T, Semeraro M, Flamment C, Caillat-Zucman S, et al. NCR3/NKp30 contributes to pathogenesis in primary Sjogren’s syndrome. Sci Transl Med. (2013) 5:195ra96. doi: 10.1126/scitranslmed.3005727

178. Chappell JD, Prota AE, Dermody TS, Stehle T. Crystal structure of reovirus attachment protein σ1 reveals evolutionary relationship to adeno virus fiber. EMBO J. (2002) 21:1–11. doi: 10.1093/emboj/21.1.1

179. Bar-On Y, Charpak-Amikam Y, Glasner A, Isaacson B, Duev-Cohen A, Tsukerman P, et al. NKp46 Recognizes the signal protein of reovirus: implications for reovirus-based cancer therapy. J Virol. (2017) 91:e01045–17. doi: 10.1128/JVI.01045-17

180. Wu M-R, Zhang T, Gacerez AT, Coupet TA, DeMars LR, Sentman CL. B7-H6-specific bispecific T cell engagers lead to tumor elimination and host antitumor immunity. J Immunol. (2015) 194:5305–11. doi: 10.4049/jimmunol.1402517

181. Peipp M, Derer S, Lohse S, Staudinger M, Klausz K, Valerius T, et al. HER2-specific immunoligands engaging NKp30 or NKp80 trigger NK-cell-mediated lysis of tumor cells and enhance antibody-dependent cell-mediated cytotoxicity. Oncotarget. (2015) 6:32075–88. doi: 10.18632/oncotarget.5135

182. Kellner C, Maurer T, Hallack D, Repp R, van de Winkel JG, Parren PW, et al. Mimicking an induced self phenotype by coating lymphomas with the NKp30 ligand B7-H6 promotes NK cell cytotoxicity. J Immunol. (2012) 189:5037–46. doi: 10.4049/jimmunol.1201321

183. Kellner C, Günther A, Humpe A, Repp R, Klauss K, Merer D, et al. Enhancing natural killer cell-mediated lysis of lymphoma cells by combining therapeutic antibodies with CD20-specific immunoligands engaging NKp42 or NKp30. Oncomunology. (2016) 6:e0158439. doi: 10.2162/21642042X.2015.1058439

184. Zhang T, Wu M-R, Sentman CL. An NKp30-based chimeric antigen receptor promotes T cell effector functions and antitumor efficacy in vivo. J Immunol. (2012) 189:2290–9. doi: 10.4049/jimmunol.1103495
195. Wu M-R, Zhang T, DeMars LR, Sentman CL. B7H6-specific chimeric antigen receptors lead to tumor elimination and host antigenic immunity. *Gene Ther.* (2015) 22:875–84. doi: 10.1038/gt.2015.29

196. Hua CK, Gacerez AT, Sentman CL, Ackerman ME. Development of unique cytotoxic chimeric antigen receptors based on human scFv targeting B7H6. *Protein Eng Des Sel.* (2017) 30:713–21. doi: 10.1093/protein/gzx051

197. Tal Y, Yaakobi S, Horovitz-Fried M, Safyon E, Rosental B, Porgador A, et al. An NCR1-based chimeric receptor endows T-cells with multiple anti-tumor specificities. *Oncotarget.* (2014) 5:10949–58. doi: 10.18632/oncotarget.1919

198. Eisenberg V, Shamalov K, Meir S, Hoogi S, Sarkar R, Pinker S, et al. Targeting multiple tumors using T-cells engineered to express a natural cytotoxicity receptor 2-based chimeric receptor. *Front Immunol.* (2017) 8:1212. doi: 10.3389/fimmu.2017.01212

199. Cervenka A, Bakker AB, McClanahan T, Wagner J, Wu J, Phillips JH, et al. Retinoic acid early inducible genes define a ligand family for the activating NKG2D receptor in mice. *Immunity.* (2000) 12:721–7. doi: 10.1016/S1074-7613(00)80222-8

200. Barrow AD, Trowsdale J. The extended human leukocyte receptor complex: diverse ways of modulating immune responses. *Immunol Rev.* (2008) 224:98–123. doi: 10.1111/j.1600-065X.2008.00653.x

201. Barrow AD, Raynal N, Andersen TL, Slatter DA, Bihan D, Pugh N, et al. OSCAR is a collagen receptor that costimulates osteoclastogenesis in DAP12-deficient humans and mice. *J Clin Invest.* (2011) 121:3505–16. doi: 10.1172/JCI45913

202. Zhou L, Hinerman JM, Blaszczyk M, Miller JLC, Conrady DG, Barrow AD, et al. Structural basis for collagen recognition by the immune receptor OSCAR. *Blood.* (2016) 127:529–37. doi: 10.1182/blood-2015-08-667055

203. Haywood J, Qi J, Chen C-C, Lu G, Liu Y, Yan J, et al. Structural basis of collagen recognition by human osteoclast-associated receptor and design of osteoclastogenesis inhibitors. *Proc Natl Acad Sci USA.* (2016) 113:1038–43. doi: 10.1073/pnas.1522572113

204. Barrow AD, Palarash Y, Bugatti M, Holehouse AS, Byers DE, Holtzman MI, et al. OSCAR is a receptor for surfactant protein D that activates TNF-α release from human CCR2+ inflammatory monocytes. *J Immunol.* (2015) 194:3317–26. doi: 10.4049/jimmunol.1402289

205. Brondijk THC, de Ruiter T, Ballering J, Wienk H, Lebbink RJ, van Ingen H, et al. Crystal structure and collagen-binding site of immune inhibitory receptor LAIR-1: unexpected implications for collagen binding by platelet receptor GPVI. *Blood.* (2010) 115:1364–73. doi: 10.1182/blood-2009-10-246322

206. Lebbink RJ, Raynal N, de Ruiter T, Bihan DG, Farndale RW, Meyvaerd L. Identification of multiple potent binding sites for human leukocyte associated Ig-like receptor LAIR on collagens II and III. *Matrix Biol.* (2009) 28:202–10. doi: 10.1016/j.matbio.2009.03.005

207. Olde Nordkamp MJM, van Eijk M, Urbanus RT, Bont L, Haagsman HP, Meyevaard L. Leukocyte-associated Ig-like receptor-1 is a novel inhibitory receptor for surfactant protein D. *J Leukoc Biol.* (2014) 96:105–11. doi: 10.1189/jlb.3AB0213-092RR

208. Bauer S, Groh V, Wu J, Steinle A, Phillips JH, Lanier LL, et al. Activation of NK cells and T cells by NKG2D, a receptor for stress-inducible MICA. *Science.* (1999) 285:727–9.

209. Eagle RA, Trowsdale J. Promiscuity and the single receptor: NKG2D. *Nat Rev Immunol.* (2007) 7:737–44. doi: 10.1038/nri2144

210. Deng W, Gowen BG, Zhang L, Wang L, Lau S, Iannello A, et al. Antitumor immunity. A shed NKG2D ligand that promotes natural killer cell activation and tumor rejection. *Science.* (2015) 348:136–9. doi: 10.1126/science.1258867

211. Eagle RA, Flack G, Warford A, Martinez-Borra J, Jafferji I, Traharne JA, et al. Cellular expression, trafficking, and function of two isoforms of human ULBP5/RAET1G. *PLoS ONE.* (2009) 4:e4503. doi: 10.1371/journal.pone.0004503

212. Barrow AD, Colonna M. Tailoring natural killer cell immunotherapy to the tumour microenvironment. *Semin Immunol.* (2017) 31:30–6. doi: 10.1016/j.smim.2017.09.001

213. Barrow AD, Colonna M. Exploiting NK cell surveillance pathways for cancer therapy. *Cancers.* (2019) 11:55. doi: 10.3390/cancers11010055

214. Chang ZL, Lorenzini MH, Chen X, Tran U, Bangayan NJ, Chen YY. Rewiring T-cell responses to soluble factors with chimeric antigen receptors. *Nat Chem Biol.* (2018) 14:317–24. doi: 10.1038/nchembio.2565

215. Hou AJ, Chang ZL, Lorenzini MH, Zah E, Chen YY. TGF-β-responsive CAR-T cells promote anti-tumor immune function. *Bioeng Transl Med.* (2018) 3:75–86. doi: 10.1002/btm2.10097

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

*Copyright © 2019 Barrow, Martin and Colonna. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.*
Author/s:
Barrow, AD; Martin, CJ; Colonna, M

Title:
The Natural Cytotoxicity Receptors in Health and Disease

Date:
2019-05-07

Citation:
Barrow, A. D., Martin, C. J. & Colonna, M. (2019). The Natural Cytotoxicity Receptors in Health and Disease. FRONTIERS IN IMMUNOLOGY, 10 (MAY), https://doi.org/10.3389/fimmu.2019.00909.

Persistent Link:
http://hdl.handle.net/11343/240630

File Description:
Accepted version

License:
CC BY