Missing self-recognition of Ocil/Clr-b by inhibitory NKR-P1 natural killer cell receptors

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Communicated by Arthur Weiss, University of California, San Francisco, CA, December 13, 2003 (received for review October 16, 2003)

The NKR-P1 family of C-type lectin-like receptors are expressed on natural killer (NK) cells and NKT cells. We report the cloning and characterization of a cognate ligand for the inhibitory mouse NK receptors (NKR)-P1B and NKR-P1D (CD161b/d). The NKR-P1B/D ligand is osteoclast inhibitory lectin (Ocil), also known as Clr-b, a member of a previously cloned group of C-type lectin-related (Clr) proteins linked to the NKR-P1 receptors in the mouse NK gene complex (NKC). Expression of Ocil/Clr-b on mouse tumor cell lines inhibits NK cell-mediated killing. Inhibition is blocked with a new mAb (4A6) specific for Ocil/Clr-b. By using 4A6 mAb, we demonstrate that Ocil/Clr-b is displayed at high levels on nearly all hematopoietic cells, with the exception of erythrocytes, in a pattern that is similar to that of class I MHC molecules. Remarkably, Ocil/Clr-b is frequently down-regulated on mouse tumor cell lines, indicating a role for this receptor–ligand system in a new form of “missing self-recognition” of tumor cells.

The natural killer (NK)-1 alloantigen was identified in 1977 (1). Further definition with the PK136 mAb (2) demonstrated expression by NK cells and certain T cell subsets. cDNA sequencing (3) revealed that NK1.1 is a member of a family of the NK receptor (NKR)-P1 family of disulfide-linked homodimeric type II transmembrane C-type lectin-like receptors initially defined in rats (4–6). The Nkr-pl cluster is located in the NK gene complex (NKC) on chromosome 6 in mice, syntenic to a similar region on chromosome 4 in rats and chromosome 12 in humans (3, 7, 8). Rat NKR-P1A was shown to confer reactivity of an NK cell line to certain mouse tumor cell lines (9), and some reports suggest a role for NKR-P1 receptors in the lysis of MHC allogeneic or semiallogeneic target cells (10–12).

An initial report (5) described three distinct murine Nkr-pl genes: Nkr-p1a, Nkr-p1b, and Nkr-p1c. A fourth Nkr-pl gene, Nkr-pld, was identified subsequently (GenBank accession nos. AF338321 and AF338322) (13). More recently, genomic sequencing identified a fifth functional gene, Nkr-plf (14). Sequence data and functional studies indicate that NKR-P1A, NKR-P1C, and NKR-P1F are stimulatory receptors and that NKR-P1A, Nkr-p1d.

The full extracellular domain or only the C-type lectin-like domain segment of mouse NKR-P1B were isolated by PCR using the following primers: P1Btet, 5’-CTCGAGCTGAAACGACATCTTCCAGGCTCAAAGATGCAGTGGCAGAAAGGACGAGATGTTCA; and P1Bcrd, 5’-CTCGAGCTGAACGACATCTTCGAGGCTC AAAAGCTGAGTGGCACAGGATGTTCA; P1Btet, 5’-CTCGAGCTGAAACGACATCTTCCAGGCTCAAAGATGCAGTGGCAGAAAGGACGAGATGTTCA; P1Bcrd, 5’-CTCGAGCTGAAACGACATCTTCGAGGCTC AAAAGCTGAGTGGCACAGGATGTTCA; P1Btet, 5’-CTCGAGCTGAAACGACATCTTCCAGGCTCAAAGATGCAGTGGCAGAAAGGACGAGATGTTCA; and P1Bcrd, 5’-CTCGAGCTGAAACGACATCTTCCAGGCTCAAAGATGCAGTGGCAGAAAGGACGAGATGTTCA.

PCR products were cloned into pET15b (Novagen) and confirmed by sequencing. The expression plasmids were transformed into BL21 Star bacteria (Invitrogen) containing an isopropyl β-D-thiogalactoside-inducible BirA vector (Avidity, Denver). On induction of protein with 1 mM isopropyl β-D-thiogalactoside at OD600 = 0.500, 50 μM D-biotin was added to allow biotinylation of proteins in vivo. Inclusion-body proteins were isolated by using BugBuster reagent and the nuclease Benzonase (Novagen), and they were refolded by using condition 6 of the FoldIt Screen kit (Hampton Research, Riverside, CA).

Equal amounts of the biotinylated C-type lectin-like domain of mouse NKR-P1B were purchased from Pharmingen or eBioscience (San Diego). Streptavidin–phycoerythrin (Molecular Probes) was used as a secondary reagent. Cells were stained as described (19). Automated magnetic cell sorting was performed by using an autoMACS sorter according to the manufacturer's instructions (Miltenyi Biotec, Auburn, CA).

Methods

Mice. C57BL/6 (B6), BALB/c, and CD1 mice were maintained in our own facility. NIH Swiss and timed-pregnant B6 mice were purchased from the Animal Production Area at Charles River Breeding Laboratories.

Cell Preparations. Adult-cell suspensions were purified by using Lympholyte–Mammal (Cedarlane Laboratories). Bone marrow macrophages were prepared as described (24). B6 total fetal cell populations (days 14–17 of gestation) include erythrocytes (no Lympholyte gradient was applied). Fetal NK-enriched populations were described as prepared (19). Fetal NK lymphokine-activated killer (LAK) cells were prepared by culturing cells in 500 units/ml rhIL-2 for 5 days.

mAbs, Flow Cytometry, and Cell Sorting. All commercial mAbs were purchased from Pharmingen or eBioscience (San Diego). Streptavidin–phycoerythrin (Molecular Probes) was used as a secondary reagent. Cells were stained as described (19). Automated magnetic cell sorting was performed by using an autoMACS sorter according to the manufacturer’s instructions (Miltenyi Biotec, Auburn, CA).

Abbreviations: LAK, lymphokine-activated killers; NK, natural killer; NKR, NK receptor; NKC, NK gene complex.

Data deposition: The sequence reported in this paper has been deposited in the GenBank database (accession no. AY320031).

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PNAS | March 9, 2004 vol. 101 no. 10 3527–3532
were prepared as described (25). For staining, cells were incubated with $4 \mu g$ of tetramer in staining buffer for 1 h on ice or at room temperature.

**Generation of CD3ζ/NKR-P1 Fusion Receptors and BWZ Reporter Cells.**

The transmembrane and extracellular domains of mouse NKR-P1B and the cytoplasmic domain of CD3ζ were isolated by PCR using the following primers: P1BF, ACCCTGGCCCGCTGCTGTGGCTGCCCTCCTCGTGCATC; P1BR, GCGGCCGCTGCTCAGGATCTTTACTGCGAGGTG; and CD3ZF, GCAAGCTATTTTTCGACGATGG. The fusion receptor was generated by gene splicing by overlap extension by using CD3ZF and P1BR, cloned into pcDNA3.1-

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Generation of the 4A6 Anti-OcI/Clr-b mAb by Immune Focusing. We injected 1-year-old Lewis rats i.p. with MNK-1, NIH 3T3, and cultured B6 bone marrow macrophages at 10-day intervals, respectively. Hybridomas were generated as described (24). For screening, hybridoma supernatants were added to MNK-1 or NIH 3T3 cells for 30 min, followed by BWZ.CD3ζ/P1B indicator cells. A rat Ig isotyping kit (Pharminen) revealed that 4A6 is a rat IgM. Produced in CELLine CL1000 vessels (Integra Biosciences, Chur, Switzerland), 4A6 mAb was purified over recombinant protein L-agarose columns (Pierce) by using EZ-link Sulfo-

Retroviral supernatants were generated by triple transfection of proviral vector with VSV-G supernatants were generated by triple transfection of proviral vector with VSV-G constructs (gift from W. Sha) into 293T cells, and 48-h supernatants were used to “spin-feet” BWZ cells for 90 min at 1,000 × g. NKR-P1A/C/D C-type lectin-like domain segments were digested and ligated into the CD3ζ/NKR-P1B fusion receptor cassette.

For plate-bound assays, mAbs were incubated on high-binding chemistry plates (Costar/Corning). For cell-based assays, $5 \times 10^4$ to $10^5$ BWZ indicator cells were cocultured overnight with $5 \times 10^4$ to $5 \times 10^5$ stimulator cells. β-galactosidase activities were determined as described (26).

**Retroviral Library Screening.** Retroviral supernatants of the pM5S-NK library (27) were generated as described above. Stable transductants ($10^7$ 293T, COS, or BWZ cells) were sorted after 3 days by using biotin-conjugated 4A6 plus phycoerythrin-conjugated streptavidin (Molecular Probes) versus FITC-conjugated 10D7 at 10 positive cells per well. Positive pools were identified by flow cytometry, and PCR of genomic DNA was performed (27). The following retroviral PCR primers were used: MSCVF2, CCCACCGCCCTCAAAGTAGACG; MSCVR, GCTTGCCAAACCTACAGGTGGGG; PMXF, GGTTGGACCATCCTCTAGACT; PMXR, CCCTTTTTCCACGACTAATAA. Transfections were performed by using Lipofectamine Plus (Invitrogen) or FuGENE 6 (Roche) reagent.

RT-PCR. RNA was prepared by using TRizol reagent (Invitrogen) and reverse-transcribed by using the Delta differential-display kit (Becton Dickinson). The following primers were used: CLRBF, ATGTGTCATCATCAGGGAATT; CLRBR, CTAGGAAGGGAAAAAAAGGAGTTT; CLRFF, ATGAATGGTCAAGGCGAAAAT; CLRFR, TRACAGATCGCTTGGTACACTT; CLRGF, ATGGCAATGTGTTGAGGAGAG; CLRGR, CTGACAGAAGGAGGAGTTTGCGAATG; G3PDHF, ACCACACTGCCATGCTAC; and G3PDRH, TCACACACCTTGTGGTGA. PCR was performed by using Expand Hi-Fidelity enzyme (Roche).

![Fig. 1. Detection of NKR-P1B ligands.](image-url)
The transiently transfected cells used as stimulator cells in expressing chimeric NKR-P1A, NKR-P1B, NKR-P1C, or NKR-P1D receptors. (Clr-b, Clr-f, or Clr-g to stimulate an equal number of BWZ indicator cells had been transiently transfected with expression plasmids encoding Ocil not Clr-f or Clr-g, is a ligand of NKR-P1B and NKR-P1D. We used 293T cells that had been transiently transfected with expression plasmids encoding Ocil/Clr-b, Clr-f, or Clr-g to stimulate an equal number of BWZ indicator cells expressing chimeric NKR-P1A, NKR-P1B, NKR-P1C, or NKR-P1D receptors. (C) The transiently transfected cells used as stimulator cells in B were stained 18 h after transfection with 4A6 mAb (1007 mAb served as a control). Markers were placed according to staining of untransfected 293T cells.

**Results**

To identify NKR-P1 ligands, we developed a rapid functional assay as well as a ligand-binding staining reagent. The functional assay was developed by transfecting the BWZ reporter cell line, which produces β-galactosidase in response to stimulation (26), with a chimeric cDNA encoding the extracellular domain of NKR-P1B fused to a modified inverted cytoplasmic domain of the stimulatory CD3ζ subunit (28). The staining reagent consisted of a tetrameric form of the soluble extracellular domain of NKR-P1B, which had been expressed in bacteria and refolded in vitro.

The BWZ-transfectants (BWZ.CD3ζ/P1B cells) responded well when stimulated on plates coated with anti-NKR-P1 mAb PKI36, but they did not respond to other Abs (Fig. 1A). The indicator cells responded well also when stimulated with the MNK-1 immature NK cell line or with NIH 3T3 fibroblasts, gave intermediate responses with the C1498 leukemic cell line or bone marrow macrophages, and gave minimal responses with several other cell lines (Fig. 1B). A separate analysis revealed strong responses with bone marrow cells and spleen cells and lower but consistent responses with thymocytes (see Fig. 6F, which is published as supporting information on the PNAS web site). Comparisons of BWZ cell lines expressing fusions of all four NKR-P1 proteins with CD3ζ showed that MNK-1 cells expressed ligands for NKR-P1B and NKR-P1D, the two inhibitory isoforms, but not NKR-P1A and NKR-P1C, two stimulatory isoforms (see Fig. 7A, which is published as supporting information on the PNAS web site). All four reporter cell lines responded when the fusion protein was crosslinked with the appropriate mAbs.

**51Cr-Release Assays.** LAK cells were generated, and lysis assays were performed as described (24).

NKR-P1B tetramers stained NIH 3T3 cells and, to a lower extent, MNK-1 cells, but they did not stain BWZ cells (Fig. 1C Upper). Notably, MNK-1 cells also express NKR-P1B receptor (13), which could interfere with tetramer binding.

To prepare an mAb specific for NKR-P1B ligands, Lewis rats were sequentially immunized with MNK-1, NIH 3T3, and B6 bone marrow macrophages to focus the Ab response on their shared antigens. Hybridoma supernatants were screened for inhibition of the response of BWZ.CD3ζ/P1B cells to ligand-expressing cells. Hybridoma 4A6 (rat IgMκ isotype), blocked stimulation of BWZ.CD3ζ/P1B cells by both MNK-1 and NIH 3T3 cells (Figs. 1D and 7B), as did PK136 (Fig. 1D). Biotinylated 4A6 stained NIH 3T3 cells brightly, MNK-1 cells less intensely, and BWZ cells only weakly, and it did not stain 293T cells (Fig. 7C), consistent with the NKR-P1B tetramer staining. Importantly, 4A6 mAb blocked staining of MNK-1 and NIH 3T3 cells by NKR-P1B tetramers (Fig. 1C Lower), providing direct evidence that 4A6 recognizes the NKR-P1B ligand.

Because we observed that NK cells stained with 4A6 mAb (see below), we transduced the pMXS-NK cell cDNA retroviral expression library into the BWZ cell line and selected 4A6+ BWZ clones. PCR of retroviral integrants from genomic DNA revealed that most of the 4A6+ clones lacked retroviral sequences and presumably expressed endogenous 4A6 antigen. However, one transductant (IB8) yielded a PCR product (see Fig. 8A, which is published as supporting information on the PNAS web site) that conferred 4A6 reactivity when transsected into 293T cells in the form of a new proviral construct (pMXS-IB8; Fig. 2A). Control DX5 mAb did not stain the transfectants. Furthermore, when virus obtained from these transsected cells was transduced into BWZ cells, we observed strong staining of
a large subset of the cells with 4A6 mAb compared with control untransfected BWZ cells (Fig. 8B). Thus, 1B8 cDNA encodes the 4A6 antigen. The 1B8 cDNA coding sequence was 100% identical to that of Oc'il isolated from B6 mice (29), and it contained only a few BALB/c-specific substitutions in the untranslated regions. Oc'il is also called Clr-b, a member of the Clr gene family, which includes Clr-f and Clr-g in B6 mice (30). Oc'il/Clr-b transcripts were up-regulated in BWZ clones with endogenously encoded (nonretroviral) 4A6 antigen expression (Fig. 8C), supporting the conclusion that Oc'il/Clr-b is the 4A6 antigen. Interestingly, the Clr genes are located in the NKC, closely linked to the Nkr-p1 genes (30).

We tested 293T cells transiently transfected with Oc'il/Clr-b, Clr-f, and Clr-g/OcilrP2 (29–31) in the BWZ assay. Oc'il/Clr-b transfectants stimulated BWZ.CD3/CD28.P1B and BWZ.CD3/CD28.P1D indicator cells but not the other indicator cells, providing direct functional evidence that Oc'il/Clr-b is a ligand for NKR-P1B and NKR-P1D but not for NKR-P1A or NKR-P1C (Fig. 2B). All of the indicator cells responded to stimulation with antireceptor Abs (Fig. 7A). Furthermore, Clr-f and Clr-g/OcilrP2 transfectants failed to stimulate any of the indicator cells (Fig. 2B). Parallel analysis showed that a fraction of the Clr-b transfectants, but not Clr-f or Clr-g/OcilrP2 transfectants, were stained with 4A6 mAb (Fig. 2C). Thus, Oc'il/Clr-b is a functional ligand for NKR-P1B and NKR-P1D but not for NKR-P1A or NKR-P1C.

Expression of Oc'il/Clr-b Resembles That of Class I MHC but Is β2m-Independent. Oc'il/Clr-b is broadly expressed by nearly all adult splenocytes, thymocytes, and lymph-node cells, as shown by 4A6 staining (Figs. 3A and B and 6). The positive cells included nearly all lymphocytes, granulocytes, and myeloid cells (Figs. 3 and 6). All four thymocyte subsets defined with CD4 and CD8 were mostly positive, although CD4−/CD8− thymocytes stained only ~1/10th as well as the other subsets (Fig. 3B). Nearly all CD4+ cells from fetal liver, fetal spleen, and fetal thymus expressed Oc'il/Clr-b, including all subsets of CD3+ CD4+ CD8− triple-negative thymocytes (Figs. 3C and 6). Notably, Oc'il/Clr-b expression diminishes during erythropoiesis, and terminally differentiated erythrocytes (CD45−/TER-119+) in the fetal liver or fetal spleen did not stain detectably with 4A6 (Figs. 3C and 6A and data not shown). In contrast, various nonhematopoietic CD45+ cells expressed high levels of Oc'il/Clr-b, as shown by 4A6 staining of NIH 3T3 and other fibroblast cell lines (Fig. 7 and see below) and mRNA analysis (30). The broad pattern of expression with lower levels on CD4−CD8− thymocytes and erythrocyte lineage cells is remarkably similar to that of classical MHC class I molecules. Furthermore, costaining with 4A6 and anti-class I (Kb) mAbs revealed a correlation between the relative levels of Oc'il/Clr-b and Kb on various ex vivo BALB/c cell populations prepared without enrichment of lymphocytes (Fig. 6G). Staining and functional analyses showed that Oc'il/Clr-b expression was normal in MHC class I-deficient (β2m−/−) mice, showing that it is not class I-dependent (Figs. 3D and 6F).

Ocil/Clr-b Is Frequently Down-Regulated in Mouse Tumor Cell Lines. Analysis of cell lines with 4A6 mAb demonstrated that numerous lymphoid tumor lines expressed dramatically lower cell-surface expression

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**Fig. 4.** Oc'il/Clr-b is frequently down-regulated on mouse tumor cells. The cell lines or cell populations indicated were stained with biotin-conjugated 4A6 and phycoerythrin-conjugated streptavidin secondary (shaded region) or phycoerythrin-conjugated streptavidin alone (region outlined in black) and analyzed by flow cytometry.
levels of Ocil/Clr-b (Fig. 4), compared with the uniformly high levels detected on their ex vivo lymphocyte counterparts (Fig. 3). Several carcinoma cell lines also expressed low levels of Ocil/Clr-b (Fig. 4). Loss of ligand for the inhibitory NKR-P1 receptors would be expected to make the tumor lines more sensitive to NK cell-mediated killing.

Expression of Ocil/Clr-b Protects Target Cells from NK Cell-Mediated Lysis. To test whether Ocil/Clr-b expression in target cells inhibits NK cell function, we compared the sorted 4A6+ variants of BWZ (BWZ.OCil+), which had up-regulated endogenous Ocil/Clr-b, with parental BWZ.36 cells, which expressed only low levels of Ocil/Clr-b (Figs. 7 and 8C). Unexpectedly, BWZ.OCil+ variants stimulated BWZ.CD3ζ/P1D indicator cells much less well than they stimulated BWZ.CD3ζ/P1B cells (Fig. 5A), possibly because of a modification of the endogenous Ocil/Clr-b (BWZ.36 cells were derived from AKR strain T lymphoma, BW5147αβ−). Presumably because B6 NK cells lack NKR-P1B and NKR-P1D fail to functionally engage Ocil/Clr-b on these target cells, B6 NK cells were not inhibited by the BWZ.OCil+ cells and lysed them equally to control BWZ cells (Fig. 5B). However, NK cells from NIH Swiss or CD1 mice, which express NKR-P1B, were inhibited by BWZ.OCil+ cells (Fig. 5B). The inhibitory effect was reversed by preincubating the effector cells with the 4A6 anti-Ocil mAb (Fig. 5C). Together, these data indicate that Ocil/Clr-b is a ligand for the inhibitory NKR-P1 receptors and that Ocil/Clr-b expression protects cells from NK cell-mediated killing.

Discussion
This article identifies Ocil/Clr-b as the cognate ligand for the inhibitory NKR-P1B and NKR-P1D receptors. As this article was in preparation, the identification of Clr family members as ligands for NKR-P1 receptors was reported by Yokoyama and colleagues (32). The NKR-P1 receptors are NKC-encoded lectin-like NKRs that are shown to recognize other lectin-like molecules. All of the other lectin-like NKRs studied have been shown to recognize class I MHC molecules or relatives of class I MHC molecules. Moreover, the fact that the Nkr-p1 genes are intermingled within the NKC with the genes encoding their ligands is remarkable. It is interesting to speculate that the NKR-P1 proteins and their Clr ligands may be descended from a common C-type lectin-like ancestral protein that engaged in homophilic interactions.

Close genetic linkage of the genes encoding receptors and ligands in this system is unprecedented for NKRs. In contrast, self-recognition systems in plants such as crucifers consist of closely linked genes for a receptor and cognate ligand, which interact to prevent pollen tube formation (33). Because both molecules are polymorphic, close genetic linkage of receptor and ligand genes enhances the likelihood that the self receptor and self ligand will be co-inherited. By analogy, close linkage of NKR-P1 and its Ocil/Clr ligands may also ensure coinheritance of cognate receptor and ligand genes.

These results demonstrate that the NKR-P1 receptors, like other lectin-like proteins encoded within the NKC, recognize proteins, in contrast to the earlier reports of oligosaccharide recognition by these receptors (20–23). It remains possible that glycans linked to Ocil/Clr-b play a role in binding by NK-R-P1. A significant finding with the 4A6 mAb was that Ocil/Clr-b is broadly expressed by hematopoietic cells, consistent with the finding that the transcripts are broadly expressed (29–31). In general, 4A6+ cells were stimulatory in our functional assay (Figs. 1, 6F, and 7). The few discrepancies may be accounted for by variable expression of accessory molecules by different cells, although it remains possible that modifications of Ocil/Clr-b occur in some cells. In contrast to our results, it was reported recently based on functional studies that Ocil/Clr-b is expressed predominantly by dendritic cells and macrophages (32). Furthermore, although it was suggested that the expression pattern of Ocil/Clr-b is dissimilar to that of MHC class I proteins (32), we found instead a remarkable similarity in their expression patterns, including the low expression on CD4+ CD8+ thymocytes and erythrocyte lineage cells and a correlation in the relative levels of expression in various ex vivo populations. The similarity in the expression patterns of ligands for NKR-P1B/D receptors and the MHC class I ligands for Ly49 inhibitory receptors suggests that low expression of ligands by these cell lineages may have significance in missing self-recognition systems. Notably, other Clr proteins, such as Clr-I and Clr-g, exhibit a much more restricted expression pattern (30), suggesting a different function for these proteins.

One of our most interesting findings was that many of the tumor cell lines we examined expressed notably low levels of Ocil/Clr-b on the cell surface. Comparisons of Ocil/Clr-b levels on lymphomas and other hematopoietic tumors with those on corresponding nontransformed lymphohematopoietic cell types...
suggested that Ocil/Clr-b down-regulation on tumor cell lines is often dramatic. These findings are consistent with the notion that down-regulation of Ocil/Clr-b on tumor cells contributes to their sensitivity to lysis by NK cells. Indeed, we observed reduced lysis of cells expressing Ocil/Clr-b that could be reversed with the 4A6 mAb. Hence, the NKR-P1B/D-Ocil/Clr-b receptor-ligand system appears to represent an independent mode of missing self-recognition that functions in parallel with missing self-recognition of class I MHC molecules.

The adaptive value to the animal of missing self-recognition of the Ocil/Clr-b molecules remains unclear, but the presence of the consensus casein kinase 2 (CK2) phosphorylation site in the cytoplasmic domain of Ocil/Clr-b (31) raises an interesting possibility. CK2 overexpression is common in tumor cell lines (34, 35), and active CK2 is known to regulate the NF-κB pathway, caspase activation, and Bel-2(X,L) levels (36), leading to lymphoproliferation and accelerated lymphomagenesis (34, 37). Furthermore, endocytosis of certain other receptors that contain CK2 phosphorylation sites, such as the transferrin receptor and possibly the polymeric Ig receptor, depends on CK2-mediated receptor phosphorylation (38, 39). Hence, it is possible that up-regulation of CK2 in tumor cells leads to Ocil/Clr-b endocytosis from the cell surface and, consequently, enhances the sensitivity of the tumor cells to NK-mediated lysis.

It is possible also that, under certain circumstances, expression of Ocil/Clr-b favors apoptosis over uncontrolled growth. For example, although we were successful in expression cloning of the Ocil/Clr-b cDNA by using a single cycle of cell sorting after transfection, these cells rapidly lost retroviral integrations when they were resorted (data not shown), possibly because the transduced cells died. Moreover, on transduction of the cloned IBS cDNA into BWZ cells, 4A6 staining waned over time, and a dramatic loss in viability of the 4A6+ (but not 4A6−) cells was observed during the 4A6 sorting procedure, as monitored by propidium iodide uptake (data not shown). Also, transient expression of Ocil/Clr-b in certain cell lines, such as COS cells, was inefficient, possibly because of induced cell death (data not shown). If Ocil/Clr-b expression normally favors the death of certain tumor cells, it is conceivable that loss of these molecules would be selected for during the evolution of a tumor. This mechanism could provide a rationale for missing self-recognition of Ocil/Clr-b molecules by NK cells. It remains a mystery why some cells, such as normal cells or BWZ cells expressing endogenous Ocil/Clr-b, display no ill effects.

In conclusion, we have demonstrated that the inhibitory NKR-P1B/D orphan NK cell receptors functionally interact with Ocil/Clr-b ligand and can mediate missing self-recognition of tumor cells by mouse NK cells. A recent study (40) identified a gene in the tunicate Botryllus schlosseri that encodes a lectin-like protein that is more similar in its extracellular domain to NKR-P1 than to other lectin-like genes. The possibility that the tunicate gene exhibits a binding specificity or function similar to that of the mouse NKR-P1 proteins remains to be explored.

We thank Jay Ryan for the 10A7 mAb, advice, and sharing unpublished work; J. C. Züniga-Pflücker for helpful suggestions and support; Eric Jensen for help with rat i.p. injections; and Hector Nolla for assistance with fluorescence-activated cell sorting. This research was supported by grants from the National Institutes of Health (to D.H.R.) and a Long-Term Fellowship from the Human Frontier Science Program (to J.R.C.).