Tumor cells release NKG2D ligands to evade NKG2D-mediated immune surveillance. The purpose of our investigation was to explore the cellular mechanisms of release used by various members of the ULBP family. Using biochemical and cellular approaches in both transfectant systems and tumor cell lines, this paper shows that ULBP1, ULBP2, and ULBP3 are released from cells with different kinetics and by distinct mechanisms. Whereas ULBP2 is mainly shed by metalloproteases, ULBP3 is abundantly released as part of membrane vesicles known as exosomes. Interestingly, exosomal ULBP3 protein is much more potent for down-modulation of the NKG2D receptor than soluble ULBP2 protein. This is the first report showing functionally relevant differences in the biochemistry of the three members of the ULBP family and confirms that in depth study of the biochemical features of individual NKG2D ligands will be necessary to understand and manipulate the biology of these proteins for therapy.

NKG2D is an activating immune receptor that can be expressed by most cytotoxic lymphocytes, including NK and CD8+ T cells (1). Engagement of NKG2D by its ligands leads to the activation or co-stimulation of lysis and cytokine secretion (for review, see Ref. 2). In humans, NKG2D ligands (NKG2D-L)5 occur in two families of proteins: the polymorphic family of MHC-I-related chain A/B (MICA/B) and the multigene family of UL16-binding proteins (ULBPs, also known as RAET1A–E). In total, 10 members of this gene family have been described, of which six can be expressed as functional proteins (3). Two members of the ULBP family have a transmembrane region (ULBP4 and -5), like MICA/B, whereas the other ULBP molecules are linked to the cell membrane via glycosylphosphatidylinositol (GPI) anchors. The existence of such a large number of ligands for a single receptor is not fully understood but may reflect a differential role for different ligands in immune surveillance or an evolutionary response to selective pressures exerted by pathogens or cancer.

In general, NKG2D-L are not expressed ubiquitously; instead, they are expressed in response to several types of cellular stress, such as pathogen infection (4), DNA damage (5), proteasome inhibition (6), and tumor transformation (7). For example, MICA/B are expressed in epithelial tumors, melanoma, neuroblastoma, various hematopoietic malignancies, and carcinomas; ULBPs are found in leukemia, gliomas and melanomas. An additional complication is that mRNA can be found in many cells that do not express protein suggesting post-transcriptional regulation of NKG2D-L expression (8–10).

Mice deficient in NKG2D expression show an enhanced susceptibility to the development of tumors (11). However, shedding NKG2D-L as soluble molecules allows tumor cells to evade NKG2D surveillance. Apart from reducing NKG2D-L expression on the tumor cell surface, the release of soluble molecules may also impair immune surveillance by promoting down-regulation of NKG2D (12, 13). In fact, the sustained presence in vivo of NKG2D-L down-modulates the receptor (14, 15), and at least in mouse models, blocking shedding of NKG2D-L can prevent tumor formation (16). In patients with colorectal or prostate cancer, hepatocellular carcinoma, and neuroblastoma, decreased NKG2D expression and impaired activation of NK cells was associated with high levels of soluble MICA in serum (17–20). Importantly, therapy of patients with chronic myeloid leukemia led to a substantial decrease of soluble MICA levels, accompanied by restored NKG2D expression on CD8+ T cells and NK cells (8). Overall, the release of soluble NKG2D-L by tumor cells has a negative impact on NKG2D-dependent immune surveillance of cancer and suggest that a better understanding of this process may lead to the identification of useful targets for therapy.

Recently, members of the ADAM (a disintegrin and metalloprotease) family have been identified as key proteases involved in the shedding of some alleles of MICA and MICB (21, 22), and a member of the disulfide isomerase family, ERp5, has also been proposed to play a role in the shedding of MICA (23). ADAMs 10 and 17 have been shown to be involved in proteolytic cleavage of ULBP2 (22), but nothing else is known about the shedding of other ULBPs. Indeed, in general, little is known about the biochemistry and cell biology of the ULBPs other than that they have signals for a GPI anchor (24, 25) and
that ULBP3 can associate with microdomains of the membrane rich in sphingolipids and cholesterol (detergent-resistant membranes) (26).

We have analyzed the biochemical features of the ULBPs released to the extracellular media of transfected and tumor cell lines and here report interesting differences in the kinetics and mechanisms of shedding of the ULBPs. Although ULBP1 was shed at low levels, and ULBP2 was abundantly shed as a soluble protein, ULBP3 was shed with slower kinetics, and, surprisingly, much of this released ULBP3 was found in the membrane of small vesicles known as exosomes. Interestingly, the ULBP released in exosomes potently down-modulated the NKG2D receptor. Overall, these data provide the first evidence of functionally relevant biochemical differences between the three members of the ULBP family linked through a GPI anchor and suggest new approaches to understand the diversity of NKG2D-L.

**EXPERIMENTAL PROCEDURES**

**Cells and Reagents**—ULBP1, -2, and -3 constructs were obtained from Dr. Richard Apps (27). The kidney monkey cell line CV1 and 293T cells (ATCC) were maintained in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal calf serum, 1-glutamine, and antibiotics. Chinese hamster ovary (CHO) cells were maintained in Hams F12 medium with the same supplements. CV1 cells were transfected as described (28). CHO cells were transfected using Lipofectamine 2000 and ULBP expression plasmid mixed (9:1 ratio) with a vector containing the same supplements. CV1 cells were transfected with puromycin (Calbiochem). Antibodies directed against ULBPs were purchased from R&D Systems (Abingdon, UK). BB94 (Batimastat) was a kind gift from Dr. M. Marsh (clone eh1C9b, generated by S. Schmid). Anti-hamster CD63 hybridoma was provided by Dr. G. Fishwild (gift of British Biotech). Leupeptin, pepstatin A, 1,10-phenanthroline, and 1H-[1,2,3]triazolo[4,5-b]pyridine (Triton X-100) were obtained from Calbiochem. Anti-hamster CD63 hybridoma was provided by Dr. G. Fishwild (gift of British Biotech). Leupeptin, pepstatin A, 1,10-phenanthroline, and 1H-[1,2,3]triazolo[4,5-b]pyridine (Triton X-100) were obtained from Calbiochem.

**Flow Cytometry**—10^5 cells were preincubated in PBS containing 1% bovine serum albumin, 0.1% sodium azide (PBA). Cells were then incubated with mouse monoclonal antibodies and bound antibody was visualized using either phycoerythrin- or fluorescein isothiocyanate-labeled F(ab’)_2 fragments of goat anti-mouse Ig (Dako). Samples were analyzed using a FACScan II flow cytometer (Becton Dickinson). Dead cells were excluded from all analyses by staining with propidium iodide.

**ELISA**—For detection of soluble proteins, cells were incubated for 16–24 h in medium in the absence of serum. Detection of sULBP was performed using a sandwich ELISA procedure. Plates were coated for 16 h at 4 °C with the appropriate polyclonal anti-ULBP1, -2, and -3 antibody (R&D Systems) (5 μg/ml) and blocked with 2% bovine serum albumin-PBS for 2 h at 37 °C. Tissue culture supernatant was added for 2 h at 37 °C. Bound ULBP protein was detected using the appropriate biotinylated goat anti-ULBP (R&D Systems) followed by streptavidin-horseradish peroxidase (Amersham Biosciences), and the assay was developed using the peroxidase substrate system (ABTS, Roche Applied Science). Absorbance was measured at 410 nm with a reference wavelength of 490. Samples were analyzed in duplicates. Under these conditions, the cut-off for detection of recombinant soluble ULBP-Fc constructs (R&D Systems) was ~1 ng/ml, and the ELISA absorbance values were directly proportional to the concentration of soluble ULBP protein over the range 1 to 100 ng/ml.

**Exosome Purification**—Exosomes were prepared by sequential centrifugation as in Raposo et al. (31). Exosomes pellet after centrifugation at 100,000 × g for 2 h were resuspended either in 100 μl of PBS for electron microscopy or directly in sample buffer for Western blot. The amount of protein in exosomes was estimated using Bio-Rad protein assay. The resulting supernatant (soluble fraction) was recovered by precipitation with trichloroacetic acid. Where indicated, further purification of exosomes by flotation on a discontinuous sucrose gradient was performed. A step sucrose gradient (4 ml of 60% sucrose, 3 ml of 30%, and 1 ml of 5%) was layered on top of the exosome suspension, mixed with 2.5 volumes of 85% sucrose in TNE buffer (20 mM Tris, pH 7.5, 150 mM NaCl, 5 mM EDTA). Gradients were centrifuged for 20 h at 100,000 × g. 1-ml fractions were collected, and proteins were further solubilized by adding deoxycholate (final concentration 0.2%).

**Western Blot**—Cell lysates were prepared by incubation in TNE buffer containing 1% Nonidet P-40 and the protease inhibitors leupeptin, pepstatin, and 1,10-phenanthroline for 30 min at 4 °C. Nuclei were eliminated by centrifugation at 13,000 × g. Lysates, supernatants, and exosome preparations were run on 12% SDS-PAGE gels and transferred to Immobilon-P (Millipore) membrane. The membrane was blocked using PBS containing 0.1% Tween 20 (PBS-T) and 5% nonfat dry milk. Detection of ULBP was performed by incubation with biotinylated goat polyclonal anti-ULBP antibody, followed by horseradish peroxidase-conjugated streptavidin. Proteins were visualized using the ECL system (Amersham Biosciences). Quantitative analysis of the Western blot data was done using NIH ImageJ software.

**Electron Microscopy**—Samples of exosomes for examination by electron microscopy were prepared by floating a carbon-coated 400 mesh Formvar electron microscopy grid on top of one drop of freshly prepared exosomes for 15 s. The grid was then briefly washed with deionized water and floated on a drop of 2% phosphotungstic acid, pH 7.0. Samples were examined using a Philips CM100 operating at 60 or 80 keV.

**Confocal Microscopy**—Cells were fixed with 4% paraformaldehyde at 4 °C for 15 min, permeabilized by incubation with 0.1% saponin at room temperature for 10 min, stained with polyclonal anti-ULBP antibodies (R&D Systems), and analyzed by confocal microscopy as described previously (32). Fluorescence images were obtained using a confocal microscope (Leica TCS-NT-UV confocal laser scanning microscope). Images of fixed cells were taken using a 63 × 1.32 objective with the confocal pinhole set to one airy unit. Images were obtained by scanning series of single focal planes across the cell using Leica TCS software. To explore the whole intracellular area, series of sections (total interval z = 2–4 μm) were acquired.

**NKG2D Down-modulation**—Primary human NK cells, 3–5 days after stimulation with feeders and interleukin-2, were cul-
Although ULBP2 and -3 were detected in large amounts, and -3 expressed in the same cellular background was observed. transfection. Interestingly, differential shedding of ULBP1, -2, and -3 in stably transfected CHO cells revealed that the kinetics of accumulation of ULBP2 and ULBP3 in the supernatant were different; large amounts of ULBP2 were detected after only brief incubations, whereas the accumulation of ULBP3 occurred at a slower rate (Fig. 1, C and D).

**Metalloproteases and the Cleavage of ULBP2 and ULBP3**—It has been reported that the release of ULBP2 is inhibited by metalloprotease inhibitors (34) but that metalloprotease inhibition had no effect on surface expression of ULBP1 or ULBP3. We extended these studies using a panel of protease inhibitors to explore the ability of inhibitors of different classes of proteases to affect ULBP2 and ULBP3 shedding in time course experiments (Fig. 2). Leupeptin is an inhibitor of serine and cysteine proteinases; pepstatin is a potent inhibitor of various aspartic proteinases; BB94 inhibits both matrix metalloproteases and some members of the ADAM family of proteases, which have been shown to affect ULBP2 release. Although leupeptin and pepstatin did not affect the shedding of ULBP2 or -3, BB94 dramatically decreased the shedding of ULBP2 from both CHO cells (Fig. 2A) and CV1 cells (data not shown). At short times of incubation, BB94 weakly inhibited ULBP3 release, but no clear effect was apparent after 16 h of incubation. These results confirmed that metalloproteases mediate shedding of ULBP2 and suggest that the release of ULBP3 depends only partially on metalloproteases.

Analysis of the effect of pharmacological stimulation on the release of ULBP2 and -3 revealed more differences between the release of these molecules. Phorbol esters (PMA) activate ADAM17-mediated shedding of proteins (known to be involved in ULBP2 shedding) (22, 35). Ca\(^{2+}\)-ionophores (ionomycin) activate ADAM10 (36) and, at long incubation times, exosome release (37). PMA treatment increased ULBP2 shedding, although the effect disappeared at longer time points but had little effect on ULBP3 release. Ionomycin weakly stimulated the release of the ULBPs only at longer times (Fig. 2B).

These results are consistent with published data, suggesting the involvement of ADAM proteases in release of ULBP2 (22). Overall, these data also imply that the release of ULBP2 and -3 depend on distinct cellular pathways.

**ULBP3 Is Released in Exosomes**—To understand the differences in the shedding of the various ULBPs, a biochemical characterization of the molecules released to the supernatant was performed using both transfected cells and tumor cell lines. Supernatants were fractionated to separate proteins present as soluble monomeric forms from those present in microparticles called exosomes. Exosomes are small vesicles (30–100 nm) secreted to the extracellular medium after...
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Both ULBP2 and ULBP3 are GPI-anchored proteins and reside in regions of membranes rich in sphingolipids and cholesterol, known as detergent-resistant membranes. Recruitment to detergent-resistant membranes is a mechanism described for incorporation of proteins into exosomes (40); thus, why would ULBP3 but not ULBP2 be preferentially included into exosomes? We hypothesized that ULBP2 is not excluded from exosomes but rather is more susceptible to metalloprotease attack and is therefore lost by proteolysis before inclusion into exosomes. To test this idea, the soluble and exosome fractions of supernatants of ULBP2 transfectants, untreated or treated with metalloprotease inhibitors, were analyzed for the presence of ULBP2 by Western blot. Fig. 4A shows that treatment with BB94 augments the amount of ULBP2 present in exosomes, confirming this hypothesis: whereas in untreated cells, only 2.77% of the released ULBP2 protein was present in exosomes, in BB94-treated cells, this percentage augmented to 31.89%, and a 30% reduction of soluble protein could be observed (from 97.23% in untreated to 68.11% in BB94). Treatment of ULBP3 transfectants with BB94 produced a decrease in the amount of shed soluble ULBP3 but not of the exosomal protein (data not shown). In confocal microscopy, the pattern of distribution within the cell was different for ULBP2 and ULBP3, suggesting that these molecules may also differ in intracellular trafficking (Fig. 4B and supplemental Fig. 2).

Both Soluble ULBP2 and Exosomal ULBP3 Can Induce Down-regulation of NKG2D—The release of soluble MICA, as well as promoting immune evasion by reducing the cell surface expression of NKG2D-L on tumors, has also been reported to trigger a systemic down-regulation of the NKG2D receptor on NK cells and CD8+ T cells (12, 17, 41). Thus, it was of interest to compare the effect of the addition of supernatants containing ULBP2 (mainly soluble) and ULBP3 (released in exosomes) on cell surface expression of NKG2D on primary human NK cells. To avoid the confounding effects of factors such as transforming growth factor-β on lymphocyte proliferation and NKG2D expression (42–44), supernatants from untransfected CHO cells and CHO transfectants expressing either ULBP2 or ULBP3 were used for these experiments. A representative example of this type of experiment is shown in

6 L. Fernandez-Messina, unpublished data.
Fig. 5. Both ULBP2- and ULBP3-containing supernatants could induce decreased cell surface NKG2D expression. A t test of the data from multiple experiments shows that these effects are statistically significant. It is interesting to note that the effect of ULBP3 is consistently stronger than that of ULBP2, especially because the amount of ULBP2 in the supernatants was always higher than that of ULBP3 (Fig. 5C).

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A. Exosome purification after treatment with metalloprotease inhibitor

| Untreated | +BB94 |
|-----------|-------|
| SOLUBLE   | EXOSOMES | LYSATE | SOLUBLE | EXOSOMES | LYSATE |
| 1 2 3 4 5 6 7 8 9 | 1 2 3 4 5 6 7 8 9 |
| ULBP2    | CD63   | ULBP2    | CD63   |
| 50       | 75     | 50       | 75     |
| 37       | 50     | 37       | 50     |

B. Confocal microscopy

ULBP2

ULBP3

D. Western blot in 293T cells (endogenous expression)

ULBP2

ULBP3

FIGURE 3. ULBP3 is released in exosomes both in transfectants and the tumor cell line 293T. A, the exosome fraction and soluble proteins from CHO-ULBP2 and -3 transfectants were purified after 24 h in culture as described under “Experimental Procedures” and compared with total lysate from the same cells in Western blot analysis. Similar results were obtained on analysis of CV1-ULBP3 cells (supplemental Fig. 1). ULBP2 was shed mainly as a soluble protein, whereas ULBP3 was released both as a soluble protein and in exosomes. B, fractionation of exosomes in a sucrose gradient shows co-migration with CD63. C, exosomes from CHO-ULBP3 cells were negatively stained with 2% phosphotungstic acid and analyzed by electron microscopy. As expected, nano-sized vesicles from 30 to 120 nm were observed. Bar, 100 nm. D, analysis of tissue culture supernatant from the 293T cell line, which expresses ULBP2 and ULBP3 endogenously, confirmed the results obtained in the transfectant system. Soluble and exosome fractions were analyzed by Western blot. ULBP2 is mainly shed as a soluble protein, whereas ULBP3 is mainly released in exosomes.

Fig. 5. Both ULBP2- and ULBP3-containing supernatants could induce decreased cell surface NKG2D expression. A t test of the data from multiple experiments shows that these effects are statistically significant. It is interesting to note that the effect of ULBP3 is consistently stronger than that of ULBP2, especially because the amount of ULBP2 in the supernatants was always higher than that of ULBP3 (Fig. 5C). The simplest interpreta-
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A. NKG2D expression on NK cells preincubated with supernatant from CHO or CHO/ULBP cells

B. NKG2D downregulation

C. Soluble ULBP in supernatant

To confirm that the down-modulation of NKG2D receptor, observed in the experiments above, was due to the effect of exosomal protein and not to any soluble protein present in the supernatant, it was important to test whether purified ULBP3-containing exosomes could affect cell surface expression of NKG2D on NK cells. Fig. 6 shows that exposure to exosomes purified from ULBP3 transfectants triggered a marked reduction in cell surface NKG2D expression. Because purified exosomes from untransfected CHO cells had no effect on receptor expression, down-modulation of NKG2D depended specifically on the presence of exosomal ULBP3.

Exosomal ULBP3 Can Compromise NKG2D-mediated Cell Cytotoxicity—The fact that exosomal protein potently down-modulated NKG2D suggested that incubation with ULBP3-containing exosomes might impair cytotoxic responses. To test this hypothesis, exosomes purified from ULBP3 transfectants and untransfected control cells were incubated with primary NK cells before including these effectors in cytotoxicity assays. As expected, human NK cells lysed MICA-transfected CHO cells in an NKG2D-dependent manner, while untransfected CHO cells were not efficiently lysed (Fig. 7A). Preincubation with ULBP3-containing exosomes resulted in a clear cut reduction of the NKG2D-mediated lytic activity of NK cells against cells expressing MICA (Fig. 7B). These results demonstrate that ULBP3-containing exosomes but not other exosomes devoid of NKG2D-L inhibit NKG2D-mediated responses.

DISCUSSION

A recurrent question in the field of the NKG2D system is why are there so many ligands for one, invariant receptor. The existence of such a large number of ligands for a single receptor is
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from tumor cells. Striking differences in both the kinetics of release and the species of protein produced were observed: truncated soluble protein versus membrane-bound. These data constitute the first demonstration of differences in the biochemistry of the GPI-linked members of the ULBP family and could be of crucial importance to understand the process of escape from the immune system by stressed cells up-regulating NKG2D-L, such as pathogen-infected and cancer cells.

ULBP2 is known to be released by the action of metalloproteinases (34) including members of the ADAM family (22). Using pharmacological agents to stimulate or inhibit different classes of protease, we reproduced the published data with ULBP2 and showed that inhibition of multiple classes of proteases including metalloproteases did not significantly affect ULBP3 release. Furthermore, the kinetics of ULBP3 shedding were not comparable to those of ULBP2, and the release of ULBP3 was much slower. Moreover, an increase in ULBP3 shedding could be observed after a long exposure to ionomycin. As this increase did not occur at short incubation times, this suggested not an effect on ADAM10 but on the process of exosome formation and release.

Analysis of the biochemistry of released ULBP2 and ULBP3 gave a definitive response; ULBP3 was found to be released in exosomes. This was observed in various transfectants and in cells endogenously expressing the protein; thus, we infer that this is due to the characteristics of the protein and not to the cell line. In the transfected cells, a small proportion of ULBP3 can also be observed in the soluble fraction; thus, a truncated monomeric form is generated in addition to the membrane-bound ULBP3 molecule. More work needs to be done to evaluate the spatio-temporal characteristics of these two processes. The observation that ULBP2 can be included in exosomes, but it is more susceptible to metalloprotease attack is consistent with the sequence differences between ULBPs in the stalk region and further study is needed to define the proteolytic site for ULBP2.

The expression of a particular NKG2D-L does not always imply cytotoxic attack toward the cell. Instead, the fate of the target cell depends on the post-transcriptional and post-translational modifications that direct the protein for surface expression, retention, or shedding. Some NKG2D-L do not even reach the surface after translation (46); in other cases, the protein recycles into endosomal compartments (47); and finally, NKG2D-L are known to be released to the surrounding medium (12, 34). In light of our data, it seems plausible to speculate that expression of a particular ULBP would lead to different amounts of protein shed to the supernatant and as different protein species, more or less potent in NKG2D down-modulation. The functional consequences of this differential release would result in the complex spectrum of situations that are observed in cancer patients, some expressing large amounts of NKG2D-L, some not expressing any. For this reason, it will be of crucial importance to study the particular cell biological properties of the individual NKG2D-L.

Elevated levels of soluble MICA have been detected in the sera of patients suffering from various types of cancer and often correlate with a poor prognosis for the patient (48). This relationship is likely to be related to the known effects of MICA/B...
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... shedding: a reduction in cell surface density of NKG2D ligands leading to a reduced susceptibility to NKG2D-mediated cytoxicity and systemic down-regulation of NKG2D on NK cells and CD8+ T cells in cancer patients. The occurrence and significance of serum ULBP molecules has not been well studied, but soluble ULBPs have been detected in the serum of patients with hematological malignancies (34) and colorectal cancer. The data presented here show that incubation with supernatants containing either soluble ULBP2 or exosomal ULBP3 molecules also leads to significant down-regulation of cell surface NKG2D expression. Interestingly, incubation of NK cells with the ULBP3-containing supernatant triggers significantly more NKG2D down-regulation (Fig. 5) than the ULBP2 culture supernatant despite containing much less ULBP protein. A goal of future research will be to investigate whether this might be related to differences in how the two ULBP species exist in the supernatant: soluble, probably monomeric ULBP2 versus multivalent ULBP3 molecules in exosome membranes and whether this is also true in patient sera.

The ability of soluble NKG2D-L to provoke NKG2D down-regulation and compromised NK and cytotoxic T lymphocytes function suggests that blockade of NKG2D-L release might be a useful addition to immunological approaches for cancer therapy, but an understanding of the mechanisms involved in the release of soluble NKG2D-L from tumor cells is crucial for the development of effective strategies to block the shedding of these proteins. Metallloproteases are key enzymes mediating proteolytic cleavage of both MICA/B and ULBP2 molecules (22, 47). However, the data in this manuscript now show that ULBP2-L molecules can be released from a tumor cell in more than one way. ULBP3 is not released from cells by proteolysis but rather as a full-length molecule in exosomes. Thus, effective blockade of the accumulation of soluble NKG2D-L in patient sera will require the use of multiple strategies.

In conclusion, we have demonstrated here that different ULBP proteins are released in diverse amounts as different species and that these released ULBP proteins can trigger NKG2D down-regulation. These findings offer the possibility of starting novel approaches to try to understand the regulation of the fate of a cell expressing NKG2D-L. For example, the study could be extended to analyze the effect of different stimuli that up-regulate particular NKG2D-L (6) on the release of that molecule.

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REFERENCES

1. Raulet, D. H. (2003) Nature 427, 781–790
2. Moretta, A., Bottino, C., Vitale, M., Pende, D., Cantoni, C., Mingari, M. C., Biassoni, R., and Moretta, L. (2001) Annu. Rev. Immunol. 19, 197–223
3. Radosavljevic, M., Cuillerier, B., Wilson, M. J., Clément, O., Wicker, S., Gilfillan, S., Beck, S., Trowsdale, J., and Bahram, S. (2002) Genomics 79, 283–292
4. Jafferi, L. Fernandez-Messina, S. Agüera-González, M. Vales-Gomez, R. McGilvray, L. G. Durrant, J. Trowsdale, and R. A. Eagle, manuscript in preparation.
34. Waldhauer, I., and Steinle, A. (2006) Cancer Res. 66, 2520–2526
35. Hooper, N. M., Karran, E. H., and Turner, A. J. (1997) Biochem. J. 321, 265–279
36. Horiuchi, K., Le Gall, S., Schulte, M., Yamaguchi, T., Reiss, K., Murphy, G., Toyama, Y., Hartmann, D., Saftig, P., and Blobel, C. P. (2007) Mol. Biol. Cell 18, 176–188
37. Savina, A., Furlán, M., Vidal, M., and Colombo, M. I. (2003) J. Biol. Chem. 278, 20083–20090
38. Stoorvogel, W., Kleijmeer, M. J., Geuze, H. J., and Raposo, G. (2002) Traffic 3, 321–330
39. Escola, J. M., Kleijmeer, M. J., Stoorvogel, W., Griffith, J. M., Yoshie, O., and Geuze, H. J. (1998) J. Biol. Chem. 273, 20121–20127
40. de Gassart, A., Geminard, C., Fevrier, B., Raposo, G., and Vidal, M. (2003) Blood 102, 4336–4344
41. Vetter, C. S., Lieb, W., Bröcker, E. B., and Becker, J. C. (2004) Br. J. Cancer 91, 1495–1499
42. Castriconi, R., Cantoni, C., Della Chiesa, M., Vitale, M., Marcenaro, E., Conte, R., Biassoni, R., Bottino, C., Moretta, L., and Moretta, A. (2003) Proc. Natl. Acad. Sci. U.S.A. 100, 4120–4125
43. Clayton, A., Mitchell, J. P., Court, J., Mason, M. D., and Tabi, Z. (2007) Cancer Res. 67, 7458–7466
44. Clayton, A., Mitchell, J. P., Court, J., Linnane, S., Mason, M. D., and Tabi, Z. (2008) J. Immunol. 180, 7249–7258
45. Eagle, R. A., and Trowsdale, J. (2007) Nat. Rev. Immunol. 7, 737–744
46. Borchers, M. T., Harris, N. L., Wesselkamper, S. C., Vitucci, M., and Cosman, D. (2006) Am. J. Physiol. Lung Cell Mol. Physiol. 291, L222–231
47. Aguera-González, S., Boutet, P., Reyburn, H. T., and Valéz-Gómez, M. (2009) J. Immunol. 182, 4800–4808
48. Holdenrieder, S., Stieber, P., Peterfi, A., Nagel, D., Steinle, A., and Salih, H. R. (2006) Cancer Immunol. Immunother. 55, 1584–1589