Inhibition of IgE-mediated mast cell activation by the paired Ig-like receptor PIR-B

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The potential of the paired Ig-like receptors of activating (PIR-A) and inhibitory (PIR-B) types for modifying an IgE antibody-mediated allergic response was evaluated in mouse bone marrow–derived mast cells. Although mast cells produced both PIR-A and PIR-B, PIR-B was found to be preferentially expressed on the cell surface, where it was constitutively tyrosine phosphorylated and associated with intracellular SHP-1 protein tyrosine phosphatase. PIR-B oligation with the IgE receptor (FcεRI) inhibited IgE-mediated mast cell activation and release of serotonin. Surprisingly, the inhibitory activity of PIR-B was unimpaired in SHP-1–deficient mast cells. A third functional tyrosine-based inhibitory motif, one that fails to bind the SHP-1, SHP-2, and SHIP phosphatases, was identified in parallel studies of FcεRI-bearing rat basophilic leukemia (RBL) cells transfected with constructs having mutations in the PIR-B cytoplasmic region. These results define the preferential expression of the PIR-B molecules on mast cells and an inhibitory potential that can be mediated via a SHP-1–independent pathway.


Introduction

The paired Ig-like receptors of activating (PIR-A) and inhibitory (PIR-B) types were originally identified in mice on the basis of limited homology with the human IgA Fc receptor (FcαR) (1, 2). Their human counterparts are considered to be the activating and inhibitory types of leukocyte Ig-like receptors/CD85 (3–6). PIR-A and PIR-B have been defined as cell surface glycoproteins with similar extracellular regions (>92% homology) containing six Ig-like domains, and distinctive transmembrane and cytoplasmic regions. PIR-A isoforms with slightly different sequences are encoded by the six Pir genes, whereas the invariant PIR-B molecule is encoded by a single Pirb gene (1, 2, 7, 8). The Pir-A proteins have a short cytoplasmic tail and a charged arginine residue in their transmembrane domain that facilitates noncovalent association with a transmembrane adaptor molecule, the Fc receptor common γ chain (FcγRc), to form a cell activation complex (9–12). The PIR-B molecule contains an uncharged transmembrane segment and four potential immunoreceptor tyrosine-based inhibitory motifs (ITIMs) in the cytoplasmic tail. Two of the ITIM regions of PIR-B, when tyrosine phosphorylated, can recruit the protein tyrosine phosphatase SHP-1, and possibly SHP-2 as well, to inhibit cell activation (10, 13–15), but these carboxy terminal ITIMs do not appear to account for all of the PIR-B inhibitory activity (13, 14).

PIR-A and PIR-B are expressed by many types of hematopoietic cells, including B lymphocytes, dendritic cells, monocyte/macrophages, granulocytes, megakaryocytes/platelets, and mast cells (11, 16). Interestingly, the PIR-B molecules on freshly isolated B lymphocytes and macrophages have been found to be constitutively tyrosine phosphorylated, but they are rarely tyrosine phosphorylated on corresponding cell lines before their ligation by antibody (17). The reduced levels of PIR-B tyrosine phosphorylation found in β2 microglobulin-deficient mice suggest that MHC class I or class I–like molecules may serve as natural PIR ligands (17).

Mast cells are important mediators of allergic responses. They are generated in the bone marrow, circulate as immature precursors, and migrate into various tissue...
sites where they undergo terminal differentiation. Basophils also develop in the bone marrow, but they circulate as fully functional granulated cells that migrate into tissues in response to inflammation. Both types of cells contain metachromatic granules loaded with histamine, serotonin, and other biologically active products. They express high-affinity IgE receptors (FcεRI) and low-affinity IgG receptors (FcγRIII), as well as receptors for multiple cytokines and growth factors. Upon activation by contact with allergens, the IgE antibody-sensitized mast cells release the pharmacologically active mediators stored in their granules, resulting in clinical manifestations of type I hypersensitivity (18).

Information about the basic biology of mast cells and basophils has been gained largely through studies of bone marrow–derived mast cells (18) and the rat basophilic leukemia cell line (RBL-2H3). The RBL-2H3 cell line has been particularly useful in analyzing the activating and inhibitory potential of PIR-A and PIR-B in transfection studies using chimeric constructs (10, 13), but the biochemical nature and functional properties of native PIR molecules on the mast cells have not been examined previously. These issues have been addressed in the present studies of cultured mast cells of bone marrow and splenic origin. In parallel studies, the RBL-2H3 cell line was used to refine the definition of PIR-B–inhibitory motifs.

Methods

Mice. Four- to 8-week-old C57BL/6 (H-2b), C3H/HeJ (H-2k), and BALB/cJ mice were purchased from The Jackson Laboratory (Bar Harbor, Maine, USA). C3H/HeJ mice heterozygous for motheaten mutation (me/+), kindly provided by K.A. Siminovitch (Mount Sinai Hospital, Toronto, Ontario, Canada), were bred to generate me homozygous mice (me/me). The me mutation status was identified by genomic PCR using diagnostic primers as described previously (19).

IL-3–induced mast cell cultures. Bone marrow cells were obtained from the femurs of adult C57BL/6, C3H/He, and BALB/cJ mice. Splenic cells were obtained from neonatal motheaten (me/me) and wild-type (+/+ ) littermate mice. After lysing the erythrocytes with 0.15 M ammonium chloride solution, bone marrow and splenic cells were cultured at a cell concentration of 1 × 10^6/ml in RPMI 1640 medium containing 10% FCS (HyClone Laboratories, Logan, Utah, USA), penicillin (100 U/ml), streptomycin (100 µg/ml), and 2% SDS, resolved on SDS-PAGE (10% acrylamide) gels, and transferred to PVDF membranes. Membrane-bound proteins were detected with specific antibodies and visualized with chemiluminescence (Pierce, Rockford, Illinois, USA). Membrane-bound proteins were stained with anti-c-kit (ACK-2, µg/ml; ref. 21), a gift of S. Nishikawa (Kyoto University, Kyoto, Japan), and for CD40 (FGK45, IgGκ; ref. 22), a gift of J. Anderson (Basel Institute for Immunology, Basel, Switzerland). All experiments were performed in triplicate, and representative gels are shown.

Immunoprecipitation analysis of cell surface and intracellular proteins. Viable cells (3 × 10^7) were surface radiolabeled with 1 mCi of Na^125I by lactoperoxidase catalysis and solubilized in ~500 µl of 1% NP-40 lysis buffer containing various protease inhibitors as described elsewhere (11, 17). The ultracentrifuged membrane lysates were examined by a solid-phase immunosolulotion technique. In brief, microtiter plates precoated with goat anti-rat Ig antibodies (Southern Biotechnology Associates, Birmingham, Alabama, USA). Stained cells were analyzed with a FACSCalibur flow cytometer (Becton Dickinson Immunocytometry Systems, San Jose, California, USA).

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Protein blot analysis. Cells (3 × 10⁷) were solubilized in lysis buffer supplemented with 2 mM sodium orthovanadate and 50 mM sodium fluoride (17). Proteins in the cell lysates that were immunoadsorbed with anti-PIR or control mAb’s or with streptavidin-coupled beads containing the biotinylated PIR-B phosphopeptides were resolved on SDS-PAGE and transferred onto nitrocellulose membranes by electroblotting. After soaking in 3% BSA in PBS containing 0.05% Tween-20, membranes were incubated with HRP-labeled mouse anti-phosphotyrosine mAb (4G10; Upstate Biotechnology Inc., Lake Placid, New York, USA) or with rabbit antibodies specific for protein tyrosine phosphatase SHP-1 (Upstate Biotechnology Inc.) or SHP-2 (Santa Cruz Biotechnology, Santa Cruz, California, USA) or for inositol polyphosphate 5-phosphatase SHIP (Santa Cruz Biotechnology). In some experiments, mouse anti-SHP-2 mAb (Transduction Laboratories, Lexington, Kentucky, USA) and HRP-labeled goat anti-mouse IgG antibody (Southern Biotechnology Associates) were used. For rabbit antibodies, HRP-labeled goat anti-rabbit Ig antibody (0.1 µg/ml; Southern Biotechnology Associates) was used as a developing reagent before visualization by enhanced chemiluminescence employing the oxidation of luminol in the presence of phenols (Amersham Life Science, Buckinghamshire, United Kingdom). Membranes were reblotted with rabbit anti-PIR antiserum (1:4,000 dilution) to detect PIR-A and PIR-B molecules.

RT-PCR analysis. Total RNA was extracted from cultured mast cells and control cell lines (10⁶ cells) by the TRIzol method (Molecular Research Center Inc., Cincinnati, Ohio, USA). Two micrograms of RNA were converted to first-strand cDNA with oligo (dT)₁₂–₁₈ primers and SuperScript RNase H– RT (Life Technologies Inc., Rockville, Maryland, USA), and amplified with a common forward primer (5′-CCCTGGAGGCTCA-CAGTCTCAG-3′), and the PIR-A specific (5′-CCCA-GAGTGTAGAACATTGAAGATG-3′) or PIR-B specific (5′-GGTTCATGGTGTCCTCCCTGGACAGCAGG-3′) reverse primers. These primers correspond to the 3′ end of D6 of PIR-A and PIR-B and to the cytoplasmic regions of PIR-A and PIR-B, respectively, and yield fragments of 252 bp for PIR-A and 470 bp for PIR-B (1). Each amplification reaction included 30 cycles of denaturation at 94°C for 1 minute, annealing at 60°C for 1 minute, and extension at 72°C for 1 minute. A final extension was performed at 72°C for 5 minutes. Amplification of actin transcripts with the primers (5′-ATCGAACATGGGATTGTGTCACC-3′ and 5′-GGCATAAGGGACAGACAGG-3′) was also performed as a control. Amplified products were electrophoresed in 1.5% agarose and stained with ethidium bromide.

Calcium mobilization analysis. Intracellular Ca²⁺ concentrations were measured after PIR and/or FcεRI cross-linkage on mast cells as described elsewhere (13). Briefly, mast cells were loaded with Indo-1/AM (1 µg/ml; Calbiochem-Novabiochem Corp., San Diego, California, USA) in the presence or absence of rat IgE anti-DNP mAb (1 µg/ml) for 30 minutes at 37°C, resuspended at 10⁶ cells/ml either in HBSS containing 5% FCS or Ca²⁺-free HBSS containing 1 mM MgCl₂ and were incubated first with F(ab)₂ fragments of the rat anti-PIR mAb (10.1 clone; γ2bκ; 20 µg/ml) and then with F(ab)’₂ fragments of rabbit anti-rat Ig antibody (50 µg/ml; Southern Biotechnology Associates) as a cross-linker. Purity of the F(ab)’₂ reagents was established by SDS-PAGE analysis. An intact rat mAb (1D6 clone; γ2aκ), against an undefined cell surface component on mouse cells, including mast cells, was used as a control for the 10.1 anti-PIR mAb. The 395/510 nm fluorescence ratio for the dye-loaded viable cells was determined with a FACStar Plus flow cytometer (Becton Dickinson Immunocytometry Systems) equipped with an Enterprise laser (Coherent Laser Group, Santa Clara, California, USA) regulated at 40 mW UV.

Serotonin-release assay. IgE-mediated serotonin release assay was performed according to methods described previously (13), with the following minor modifications. IL-3–induced mast cells (1 × 10⁶ cells/ml) were incubated with 5-[1,2–³H(N)]-hydroxytryptamine binoxalate (5 µCi/ml; 37,000 MBq; NEN Life Science Products Inc.) for 5 hours at 37°C, washed, and reincubated for an additional 1 hour at 37°C to reduce background radioactivity. Thirty microliters of ³H-serotonin–loaded mast cells (1.4 × 10⁷ cells/ml) were plated in 96-well plates that contained 10 µl of rat IgE anti-DNP mAb (25 µg/ml) and 10 µl of various concentrations (0.5–50 µg/ml) of F(ab’)₂ fragments of rat anti-PIR mAb in triplicate, incubated for 30 minutes at 4°C, washed, and resuspended in 25 µl of culture medium. The 1D6 rat mAb (intact form) was used as a control antibody. Antibody-primed mast cells were challenged with 25 µl of F(ab’)₂ fragments of rabbit anti-rat Ig antibody (40 µg/ml) for 30–60 minutes at 37°C. In other experiments, DNP-coupled human serum albumin and goat anti-rat IgG antibody were used to aggregate separately the IgE and PIR receptors. The reactions were terminated by adding 50 µl of cold HBSS and centrifugation.

³H-Serotonin in the supernatants was measured by a liquid scintillation counter. The total incorporation of ³H-serotonin was determined by lysis of unprimed cells with 1% SDS/1% NP-40, and the percent of serotonin release was defined as: 100 × (cpm of supernatants ÷ cpm of total incorporation). For cells transfected with FcγRIIB/P-PIR chimeric constructs, mouse IgE anti-TNP mAb (2682-1), rat anti-mouse FcγRIIB/III mAb (2.4G2; γ2bκ), and F(ab)’₂ fragments of goat anti-murine Ig antibodies (Immunotech, Marseille, France) were used to ligate FcγRI and FcγRIIB/P-PIR as described elsewhere (13).

Constructions. cDNA constructs generated from a pNT-Neo plasmid included a chimeric cDNA encoding the extracellular, transmembrane, and the first six amino acids (KKKVQP; one-letter code) of the cytoplasmic tail of the mouse FcγRIIB, fused with the cytoplasmic region of the mouse PIR-B (FcγRIIB/P-PIR-B) as described previously (13). The initial FcγRIIB in
characteristic for IL-3-induced, bone marrow–derived mast cells (18, 24). Importantly for our studies, the IL-3–induced mast cells from normal and FcεRIα−/− mice expressed cell surface PIR proteins at relatively high levels (Figure 1, a and b).

Mast cell cultures were also derived from neonatal splenic progenitors in SHP-1–deficient motheaten (me/me) mice and control littersmates because the me/me defect leads to early death. Like the mast cells generated from adult bone marrow, the spleen-derived mast cells expressed cell surface PIR as well as c-kit, CD13, FcγRII/III, CD69, IgE binding capacity, and intracellular metachromatic granules (Figure 1, c and d).

**Predominant expression of constitutively tyrosine-phosphorylated PIR-B on mast cells.** In a previous analysis, both PIR-A and PIR-B transcripts were found to be expressed by the P815 mast cell line, although this cell appeared to express higher levels of PIR-B transcripts than PIR-A transcripts. When a similar RT-PCR analysis was conducted for the bone marrow–derived mast cells, both PIR-A and PIR-B transcripts were found in roughly equal abundance (Figure 2a).

An immunoprecipitation analysis of radiolabeled cell surface proteins was then performed with the 6C1 anti-PIR antibody. This antibody recognizes a common epitope on the D1 and D2 Ig-like ectodomains of PIR-A and PIR-B proteins that can be distinguished on the basis of their relative molecular masses (11). To compare the array of PIR molecules expressed on the different types of cells, nonadherent mast cells and adherent macrophages in the bone marrow–derived cultures were separated and their cell surface PIR proteins were analyzed along with those expressed by splenocytes, the majority of which were B lymphocytes. Both the 120-kDa PIR-B and 85-kDa PIR-A were precipitated from membrane lysates of surface iodinated splenocytes and macrophages, whereas only PIR-B molecules were detected in the membrane lysates of mast cells (Figure 2b). Given that FcεRIγ is required for the expression of functional PIR-A, limited FcεRIγ availability due to competition with the FcεRI could account for the paucity of cell surface PIR-A expression. To test this possibility, the FcεRIα−/−–derived mast cells were examined for their surface PIR expression. The results again indicated predominant expression of PIR-B on the surface of FcεRIα−/− mast cells, thereby suggesting that a limitation in the availability of FcεRIγ is unlikely to be the basis for the paucity of surface PIR-A on mast cells (Figure 2c, upper panel). The predominance of cell surface PIR-B was also observed for mast cells derived from splenic progenitors in neonatal me/me mice and control littersmates (Figure 2c, lower panel) The PIR-B molecules on mast cells were also found to be constitutively tyrosine phosphorylated and to be associated with SHP-1 protein tyrosine phosphatase, except in mast cells from me/me mice. SHP-2 was not found to be associated with PIR-B in mast cells (data not shown), consistent with the results of an earlier
analysis of PIR-B molecules on splenic B lymphocytes and macrophages (17).

PIR-B–mediated inhibition of FcεRI-induced Ca^{2+} mobilization and serotonin release. Because mast cells preferentially express PIR-B on their cell surfaces, examination of the functional consequence of PIR-A/PIR-B ligation with a nondiscriminative antibody is a more straightforward analysis than for the B lymphocytes and myeloid cells that express both PIR-A and PIR-B molecules on their cell surface. The effects of PIR-B cross-linkage on FcεRI-mediated Ca^{2+} mobilization were examined initially as an early indicator of the cellular response. Ligation of FcεRI on mast cells induced an immediate Ca^{2+} release from intracellular pools, followed by a sustained increase that may reflect the influx of extracellular Ca^{2+}. Although PIR ligation alone had no demonstrable effect on intracellular Ca^{2+}, levels, coligation of FcεRI and PIR inhibited IgE-induced Ca^{2+} mobilization (Figure 3, upper panel). When this experiment was repeated in Ca^{2+}-free media, IgE-induced Ca^{2+} mobilization was again inhibited by PIR-B and FcεRI coligation (data not shown), thereby confirming that the PIR-B inhibition affects Ca^{2+} release from the intracellular pool.

When the effects of PIR-B cross-linkage on FcεRI-mediated serotonin release were evaluated, PIR coligation with the FcεRI inhibited this mast cell response in a dose-dependent manner (Figure 4, left panel). The inhibition was observed only when the PIR-B and FcεRI were brought into physical proximity by ligation with a common secondary reagent. In a test of the PIR-B specificity of the inhibitory effect, coligation of FcεRI with an unrelated cell surface molecule did not inhibit IgE-mediated serotonin release. Separate ligation of cell surface PIR-B and FcεRI with individual secondary reagents also did not result in inhibition of the IgE-mediated serotonin release, and ligation of PIR-B alone did not induce serotonin release (data not shown). These findings collectively demonstrate the functional potential of PIR-B as a negative regulator of IgE-mediated mast cell activation.

SHP-1 is dispensable in the PIR-B inhibitory activity. In native B cells and in transfected cell models, PIR-B has been shown to recruit the SHP-1 protein tyrosine phosphatase (10, 13–15). To test the requirement for SHP-1 in the PIR-B-mediated inhibition of FcεRI signaling, we examined mast cells from SHP-1–deficient motheaten (me/me) mice and normal littermate (+/+) controls. To our surprise, IgE antibody-mediated intracellular Ca^{2+}-
mobilization and serotonin release were inhibited as efficiently by PIR-B and FcRI coligation on mast cells from SHP-1–deficient motheaten (me/me) mice as for mast cells from control (+/+ ) littermates (Figure 3, lower panel; Figure 4, right panel). The absence of the 65-kDa SHP-1 protein was confirmed by immunoprecipitation analysis of the motheaten-derived mast cells with anti–SHP-1 antibodies as described in Methods. The standard deviation from the mean (circle) is indicated by bars only for the points at which this value (1 SD) is greater than the circle radius.

Rather, the data raise the possibility that another, yet unidentified, inhibitory signaling element can participate in the PIR-B inhibitory function.

A third tyrosine-based motif in the PIR-B cytoplasmic tail mediates inhibitory function. The PIR-B protein contains five cytoplasmic tyrosine residues, four of which, Y713 (SLYASYV), Y742 (ETYAQTV), Y794 (VTYAQL), and Y824 (SVYATL), reside within ITIM-like sequences (I/V/L/S-x-Y-x-x-V/L), whereas the other tyrosine, Y770 (TEYEQA), does not reside in an ITIM consensus sequence. In previous studies (13), phosphopeptides corresponding to the Y794 tyrosine– and Y824 tyrosine–based motifs were shown to bind SHP-1 and SHP-2 phosphatases derived from a B cell line, whereas phosphopeptides representing the other two ITIM candidates failed to bind either phosphatase. To test whether these relationships held true for mast cells, the cellular proteins in lysates of metabolically labeled and unlabeled mast cells were incubated with phosphopeptides containing the PIR-B tyrosine residues (Y713, Y742, Y770, Y794, Y824) or the FcγRIIB ITIM as a control. Although the FcγRIIB phosphopeptide bound multiple SHIP isoforms as well as SHP-1 and SHP-2, the two carboxy terminal PIR-B ITIM phosphopeptides (Y794 and Y824) again were found to bind both SHP-1 and SHP-2. Although the Y742 phosphopeptide failed to bind SHIP, SHP-1 or SHP-2, a protein of approximately 120 kDa was instead found to be associated with this ITIM candidate (Figure 5).

Previous transfection studies have indicated that the two carboxy terminal PIR-B ITIMs (Y794 and Y824) are responsible for most, but not all, of the inhibitory capacity of PIR-B (13, 14). To determine whether the

Figure 4
PIR-B inhibitory activity in FcγRI-induced serotonin release. ³H-Serotonin preloaded mast cells from wild-type neonatal spleen (left) and motheaten mutant neonatal spleen (right) were incubated with rat IgE anti-DNP mAb plus various concentrations of F(ab)² fragments of rat mAb specific for PIR (filled circles) or unknown cell surface antigen (open circles). Antibody-primed cells were then triggered with F(ab)²; fragments of rabbit anti-rat Ig antibody as a cross-linker, and the ³H-serotonin release was measured as described in Methods. The Journal of Clinical Investigation | October 2001 | Volume 108 | Number 7

Figure 5
Analysis of mast cell proteins binding to PIR-B ITIM. Left panel: Metabolically labeled mast cell proteins were solubilized in 1% NP-40 lysis buffer before incubation with streptavidin-coupled beads bearing biotinylated 15-mer peptides corresponding to the mouse FcγRIIB ITIM (tyrosine-phosphorylated, pY, or tyrosine-nonphosphorylated, Y, as controls) and each of the tyrosine-phosphorylated and biotinylated PIR-B ITIM (pY713, pY742, pY770, pY794, pY824). Bound materials were resolved by SDS-8% PAGE before electrophoresis on reducing conditions before exposure to x-ray films. Arrowhead indicates 120-kDa protein band precipitated by the pY742 phosphopeptide. Right panel: Proteins isolated from mast cell lysates with the same phosphopeptides were resolved by SDS-PAGE before transfer onto membranes and immunoblotting with antibodies specific for SHP-1, SHP-2, or SHIP before visualization by enhanced chemiluminescence.
other two tyrosine-based motifs, centered around Y713 and Y742, can also mediate PIR-B inhibitory activity, four additional chimeric constructs were prepared: (a) One encodes a short PIR-B cytoplasmic tail with a carboxy terminal deletion from five amino acids after Y742 (YF); (b) the second encodes a similar short PIR-B cytoplasmic tail with a point mutation of Y742F (YF); (c) the third includes three point mutations, Y713F, 794F, and Y824F, to allow evaluation of the role of Y742 (FYFF); and (d) the fourth encodes the shortest PIR-B cytoplasmic tail, rendered tyrosine-free by deleting the carboxy terminus from position 712 onward (Δ712). These chimeric constructs were transfected into RBL-2H3 cells, and the resultant stable transfectants were shown by immunofluorescence to express levels of FcγRIIB/PIR-B chimeric receptors comparable to that of the wild-type transfectant, except for the YF transfectant, where the FcγRIIB/PIR-B level was higher (Figure 6). The cell surface levels of endogenous FcεRI on the different transfectants were not altered.

These chimeric receptor transfectants were examined for their potential to inhibit the FcεRI-mediated serotonin release. As previously shown (13), coligation of FcεRI and the wild-type (YYYY) chimeric receptor effectively inhibited the IgE-mediated serotonin release (Figure 7). The YY-type chimeric receptor could also mediate significant inhibition over a wide range of IgE stimulation dosages, whereas the YF type chimeric receptor did not, even though the YF transfectant expressed higher levels of chimeric receptors than did the YY transfectant. The FYFF-type chimeric receptor mediated inhibition to a similar degree as did the YY-type chimeric receptor, thereby implicating the Y742-based motif in the inhibition. No inhibitory capability could be demonstrated for the Δ712-type chimeric receptor. These findings demonstrate that three of the four PIR-B ITIM candidates, namely the Y742-, Y794-, and Y824-based motifs, have inhibitory capability. The inhibitory function of the newly verified Y742-based ITIMs may involve the associated p120 molecule (see Figure 5).

**Discussion**

Several unanticipated features of the PIR-B inhibitory potential in mast cells are indicated in these studies. Although mast cells apparently produce PIR-A and PIR-B in similar amounts, the latter is preferentially expressed on the mast cell surface as a constitutively tyrosine phosphorylated molecule. Although the SHP-1 protein tyrosine phosphatase is normally associated with PIR-B molecules, surprisingly, this phosphatase appears to be entirely dispensable in the PIR-B-mediated inhibition of IgE antibody-mediated mast cell activation. A third functional PIR-B ITIM identified in this study, the Y742-based ETYAQV motif, that failed to bind either protein tyrosine phosphatases (SHP-1 and SHP-2) or inositol polyphosphate 5-phosphatase (SHIP) (8,10,13), was found to associate with another potential regulatory element, a presently unidentified protein of approximately 120 kDa. Mast cells derived from bone marrow or splenic progenitors were found to produce the PIR-B and PIR-A transcripts in comparable levels, but an analysis of their cell surface expression indicated a striking prevalence of PIR-B. PIR-A could not be detected on the mast cells, whereas PIR-B was easily and consistently detected on the cell surface. Previous studies have indicated that an association with FcγRI homodimers is required for the normal expression of PIR-A molecules on the cell surface (11). As FcγRIc is also an essential component of the high-affinity IgE receptor complex, FcεRI (25), an inability to compete efficiently with the FcεRI for a limited supply of FcγRIc dimers could explain the paucity of PIR-A molecules that reach the mast cell surface. Indeed, previous studies have suggested competition between FcεRI and FcγRIII for limiting amounts of FcγRIc (26, 27). Functional FcγRIII is not expressed on mast cells, as it is not efficiently associated with the FcγRIc and thus...
Absence of FcεRIα chain thus results in upregulation of FcγRIII-dependent mast cell responses (27). However, we found that FcεRIα chain–deficient mast cells also lack detectable cell surface PIR-A expression, thereby suggesting that other regulatory mechanisms may account for the paucity of PIR-A on mast cells. The IL-3–induced mast cells should provide an ideal cell source to determine the regulatory mechanisms affecting the equilibrium of cell surface PIR-A and PIR-B expression.

The integrity of the PIR-B–mediated inhibition of IgE triggered Ca2+ mobilization and serotonin release in SHP-1–deficient mast cells was also unanticipated, because previous studies have suggested that the inhibitory activity of PIR-B is mediated primarily through its association with SHP-1 (13, 14, 17). The possibility was therefore considered that the inhibition seen in me/me mast cells reflected artificial co-cross-linkage of PIR-A with the FcγRIIB via the intact antibodies used for receptor coligation. FcγRIIB has its own ITIM motif, through which it can interact with the SHIP inositol polyphosphate 5-phosphatase to inhibit cell activation (28–30). However, this explanation of our results was rendered unlikely by the observations that (a) intact IgG molecules could not be detected in the F(ab′)2 preparations of rat anti-PIR and rabbit anti-rat Ig antibodies that were used for receptor coligation, (b) inhibitory responses generated by PIR-B and FcγRIIB are dissimilar in that PIR-B–mediated inhibition involves a blockage of Ca2+ release from intracellular stores (see Figure 3), whereas FcγRIIB-mediated inhibition involves blockage of Ca2+ influx (28), and (c) the coligation of FcεRI and an unrelated molecule with an intact antibody failed to inhibit the IgE-mediated Ca2+ mobilization and serotonin release. The alternative possibility that the inhibitory activity of PIR-B could directly involve SHIP is rendered unlikely by the inability to identify an association between these two molecules in the present studies or in earlier analyses (13–15, 17).

Protein tyrosine phosphatase activity mediated by SHP-2 is a third possible explanation for the PIR-B inhibitory effect observed in SHP-1–deficient mast cells. In a previous analysis using a B cell line, we observed that SHP-1 and SHP-2 could bind to phosphopeptides corresponding to the two carboxy-terminal ITIM (Y794 and Y824) in the PIR-B cytoplasmic tail (13). The present studies indicate that the PIR-B Y794 and Y824 phosphopeptides also associate with SHP-1 and SHP-2 in mast cells. However, in normal mast cells, splenic lymphocytes (17), and RBL-2H3 cells transfected with chimeric receptor (extracellular FcγRIIB/PIR-B cytoplasmic tail) constructs (13), we have consistently observed the association of PIR-B with SHP-1, and not with SHP-2. Nevertheless, when Maeda et al. used PIR-B constructs to transfect DT-40 chicken B cells deficient in SHP-1, SHP-2, or both, their results suggested that both the SHP-1 and SHP-2 phosphatases may participate in PIR-B–mediated inhibition (14). Although this interpretation could also explain our results with the motheaten (me/me) mast cells, PIR-B–associated SHP-2 molecules were not demonstrable in mast cells from normal mice or the SHP-1–deficient motheaten mice. This raises the possibility that PIR-B signaling elements other than SHP-1, SHP-2, and SHIP contribute to the inhibitory function. Consistent with this hypothesis, Maeda et al. found residual PIR-B inhibitory activity in the SHP-1 and SHP-2 double-deficient DT40 cells, leading them to invoke the involvement of an additional SH2-containing protein(s) in mediating the PIR-B inhibitory signal (14).

Figure 7
Inhibitory activity of FcγRIIB/PIR-B chimeric receptors expressed by RBL transfected cells. RBL-2H3 cells transfected with the indicated FcγRIIB/PIR-B constructs were incubated with various dilutions of mouse IgE in the absence (filled circles) or presence (open circles) of 2.4G2 rat anti-FcγRII/III mAb (20 µg/ml). F(ab′)2 fragments of goat anti-murine Ig antibody (50 µg/ml) were used as a cross-linker. A representative result is shown as a mean ± 1 SD (if the SD values are greater than the radius of circles) from at least three independent experiments. Statistical analysis revealed that inhibition is significant for IgE at 1:1,000 and 1:300 dilutions in a, b, and d (data not shown).
In this regard, a third functional ITIM in the PIR-B cytoplasmic tail is indicated by the present results. In addition to the two carboxy-terminal ITIM centering around the Y794 and Y824 residues, transfection experiments with constructs differing in the integrity of their PIR-B cytoplasmic region indicate that one of the more membrane proximal ITIM (Y742: ETYAVQY) is capable of mediating inhibition of the FcεRI-induced serotonin release. While the fourth candidate ITIM (Y713, SLYASV) matches the consensus sequence (I/V/L/S-x-Y-x-x-V/L), no inhibitory activity could be demonstrated for a transfectant having an intact Y713-based motif alone. Whether or not a given cytoplasmic tyrosine residue mediates an inhibitory signal therefore cannot be predicted entirely by sequence similarity with an ITIM consensus motif. The functional Y742-based motif matches the consensus ITIM motif, except for a charged glutamic acid in position –2 to the tyrosine. This charged amino acid could alter the properties of this motif relative to those of the consensus ITIM motif. The molecular mechanism through which the Y742-based ITIM exerts its inhibitory function will be interesting to determine, especially in view of its inability to bind SHP-1, SHP-2, and SHIP (8, 10, 13). Instead, the PIR-B Y742 phosphopeptide was found to bind a protein of approximately 120 kDa, the identity of which remains to be elucidated. This analysis suggests a further surprising feature of the PIR-B ITIM motifs in that they are able to mediate inhibition on their own and do not need to be present in tandem to mediate inhibition, as has been suggested for the ITIM in the killer Ig-like receptors (23).

While the ligands for the PIR molecules are unknown, our functional analysis indicates that PIR-B oligation can inhibit IgE antibody–mediated mast cell activation. PIR-B is thus the third ITIM-containing inhibitory receptor to be identified on murine mast cells, a trio including FcγRIIB and gp49B1 (31, 32). Multiple negative regulators may be needed for mast cells to attenuate their release of potent inflammatory mediators. In this regard, mast cells from FcγRIIB-deficient mice have been found to be very sensitive to IgG-triggered degranulation (33). Moreover, these mice exhibit exaggerated anaphylactic susceptibility as well as enhanced humoral responses. It might be predicted that PIR–B−/− deficient mice will also exhibit enhanced type I hypersensitivity. Given that human basophils and mast cells express at least four ITIM-containing inhibitory receptors, the FcγRIIB (34), the mast cell function-associated antigen (35), signal regulatory protein ζ (36) and the PIR relatives, Ig-like transcripts (also known as leukocyte Ig-like receptors, monocyte/macrophage Ig-like receptors, monocyte CDNA-18, and CD85) (3–6), mast cell modulation by these negative regulators could prove to be important in allergic disorders (for a review, see ref. 37).

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