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J Immunol published online 16 February 2015
http://www.jimmunol.org/content/early/2015/02/16/jimmunol.1400817

Supplementary Material http://www.jimmunol.org/content/suppl/2015/02/16/jimmunol.1400817.DCSupplemental

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PRL-3 Mediates the Protein Maturation of ULBP2 by Regulating the Tyrosine Phosphorylation of HSP60

Wai-Hang Leung,*1 Queenie P. Vong,*1 Wenwei Lin,† David Bouck,‡ Susanne Wendt,* Erin Sullivan,* Ying Li,* Rafijul Bari,* Taosheng Chen,‡ and Wing Leung*,‡

Many malignant cells release the NKG2D ligand ULBP2 from their cell surface to evade immunosurveillance by NK cells and CD8 T cells. Although the shedding mechanism remains unclear, various inhibitors of matrix metalloproteinases have been shown to efficiently block the release of soluble ULBP2. The clinical use of these inhibitors, however, is limited because of adverse side effects. Using high-throughput screening technique, we identified a specific inhibitor of phosphatase of regenerating liver 3 (PRL-3) that could reduce the level of soluble ULBP2 in the culture supernatant of various cancer cell lines. Inhibition or gene knockdown of PRL-3 did not reduce ULBP2 shedding, but rather suppressed posttranslational maturation of ULBP2, resulting in intracellular retention of immature ULBP2. We then found that ULBP2 was constitutively associated with heat shock protein HSP60. Complete maturation of ULBP2 required tyrosine phosphorylation of HSP60 which was mediated by PRL-3.

The Journal of Immunology, 2015, 194: 000–000.

The online version of this article contains supplemental material.

Received for publication March 31, 2014. Accepted for publication January 13, 2015.

This work was supported in part by National Institutes of Health Grant P30 CA021765, the Press On Fund, the Assisi Foundation of Memphis, and the American Lebanese Syrian Associated Charities.

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Abbreviations used in this article: Endo H, endoglycosidase H; IP, immunoprecipitation; KD, knockdown; MIC, MHC class I chain-related molecule; MMP, matrix metalloproteinase; NKG2DL, NKG2D ligand; PI-PLC, phosphatidylinositol-specific phospholipase C; PNGase F, peptide-N-glycosidase F; PRL-3, phosphatase of regenerating liver 3; PRL3-I, PRL-3 inhibitor I; siRNA, small interfering RNA; UTR, untranslated region.

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www.jimmunol.org/cgi/doi/10.4049/jimmunol.1400817

The Journal of Immunology

Published February 16, 2015, doi:10.4049/jimmunol.1400817
PRL-3 REGULATES ULBP2 MATURATION

Materials and Methods

Cell lines, Abs, and reagents

HCT116, CAL27, 293T, HT29, SW480, Hela, and DU145 cell lines were obtained from American Type Culture Collection. The protease and phosphatase inhibitor library were purchased from Enzo Life Sciences. Mouse anti-ULBP1 (AUMO2) (as primary Ab for flow cytometry) and anti-ULBP2 (BUMO1) (as capture Ab for ELISA) Ab were purchased from BAMOMAB; mouse anti-ULBP2 (IgG2a, as detection Ab for ELISA), allophycocyanin-conjugated mouse anti-ULBP2 (for flow cytometry), goat anti-ULBP2, and anti-ULBP1 (for Western blot) from R&D Systems; allophycocyanin-conjugated anti-mouse IgG Ab (as secondary Ab for flow cytometry) and HRP-conjugate goat anti-mouse IgG2a (as secondary Ab for ELISA) from Jackson ImmunoResearch Laboratories; mouse anti-ubiquitin (DM1A + DM1B) and rabbit anti-HSP60 from Abcam; HRP-conjugated anti-mouse and anti-rabbit secondary Abs, rabbit anti-CD73, tubulin (DM1A + DM1B) and rabbit anti-HSP60 from Abcam; HRP-conjugated anti-mouse IgG Ab (as secondary Ab for flow cytometry) from Jackson ImmunoResearch Laboratories; mouse anti-ULBP2 (IgG2a, as detection Ab for ELISA), Mouse anti-ULBP1 (AUMO2) (as primary Ab for flow cytometry) and anti-ULBP2, and anti-ULBP1 (for Western blot) from R&D Systems; allophycocyanin-conjugated anti-mouse IgG Ab (as secondary Ab for flow cytometry) and HRP-conjugate goat anti-mouse IgG2a (as secondary Ab for ELISA) from Jackson ImmunoResearch Laboratories; mouse anti-ubiquitin (DM1A + DM1B) and rabbit anti-HSP60 from Abcam; HRP-conjugated anti-mouse and anti-rabbit secondary Ab, rabbit anti-CD73, and rabbit anti-Na,K-ATPase from Cell Signaling; HRP-conjugated anti-goat secondary Ab from Santa Cruz; HRP-conjugated anti-phospho-tyrosine Ab (4G10 Platinum) and MMP inhibitor III from EMD Milipore; mouse anti-Flag (clone M2) Ab, DMSO, PRL-3 inhibitor I, curcumin, nonactin, staurosporine, and phosphatidylinositol-specific phospholipase C (PI-PLC) from Sigma-Aldrich. Src II and GM 6001 from Tocris Bioscience; tetratemethylbenzidine ELISA substrate solution from eBioscience; and Cell Titer-Glo luminescent cell viability assay from Promega.

Flow cytometry

Cell lines were stained with isotype control or allophycocyanin-conjugated anti-ULBP2 for 20 min at 4°C. For ULBP1 staining, cells were first incubated with anti-ULBP1 and then with allophycocyanin-conjugated anti-mouse IgG Ab. Cells were washed and analyzed on a 3D flow cytometer (Accuri). Data were analyzed using FlowJo software (Tree Star).

ULBP2 ELISA

ULBP2 shed from cell surface to the culture supernatant was detected by ELISA method, as described before (19). Briefly, 100 μl (triplicate) cultured supernatant was added into a 96-well plate precoated with 1 μg/ml mouse anti-ULBP2 (BUMO1) and incubated at 4°C overnight. The plate was washed, and then incubated with mouse anti-ULBP2 (IgG2a, 100 μl/ well at 1 μg/ml) for 1 h, followed by incubation with HRP-conjugated goat anti-mouse IgG2a for 1 h at room temperature. Tetratemethylbenzidine substrate was added, and the plate was read using Victor® plate reader (Perkin Elmer).

Quantitative real-time PCR assay

Total RNA was extracted from DMSO- or PRL3-I–treated (40 μM) tumor cells by using the RNA Clean & Concentrator (Zymo Research). cDNA was generated by using the SuperScript VILO cDNA Synthesis kit (Life Technologies) and diluted 10-fold for analysis using the 7900HT Fast Real-Time PCR System (Applied Biosystems). The data were calculated as the cycle threshold of target genes normalized to the cycle threshold of GAPDH of each sample. Primer sequences of the target genes are as follows: ULBP2 forward, 5'-AAATGTCACACGGCCTG-3'; ULBP2 reverse, 5'-TGGAGG-GGTTCCTTTGGG-3'; GAPDH forward, 5'-ATGGGGAAGTGGAG-TGC-3'; GAPDH reverse, 5'-GGGTCACTTATGGCCACAAATA-3'.

Cytotoxicity assay

NK cells were isolated from health donors by autoMACS Pro Separator using NK cell isolation kit (Miltenyi Biotec) and cultured overnight with 10 U/ml IL-2 (eBioscience). Cells were treated for 18 h with DMSO or PRL-3-I (40 μM) were subjected to BATDA release assays (Perkin Elmer), according to the manufacturer’s instructions. The results were calculated as follows:

\[
\% \text{ specific lysis} = \frac{\text{Experimental release} - \text{Spontaneous release}}{\text{Maximum release} - \text{Spontaneous release}} \times 100
\]

Immunoprecipitation

Total cell lysates from ~1 × 10⁷ cells (lysed in cell lysis buffer (50 mM Tris [pH 8.0], 10 mM EDTA, 125 mM NaCl, 1% Triton X-100, proteinase inhibitor mixture (Thermo Scientific) for 10 min at 4°C) were incubated with 1 μg Ab (anti-ULBP2, anti-HSP60, anti-phosphorysorine, or anti-Flag) for 2 h at 4°C, followed by 1-h incubation with 20 μl protein G microbeads (Miltenyi Biotec). Target proteins were separated by MultiMACS M separator (Miltenyi Biotec), according to the manufacturer’s instructions.

Western blot analysis

Protein samples prepared in NuPAGE lithium dodecyl sulfate sample buffer (Life Technologies) were run onto NuPAGE 4–12% Bis-Tris gels and then transferred onto nitrocellulose membranes using the iBlot transfer system (Life Technologies). After incubation with 5% nonfat milk in TBST (10 mM Tris [pH 8.0], 150 mM NaCl, 0.5% Tween 20) for 180 min, the membranes were incubated with Abs against ULBP2 (1:1000), HSP60 (1:2000), Flag-tag (1:2000), phosphorysorine (1:800), or tubulin (1:5000) at 4°C for 16 h. Membranes were washed three times for 15 min and incubated with a 1:2500 dilution of HRP-conjugated anti-mouse, anti-goat, or anti-rabbit Secondary Abs for 1 h (except for phosphorysorine detection because the Ab, 4G10, has already been HRP conjugated). The membranes were washed with TBST five times and developed with the Western Lightning Plus ECL system (Perkin Elmer), according to the manufacturer’s instructions.

Endoglycosidase H, peptide-N-glycosidase F, and PI-PLC digestion

A total of 20 μg total proteins from the PRL3-I–treated or untreated HCT116-ULBP2 cell lysates was digested with 1500 U endoglycosidase H (Endo H) or peptide-N-glycosidase F (PNGase F) (NEB) at 37°C for 2 h, according to the manufacturer’s instruction. For experiments with PI-PLC and PNGase F double digestion, cell lysates were first incubated with 0.2 U PI-PLC at 37°C for 1 h before the addition of PNGase F. Digested samples were subjected to Western blot analysis.

Pulse-chase experiment

HCT116-ULBP2 cells treated with DMSO or 40 μM PRL3-I for 18 h were washed once with PBS and then starved in DMEM without methionine, cysteine, and glutamine for 1 h at 37°C. A final concentration of 50 μM L-azidohomoalanine (Life Technologies), 5 mM L-cysteine (Sigma-Aldrich), and 1X GlutaMAX (Life Technologies) was added for 1-h additional incubation. Cells were washed and replaced with complete DMEM supplemented with 10% FBS. Cell lysates were collected at different time points for Click-it biotinylation, according to the manufacturer’s instructions. Biotinylated proteins were isolated using streptavidin microbeads (Miltenyi Biotec) for Western blot analysis to detect the protein synthesis of ULBP2.

Subcellular fractionation

Subcellular protein fractions (plasma membrane, organelle, and cytosol) were separated using the Minute plasma membrane isolation kit (Invent Biotechnologies), according to the manufacturer’s instructions.
**Lentiviral transduction**

Lentiviral vector was cotransfected with the ViraPower lentiviral packaging mix (Life Technologies) into 293T cells, according to the manufacturer’s manual. The culture supernatant, which contained the lentiviral particles, was collected 72 h posttransfection and concentrated by ultracentrifugation. Lentiviral titers were determined (based on GFP+ cell detection by flow cytometry) after transduction by transducing 293T cells with serial dilutions of the concentrated lentiviral culture supernatant. Stable lentiviral transduced HCT116 cell lines were established by culturing 1 × 10^6 cells with the concentrated lentiviral supernatant (multiplicity of infection = 10) containing 8 μg/ml polybrene. Positive transduced cells were isolated by GFP-based cell sorting 1 wk after transduction.

**Small interfering RNA knockdown experiments**

Small interfering RNA (siRNA)-targeting PRL-3 (D-006859-04), HSP60 (D-010600-02), and Src (D-003175-05) were purchased from ThermoFisher Scientific. Negative knockdown control siRNA (4390843) was from Life Technologies. For siRNA transfection, 2 × 10^5 cells were transfected with 60 pmol siRNA molecules using lipofectamine RNAiMAX (Life Technologies), according to the manufacturer’s instructions.

**Confocal microscopy**

A total of 2 × 10^5 HCT116-ULBP2 cells was seeded on glass slides and treated with DMSO or PRL3-I (40 μM) for 18 h. Cells were fixed with 4% paraformaldehyde and permeabilized by PBS containing 0.5% Triton X-100 and then stained with anti-ULBP2 (clone 165903 from R&D Systems), DAPI, and either anti-calnexin (clone C390 from Cell Signaling) or anti-GM130 (ab31561 from Abcam). Cellular localization of ULBP2 was analyzed by using the Nikon C2plus confocal microscope. Fluorescence images were taken using 60× 1.45NA Plan Apo objective (pinhole radius = 40 μm, 1 Airy unit) with the depth of the scanned images ~500 nm. Images were analyzed using NIS-Elements Advanced Research software.

**Triton X-114 partitioning experiment**

DMSO- or PRL3-I (40 μM)-treated HCT116-ULBP2 cells were lysed in 2% Triton X-114 in TBS (prepared from the 10% Triton X-114 stock solution from ThermoFisher Scientific). Extraction was performed by incubating the cell lysate at 4˚C for 1 h with occasional mixing. Insoluble cell debris was removed by centrifugation at 10,000 × g at 4˚C for 10 min. Supernatant was collected and incubated at 37˚C for 5 min (solution became cloudy after the incubation), and phases were separated by centrifugation at 1,000 × g at 25˚C for 10 min. The upper aqueous phase and the lower detergent phase were analyzed by Western blotting.

**PRL-3 activity assay**

HCT116 cells stably transduced with a control lentiviral vector (control) or a PRL-3 overexpression construct (HCT116-PRL-3) were treated with DMSO, 40 μM PRL3-I, or 40 μM Src In 1 h. One million cells were lysed in 80 μl cell lysis buffer (PBS–1% Triton X-100) and then diluted 20-fold with 20 mM Tris-HCl (pH 8.0). One hundred microliters of the diluted lysate was incubated with 100 μl phosphatase substrate, DiFMUP (16 μM; Life Technologies) for 30 min. The fluorescence intensity was measured using the Victor2 plate reader (Perkin Elmer). PRL-3 activity in HCT116-PRL-3 cells was calculated as follows:

\[
\% \text{ PRL3 activity} = \frac{\text{HCT116-PRL3 intensity} - \text{HCT116}}{\text{DMSO-HCT116-PRL3 intensity} - \text{DMSO-HCT116}} \times 100
\]

**Results**

**High-throughput screening identified a PRL-3 inhibitor that reduced the release of soluble ULBP2 from tumor cell lines**

We generated a lentiviral construct that contained a ULBP2-knockdown short hairpin RNA expression cassette targeting the 5' untranslated region (UTR) of human ULBP2. The vector also contained a CMV promoter-driven recombinant ULBP2 that lacked the natural 5'-UTR and was fused to two successive GFP at the N terminus (ULBP2-2GFP^N-term) (Fig. 1A). The 293T cells transduced with this construct expressed a significantly lower level of endogenous ULBP2 (determined by quantitative real-time PCR using primers detecting the 5'-UTR; data not shown), but a high level of ULBP2-2GFP^N-term on the cell surface (determined by flow cytometry and fluorescence microscopy; W.-H. Leung and W. Leung, unpublished observations). With the coexpression of two GFP molecules at the N terminus, the release of the recombinant ULBP2 from the cell surface could be detected by the measurement of fluorescence intensity in the cultured supernatant. In line with the previous studies (19), the release of ULBP2-2GFP^N-term from 293T cells could be blocked by a MMP inhibitor, MMP-I III, in a dose-dependent manner (Fig. 1B). The subsequent high-throughput screening procedure is illustrated in Fig. 1C. Culture supernatants were collected 5 d after the treatment of different inhibitors (14 μM) for fluorescence detection. The viability of the cells was determined by analyzing the production of ATP in the cells. Compounds that reduced ≥20% of ULBP2 release, but affected ≤5% of cell viability, were considered potential candidates. Among 53 protease inhibitors and 33 phosphatase inhibitors that we screened, 6 compounds fulfilled these criteria, including the broad-spectrum MMP inhibitor GM 6001 that has been reported to inhibit ULBP2 shedding (29) (Fig. 1D). Another candidate was a specific phosphatase inhibitor targeting PRL-3 (PRL3-I). Confirmatory experiments showed that this PRL3-I could reduce the release of endogenous ULBP2 from cancer cell lines HCT116 and CAL27 (Fig. 1E).

**Surface expression of ULBP2 unexpectedly decreased in cancer cell lines upon PRL3-I treatment**

Effective blockage of ULBP2 shedding by GM 6001 or MMP-I III in HCT116 cells resulted in an increase in ULBP2 surface expression (Fig. 2A). Surprisingly, the inhibition of ULBP2 released from cancer cell lines upon PRL3-I treatment was not associated with an increase, but rather a decrease in ULBP2 surface expression (Fig. 2B, Supplemental Fig. 1A, 1B), even though the mRNA level of ULBP2 was increased in PRL3-I–treated cells (Fig. 2C). These results suggested that the inhibition of ULBP2 release by PRL3-I was a result of downregulation of ULBP2 surface expression rather than blockage of shedding. The surface expression of other NKG2D ligands such as ULBP1 (Fig. 2B) or MICA (W.-H. Leung and W. Leung, unpublished observations) remained unchanged with PRL3-I treatment; thus, the specific reduction of surface ULBP2 in PRL3-I–treated HCT116 and CAL27 cells resulted in only a moderate decrease in susceptibility to NK cell cytolysis (Fig. 2D).

**PRL3-I suppressed ULBP2 protein maturation, resulting in retention in endoplasmic reticulum**

To investigate how PRL3-I affected the surface expression of ULBP2, we first performed Western blot analysis on cell lysate prepared from HCT116 cells treated with various concentrations of PRL3-I (Fig. 3A). A 37-kDa band representing the mature form of ULBP2 was detected in the lysate from HCT116 cells without PRL3-I treatment. An additional band ~32 kDa began to appear in samples treated with 5 μM PRL3-I. Increasing the concentration of PRL3-I (up to 40 μM) increased the abundance of the 32-kDa band, alongside with a gradual reduction of the 37-kDa mature ULBP2. Similar results were observed in CAL27 cells upon PRL3-I treatment. To confirm the ULBP2 identity of the 32-kDa band, we generated a stable transduced HCT116 cell line that specifically overexpressed ULBP2 (HCT116-ULBP2). In the absence of PRL3-I, a strong 37-kDa band and a faint 32-kDa band were detected in the HCT116-ULBP2 cell lysate. Consistent with the results of endogenous ULBP2, PRL3-I treatment increased the abundance of ULBP2 in the 32-kDa form in HCT116-ULBP2 cells (Fig. 3B). This particular switch in molecular size of ULBP2...
accompanied a decrease in surface expression of ULBP2 (Fig. 3C). In contrast, there was no difference in protein size and surface expression of ULBP1 in PRL3-I–treated HCT116 cells overexpressing ULBP1 (HCT116-ULBP1).

Newly synthesized ULBP2 consists of 246 aa, which has a molecular mass of 27 kDa. The protein is then posttranslationally modified by glycosylation to become a mature 37-kDa glycoprotein (30). The 32-kDa ULBP2 appeared upon PRL3-I treatment might therefore represent either an immature form of ULBP2 or a degradation product from the mature protein. We treated the cell lysates from HCT116-ULBP2 cells cultured overnight with DMSO (solvent control) or 40 μM PRL3-I with either Endo H or PNGase F. Endo H preferentially cleaves immature high mannose-containing N-linked oligosaccharides, whereas PNGase F removes all N-linked glycans regardless of their level of processing as immature or mature proteins. We found that the 32-kDa bands in both DMSO- and PRL3-I–treated cell lysates were sensitive to Endo H digestion, whereas PNGase F treatment converted both 32- and 37-kDa bands to the 27- to 28-kDa unglycosylated ULBP2 (the appearance of slightly different sizes of ULBP2 upon DMSO and PRL3-I treatment was further investigated in the following section) (Fig. 4A). Similar results were also observed in PRL3-I–treated HCT116 cell lysates (Supplemental Fig. 2). In addition, we also performed a pulse-chase experiment to track ULBP2 synthesis in HCT116-ULBP2 cells (Fig. 4B). In the absence of PRL3-I, both 32- and 37-kDa bands appeared during early ULBP2 synthesis. The abundance of the 37-kDa mature ULBP2 gradually increased, which eventually predominated over the 32-kDa immature form. Upon PRL3-I treatment, the synthesis of ULBP2 was completely halted at the immature stage. Collectively, these results suggest that the 32-kDa band corresponded to an immature form, rather than a degradation product of ULBP2, and that the complete maturation of ULBP2 required the activity of PRL-3.

We next performed subcellular fractionation of PRL3-I–treated or -untreated HCT116-ULBP2 cells for Western blot analysis. The 37-kDa mature ULBP2 located primarily in the plasma membrane fraction, whereas the 32-kDa immature form largely remained in the organelle fraction (Fig. 4C). Similarly, confocal microscopy confirmed that the majority of ULBP2 proteins expressed on the cell surface of DMSO-treated control cells (Fig. 4D). Upon PRL3-I treatment, large amounts of ULBP2 were retained in the endoplasmic reticulum (Fig. 4E). Collectively, these results suggest that PRL3-I suppresses the posttranslational maturation of ULBP2 proteins, which, in turn, affected their cell surface expression.
PRL3-I inhibited GPI anchor formation of ULBP2

It has been suggested that the level of ULBP2 surface expression could be influenced by the GPI anchor formation of the proteins (31). We generated a mutated ULBP2 construct (Fig. 5A) that had the amino acid sequence 216SSG218 substituted with 216TPV218 (ULBP2-DGPI) so that it lacked the GPI anchor binding site (Supplemental Fig. 3A). Consistent with previous findings, the surface expression of ULBP2 was lower in HCT116-ULBP2-DGPI cells compared with that in HCT116-ULBP2 cells (Fig. 5B). Importantly, we found that the overexpressed ULBP2-DGPI proteins were in the 32-kDa immature form (Fig. 5C). Treatment with PRL3-I did not further affect the protein level (Fig. 5C) and surface expression of ULBP2-DGPI (Fig. 5D). Furthermore, the ULBP2-DGPI proteins demonstrated an intracellular endoplasmic reticulum retention pattern that was the same as the wild-type ULBP2 proteins upon PRL3-I treatment (Figs. 4E, 5E).

FIGURE 2. PRL3-I reduces the surface expression of ULBP2 on tumor cells. (A) Cell surface expression of ULBP2 was analyzed by flow cytometry in HCT116 cells treated with DMSO (solvent control), 40 μM GM 6001, or MMP-1 III for 18 h. Data are representative of two independent experiments. (B) Cell surface expression of ULBP1 and ULBP2 was analyzed by flow cytometry in different cancer cell lines treated with DMSO or PRL3-I (40 μM) for 18 h. Data are representative of three independent experiments. (C) ULBP2 mRNA expression in HCT116 and CAL27 cells treated with DMSO or PRL3-I (40 μM) for 18 h was analyzed by quantitative real-time PCR analysis. Data are normalized to GAPDH mRNA levels and are presented as fold-change relative to the expression in DMSO-treated cells (n = 4). (D) NK cell cytotoxicity on HCT116 and CAL27 cells treated with DMSO or PRL3-I (40 μM) for 18 h was determined by a BATDA release assay using IL-2 (10 U/ml)-primed primary NK cells isolated from healthy donors at various E:T ratios (n = 4). *p < 0.05, **p < 0.01, ***p < 0.005, ****p < 0.0001.

FIGURE 3. Treatment with PRL3-I associated with the appearance of low-molecular mass ULBP2. (A) HCT116 cells were treated with increasing concentration (0–40 μM) of PRL3-I, whereas CAL27 cells were treated with either DMSO or 40 μM PRL3-I for 18 h. Total cell lysates were collected for Western blot analysis to detect ULBP2 expression. Data shown are representative of two independent experiments. (B) HCT116 cells overexpressing ULBP2 (HCT116-ULBP2) or ULBP1 (HCT116-ULBP1) were treated with DMSO or PRL3-I (40 μM) for 18 h. Total cell lysates were collected for Western blot analysis to detect ULBP2 or ULBP1 expression. Data shown are representative of three independent experiments. (C) HCT116-ULBP2 and HCT116-ULBP1 cells were treated with DMSO or PRL3-I (40 μM) for 18 h. Cell surface expression of ULBP2 or ULBP1 was analyzed by flow cytometry. Results are representative of three independent experiments.
To gain more insight into the correlation between GPI anchor formation of ULBP2 and the effect of PRL3-I on ULBP2 expression, we generated a lentiviral construct expressing a chimeric GFP protein that possessed the ULBP2 signal sequence at the N-terminal and the cytoplasmic tail of ULBP2 carrying the GPI anchor sequence at the C-terminal (GFP-ULBP2 210–246) (Fig. 5F). Because GFP is not a glycoprotein, its expression, protein folding, and fluorescent property are not affected by the posttranslational glycosylation. With the GPI anchor sequence of ULBP2, GFP-ULBP2210–246 was able to express on the cell surface of HCT116 cells (detected by flow cytometry using Alexa Fluor-647–conjugated anti-GFP Ab; Fig. 5G). Upon PRL3-I treatment, the surface expression of GFP-ULBP2 210–246 was greatly reduced, but the total GFP protein level was not affected (Fig. 5G). We also performed Triton X-114 phase partitioning experiments to detect the soluble immature ULBP2 proteins appeared upon PRL3-I treatment. With this method, membrane proteins and GPI-anchored proteins are partitioned into the detergent phase, whereas soluble proteins remain in the aqueous phase (32). As shown in Fig. 5H, mature ULBP2 was preferentially partitioned into the detergent phase. However, in the presence of PRL3-I, a significant amount of immature form of ULBP2 (but not the control GPI protein CD73) appeared in the aqueous phase, indicating that some of the immature ULBP2 proteins became soluble upon PRL3-I treatment.

Because we observed a molecular mass difference of PNGase F–digested ULBP2 proteins between DMSO (28 kDa)– and PRL3-I (27 kDa)–treated samples (Fig. 4A), we speculated that the slightly larger ULBP2 from DMSO control was due to the presence of GPI anchor. To verify the existence of GPI anchor in the 28-kDa deglycosylated ULBP2 proteins, we performed PNGase F digestion with or without PI-PLC treatment. Previous studies have shown that PI-PLC–treated GPI proteins migrate more slowly on SDS-PAGE (33, 34). As shown in Fig. 5I, the 28-kDa deglycosylated ULBP2 only appeared in DMSO control, but not in PRL3-I–treated sample or ULBP2-DΔGPI upon PNGase F digestion. With the PI-PLC digestion, ULBP2 proteins remain in the aqueous phase (32). As shown in Fig. 5H, mature ULBP2 was preferentially partitioned into the detergent phase. However, in the presence of PRL3-I, a significant amount of immature form of ULBP2 (but not the control GPI protein CD73) appeared in the aqueous phase, indicating that some of the immature ULBP2 proteins became soluble upon PRL3-I treatment.

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treatment, the 28-kDa band migrated more slowly on SDS-PAGE, which became ∼29 kDa. These results suggested that ULBP2 protein in DMSO- but not PRL3-I–treated cells contained GPI anchor.

We further examined the surface expression of other transmembrane (Na,K-ATPase and HLA-E) or GPI-anchored proteins (CD73) on HCT116 cells. Unlike ULBP2, none of them was affected upon PRL3-I treatment (Supplemental Fig. 3B). Collectively, we conclude that GPI anchor formation is essential for the complete maturation and surface expression of ULBP2 and that this process is specifically inhibited by PRL3-I.

PRL-3 was directly involved in protein maturation and cell surface expression of ULBP2

It has been reported that PRL3-I can inhibit other protein tyrosine phosphatases in addition to PRL-3 (35). To confirm that the suppression of PRL-3 by PRL3-I directly gave rise to the altered ULBP2 expression, we treated HCT116 cells with curcumin or PRL-3–knockdown (KD). As shown in Fig. 6A, both curcumin and PRL-3-KD treatment reduced the surface expression of ULBP2 in HCT116 cells. In line with the results of PRL3-I treatment, the reduction of PRL-3 expression by curcumin or PRL-3-KD in HCT116-ULBP2 cells was associated with predominant expression of 32-kDa ULBP2 (Fig. 6B).

Previous reports suggested that Src mediates the tyrosine phosphorylation of PRL-3, which is required for PRL-3 functions (36). Treatment of HCT116 cells overexpressing PRL-3 (HCT116-PRL-3) with a Src kinase inhibitor, Src I1, significantly reduced the phosphatase activity of PRL-3, which was comparable to the effect of PRL3-I (Fig. 6C). Moreover, Src I1 treatment reduced the surface expression of ULBP2 in HCT116.

FIGURE 5. PRL3-I suppressed cell surface expression of ULBP2 by the inhibition of GPI anchor formation. (A) Schematic representation of the overexpression construct of wild-type (ULBP2) and mutant ULBP2 that is not able to form GPI anchor (ULBP2-DGPI). (B) Flow cytometry was performed to compare the surface expression of ULBP2 between the untransduced and transduced HCT116 cells with either ULBP2 (HCT116-ULBP2) or ULBP2-DGPI (HCT116-ULBP2-DGPI) expression construct. Results are representative of three independent staining. (C) Total cell lysates from HCT116-ULBP2 and HCT116-ULBP2-DGPI cells treated with DMSO or PRL3-I for 18 h were collected for the detection of ULBP2 expression by Western blot analysis. Data shown are representative of three independent experiments. (D) The surface expression of ULBP2 in DMSO- or PRL3-I–treated HCT116-ULBP2-DGPI cells was determined by flow cytometry. Results are representative of three independent experiments. (E) HCT116-ULBP2-DGPI cells seeded on glass slides were fixed, permeabilized, and stained with anti-ULBP2, DAPI, and either anti-calnexin (endoplasmic reticulum staining) or anti-GM130 (Golgi staining) for the detection of ULBP2 localization by confocal microscopy. Results are representative of two independent experiments. Scale bars, 5 μm. (F) Schematic representation of the lentiviral construct expressing a chimeric GFP protein (GFP-ULBP2210–246) that was fused with the signal sequence (N-terminal) and the cytoplasmic tail of ULBP2 (C-terminal). (G) HCT116-GFP-ULBP2210–246 cells were treated with DMSO or 40 μM PRL3-I for 18 h. Flow cytometry was performed to detect the total GFP proteins, and the cell surface expressed GFP proteins that were stained with the Alexa Fluor-647-conjugated anti-GFP Ab. Results are representative of three independent experiments. (H) Triton X-114 partitioning experiments were performed to detect the GPI anchor formation on ULBP2 with or without PRL3-I treatment. The GPI-anchored protein CD73 was detected for the partitioning control. Results are representative of two independent experiments. (I) Total cell lysates collected from DMSO- or PRL3-I–treated HCT116-ULBP2 cells (18 h) or from HCT116-ULBP2-DGPI cells were untreated or treated with Endo H or PNGase F (with or without PI-PLC) for Western blot analysis to detect ULBP2 expression. Results are representative of two independent experiments.

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and CAL27 cells (Fig. 6D). Likewise, siRNA-mediated knockdown of Src in HCT116 cells resulted in predominant expression of 32-kDa ULBP2 (Fig. 6E) and decreased ULBP2 surface expression (Fig. 6F). Collectively, these results suggest an absolute requirement of Src-dependent PRL-3 activity in modulating the protein maturation and cell surface expression of ULBP2.

HSP60 interacted constitutively with ULBP2 and PRL-3 regulated HSP60 tyrosine phosphorylation

Recent studies have shown that PRL-3 regulates the tyrosine phosphorylation of various molecules such as ERK1/2 and integrin β1 (37). Because ULBP2 is not a tyrosine-phosphorylated protein and there is no direct protein–protein interaction between ULBP2 and PRL-3 (W.-H. Leung and W. Leung, unpublished observations), we speculated that PRL-3 controls ULBP2 maturation through the regulation of phosphorylation in other tyrosine phosphoproteins. Lysates of HCT116-ULBP2 cells treated with different concentrations of PRL3-I were subjected to immunoprecipitation (IP) with anti-ULBP2 Ab and were analyzed by Western blotting using anti-phosphotyrosine Ab (4G10). In the absence of PRL-3-I, a 60-kDa tyrosine-phosphorylated protein was coimmunoprecipitated with ULBP2. However, the intensity of the tyrosine-phosphorylated protein band gradually decreased when the concentration of PRL3-I increased (Fig. 7A). Using 4G10 for IP of cell lysates from DMSO- or PRL3-I–treated HCT116-ULBP2 cells, we observed a bias in favor of the mature ULBP2 (37 kDa) over the immature form (32 kDa) to be coimmunoprecipitated by the Ab (Fig. 7B).

The 60-kDa member of the heat shock protein family, HSP60, has been known to act as a chaperone to regulate posttranslational modification (38). It has also been shown that tyrosine phosphorylation is required for HSP60 to be expressed on cell membrane (39). To examine whether the 60-kDa tyrosine phosphoprotein was HSP60, we performed IP of both ULBP2 and HSP60 from the HCT116-ULBP2 cell lysates. As shown in Fig. 7C, anti-ULBP2 Ab could pull down HSP60, whereas anti-HSP60 Ab coimmunoprecipitated both HSP60 and ULBP2. In both cases, the HSP60 bands were colocalized with the 60-kDa tyrosine phosphoproteins.
on the immunoblots and that PRL3-I treatment consistently reduced the intensity of the 60-kDa tyrosine phosphoprotein, but did not affect the co-IP of HSP60 and ULBP2 (both mature 37-kDa and immature 32-kDa forms). It has been suggested that HSP60 can be tyrosine phosphorylated at positions 227 and 243 (40, 41). Accordingly, we generated lentiviral constructs expressing Flag-tagged HSP60 with or without the substitution of tyrosine with phenylalanine at both positions 227 and 243 (Fig. 7D). As shown in Fig. 7E, the level of tyrosine phosphorylation was lower in the HSP60 mutant (F-HSP60-YY-mu) compared with that in the wild-type protein (F-HSP60). PRL3-I treatment caused a reduction (∼2.8-fold) of phosphorylation intensity on F-HSP60, but exerted minimal effect on F-HSP60-YY-mu. These results suggested that the tyrosine phosphorylation of HSP60 was regulated by PRL-3.

Protein maturation and cell surface expression of ULBP2 required tyrosine-phosphorylated HSP60

We next investigated whether tyrosine phosphorylation of HSP60 is required for the maturation and surface expression of ULBP2. We transduced HCT116-ULBP2 cells with F-HSP60 or F-HSP60-YY-mu. Cell lysates were subjected to IP with anti–Flag-tag Ab for Western blot analysis. A much higher amount of mature ULBP2 was coimmunoprecipitated with F-HSP60 than with F-HSP60-YY-mu (Fig. 8A). We further validated the role of HSP60 in ULBP2 maturation by treating HCT116 cells with nonactin or a siRNA-specific targeting HSP60 (HSP60-KD). Nonactin is an ionophore that blocks the mitochondrial import and consequently inhibits the maturation of HSP60 (42). Consistent with previous studies, nonactin treatment inhibited the surface expression of HSP60 in HCT116 cells (Fig. 8B). Tyrosine phosphorylation of HSP60 was also reduced upon nonactin treatment (Fig. 8C). As expected as a consequence of reduced HSP60 phosphorylation, the maturation (Fig. 8D), together with the surface expression (Fig. 8E), of ULBP2 was inhibited by nonactin treatment. Similar results were observed by direct HSP60-KD (Fig. 8F, 8G). However, neither knockdown nor overexpression of HSP60 affected the mRNA level of ULBP2 (data not shown). Also, there was no change in the surface expression of ULBP2 in HSP60-overexpressing HCT116 cells (Supplemental Fig. 4), suggesting that the gene expression of ULBP2 was not affected by the expression level of HSP60. Collectively, these data suggest that the association with tyrosine-phosphorylated HSP60 is essential for the protein maturation and surface expression of ULBP2.

Discussion

This study reveals a novel role of PRL-3 in regulating the post-translational maturation of ULBP2. Either decreased expression or functional inhibition of PRL-3 reduced both the soluble release

FIGURE 7. ULBP2 associated with HSP60, and the tyrosine phosphorylation of HSP60 was regulated by PRL-3. (A) HCT116-ULBP2 cells were treated with increasing concentration of PRL3-I for 18 h. Total cell lysates were collected for IP of ULBP2. Western blot analysis was performed to detect tyrosine-phosphorylated proteins in the immunoprecipitated fractions. Data shown are representative of three independent experiments. (B) Total cell lysates from HCT116-ULBP2 cells treated 18 h with DMSO or PRL3-I (40 μM) were collected for IP of tyrosine-phosphorylated proteins or ULBP2. Western blot analysis was performed to detect ULBP2 in the immunoprecipitated fractions. Results are representative of three independent experiments. (C) Reciprocal IP of ULBP2 and HSP60 was performed from the total cell lysates of HCT116-ULBP2 cells treated with DMSO or PRL3-I for 18 h. Western blot analysis was performed to detect tyrosine-phosphorylated proteins, HSP60 and ULBP2. Results are representative of three independent experiments. (D) Schematic representation of the overexpression construct of the Flag-tagged wild-type (F-HSP60) or the tyrosine phosphorylation mutant HSP60 (F-HSP60-YY-mu). (E) HCT116 cells stably transduced with either of these constructs were treated with DMSO or PRL3-I for 18 h. The Flag-tagged HSP60 proteins were immunoprecipitated for the detection of their tyrosine phosphorylation status. Results are representative of two independent experiments.
and the cell surface expression of ULBP2. Although there was no significant reduction in the total protein level of ULBP2, inactivation of PRL-3 significantly affected the glycosylation and GPI anchor formation of the protein, rendering the production of an immature form of ULBP2 that did not favor cell surface expression, which, in turn, reduced the soluble release of ULBP2.

Although PRL-3 possesses tyrosine phosphatase activity, inhibition of PRL-3 greatly decreased, rather than increased, the tyrosine phosphorylation of HSP60. In fact, several studies have shown that PRL-3–expressing cells exhibit a pronounced increase in protein tyrosine phosphorylation (43). Src kinase has been shown to regulate PRL-3 activity (36). However, the tyrosine phosphorylation of Src is indirectly dependent on PRL-3 through its downregulation of Csk, a negative regulator of Src (44). Little is known about the mechanism controlling the tyrosine phosphorylation of HSP60. In this study, we showed that treatment with PRL3-I or nonactin significantly reduced the level of tyrosine phosphorylation of HSP60 and that both of the treatments gave rise to a very similar alteration in the expression of ULBP2. Whereas nonactin is well known to affect the mitochondrial transportation and maturation of HSP60 (42), it is possible that PRL-3 may directly regulate the tyrosine phosphorylation mechanism of HSP60 or indirectly through the regulation of HSP60 maturation process. Further investigation is necessary to understand the regulatory role of PRL-3 in the tyrosine phosphorylation of HSP60.

Although we have demonstrated an absolute requirement of PRL-3 to produce the 37-kDa complete mature form of ULBP2, treatment with PRL3-I in either HCT116 or HCT116-ULBP2 cells could never completely inhibit the surface expression of the endogenous or the overexpressed ULBP2 proteins, respectively. Moreover, overexpression of ULBP2-ΔGPI in HCT116 cells could also moderately enhance the surface expression of ULBP2 (Fig. SB), suggesting that the immature form of ULBP2 can somehow inefficiently express on the cell surface in the absence of a GPI anchor. These results are consistent with a previous report in which the...
The authors have no financial conflicts of interest.

Acknowledgments
We thank Barbara Rooney, James Houston, and Martha Holladay for assistance with flow cytometry and cell sorting and Jimmy Cui for high-throughput library screening.

Disclosures
The authors have no financial conflicts of interest.

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