A 60 kDa prolactin variant secreted by cervical cancer cells modulates apoptosis and cytokine production

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Abstract. Prolactin (PRL) is associated with different types of cancer, such as cervical cancer. Recombinant PRL has anti-apoptotic effect on cervical cancer cells, and it can also induce cytokine production on macrophages. A 60 kDa variant of PRL is produced by cervical cancer cells. The aim of the present study was to evaluate this variant's bioactivity, to test its effect on cervical cancer cell apoptosis, and to assess its ability to induce cytokine production on THP-1 macrophages. First, 60 kDa PRL was isolated and used to stimulate Nb2 cells. Later, apoptosis was measured after exposure to 60 kDa PRL. Finally, cytokines were measured on THP-1 stimulated supernatants. Our results show that 60 kDa PRL increased Nb2 cell proliferation. Apoptosis was decreased after stimuli with 60 kDa PRL in cervical cancer cells. IL-1β and TNF-α are produced by THP-1 macrophages after stimuli. These results suggest that 60 kDa PRL produced by cervical cancer cells is able to reduce apoptosis in HeLa, SiHa and C-33A cells and induce IL-1β and TNF-α production by THP-1 macrophages.

Introduction

Prolactin (PRL) is a peptidic hormone that has been shown to play multiple biological functions. In addition to its clear role in lactation, PRL participates in some cellular processes such as proliferation, growth and differentiation (1). PRL can modulate the immune system, being involved in alterations of both cellular and humoral responses (2,3). For instance, PRL is able to influence different immunological processes such as proliferation (4), antigen presentation (5) and immunoglobulin production (6). Human PRL, besides being expressed in the pituitary gland, is independently and differentially expressed by different tissues such as breast, prostate, endometrium and immune cells (7).

Several studies have described the modulating effects of PRL involving disturbances in chronic inflammatory processes such as autoimmune diseases (8). Low doses of PRL are able to induce the production of proinflammatory cytokines such as TNF-α, IL-1β, IFN-γ in murine peritoneal macrophages; however, higher doses of PRL (1,000 ng/ml) induced IL-10 synthesis with significant inhibition of proinflammatory cytokine production in the same cells (9,10). One study showed that proinflammatory cytokines induced in vitro PRLR expression in pulmonary fibroblasts (11). Increased PRL levels have been shown in autoimmune diseases, such as systemic lupus erythematosus (SLE), rheumatoid arthritis (RA), systemic sclerosis and Sjögren-sindrome (12-14). Proinflammatory cytokines have the ability to induce pituitary PRL secretion (15), which in turn may contribute to circulating pool of the hormone. Several studies have focused on the blockade of proinflammatory cytokines to prevent autoimmune diseases in murine models (16-18), which may be a possible therapeutic strategy.

The PRL protein can undergo post-translational modifications that include glycosylation, phosphorylation, polymerization and proteolytic cleavage, which can influence its biological activities (19). Pituitary PRL is a 23 kDa polypeptidic hormone; however, other forms of PRL called

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macrolactam (over 100 kDa) and big PRL (40-60 kDa) have been identified in human serum (19,20). The 60 kDa PRL corresponds to the big PRL form, which is a dimer that may be an aggregate of glycosylated subunits (25 kDa) (21), and this modification may be involved in mechanisms of biosynthesis, secretion, and clearance of the hormone (19).

There is also evidence of the association between PRL/PRLR and tumorigenesis in clinical samples. A previous study of our research group showed the relation of high PRLR expression and malignant laryngeal tumors (22).

In a previous study we provided evidence that the cell lines and tissues of cervical cancer synthesize a 60 kDa PRL variant unlike the non-tumorigenic keratinocytes (HaCaT) (23,24). This PRL variant has been previously identified in peripheral blood mononuclear cells (PBMC) from patients with SLE and THP-1 monocytes (24,25).

The diverse activities of PRL are mediated by PRLR, and this complex may activate several signaling pathways including Jak-STAT (26,27), MAPK (28) and PI3K (29,30). However, in a previous study, our group reported that STAT3 is an important transcription factor for recombinant hPRL (rec-hPRL) to carry its effects (31).

There is an association between PRL and its receptor's overexpression with the development of different types of cancer such as breast, laryngeal, prostate, colon and cervical cancer (22,23,32-36). The role of circulating PRL in tumorigenesis, mainly in breast cancer, has been a topic of debate for more than 20 years since various epidemiological studies were unable to reach unified conclusions on the correlation between circulating PRL levels and risk for cancer (37,38). However, there is no evidence that this 60 kDa PRL, which may be acting in an autocrine manner, may cooperate with the development of cancer.

Cervical cancer is still a major public health problem and the third most common type of cancer in women worldwide (39). Persistent high-risk human papillomavirus infection is the main risk factor for cervical cancer; however, there are other important cofactors for this outcome. Previous results of our group show that a long PRLR isoform is mainly expressed in cervical cancer tissues (40), and its overexpression is associated to cell survival by inhibition of apoptosis (23). Moreover, we found the presence of a 60 kDa PRL isoform that is present in cervical cancer cells (23).

There are no studies that evaluate the biological effect of PRL produced by tumor cells and the different responses due to the existence of different isoforms. The aim of the present study was to obtain the 60 kDa PRL produced by cervical cancer cells and to evaluate some aspects of its functionality. We tested whether the 60 kDa PRL is bioactive in Nb2 cells and its effect on apoptosis in cervical cancer cells. In addition, we assessed its impact on the induction of proinflammatory cytokines in THP-1 macrophages.

Materials and methods

Reagents. Rec-hPRL (L-4021) and 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium reagent (MTT) were obtained from Sigma-Aldrich® (St. Louis, MO, USA). Polyvinylidene difluoride (PVDF) membranes, enhanced chemiluminescence (ECL), and western blotting detection kit (WBKLS0500) were purchased from Merck Millipore (EMD Millipore Corporation Billerica, MA, USA) as well as Amicon® Ultra 0.5 ml centrifugal filters. Monoclonal antibody anti-PRL (E-9) sc-48383 was obtained from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA, USA). Dulbecco's modified Eagle's medium (DMEM), DMEM advanced, charcoal stripped fetal bovine serum (FBS) and antibiotic/antimicotic were purchased from Gibco Life Technologies (Carlsbad, CA, USA). Polyclonal antibodies anti-pSTAT3 (Ser 727), sc-13564, anti-bcl-xl (H-5) sc-8392 and anti-bcl-2 (C-2) sc-7382 were obtained from Santa Cruz Biotechnology. Polyclonal antibody anti-survivin AF886 was purchased from R&D Systems (R&D Systems, Inc., Minneapolis, MN, USA). Inhibitor for JAK/STAT signaling pathway α-cyano-(3,4-dihydroxy)-N-benzylcinammide (AG490) was dissolved in dimethyl sulfoxide (DMSO) and stored at -20°C as recommended by the manufacturer. Phorbol 12-myristate 13-acetate (PMA) was obtained by Sigma (P8139). μMACSTM Protein G MicroBeadsMultiMACSTM Protein G kit was provided by MiltenyiBiotec. Finally, multiple cytokine magnetic bead MIHCYTOMAG-60K-05 kit was manufactured by MILLIPLEX® MAP.

Cell culture. Cervical cancer derived cells (HeLa, SiHa and C-33A), non-tumorigenic immortalized keratinocytes (HaCaT), as a negative control, THP-1 monocytic cell line, and Nb2 rat lymphoma-derived cells were used for stimulation with PRL assays. All the cell lines were obtained from American Type Culture Collection (ATCC; University Boulevard Manassas, VA, USA) and were grown with DMEM, DMEM advanced or RPMI-1640 medium supplemented with 5% of charcoal stripped FBS, penicillin G, streptomycin and amphotericin B.

PMA was used to differentiate THP-1 monocytes into macrophages, and its concentration was 200 nM for 72 h.

Cells were cultured in a jacket-water incubator at 37°C with an atmosphere of 5% CO₂. The cells were grown to reach 80% confluence, thus they could be used for assays.

Western blotting. Protein (40 µg) from supernatants of cervical cancer cells were mixed with loading buffer and denatured at 95°C for 5 min. Afterwards, they were loaded on 10% SDS polyacrylamide gels to be resolved. Protein transference was carried out in PVDF membranes (Bio-Rad Laboratories, Hercules, CA, USA). Blocking solution was prepared with 5% of Blotting Grade Blocker (Bio-Rad Laboratories), and membranes were incubated in this solution for 2 h. Solutions with primary antibodies were prepared at a dilution of 1:500, and membranes were kept overnight followed by the secondary antibody solutions (diluted 1:5,000). The process was developed with a chemiluminescence system (Immobilon, Merck Millipore). MicroChemi 6.0 was also used to reveal membranes and measure optical density.

Immunoprecipitation with magnetic beads. Immunoprecipitation was carried out to isolate the 60 kDa isoform of PRL. Supernatants of different cervical cancer cell cultures were tested by western blotting to evaluate whether the isoform was present. Positive samples (250 µl) were mixed with...
10 µl of anti-PRL and 50 µl of microbeads using µMACSTM Protein G MicroBeadsMultiMACSTM Protein G kit. The mixture was kept on ice for 30 min, and then it was transferred into magnetic columns followed by four rinse steps and a final one to detach the protein with an acid glycine solution. After testing the correct PRL isolation, the protein was quantified to proceed with stimuli.

**Molecular weight cut-off filtration.** The PRL isolated from cervical cancer cell supernatants was filtered to purify the 60 kDa isoform using Amicon® Ultra 0.5 ml centrifugal filters setting up a molecular weight cut-off at 50 kDa. Filtration conditions were 14,000 x g for 30 min for concentration spin and 1,000 x g for 2 min for reverse spin.

**Quantification of 60 kDa PRL.** Lowry method was used to measure the isolated and purified 60 kDa PRL, having a bovine serum albumin standard curve and reading samples by triplicate. The plates were read in an iMark microplate reader Bio-Rad version 6.1.

**Silver nitrate staining.** Polyacrylamide gels (10%) were used to determine the presence of PRL after isolation assays and to determine the correct 60 kDa isoform isolation after molecular weight cut-off filtration. After electrophoretic running, a silver nitrate staining was performed.

**Nh2 cell proliferation assay.** Nh2 cells were cultured with complete medium (10% of horse serum and 10% of FBS) and 24 h before the assay cells were transferred to maintenance medium (1% FBS). Finally, assay medium (0.1% FBS) was used to carry out the test. Cells were grown for 60 h with no stimulus, rec-hPRL (200 ng/ml), or 60 kDa PRL (200 ng/ml). MTT (5 mg/ml) was added at a proportion of one tenth of final medium volume and incubated for 4 h, according to the protocol of the manufacturer. Blue formazan crystals were solubilized with acidified isopropanol, and absorbances were read at a wavelength of 570 nm in an Epoch Microplate Spectrophotometer (BioTek Instruments, Inc., Winooski, VT, USA).

**Apoptotic assay (TUNEL assay).** The kit APO-BrdU (Invitrogen, Carlsbad, CA, USA) was used to carry out TUNEL assay. Cells were grown in 8-well chamber slides seeded with 5x10^4 cells/well, and treated with etoposide alone or in combination with the 60 kDa PRL for 24 h and incubated at 37°C. The slides were washed with PBS and fixed with 4% paraformaldehyde for 30 min at room temperature. Fixed cells were washed and permeabilized using 0.2% Tween-20 for 10 min and then incubated with terminal deoxynucleotidyl transferase and BrdU for 1 h at 37°C. After rinsing with PBS, cells were treated with Alexa Fluor 488 dye-labeled anti-BrdU antibody at 37°C for 30 min and mounted with a glass coverslip. Staining of DNA fragmentation was observed with ultraviolet fluorescent microscope (Carl Zeiss, Jena, Germany) counting at least 200 cells/well. Semi-automatic quantification was performed using the cell counter function of Image-pro Plus 6.0 software. The color intensity of the positive objects (green) was manually preset for each pattern (pixel/pixel) based on the hue-saturation-intensity (HSI) histogram. The final option of the ‘count/size’ command allows to obtain the mean optical densities of 5 fields/sample.

**Human cytokine magnetic bead panel assay.** Cytokines secreted by THP-1 cells differentiated into macrophages were quantified using a multiple cytokine magnetic bead kit (MILLIPLEX® MAP, MHCYTOMAG-60K-05; Millipore) following the instructions of the manufacturer. This kit was targeted to TNF-α, IL-1β, IL-6, IL-12 and IL-10.

**Statistical analysis.** Data were analyzed using (GraphPad Prism software (GraphPad version 6.01). Significant effects were determined using ANOVA. Statistically significant differences were considered for P-values <0.05.

**Results**

**Isolation and purification of 60 kDa PRL.** After demonstrating the presence of 60 and 80 kDa isoforms of PRL in total protein extracts from cervical cancer cell lines HeLa, SiHa and C-33A (23), it was decided to evaluate whether both isoforms were also present in these cell culture supernatants. The 60 kDa variant of PRL was present in all of the 18 samples of cervical cancer cell supernatants evaluated; in contrast, the 80 kDa variant was absent (Fig. 1A). Additionally, the supernatants of non-tumorigenic immortalized keratinocytes (HaCaT) were also evaluated and the 60 kDa variant was not found (data not shown). Hence, compared with cervical cancer cell lines, HaCaT does not produce the 60 kDa PRL variant, and do not express long isoforms of PRLR (23).

Due to its absence in HaCaT cell line, the PRL isoform of 60 kDa was isolated only from cervical cancer cells (particularly from HeLa) and was collected using magnetic bead immunoprecipitation assays. After a second elution, PRL was not recovered, which means that one rinse was enough to recover the total immunoprecipitated PRL (Fig. 1B).

To identify the optimal conditions for the best purification performance, different volumes of the supernatants (250, 500 and 1,000 µl) and the antibody (1 and 2 µg) were assayed. The highest amount of protein obtained was accomplished with 250 µl of supernatant (Fig. 1B) and 2 µg of antibody (Fig. 1C).

During the immunoprecipitation assays, positive and negative controls were used to test the correct PRL isolation. In Fig. 1D it is observed that when using PBS buffer (negative control) there are no bands present, which means no protein was recovered. In contrast, when using rec-hPRL (positive control), we observed a band that shows immunoprecipitation is carried out correctly.

To confirm the correct isolation of PRL and exclude the presence of other interfering proteins, a silver staining assay in polyacrylamide gels was performed, and we detected the presence of PRL isoform of 60 kDa and other lower molecular weight bands, which may be degraded proteins (Fig. 1E). These fragments were eliminated by molecular weight cut-off filtration and the unique presence of the 60 kDa band was observed by a silver staining assay in polyacrylamide gels (Fig. 1F).

**Bioactivity of 60 kDa PRL.** Nb2 cell proliferation assay was used to determine whether the autocrine 60 kDa PRL is
bioactive. Nb2 cell proliferation assay is the gold standard to evaluate PRL bioactivity, based on the studies carried out by Noble et al. In these studies, they used a broad panel of hormone stimuli and identified that the only ones that induced proliferation on the Nb2 cell line were the members of the PRL family, particularly PRL, whose effects were observed to be in a more sensitive manner (even to 1 ng/ml). This is the reason why we decided to use this model to evaluate 60 kDa PRL bioactivity (41,42).

Nb2 cells were stimulated either with rec-hPRL or 60 kDa PRL (200 ng/ml for 60 h), and then compared to non-stimulated cells. The concentration of hormones and time of stimulus had been standardized in previous studies published by our group (24).

As observed in Fig. 2, 60 kDa PRL strongly induced proliferation in Nb2 cells almost in a 2-fold basis and this behavior is similar to that observed with rec-hPRL.

Antiapoptotic effect of 60 kDa PRL in cervical cancer cells. Our research group previously showed that rec-hPRL has an important effect on cervical cancer cell survival by inhibition of apoptosis (23) and the optimal concentration of rec-hPRL and etoposide was tested in previous studies for apoptosis assays (23,31). This is the reason to evaluate whether the 60 kDa PRL has the same capacity. The effect of the 60 kDa PRL on a model of apoptosis induced by etoposide in cervical cancer cells was analyzed. Treatment with etoposide (30 µg/ml) augmented the number of cells with fragmented DNA in all cell lines after 24 h of incubation, as expected. 60 kDa PRL (200 ng/ml) costimulus significantly decreased the number of cells with DNA fragmentation. As observed in optical analysis, HeLa, SiHa and C33A reduced its apoptotic cells when they were stimulated with 60 kDa PRL (from 698 to 248.6 for HeLa, from 642 to 302.4 for SiHa, and from 695 to 407.6 for C33A; Fig. 3A). Representative images of immunofluorescence from TUNEL assays in HeLa cell line are shown (Fig. 3B).
Since rec-hPRL is able to induce phosphorylation of STAT3, and this pathway is required for antiapoptotic effects of PRL in cervical cancer cells (31), we decided to evaluate whether autocrine 60 kDa PRL has the same capacity. The 60 kDa PRL augmented STAT3 phosphorylation in HeLa, SiHa and C33A; however, in HaCaT cells was not observed (Fig. 3C).

The antiapoptotic effect the autocrine 60 kDa PRL exerted was abolished when Jak-STAT signaling pathway was inhibited with AG490 inhibitor (Fig. 3A and B). This behavior was similar on the three cervical cancer cell lines used and similar to that observed in previous studies of our group analyzing apoptosis of cervical cancer cells and STAT3 activation in response to rec-hPRL (31).

Since a similar behavior was observed regarding the bioactivity and the antiapoptotic effect exerted by 60 kDa PRL compared to rPRL in cervical cancer cells, we decided to evaluate how the expression of antiapoptotic proteins was modulated.

In HeLa and C-33A cells, the expression of Bcl-xL and Survivin slightly increased after stimuli with 60 kDa PRL; as expected protein levels decreased when STAT3 pathway is blocked. For Bcl-2 these changes were not observed. However, in SiHa the expression of these antiapoptotic proteins were not modified (Fig. 3D).

**Figure 3.** Effect of 60 kDa PRL on apoptosis induced by etoposide in cervical cancer cells. (A) HeLa, SiHa and C-33A were stimulated with 60 kDa PRL for 24 h. Apoptosis was measured by TUNEL assays and optical analysis was performed with the software Image-pro Plus 6.0 (**P<0.01, *P<0.05). (B) Representative images of immunostaining from TUNEL assays in HeLa cells, apoptotic cells (green B, D, F, H) and DAPI staining (blue A, C, E, G) are shown. (C) Phosphorylation of STAT3 is shown after 30 min of stimulus with 60 kDa PRL in HeLa, SiHa and C33A. (D) Expression of antiapoptotic proteins after stimuli with 60 kDa PRL for 48 alone or in combination with AG490 in HeLa, SiHa and C-33A cells. 1, not stimulus; 2, with 60 kDa PRL; 3, with 60 kDa PRL plus AG490.

The 60 kDa PRL induces TNF-α and IL-1β production in THP-1 macrophages. The THP-1 monocytes were differentiated into macrophages. After 48 h of stimulation with the 60 kDa PRL (200 and 300 ng/ml), the concentration of cytokines on macrophage supernatants was measured by MILLIPLEX® MAP, MIHCYTOMAG-60K-05 kit (*P<0.01, **P<0.001).

**Figure 4.** Proinflammatory cytokine production by THP-1 macrophages in response to 60 kDa PRL. (A) TNFα and (B) IL-1β production by THP-1 macrophages stimulated with 200 and 300 ng/ml of 60 kDa PRL for 48 h was measured using MILLIPLEX® MAP, MIHCYTOMAG-60K-05 kit (*P<0.01, **P<0.001).
and 215.40 pg/ml, respectively) and IL-1β (10.08 and 10.39 pg/ml, respectively) (Fig. 4). The production of IL-12, IL-6 and IL-10 was not detected under the conditions set for this assays.

**Discussion**

Prolactin (PRL) activates signaling pathways that regulate cell proliferation, migration, differentiation and apoptosis; therefore, it has been implicated in the etiology and progression of cancer (1,37,43).

Even though pituitary gland is the main source of this hormone, there are several cells and tissues that can produce it such as brain, decidua, spleen, adipocytes and breast cancer cells among others. In the same context, PRLR is also expressed, which suggests an autocrine/paracrine mechanism (7).

However, opposite effects have been found on prostate cancer cell lines: rec-hPRL, induces apoptosis in LNCaP cells, but not in PC3 cells (44). This demonstrates that PRL can modulate different actions depending on the type of tumor.

Previous studies of our group demonstrated that rec-hPRL induces apoptosis inhibition in cervical cancer cells (23). Similar effects were observed in ovarian carcinoma, where the rec-hPRL did not affect proliferation, but it decreased apoptosis (45).

The molecular heterogeneity of PRL has been previously described (46), and we have provided evidence that the cell lines and tissues of cervical cancer synthesize a 60 kDa PRL variant, unlike the non-tumorigenic keratinocytes (HaCaT) that do not express this isoform (23,40). This PRL variant has been previously detected in PBMCs from patients with SLE and THP1 monocytes (25,32).

Consequently, we hypothesize that the expression of an autocrine/paracrine loop of PRL may play an important role in the tumoral microenvironment in cervical cancer that can lead to changes in cell proliferation, apoptosis and immune response, among other functions.

PRL is a pleiotropic hormone whose effects can affect cellular processes that may favor the development of cancer. Once the presence of PRL of 60 kDa was identified in the supernatant of the 3 cell lines derived from cervical cancer, we considered it may be important to determine whether the 60 kDa PRL was bioactive and whether it was performing some functions in these cells.

After the isolation of 60 kDa PRL from supernatants of HeLa cells, its bioactivity was tested. In the present study, a high proliferative bioactivity of the PRL of 60 kDa was demonstrated on Nb2 cells. These results are in concordance with the activity of PRL derived from PBMCs of SLE patients in the same cells (25).

Unlike the study of Larrea et al, in which stimuli were performed with the complete supernatant from PBMCs (25), in the present study purified 60 kDa PRL from cervical cancer cell supernatants was used to avoid masked effects of other molecules contained in the culture medium.

To prevent any interference due to hormones present in the FBS, charcoal stripped serum was used and; moreover, we corroborated that there was no presence of PRL or the levels are so low they cannot be detected.

Subsequently, we determined that 60 kDa PRL decreases apoptosis in HeLa, SiHa and C-33A cells, and it is able to phosphorylate STAT3 in a similar way to that observed in response rec-hPRL (31).

The high STAT3 expression has been related to many types of cancer such as prostate (47), breast (48,49), skin (50,51) and gliomas (52,53). In a recent study of Shukla et al, a positive correlation between STAT3 and increased E6/E7 expression as well as a diminished p53/pRB were shown, which opens the possibility to focus on therapeutic targets blocking STAT3. In the latter, curcumin is proposed as a potential candidate (54). In the same regard, there are other molecules proposed as STAT3 blockers such as SC99, proved in an antymyeloma model with ability to induce apoptosis in a selective STAT3 downregulation manner (55); however, on cervical cancer there is only one study involving an in vitro model where IL-37 inhibits STAT3 and supressed proliferation of cervical cancer cells (56).

The latter effects may be carried out by expression of antiapoptotic genes, such as bcl-2, bcl-xl and survivin. In preliminary experiments we observed that in HeLa cells there was an increase of Bcl-xL and survivin in response to etoposide, and these proteins showed a decrease when 60 kDa PRL was added. Whereas in SiHa cells, Bcl-xl and survivin do not show this tendency. These results may be explained since it is well known that antiapoptotic proteins are redundant in their functions (57-59), and this is why it may occur that one or another protein is expressed and in some cases both; however, more experiments must be performed in order to conclude with more certainty.

It may be interesting to use siRNAs to block the translation of 60 kDa PRL, however, it is necessary to characterize first the mRNA sequence to design a specific probe to carry out this assay. Nevertheless, this is a perspective our group is eager to investigate in future studies.

Regarding the immune system in cervical cancer, it is known that in early stages of precancerous cervical lesions there is a proinflammatory cytokine profile in the cervical microenvironment; however, in late phases, or cancer, this profile is shifted to an anti-inflammatory one, characterized by immunosuppressor cytokines (60,61).

One limitation of the present study was not blocking the 60 kDa PRL. However, it is necessary to characterize first the sequence of the protein, in order to use a specific antibody to directly inhibit 60 kDa PRL. This is an ongoing project we are currently working on.

We demonstrate that the 60 kDa PRL has the capacity to induce production of proinflammatory cytokines (IL-1β and TNF-α) in THP-1 macrophages at the concentration used. However, no IL-6, IL-12 or IL-10 production was detected. Previous studies show that rec-hPRL induces expression of TNF-α, IFN-γ, IL-12 and IL-1β in THP-1 macrophages using the concentration 100 ng/ml, and, as for the case of IL-10, its induction requires a higher concentration (1,000 ng/ml) (9,10). Further analysis must be performed to evaluate the complete immune role of the 60 kDa PRL on THP-1 macrophages or other cells.

In conclusion, we have shown that cervical cancer cells synthesize and secrete a 60 kDa PRL isoform that can inhibit apoptosis, and it can activate STAT3. In addition, this PRL
form induces IL-1β and TNF-α production in THP-1 macrophages. It may be interesting to evaluate a broader panel of cytokines to establish PRL autocrine/paracrine effects on the inflammatory response in cervical cancer.

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Competing interests

The authors declare that they have no competing interests.

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