Tumor Necrosis Factor-α Induced Cellular Stress on Trophoblastic Cells: NF-κB Signaling Could be a Potential Therapeutic Target

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ABSTRACT

Objective: The development of the human placenta depends on proliferation and differentiation of trophoblastic cells. Deficiencies in trophoblastic functions are known to have a critical role in the progression of placental pathologies such as preeclampsia. Therefore, in this research, it was aimed to evaluate the responses of trophoblastic cells to tumor necrosis factor-α (TNF-α) mediated cellular stress.

Materials and Methods: In this study, the cellular stress model was set up by treating JAR cells with 100ng/ml TNF-α for 1, 6, 12 and 24 hour long periods. In this model, the effects of TNF-α on the proliferation capacity and apoptotic activity of JAR trophoblastic cells were investigated by immunocytochemistry. The nuclear and total expression levels of nuclear factor-κB (NF-κB) was evaluated with immunocytochemistry and Western blot, respectively.

Results: It was shown that 100 ng/ml TNF-α treated cells had a reduced proliferative capacity and increased apoptotic activity by immunocytochemical staining of PCNA and caspase-8 proteins respectively. In this respect, the NF-κB signaling pathway plays a critical role in TNF-α induced processes. So that, it was shown that the TNF-α treated group had increased nuclear and total NF-κB expressions compared to the untreated one.

Conclusion: Our findings showed that TNF-α has a significant role as a cellular stress source in JAR cells. TNF-α stimulated cellular response could be defined as decreased proliferative capacity, increased apoptotic activity and NF-κB signaling in JAR syncytiotrophoblastic cell lines. Therefore, investigation of TNF-α related cellular responses especially NF-κB signaling is further required for the understanding of the mechanism of placental pathologies which is crucial for the development of therapeutic approaches.

Keywords: Cellular stress, NF-κB signaling pathway, placenta, TNF-α, trophoblastic cells

INTRODUCTION

The development of the human placenta depends on the proliferation and differentiation of trophoblastic cells. In this respect, dysfunction of trophoblasts plays an important role in the development of placental pathologies (1). Therefore, an aberrant development and differentiation of the villous syncytiotrophoblast risks the integrity of the placental barrier and causes the release of necrotic and apoptotic trophoblast fragments (2). Histological evidences suggest a role for trophoblasts in remodeling of the uterine spiral arteries. The disruption of trophoblastic invasion and incomplete remodeling result in reduction of uteroplacental perfusion, which in turn could cause ischemia of the placenta. The alterations resulted from ischemic placenta lead to increased production of oxidative stress and stimulation of proinflammatory cytokine secretion. In this connection, it was found that production of proinflammatory cytokines such as tumor necrosis fac-
tor alpha (TNF-α) was increased in placent al pathologies including preeclampsia (3,4). TNF-α regulates the expression of genes associated with inflammation, cell survival, proliferation and differentiation mainly through the activation of the nuclear factor κB (NF-κB) signaling pathway (5,6). TNF-α is associated with apoptotic cell death via two distinct caspase-8 activation pathways. One of the pathways is regulated by cIAP1 (cellular inhibitor of apoptosis protein1) and cIAP2, the two of which join to form the signaling complex referred to as complex I that leads to the activation of the NF-κB pathway (7). On the other hand, pentoxifylline (PTX), which is a pharmacologic agent used for improving the circulation, has been reported to have various effects at the cellular level including inhibition of TNF-α (8,9).

TNF-α gene expression was demonstrated at endometrial cells, decidual cells and trophoblastic cells during the trimesters of pregnancy (10). Moreover a group of pregnancy pathologies was associated with increased maternal TNF-α, which was suggested to influence fetal-maternal crosstalk during pregnancy. In this respect, because it is difficult to elucidate the role of TNF-α in such a complex process in in vivo, in vitro experiments with cell lines treated with recombinant TNF-α could be illuminating. Therefore, in this research, it was aimed to examine the response of trophoblasts to TNF-α mediated cellular stress in the JAR cell line by the evaluation of proliferative, apoptotic indexes and expression levels of NF-κB which is a key signaling molecule and to assess the therapeutic potential of PTX in TNF-α induced interactions.

MATERIALS AND METHODS

Cell Culture

JAR human choriocarcinoma cell line was purchased from the American Type Culture Collection. The cells were cultured in Dulbecco's modified Eagle's medium/F12 medium with 10% heat inactivated fetal bovine serum under the humidified atmosphere with 5% CO₂ at 37 °C. First of all, the cells grown on coverslips were cultured with experimental doses of TNF-α (PI-RP-10921, Thermo Fisher, MA, USA) ranging from 0.1 to 1000 ng/ml; the optimal dose of TNF-α was determined in accordance with evaluation of NF-κB expressions preliminarily. Then experimental groups were organized at 1, 6, 12 and 24 h-long 100 ng/ml TNF-α treatment. Afterwards, experimental groups were set up, treating the cells with 1, 10 and 20 mM doses of PTX (prepared from 100 mg per injection ampule) in the presence of TNF-α or not for 1h.

Immunocytochemistry

JAR cells were fixed with cold methanol. Following the incubation with blocking serum at room temperature, primary antibodies against proliferating cell nuclear antigen (PCNA) (MA1-16827, Thermo Fisher, MA, USA), caspase-8 (PI-MA1-91442, Thermo Fisher, MA, USA) and NF-κB p65 (Sc-109, Santa Cruz, CA, USA) were applied overnight at 4°C. After washing with phosphate buffered saline (PBS), biotinylated secondary antibodies and horseradish peroxidase (HRP) conjugated streptavidin were applied in order. Finally, after treatment with aminoethyl carbazole, the cells were investigated with an Olympus BX-61 bright field microscope. Proliferation indexes were calculated by taking the averages of the values obtained by dividing the number of PCNA positive cells by the total number of cells in each one of the 5 different areas. The intensities of immunocytochemical stainings for caspase-8 and NF-κB were semi-quantitatively scored in accordance with the following grading system: 0 (no staining), 1+ (weak, but detectable staining), 2+ (moderate or distinct staining), and 3+ (intense staining). Experiments were repeated three times and histological scores (HSCORE) were obtained for each slide. HSCORE = Σ Pi (i + 1), where i represents the intensity score, and Pi is the corresponding percentage of the cells (11).

DAPI Staining

The cells cultured on lamellas were treated with 100 ng/ml TNF-α for 24 h and then fixed with 4% of parafomaldehyde. After two times of washing with phosphate buffered saline (PBS) for 5 min, 4',6-diamidino-2-phenylindole dihydrochloride (DAPI) stain was applied to detect apoptotic cells with their characteristic nucleus morphologies (nuclear compaction, fragmentation or semilunar appearance). The samples were investigated under an Olympus BX-61 florescence microscope and apoptotic indexes were calculated by dividing the number of apoptotic cells by the total cell number.

Western Blot Analysis

JAR cells grown in different experimental conditions were washed with ice-cold PBS and scraped from culture flasks and then lysed with the cell lysis buffer containing a protease inhibitor cocktail to extract the total protein. The collected samples were subjected to electrophoresis on SDS-PAGE gels and transferred to nitrocellulose membranes. The membranes were blocked with 5% non-fat dried milk and then incubated with the primary antibodies against proliferating cell nuclear antigen (PCNA) (PA-109, Thermo Fisher, USA) and NF-κB p65 (Sc-109, Santa Cruz, CA, USA) dispersed in 1% BSA PBS for 1 h at room temperature. The membranes were washed with tris-buffered saline containing 0.1% Tween 20 for 15 min, then exposed to HRP-conjugated anti-rabbit secondary antibody (PI-31460 Thermo Fisher, USA) for 1 h at room temperature. Then, following a second washing step, the membranes were incubated with HRP-conjugated β-actin primary antibody (Sc-47778, Santa Cruz, CA, USA) as a loading control. The protein bands were visualized by using 3,3′-diaminobenzidine. Experiments were repeated three times and band intensities were quantified by densitometric analysis (Adobe Photoshop CS5) and normalized to β-actin readings.

Statistical Analyses

Statistical analyses were performed with Sigma Plot 12.0 software packages for the immunocytochemical and Western blot analyses. The data were presented as mean ± standard error (SE). Analysis between the groups were performed with One Way ANOVA test followed by Student t-test and non-parametric Kruskal Wallis-H tests for the immunocytochemistry and Western blot scores respectively. A value of p<0.05 was considered statistically significant.
RESULTS

In accordance with our preliminary studies, 100 ng/ml TNF-α was determined as an optimal dose for cellular stress induction in JAR cells. In this respect, the effects of cellular stress over the proliferation capacity of JAR cells were evaluated at the end of the 6, 12 and 24 h long incubations with TNF-α through the PCNA immunocytochemical analysis. The percentages of PCNA expressing cells after TNF-α treatment were statistically lower than the control and PCNA expression tended to increase with higher incubation time from 6th hr (Figure 1).

Second of all, cellular stress related apoptotic activity in JAR cells was investigated by evaluation of the immune reactivity of caspase-8 at the end of the 1, 6, 12 and 24 h of treatment with TNF-α. The caspase-8 immune reactivities of all the TNF-α applied groups were significantly higher than the control group and 1 h of TNF-α treatment brought about the highest caspase-8 immune reactivity (Figure 2A). Furthermore, cells treated with TNF-α for 24 h were compared with the non-treated ones by DAPI staining which is specific for DNA in order to show apoptosis related morphologic changes (Figure 2B-D). The number of cells having characteristic apoptotic nuclear morphology in the TNF-α treated group was significantly higher than the ones in the control (Figure 2B).

NF-κB signaling is one of the key pathways for the regulation of expressions of genes related to cell survival and proliferation, and activated by TNF-α (5,6). Therefore, nuclear NF-κB expression levels of the JAR cells were evaluated at the end of the 1-, 6-, 12- and 24 h long incubation with TNF-α through the immunocytochemical (Figure 3A-E and Table 1) and Western blot analysis (Figure 3F). It was found that 1 h long induction with TNF-α was effective to induce NF-κB expression significantly (Table 1). At this point, we assessed the effects of dif-
different doses of PTX, which has been reported as an inhibitory molecule for TNF-α on NF-κB expression levels. NF-κB immunoreactivities disappeared in all of the nuclei of cells treated with only 1 mM, 10 mM and 20 mM PTX. However, it was found that only 10 mM PTX was able to completely abolish the basal level of nuclear NF-κB expression, and significantly reduced the nuclear NF-κB expression induced by 1-h long incubation with 100 ng/ml TNF-α (Figure 4, Table 2).

Table 1. H scores for nuclear expressions of NF-κB average±SD in the control and TNF-α groups with different time intervals

<table>
<thead>
<tr>
<th>Groups</th>
<th>Average±SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ctrl</td>
<td>169±7</td>
</tr>
<tr>
<td>TNF-α (1h)</td>
<td>329±7</td>
</tr>
<tr>
<td>TNF-α (6h)</td>
<td>219±9</td>
</tr>
<tr>
<td>TNF-α (12h)</td>
<td>302±11</td>
</tr>
<tr>
<td>TNF-α(24h)</td>
<td>224±7</td>
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</tbody>
</table>

SD: standard deviation, aP<0.05 vs. control (ctrl) group, bP<0.05 vs. 6 h and 24 h long TNF-α treated groups.

Table 2. H scores for nuclear expressions of NF-κB average±SD in the control, TNF-α (1h) and PTX groups.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Average±SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ctrl</td>
<td>169±7</td>
</tr>
<tr>
<td>TNF-α (1h)</td>
<td>329±7</td>
</tr>
<tr>
<td>10 mM PTX</td>
<td>0</td>
</tr>
<tr>
<td>10 mM PTX + TNF-α (1h)</td>
<td>180±10</td>
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</table>

SD: standard deviation, aP<0.05 vs. TNF-α (1h) treated group.
Furthermore, Western blot analysis showed that total NF-κB expression was slightly increased with the TNF-α treatment and PTX application reduced total NF-κB expressions significantly compared to the control and only TNF-α treated ones (Figure 5).

![Figure 5. Graphical representation of average densitometric ratios of NF-κB to β-actin. *P<0.05 vs control group, **P<0.01 vs TNF-α group.](image)

**DISCUSSION**

Invasiveness of syncytiotrophoblasts is required for remodeling of spiral arteries during pregnancy. Inadequate invasion of trophoblasts leads to a deficient rupture of spiral arteries in the muscular layer, which in turn leads to disruption of utero-placental circulation (12,13). Interrupted arterial blood supply results in an increased generation of inflammatory cytokines like TNF-α (14,15). In this respect, determination of increased level of TNF-α in the plasma of preeclamptic pregnant women has shown that TNF-α could be used as an inducing agent for the establishment of experimental models (16). TNF-α takes part in induction of rapid transcription of genes associated with the regulation of proliferation, cell survival, inflammation and differentiation mainly through activation of the NF-κB pathway (6). For those reasons, it was aimed to evaluate the effects of TNF-α as a potential cellular inducer on JAR trophoblastic cells. In addition to all of this, PTX could be a potential therapeutic agent for the treatment of TNF-α related placental pathologies. In our experiments, we showed that a 1 h long induction with TNF-α brought about a significant increase in the total and nuclear expression of NF-κB. Even though the nuclear expression of NF-κB was fluctuating with increasing time of TNF-α stimulation, the decrease in proliferative capacity and increase in apoptotic activity are quite consistent with longer incubation time. Similarly, increased NF-κB expression induced with TNF-α was demonstrated in ED27 cells, which are immortalized trophoblast-like cells (23). Furthermore, increased NF-κB expression was also shown for syncytiotrophoblastic cells of preeclamptic placental tissues (24). As a result, we could define a positive feedback loop between TNF-α and NF-κB expressions in JAR trophoblastic cells which possibly participates in progressive aggravation of inflammation in the placenta.

PTX, which is a methylxanthine derivative and a non-specific inhibitor of cAMP phosphodiesterase, is generally applied as a pharmacologic agent for improvement of circulation in peripheral vascular disorders (8,25). Moreover, possible therapeutic effects of PTX as an inhibitor of TNF-α synthesis have been investigated in various diseases (26,27). Importantly, it was noted that PTX treatment has a reducing effect on plasma levels of proinflammatory cytokines including TNF in addition to its antioxidant effects (9). In our study, we investigated the optimal dose of PTX for inhibition of TNF-α induced NF-κB expression and it was found that 10 mM PTX application was quite effective in the reduction of TNF-α stimulated nuclear expression of NF-κB. It was reported that NF-κB translocation to nuclei was blocked by PTX application in TNF-α stimulated vascular smooth muscle cells (28). Therefore, PTX could be a potential therapeutic agent for the treatment of TNF-α related placental pathologies.

**CONCLUSION**

In summary, the present study showed that incubation with TNF-α leads to a decrease in proliferation capacity and an increase in apoptotic activity and NF-κB signaling in JAR syncytiotrophoblastic cell lines. In addition to all of this, PTX could be a potential regulatory agent for TNF-α and NF-κB signaling in JAR syncytiotrophoblastic cell lines. Therefore, we wanted to know the role of the NF-κB signaling pathway in those processes. In the absence of inducing stimuli, NF-κB molecules are in an inactive state in the cytoplasm. Upon activation by TNF-α, NF-κB is transferred to the nucleus, and regulates the associated genes (21,22). In this respect, the effects of TNF-α treatment on the nuclear NF-κB expression levels in different time intervals were evaluated. In our experiments, we showed that a 1 h long induction with TNF-α brought about a significant increase in the total and nuclear expression of NF-κB. Even though the nuclear expression of NF-κB was fluctuating with increasing time of TNF-α stimulation, the decrease in proliferative capacity and increase in apoptotic activity are quite consistent with longer incubation time. Similarly, increased NF-κB expression induced with TNF-α was demonstrated in ED27 cells, which are immortalized trophoblast-like cells (23). Furthermore, increased NF-κB expression was also shown for syncytiotrophoblastic cells of preeclamptic placental tissues (24). As a result, we could define a positive feedback loop between TNF-α and NF-κB expressions in JAR trophoblastic cells which possibly participates in progressive aggravation of inflammation in the placenta.
REFERENCES


