TNFα-induced IKKβ complex activation influences epithelial, but not stromal cell survival in endometriosis

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STUDY QUESTION: Can the activity of the IkB kinase (IKKβ) complex in endometriotic cells contribute to endometriotic lesion survival?

SUMMARY ANSWER: There is a constitutive activity of the IKKβ catalytic complex in peritoneal and deeply infiltrating lesions that can influence epithelial, but not stromal cell viability.

WHAT IS KNOWN ALREADY: Endometriotic lesions exist in an inflammatory microenvironment with higher local concentrations of cytokines, such as tumour necrosis factor α (TNFα). TNFα stimulates the activation of the IKKβ complex, an important nodal point in multiple signalling pathways that influence gene transcription, proliferation and apoptosis. However, few data on the regulation of IKKβ in endometriotic tissue are currently available.

STUDY DESIGN, SIZE, DURATION: A retrospective analysis of endometriotic tissue from peritoneal, ovarian and deeply infiltrating lesions from 37 women.

PARTICIPANTS/MATERIALS, SETTING, METHODS: Basal and activated (phosphorylated) IKKβ concentrations were analysed by western blotting and immunohistochemistry. The relationship between the expression and activation of these proteins and peritoneal fluid (TNFα) concentrations, measured via ELISA, was examined. A subsequent in vitro analysis of TNFα treatment on the activation of IKKβ and the effect on epithelial and stromal cell viability by its inhibition with PS1145 was also performed.

MAIN RESULTS AND ROLE OF CHANCE: Levels of the phosphorylated IKKβ complex in endometriotic lesions had a significant positive correlation with peritoneal fluid TNFα concentrations. Phosphorylated IKKβ complex was more prevalent in peritoneal and deeply infiltrating endometriosis lesions compared with ovarian lesions. IKKβ was present in both epithelial and stromal cells in all lesions but active IKKβ was limited to epithelial cells. TNFα stimulated an increased expression of phosphorylated IKKβ and the inhibition of this kinase with PS1145 significantly influenced ectopic epithelial cells viability but not eutopic epithelial cells, or endometrial stromal cells.

LIMITATIONS, REASONS FOR CAUTION: In vitro analysis on epithelial cells was performed with immortalized cell lines and not primary cell cultures and only low sample numbers were available for the study.

WIDER IMPLICATIONS OF THE FINDINGS: The regulation of aberrant signalling pathways represents a promising yet relatively unexplored area of endometriosis progression. The IKKβ complex is activated by inflammation and is critical nodal point of numerous downstream kinase-signalling pathways, including NFkB (nuclear factor κB), mTOR (mammalian target of rapamycin) and BAD (Bcl2-antagonist of cell survival)
Introduction

Endometriosis is characterized by the growth of endometrial epithelial and stromal cells outside the uterine cavity. It is an extremely prevalent disease occurring in 10–20% of women of reproductive age and is accompanied by chronic pelvic pain and subfertility. Although the exact pathogenesis is not yet clear, Sampson’s theory of transplantation is commonly accepted (Sampson, 1928). This theory proposes that viable endometrial epithelial and stromal cells are refluxed back through the Fallopian tube into the peritoneal cavity during menstruation. Once in this ectopic environment these cells avoid immune detection, invade the underlying mesothelial layer and stimulate a chronic inflammatory response.

Numerous inflammatory cytokines and chemokines (Borrelli et al., 2013, 2014) are increased in the peritoneal fluid of women with endometriosis, which occurs through the coordinated interaction of the refluxed endometrial and peritoneal immune cells. Refluxed endometrial cells produce and secrete chemokines (Hornung et al., 1997) that attract leukocytes and activated peritoneal macrophages (Halm et al., 1983). The activated macrophages produce inflammatory cytokines, which in turn further stimulate cytokine production by the endometrial cells, creating a feed forward regulatory loop (Lebovic et al., 2001) and the chronic inflammatory environment. This inflammatory environment has the potential to both influence symptomology (McKinnon et al., 2015) and disease progression (Bruner-Tran et al., 2013).

A chronic inflammatory environment can contribute to endometriotic lesion progression through the activation of a series of intracellular kinase-signalling pathways (McKinnon et al., 2016). The IκB kinase (IKK) complex represents a significant, early nodal point in many of the kinase-signalling pathways. In the nuclear factor (NF)κB signalling pathway, the IKK complex removes the inhibitory IκB protein from NFκB allowing translocation into the nucleus and gene transcription (Bonizzi and Karin, 2004) subsequently influencing the gene expression of many cytokine and chemokines, immunoreceptors, cell adhesion molecules, stress response genes and growth factors (Pahl, 1999). IKKβ interacts with the tubular sclerosis (TSC2) protein in the mammalian target of rapamycin (mTOR) pathway influencing cellular proliferation (Lee et al., 2007) and phosphorylates Bcl2-antagonist of cell death (BAD) pathway suppressing apoptosis (Yan et al., 2013).

The IKKβ complex consists of two catalytic subunits, IKKα and IKKβ and one regulatory subunit (IKKγ) (Hinz and Scheidereit, 2014). The binding of extracellular tumour necrosis factor (TNF)α to its cell membrane receptor TNF receptor (TNFR) (Haider and Knöll, 2009) stimulates the phosphorylation of both IKKα and IKKβ and activation of the IKKβ complex. TNFα concentrations are increased in the peritoneal fluid of endometriosis (Harada et al., 1999) and are correlated with the severity of the disease (Bedaiwy et al., 2002); and thus, TNFα-stimulated IKKβ activity may have a significant influence on the endometriotic lesions.

At present, very little information is available on the expression and activity of this important upstream nodal kinase in endometriotic cells. In this study, we used a combination of clinical and in vitro experiments to determine the presence and importance of IKKβ in endometriosis. We found that peritoneal fluid TNFα concentrations had a significant positive correlation with the activated IKKβ complex and that this was most likely due to epithelial cell expression. Furthermore, IKKβ activity was important in regulating epithelial cell, but not stromal cell survival. These results therefore suggest a significant role of IKKβ in endometriotic epithelial cells that deserves further attention.

Materials and Methods

Patient samples

Prior to surgery, the relevant institutional review board granted ethical approval and informed consent was obtained from all patients. During surgery performed for suspected endometriosis samples of endometrium, peritoneal fluid and endometriotic lesions were collected. Endometrial biopsies were collected via soft curette (Pipelde de Cornier, Laboratoire CCD, France) and stored in RNaLater (Thermo Fischer Scientific, Waltham, MA, USA) at −80°C as described previously (Santi et al., 2011). Peritoneal fluid was collected during the laparoscopic procedure from the cul-de-sac and centrifuged to remove blood cells, aggregates and debris. Samples were excluded if blood remained in the samples. The pelvic cavity was examined and any endometriotic lesions were removed and the patient staged (no endometriosis, or stages I–IV) according to the revised American Fertility Society staging system (rAFS) (American Society of Reproductive Medicine, 1997). The lesions were recorded as peritoneal, ovarian or deeply infiltrating endometriosis (DIE). All surgeries were performed during the proliferative phase of the menstrual cycle and endometriosis was confirmed by histological analysis.

Endometrial biopsies were collected from both women with (n = 8) and without endometriosis (n = 7) and used for the isolation of primary cells via collagenase digestion and size exclusion, as described previously (McKinnon et al., 2012). Strong yields were obtained for the primary endometrial stromal cells (ESC) from all women except one without endometriosis; however, only limited amounts of epithelial cells could be successfully isolated. As endometrial epithelial cells are terminally differentiated and do not propagate immortalized epithelial cells were acquired. Matching peritoneal fluid of sufficient quality was not always available for cytokine measurement; therefore, if peritoneal fluid was available then the ectopic lesions were immediately frozen and stored for subsequent Western blot analysis. If peritoneal fluid was not available, then they were formalin fixed and paraffin embedded for immunohistochemistry analysis.

STUDY FUNDING/COMPETING INTEREST(S): The study was funded by the Swiss National Science Foundation (Grant Number 320030_140774). The authors have no conflict of interest to declare.
Cytokine measurement in peritoneal fluid

TNFα was measured by an enzyme-linked immunoabsorbent assay (ELISA) kit (R&D Systems, Abingdon, England) using a high-sensitivity NADH cascade ampliﬁed alkaline phosphatase with antigen–antibody incubations at 28°C in a dry incubator and at a dilution of 1:2 in the diluent provided. Peritoneal ﬂuid progesterone concentrations were also measured via a radioimmunoassay (Coa-tect, DPC; Buhlmann Laboratories, Allschwil, Switzerland) to conﬁrm the patient cycle phase (McKinnon et al., 2014).

Protein isolation and quantitation in ectopic lesions

Approximately 30 mg of fresh frozen ectopic tissue was used to prepare whole-cell extracts via homogenization with the FastPrep 120 tissue homogenizer (30 s at 4.0 m/s) in radioimmunoprecipitation assay buffer (RIPA; 50 mM Tris-Cl, pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% v/v Triton X-100, 1% w/v sodium deoxycholate, 0.1% w/v sodium dodecyl sulphate and 1% v/v protease and phosphatase inhibitor cocktail; Cell Signalling Technology, Danvers, MA). Final protein concentrations were determined with the bicinchoninic acid assay (QuansPro BCA; Sigma-Aldrich, St Louis, MO, USA).

Proteins separation was performed by heating 20 μg of total protein into LDS buffer (Invitrogen) to 70°C for 10 min and running it on a 4–15% Novex NuPAGE Bis/Tris gel (Invitrogen). Proteins were transferred onto a 0.45-μm nitrocellulose membrane in 4-morpholinepropanesulfonic acid buffer (MOPS; Invitrogen) pH 7.7 for 1 h. Non-speciﬁc binding was blocked by incubation with 5% bovine serum albumin (BSA) in phosphate-buffered saline (PBS) containing 0.1% Tween-20 (PBST). Membranes were probed with rabbit anti-IKKβ antibody (1:1000) (Cell Signalling Technology), and with rabbit anti-pIKKα/β (176/180) (1:1000) antibody (Cell Signalling Technology), or mouse anti-actinβ antibody (Abcam, Cambridge, UK) 1:5000. Secondary anti-rabbit (GE Healthcare, Opﬁkon, Switzerland) and anti-mouse (Sigma) antibodies conjugated to horseradish peroxidase were diluted 1:50 000 and 1:200 000, respectively. Immuno-reactivity was determined with the SuperSignal West Femto kit (Pierce; Thermo Scientiﬁc) using the Chemi-Doc XRS+ system (Bio-Rad Laboratories, AG, Cressier, Switzerland). Band densitometry was quantiﬁed with the Quantity One software and in each western blot a calibrator sample with strong IKKβ, and pIKKα/β expression was included to normalize concentrations across gels and determine protein concentrations relative to the calibrator sample. The relative expression of each sample was expressed as a percentage of the calibrator. Actinβ was used as a loading control.

Immunohistochemistry

Immunohistochemistry was performed using serial sections of 4 μm mounted onto glass slides (Superfrost, Braunschweig, Germany), dewaxed in xylene and rehydrated through a series of decreasing ethanol concentrations. Epitope retrieval was performed with 10 mM citrate buffer, pH 5.5 for 5 min in a 450-W microwave. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide (H2O2) and a blocking step performed with 3% BSA for 30 min in Tris-buffered saline (TBS; Tris 100 mM, NaCl 0.15 M; pH 7.4). Rabbit anti-IKKβ antibody (1:100) and rabbit anti-pIKKα/β (176/180) antibody (1:100) were diluted in 3% BSA in TBS and incubated at 4°C overnight in a humididied chamber. Slides were washed with TBS and 0.1% Tween-20 (TBST) prior to incubation with an afﬁnity-puriﬁed, biotin-conjugated goat anti-rabbit antibody (Dako, Glastrup, Denmark) for 90 min at room temperature. After a ﬁnal wash, slides were incubated with an avidin–biotin HRP complex ( Vectastain, ABC Kit, Vector Laboratories, Burlingame, CA, USA) for 45 min. The antigen–antibody complex was detected by incubation with 3,3′- diaminobenzidine substrate and slides were counterstained with hematoxylin and mounted in Aquatex (Merck). The primary antibodies were excluded for the negative controls. Images were photographed with a Nikon Eclipse E800 microscope (Nikon, Japan). Semi-quantitative analysis of antibody staining in the epithelial and stromal cells of the endometriotic lesions was determined by the allocation of scores between 0 and 3; 0 (negative), 1 (weak), 2 (moderate) and 3 (strong) based on the intensity of staining. The percentage of cells with positive immuno-reactivity was also determined and allocated a score between 0 and 6 as described % = 0, 1–10% = 1, 11–30% = 2, 31–50% = 3, 51–70% = 4, 71–90% = 5; >91% = 6 in each cell type of the endometriotic lesions. For the ﬁnal immuno-reactive score, the staining intensity and percentage of positive cells was multiplied, as described previously (Samartzis et al., 2012).

Cell culture and TNFα treatment

Isolated primary ESC were maintained in Iscoves’s modiﬁed Eagle medium (IMEM) (Invitrogen) supplemented with 10% fetal calf serum (FCS) (Invitrogen) and 1% antibiotic/antimycotic (Invitrogen). The immortalized epithelial cell lines, EM E6/E7 and EM’Osis, were provided by Professor Kyo, Kanazawa, Japan and were isolated from eutopic endometrium (Kyo et al., 2003) and an ectopic endometrioma (Bono et al., 2012), respectively. These cells were maintained in Dulbecco’s modiﬁed Eagles medium (DMEM) (Invitrogen) with 10% FCS and 1% antibiotic/antimycotic. The 12Z cells were provided by Professor Starzinski-Powitz, Goethe University and were originally isolated from a peritoneal endometriotic lesion (Zeitvogl et al., 2001). These cells were also maintained in complete DMEM.

To determine the inﬂuence of TNFα on pIKKα/β activity in all cell types, the cells were seeded into 6-well plates at ~3 x 10^5 cells/well. After reaching ~80% confluence, the media was changed to 0.5% FCS for overnight incubation prior to treatment. Cells were treated with either control media (0.5% FCS) or media plus 10 ng/ml and 100 ng/ml recombinant human TNFα (R&D Systems, Minneapolis, MN, USA) for 6 h. At the end of the treatment period, the cells were rinsed and collected in RIPA buffer.

MTS assay and treatment with PS1145

PS1145 is a small molecular weight compound that is a speciﬁc inhibitor of IKKβ activity (Lam et al., 2005). Inhibition of IKKβ activity with PS1145 was performed in 96-well plates seeded at a density of 6 x 10^4/well. Twenty-four hours prior to treatment, the cells were changed into serum-free media and treatment media prepared by diluting PS1145 into either serum-free media at a ﬁnal concentration of 1 μM. Subsequent concentrations were prepared by a 1:3 serial dilution (333.3, 111.1, 37.0, 12.3 and 4.12 nM). Cell viability was measured after 72 h by the CellTiter 96 Aqueous One Solution Cell Proliferation Assay (Promega, Southhampton, Hampshire, UK). Triplicate wells were used for each cell type and experiment. For the immortalized epithelial cell lines the experiment was repeated three times and for the primary ESC an experiment on each of the eight endometriosis and seven non-endometriosis preparations was performed separately. A control (without PS1145) was included for each experiment and designated as 100% viability and subsequent values expressed as a percent of control.

Statistical analysis

All statistical analyses were performed with Graphpad Prism version 6.0. The correlation between the peritoneal ﬂuid cytokines and IKKβ and pIKKα/β expression was performed using the non-parametric Spearman’s rank correlation coefﬁcient. Two-group comparisons were performed with a non-parametric Mann–Whitney U test and the comparison of three or more groups with the non-parametric Kruskal–Wallis one-way analysis (ANOVA) and Dunn’s multiple comparison post hoc test. The interaction between two variables was determined via a two-way ANOVA with a post
Results

Patient data and characteristics

In total, endometriotic tissue was removed from 37 endometriosis patients and endometrial biopsies from 8 women without endometriosis. Of these 37 women, we collected accompanying peritoneal fluid from 21 in order to compare peritoneal fluid TNFα and ectopic IKKβ. Of the 21 samples, 14 were collected from women without any hormonal treatment, 4 were using combined oral contraceptives (COC) and 3 were using GnRH analogues. No significant variation in TNFα, IKKβ or pIKKα/β based on hormonal use (Table I) was identified. Five of the lesions were peritoneal, eleven ovarian and five DIE.

The remaining 16 samples without accompanying peritoneal fluid were kept for immunohistochemistry. Of these, six women had no history of hormonal treatment, five were using COC and five were using GnRH analogues. Three lesions were peritoneal, seven ovarian and six DIE. No significant variation was observed between either pIKKα/β or IKKβ expression in both the epithelial and stromal cells based on hormonal treatment (Table II).

IKKβ expression and activation in endometriotic tissue and its relationship to peritoneal fluid TNFα concentrations

Comparison of peritoneal fluid TNFα and endometriotic lesion IKKβ and pIKKα/β showed a significant positive correlation between TNFα and pIKKα/β (r = 0.6268, n = 21, P = 0.0024) (Fig. 1A), but not IKKβ (r = 0.4216, n = 21, P = 0.0570) (Fig. 1B), as determined by semi-quantitative western blot (Fig. 1C). A significant variation in pIKKα/β concentrations between lesions from different locations (P < 0.05) was observed with a post hoc analysis confirming a significantly lower expression in ovarian lesions (50 ± 8.7, n = 11) compared with the peritoneal lesions (99 ± 9.7, n = 5) (P = 0.041) (Fig. 1D). No significant difference was observed with the DIE lesions (91 ± 21.6, n = 5). In contrast, IKKβ expression showed no variation between lesion types (P = 0.4905) with similar expression in the peritoneal (138 ± 29.3, n = 5), ovarian (128 ± 26.7, n = 11) and DIE lesions (80 ± 26.9, n = 5) (Fig. 1E). The active to inactive ratio of IKKβ (pIKKα/β:IKKβ) varied significantly between lesions (P = 0.0123) with both peritoneal (1.0 ± 0.39, n = 5) and DIE (1.7 ± 0.54, n = 5) higher than ovarian lesions (0.5 ± 0.09, n = 11) with a post hoc analysis showing a significant difference between DIE and ovarian lesions (P = 0.0168).

Cell-specific activation of IKK in endometriotic lesions

Immunohistochemistry identified a low but uniform expression of IKKβ in peritoneal (Fig. 2A), ovarian (Fig. 2B) and DIE lesions (Fig. 2C) in both stromal (red arrows) and epithelial cells (black arrows). In contrast, pIKKα/β in peritoneal (Fig. 2D), ovarian (Fig. 2E) and DIE (Fig. 2F) lesions was predominantly epithelial (black arrows), with significantly less stromal cells expression (red arrows). Negative controls showed no expression in peritoneal (Fig. 2G), ovarian (Fig. 2H) or DIE (Fig. 2I) lesions. No statistically significant difference was observed in IKKβ expression between either lesion location (P = 0.2420) or cell

Table I Comparison of endometriotic lesion and peritoneal fluid protein expression in samples removed from women subject to different hormonal treatments.

<table>
<thead>
<tr>
<th>Hormone treatment</th>
<th>No hormone (n = 14)</th>
<th>Combined oral contraceptive (n = 4)</th>
<th>GnRH analogue (n = 21)</th>
</tr>
</thead>
<tbody>
<tr>
<td>pIKKα/β (%)</td>
<td>66 ± 9.8</td>
<td>77 ± 27.8</td>
<td>92 ± 17.5</td>
</tr>
<tr>
<td>IKKβ (%)</td>
<td>120 ± 23.5</td>
<td>90 ± 27.7</td>
<td>151 ± 27.7</td>
</tr>
<tr>
<td>TNFα (pg/ml)</td>
<td>2.1 ± 0.61</td>
<td>1.2 ± 0.53</td>
<td>1.5 ± 0.54</td>
</tr>
</tbody>
</table>

Data are mean ± SEM. IKKβ and pIKKα/β protein values were determined via semi-quantitative western blot densitometry analysis and expressed as a percentage of a standardized control sample included in each western blot, as described in detail in the Materials and methods section. TNFα values represent peritoneal fluid TNFα concentrations determined by ELISA and expressed as pg/ml. Analysis of significance was performed via the non-parametric one-way ANOVA test (Kruskal–Wallis) and a post hoc comparison of all groups (Dunn’s multiple comparison test) significance P < 0.05.

Table II Comparison of protein expression in formalin fixed paraffin embedded endometriotic tissue according to hormonal treatment.

<table>
<thead>
<tr>
<th>Hormonal use</th>
<th>No hormone (n = 6)</th>
<th>Combined oral contraceptive (n = 5)</th>
<th>GnRH analogue (n = 5)</th>
<th>Total (n = 16)</th>
<th>*P</th>
</tr>
</thead>
<tbody>
<tr>
<td>pIKKα/β</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Epithelial</td>
<td>1.7 ± 0.56</td>
<td>3.4 ± 1.78</td>
<td>6.2 ± 2.27</td>
<td>3.6 ± 0.98</td>
<td>0.108</td>
</tr>
<tr>
<td>Stromal</td>
<td>0.3 ± 0.33</td>
<td>0.8 ± 0.37</td>
<td>0.6 ± 0.60</td>
<td>0.5 ± 0.24</td>
<td>0.448</td>
</tr>
<tr>
<td>IKKβ</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Epithelial</td>
<td>2.3 ± 0.95</td>
<td>2.2 ± 1.11</td>
<td>3.0 ± 1.76</td>
<td>2.5 ± 0.70</td>
<td>0.949</td>
</tr>
<tr>
<td>Stromal</td>
<td>0.7 ± 0.42</td>
<td>2.4 ± 1.75</td>
<td>0.0 ± 0.00</td>
<td>1.00 ± 0.58</td>
<td>0.327</td>
</tr>
</tbody>
</table>

Data are mean ± SEM. Values represent the scores derived from the semi-quantitative IHC analysis, described in detail in the Materials and methods. Analysis of significance was performed via the non-parametric one-way ANOVA test (Kruskal–Wallis) and a post hoc comparison of all groups (Dunn’s multiple comparison test).

*Significance is P < 0.05.
type \((P = 0.1972)\) (Fig. 2J), although this could be due to a lack of power. \(pIKK_{\alpha/\beta}\) expression was significantly different in cell types \((P = 0.0198)\), but no statistically significant difference could be observed between lesion type \((P = 0.3402)\) (Fig. 2K) possibly again due to lack of power.

**IKKβ activity after TNFα treatment**

Western blot analysis of \(pIKK_{\alpha/\beta}\) after TNFα treatment of epithelial cell cultures confirmed a low but positive expression in all cell lines examined (Fig. 3A). Semi-quantitation of band densitometry indicated that the 12Z cells showed the strongest basal expression (no TNFα), but this did not vary after TNFα treatment \((P = 0.2320)\). The EM’E6/E7 cells showed lower basal expression and also no significant variation after TNFα treatment \((P = 0.4475)\). In contrast, TNFα treatment of EM’osis cells significantly increased \(pIKK_{\alpha/\beta}\) above the no treatment control \((46 \pm 5.0, n = 3)\) at concentrations of both 10ng/ml \((141 \pm 34.2, n = 3, P = 0.0173)\) and 100 ng/ml \((197 \pm 23.3, n = 3, P = 0.0085)\) (Fig. 3B). Similar western blots were performed on protein lysate isolated from stromal cells; however, no protein expression could be observed in these preparations.

**Cell-specific influence of IKKβ inhibition on viability**

In the epithelial cell cultures, there was a significant influence of PS1145 on cell viability based on cell type \((P < 0.0001)\) (Fig. 4A). No significant effect of PS1145 was observed on the EM’E6/E7 at any concentration. For EM’osis cells, there was a significant increase in cell viability at the lowest concentrations \((PS1145 2.43 nM;\)
123 ± 0.2, n = 3, P < 0.001) that was gradually diminished as concentrations increased (PS1145 1 μm; 106 ± 6.3 P > 0.05), whereas I2Z cell viability was significantly decreased at the lowest concentrations (PS1145 2.43 nM; 70 ± 4.6, n = 3, P < 0.0001) and remained significantly reduced through to the highest concentration (PS1145 1 μm; 73 ± 2.6, n = 3, P < 0.0001). In contrast, primary ESC isolated from women with and without endometriosis showed no significant variation based on either PS1145 (P = 0.8868) or cell type (P = 0.3516) (Fig. 4B).

**Discussion**

In this study, we examined the expression of the IKKβ protein kinase complex and the activation of its catalytic subunits pIKKα/β in both...
endometriotic lesions, as well as its influence on cell survival in in vitro models. The results show a significant relationship between the phosphorylation of the IKKβ complex and peritoneal fluid TNFα. Subsequent immunohistochemistry staining showed that although no statistically significant difference in IKKβ expression was observed across all cells and lesion types the phosphorylated IKKβ complex was predominantly epithelial. These data were supported by the in vitro studies that confirmed pIKKα/β expression in epithelial cell culture models, but not primary stromal cells and that inhibition of IKKβ activity significantly influenced endometriotic epithelial cell viability, but not eutopic epithelial cell viability, nor the viability of endometrial stromal cells from women with and without endometriosis. These results therefore suggest that a dysregulation of the IKKβ kinase occurs in ectopic epithelial cells that may be related to the inflammatory microenvironment. Given the role of IKKβ in transmitting extracellular signals into cell survival via kinase-signalling pathways, it may represent a significant molecule in endometriosis pathogenesis.

At present, there is very little known about the role of IKKβ in endometriosis. The results from our clinical samples suggest that the constitutive IKKβ activity is significantly different between peritoneal and DIE lesions compared with ovarian lesions. This difference of expression was supported by our in vitro results that showed TNFα stimulated an increase in pIKKα/β expression and that inhibition of IKKβ activity increased the Em’osis cell viability. In contrast in the peritoneal-

![Figure 3](image3.png)

Figure 3  pIKKα/β after TNFα treatment in endometriotic epithelial cells. (A) Western blot analysis confirmed the expression of pIKKα/β in the 12Z, EM E6/E7 and EM’osis cell lines both with and without TNFα treatment. (B) Semi-quantitative analysis indicated TNFα did not significantly influence either 12Z or EM E6/E7 expression. There was however a significant increase in pIKKα/β after both 10 ng/ml and 100 ng/ml TNFα in the EM’osis cell line.

![Figure 4](image4.png)

Figure 4  Influence of IKKβ inhibition on epithelial and stromal cell viability. (A) Inhibition of IKKβ activity with increasing concentrations of PS1145 had no influence on the eutopic-derived epithelial EM E6/E7 cells, significantly increased the cell viability of the ovarian-derived EM’osis cells, significantly reduced the viability of the peritoneal-derived 12Z cells. (B) PS1145 had no influence on stromal cells from women with and without endometriosis. Analysis on the influence between cell type and PS1145 concentrations on cell viability performed with a two-way ANOVA test with a post hoc Tukey’s multiple comparison test. **P < 0.01, ***P < 0.001, ****P < 0.0001.
derived epithelial cells TNFx had a limited influence of plIKKα/β expression and inhibition of IKKβ activity decreased cell viability. Unfortunately, a DIE-derived cell line was not available.

Endometriosis is a significantly heterogeneous condition, although whether these lesions have different pathologies (Nisolle and Donnez, 1997), or represent a continuum of the same disease (Somigliana et al., 2004) is still debated. These data suggest a varied cellular response to inflammation may occur in different lesions. It has previously been shown that rectovaginal septum lesions have a distinctly inflammatory phenotype (Bertschi et al., 2013) and that concentrations of inflammatory mediators are stronger in the peritoneal fluid of DIE compared with lesions from other locations (Santulli et al., 2012). As IKKβ can be associated with TNFx both in ours and other studies (Lee et al., 2007), it is possible that the increased production of inflammation associated with DIE lesions is related to the higher IKKβ activity.

The identification of a TNFx influenced activation of the IKKβ in endometriotic tissue is significant because of the multiple downstream pathways it regulates (Fig. 5) and the influence this can have on gene transcription, protein translation and both cellular proliferation and apoptosis. Activation of IKKβ stimulates NFkB gene transcription and a constitutive activation of NFkB has been observed in peritoneal endometriosis (González-Ramos et al., 2007). Multiple factors present in the peritoneal fluid of women with endometriosis including cytokines and iron overload (Alvarado-Díaz et al., 2015) may lead to this constitutive activation. Furthermore, an increased NFkB activity has been linked to recurrence of ovarian endometrioma (Shen et al., 2008). Neither of these studies, however, examined IKKβ expression directly. In the immortalized epithelial I2Z cells IKKβ inhibition attenuated inflammatory cytokine secretion (Grund et al., 2008) and in ectopic endometrial stromal cells mir200a suppresses IKKβ (Dai et al., 2012), raising the possibility of suppressed IKKβ activity in stromal cells occurs via an epigenetic regulation. In contrast to its role in inflammation via the NFkB pathway, IKKβ can also regulate cellular proliferation and apoptosis through the mTOR and BAD pathways (Yan et al., 2013; Dunlop and Tee, 2014). A dysregulation of mTOR has previously been implicated in endometriosis pathogenesis of DIE lesions leading to increased proliferation (Lecorte et al., 2011), as has a role for mTOR-mediated autophagy (Choi et al., 2014) and BAD activation in ovarian endometriomas (Stickles et al., 2015). Together, this suggests that TNFx has the potential to modulate all of these activities via IKKβ activation.

We found that the constitutive activation and influence on cell survival was largely restricted to epithelial cells. Endometriotic lesions are a combination of epithelial and stromal cells and an interdependency between the cells types is required for endometriotic lesions to continue proliferating as tissue integrity of refluxed endometrial matter is essential to endometrial tissue implantation (Nap et al., 2003). We have also previously shown that the stromal cells produce significantly more inflammatory cytokines than epithelial cells in response to stimulation (Bersinger et al., 2008). It could therefore be postulated that a paracrine regulation occurs in the lesions through the stromal cells production of cytokines stimulating a constitutive activation of the IKKβ complex in epithelial cells, which ultimately contributes to cell survival. More research however is required to explore this hypothesis.

Furthermore, the activity of IKKβ in other cell types other than endometriotic cells was not directly addressed in this study, but may give further insight into this mechanism in normal tissue. For endometriosis, however, whether this mechanism also happens in healthy eutopic epithelial cells may be of less consequence. This is because epithelial cells will only be present in the peritoneal cavity when endometriosis is present, and when endometriosis is present there is a constant inflammatory environment. We believe it is this influence of ectopic epithelial cells and constant inflammation that makes the contribution of TNFx-stimulated IKKβ activity significant. It may be such that this is a characteristic not inherent in the endometrium, but rather acquired during the life of the lesion and contribute more to progression through a constant stimulation of the inflammatory cascade. Further study on whether there is a significant difference between the activation of IKKβ in the eutopic endometrium of women with and without endometriosis would be an interesting follow-up.

Whether other cell types also show a constitutive activity of IKKβ in the presence of inflammation would also be interesting. In this study, the images in Fig. 2 indicate that cells proximal to the endometriotic lesion are largely negative for plIKKα/β expression, providing circumstantial evidence for the preferential activation of plIKKα/β in endometriotic epithelial cells. Previous studies suggest that cells proximal to the lesion may have different characteristics to cells distal to the lesions (Young et al., 2014) and these cells thus may also be interesting to study; however, we were unable to collect this tissue due to our
current ethical approvals. Future studies on this topic may however be warranted.

Limitations of this study were the inclusion of women with hormonal treatment. Previous research however has suggested there was no significant influence on NFκB activation by oral contraceptives (González-Ramos et al., 2007). We also observed no statistically significant difference for IKKβ in this study, although the power of this analysis was limited by the small sample size. It is possible that this may introduce a variability in peritoneal fluid cytokine concentrations in endometriotic women, as GnRHα analogues have been shown to have an influence on the inflammatory environment (Nirgianakis et al., 2013), however we postulate that a reduction in inflammatory cytokines by hormonal treatment would also be reflected by a subsequent reduction in IKKβ activation, maintaining any correlation between the extra and intracellular environment. A further limitation of this study is the small sample size. An expansion of the sample number would provide more definitive information on both the influence of TNFα, as well as hormonal treatments, on IKKβ activity. It is difficult to draw direct conclusions on the contribution of IKKβ to cell survival in different lesions types as immortalized cell lines were used, however given the similarity observed in the in vitro results to the clinical samples we can be confident that IKKβ has a role in epithelial endometriotic cells.

In conclusion, we have observed a significant relationship between TNFα and the activation of IKKβ complex in the endometriotic microenvironment and that this activation occurs predominantly in the epithelial cells of peritoneal and DIE lesions. Furthermore, IKKβ inhibition in vitro significantly influenced epithelial cell, but not stromal cell behaviour. The regulation of kinase-signalling pathways is a significant, but under explored area of endometriosis pathogenesis and progression and may represent potential non-hormonal treatment targets for endometriosis (McKinnon et al., 2016). Given the ability of the extra-cellular inflammatory environment to influence IKKβ activity and its subsequent affect on downstream pathways, this kinase may be of significant interest in endometriosis.

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Authors’ roles

V.K. performed experiments, cell culture and assisted with sample collection. C.W. performed immunohistochemistry and analysis. G.G. assisted with cell culture. N.A.B. assisted with sample collection and intellectual development of the project. M.D.M. assisted with sample collection and intellectual development of the project. B.D.M. conceived project, performed experiments and prepared the manuscript.

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Conflict of interest

None declared.

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