Betaine Attenuates The Expression of Vasoactive Mediators and Histological Alterations Associated with Ovarian Hyperstimulation Syndrome in Rats

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Abstract

Objective: Ovarian hyperstimulation syndrome (OHSS) is one female reproductive disorder that can occur after administration of injectable hormonal drugs to stimulate ovulation. Betaine (BET) is an intracellular biomolecule with anti-inflammatory and tissue protective effects. There is no information about its effects in an experimental model of OHSS. The current study aims to investigate the possible effects of BET on abnormal expressions of vasoconstrictor proteins and ovarian histological changes in an experimental OHSS rat model.

Materials and Methods: In this experimental study, 30 adult female rats (two months old) were randomly divided into six groups (n=5 per group): i. Control, ii. OHSS [10 IU sc equine chorionic gonadotropin (eCG) for 4 days followed by 30 IU sc human chorionic gonadotropin (hCG) on the fifth day], iii. OHSS+BET (200 mg/kg/day, orally for seven days), iv. OHSS+Cabergoline (CAB, 100 mg/kg/day, orally for six days), v. BET, and vi. CAB. Expression levels of vascular endothelial growth factor (VEGF), cyclooxygenase-2 (COX-2), and blood levels of oestradiol (E2) and progesterone (P4) were measured at the end of the experiment. The ovaries were studied for histomorphological changes.

Results: Induction of OHSS altered tissue histology, including an increase in the number of corpora lutea and atretic follicles, and decreased the number of follicular reserves. In this group, we observed increased expressions of the VEGF and COX-2 proteins, and increased serum E2 and P4 levels. Administration of CAB and BET significantly attenuated all molecular and histological alterations observed in the OHSS animals.

Conclusion: Our findings, for the first time, indicate the beneficial effects of BET to reduce OHSS complications in patients by reducing the expressions of vasoactive proteins and improving changes to the ovarian tissues. The findings are similar to CAB and can be a new avenue for future research on BET.

Keywords: Angiogenesis, Betaine, Cabergoline, Ovarian Histomorphology, Ovarian Hyperstimulation Syndrome


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Introduction

The ovaries are paired structures that contain immature ovum that can develop into mature eggs and acquire the ability to fertilise (1). Approximately 10% of patients who undergo in vitro fertilization (IVF) cycles, embryo transfer, stimulation of ovulation, or intrauterine insemination develop ovarian hyperstimulation syndrome (OHSS) (2-4). Clinical manifestations of this complication include extensive accumulation of fluid in the extracellular spaces and condensation of blood due to leakage of fluid from the arteries, and this can be associated in advanced-stage renal impairment, hypervolemic shock, thromboembolism, and respiratory distress (5, 6). This syndrome is caused by increased expression and secretion of vascular endothelial growth factors (VEGF) and cyclooxygenase 2 (COX-2) (5, 7-10). Therefore, the use of compounds that inhibit VEGF secretion can play an effective role in the prevention of OHSS (8,11,12) Cabergoline (CAB), a low-dose dopamine receptor (Dp-R2), inhibits vascular permeability and can be used as a prophylactic treatment for OHSS syndrome (12-16). Trimethylglycine, also known as betaine (BET), is synthesised in the inner mitochondrial membrane and functions as an osmoprotectant and methyl group donor in various cellular pathways in mammals (17). BET may have a functional role in female reproductive health and fertility. Recent findings indicate that BET can inhibit retinal vascularisation by suppression of VEGF and COX-2 expressions (17, 18). Currently, the effects of BET on
the molecular and histological changes observed in animal models of OHSS are unknown. The current study aims to investigate the effects of BET on the expressions of vasoactive proteins and histological changes induced by experimental OHSS in rat ovaries and compare its effects with CAB, an effective drug used to manage OHSS complications.

Materials and Methods

Animals and treatment assignments

A total of 30 female, two-month-old unmated Wistar rats (weights: 150-200 g) were acquired from the Animal House of the Faculty of Veterinary Medicine, Shahid Chamran University, Ahvaz, Iran. The animals were kept under a constant room temperature of 20 ± 1°C, humidity of 40-50%, and a 12-hour light/dark cycle. The rats were fed pelleted food (Pars, Tehran, Iran) for one week to allow them to adjust to their new environment.

The animals were weighed and randomly divided into six groups (n=5 per group), as follows: group I (control) rats were given 0.9% normal saline (0.1 ml/kg, sc) for five days. From day 0 of the assessment, the rats received a daily oral dose of 1 mg/kg normal saline for ten days. Group II (OHSS) rats received sc injections of equine chorionic gonadotropin (eCG, IBSA, Montagnola, Switzerland) dissolved in 0.1 ml of 0.9% normal saline (10 IU) for 4 days. On the fifth day, they received an sc injection of 30 IU human chorionic gonadotropin (hCG; Darou Pakhsh, Tehran, Iran) (19). Group III (OHSS+BET) OHSS rats were given 200 mg of oral BET (betaine anhydrous, Sigma, USA) two days before the eCG administration, for seven days. Group IV (OHSS+CAB) OHSS rats, from the first day of the test, were given 100 µg/kg/d oral CAB (Shahredaru, Iran) dissolved in 5% glucosamine for six days (19). Group V (BET) rats were given 200 mg of oral BET for seven days. Group VI (CAB) rats received 100 µg/kg, oral CAB dissolved in 5% glucosamine for five days.

As described previously, OHSS induction was confirmed by an increase in the delta value of body weight and elevation of the oestradiol (E2) and progesterone (P4) hormones.

This research was approved by the Ethics Committee of Shahid Chamran University, Ahvaz, Iran (EE/99.3.02.38347/scu.ac.ir). Figure 1 presents the research design.

At the end of trial period, the rats in all groups were weighed and euthanised with ketamine and xylazine (100 mg/kg+10 mg/kg). Serum samples were taken by heart puncture and stored at -20°C until hormone analysis. Both ovaries of each rat were removed and weighed. One ovary was fixed in 10% formalin buffer for the histological analyses, and other ovary was stored at -70°C for molecular analysis.

Histological evaluations

The tissues were cut into 5 µm sections and stained with haematoxylin and eosin (H&E, Merck, Germany) (20). Histomorphometric studies were performed using a light microscope (Olympus BH-2, Japan) equipped with a Dino-Eye lens (AM7023, Taiwan) and DinoCapture 2.0 imaging software at 200× magnification.

Ovarian histological changes to the ovarian diameter (mm), the follicular reserve count, number and diameter of the corpora lutea (µm), number of atretic follicles and vascular reserve were evaluated, in addition to assessments of hyperaemia, haemorrhage, mononuclear cell infiltration, and interstitial tissue oedema. Counts were performed in 10 microscopic fields of view from each ovary and given a score between 0 and 3 for the ovarian tissue changes: 0 (unchanged), 1 (lowest), 2 (lowest moderate), and 3 (most change) (21).

![Fig.1: Schematic diagram of the treatment protocol. BET; Betaine, CAB; Cabergoline, VEGF; Vascular endothelial growth factor, COX-2; Cyclooxygenase-2, and OHSS; Ovarian hyperstimulation syndrome.]

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Biochemical analyses

The serum levels of P4 (ng/ml) and E2 (pg/ml) were measured by an ELISA Kit (Alpine, USA) according to the manufacturer’s instructions.

Evaluation of vascular endothelial growth factor and cyclooxygenase 2-expressions

The protein expressions of VEGF and COX-2 in the ovaries of the experimental animals were determined by Western blot analysis. The frozen tissues were lysed in 200 μl lysis buffer (50 mM tris-HCl, 150 mM NaCl, 0.1% Triton X-100, 1 mM NaF) supplied with protease inhibitor cocktails (Sigma-Aldrich, MO, USA) for 30 minutes on ice. Primary antibody VEGF (Abcam, Cambridge, UK, Art No: ab46154) and COX-2 (Abcam, Cambridge, UK, Art No: ab179800) were incubated with an appropriate secondary antibody [goat anti-rabbit IgG, Abcam: ab133470 (618)]. β2-microglobulin (B2M, Abcam, Cambridge, UK, Art No: ab214769) was used as the calibrator protein. ImageJ software (National Institutes of Health, Bethesda, MD, USA) was used to analyse the optical density of the protein bands. A control group was the calibrator group (21).

Statistical analysis

Data were analysed by Graphpad Prism software (version 5.0.4, Graphpad Software, Inc., San Diego, CA, USA). Data are presented as mean ± SEM. Bilateral ANOVA was performed and Tukey’s post hoc test was used for multiple comparisons. Statistically significance differences between the different experimental groups were determined, as follows: *; P<0.05, **; P<0.01, ***; P<0.001, and ****; P<0.0001.

Results

Analysis of body weight changes

Table S1 (See Supplementary Online Information at www.celljournal.org) displays the weight changes in the groups. Statistical analysis revealed that the delta value of body weight (final weight-initial weight) showed a significant increase in the OHSS group compared to the control group. The delta body weight in the OHSS group treated with CAB showed a significant decrease compared to the OHSS group treated with BET (P<0.01). The weight changes in the BET and CAB groups were similar to the control group.

Analysis of hormones in serum

Figure 2 illustrates the serum concentrations of E2 and P4 in the study groups. There was a significant increase in serum E2 and P4 levels in the OHSS group compared with the control group (P<0.0001). The CAB treated group had a decrease in E2 and P4 levels compared to the BET treated group (P>0.05).

Analysis of protein expressions in the ovaries

Figure 3 shows the VEGF and COX-2 protein expressions in the ovarian tissues for all of the study groups. Our results showed a significant increase in VEGF and COX-2 protein expression in the OHSS group compared with the control group (P<0.0001). The OHSS+CAB group showed a significant decrease in VEGF (P<0.001) and COX-2 (P<0.001) expressions.

Analysis of ovarian morphometry

Our results revealed that ovarian weight and diameter significantly increased in the OHSS group compared to the control group (P<0.0001). The ovarian diameters after treatment with BET (P<0.001) and CAB (P<0.001) were significantly lower compared to the OHSS group. Our results indicated that the ovarian weight was lower in the BET-treated group compared with the CAB-treated group (P<0.01). Figure 4 shows the results of ovarian anatomical analysis.
Fig. 3: Quantitative results of VEGF and COX-2 protein expressions in ovarian tissues of the studied groups. VEGF; Vascular endothelial growth factor, COX-2; Cyclooxygenase-2, OHSS; Ovarian hyperstimulation syndrome, BET; Betaine, CAB; Cabergoline, ****; P<0.0001, ####; P<0.0001, &; P<0.05, &&; P<0.01, $$; P<0.01, and $$$; P<0.001.

Fig. 4: Comparison of the protective effect of BET and CAB on morphometrical changes in the study groups. OHSS; Ovarian hyperstimulation syndrome, BET; Betaine, CAB; Cabergoline, ***; P <0.001, ###; P <0.001, &; P<0.05, &&; P<0.01, $$; P<0.01, and $$$; P<0.001.
Analysis of histological parameters

Histological analysis revealed that the control group showed a normal ovary with normal follicles (Fig.5A). OHSS induction caused the tunica albuginea tissue around the ovary to become thinner. In the OHSS group, there was an increase in the atretic follicles, and the number and diameter of corpora lutea in the ovarian cortex, and a decrease in the number of follicular reserves. Also, in the OHSS group, the ovarian stroma and lutein cells of the corpus luteum had numerous blood vessels, and the nuclei of the lutein cells of the corpus luteum were dense and pyknotic. The medullary part of the ovary of the OHSS group had hyaline casts and numerous dilated blood vessels. Oedema, hyperaemia, and haemorrhage were also observed (Fig.5B). These changes were reduced after treatment with CAB or BET compared to the OHSS group (Fig.5C, D). Ovaries in the BET and CAB groups showed a tissue structure similar to the control group (Fig.5E, F).

The results of the histomorphometric analysis of the ovaries are presented in Figure 6. The numbers and diameter of the corpora lutea and numbers of the atretic follicle in the OHSS group were higher, and the follicular reserve count was lower than the control group (P<0.001). Treatment with CAB or BET significantly reduced all histomorphometric changes observed in the OHSS group.

Fig. 5: These figures show the histological images of ovaries in the study groups (H&E stain, 100× magnification, scale bar: 50 μm). A. The control group, B. OHSS, C. OHSS+BET, D. OHSS+CAB, E. BET, and F. CAB groups. Superficial lining tissue of the ovary (arrowhead), primordial follicle (white arrow), primary follicle (black arrow), thickening of the connective tissue wall around the follicle (green arrow), hyperaemia in the corpus luteum (red arrow), and hyperaemia in the ovarian stroma (star) are shown. G. Histomorphometric changes. Data are presented as mean ± SEM from 10 sections per rat and five animals per group. H&E; Haematoxylin and eosin, CL; Corpus luteum, OHSS; Ovarian hyperstimulation syndrome, BET, Betaine, CAB; Cabergoline, ****; P<0.0001, ###; P<0.001, &&&; P<0.001, and $$; P<0.01.
Discussion

Increasing access to *in vitro* fertilisation has assisted numerous women worldwide who otherwise would not be able to achieve a normal pregnancy. However, an increased risk of OHSS is a potential side effect of this treatment (4, 22-25). OHSS syndrome increases the expression and secretion of VEGF, and the expression of VEGF receptors from stimulated ovaries; this, in turn, creates increased vascular permeability and ovarian disorders (5, 9). Women with OHSS have evidence of significantly increased body weight, and ovarian weight and diameter (4, 7, 26, 27). The present study showed a significant increase in delta values of body weight, and ovarian weight and diameter in the OHSS group. We observed that BET and CAB treatment attenuated the elevated levels of body weight, and ovarian weight and diameter in the OHSS group. We observed that BET and CAB treatment attenuated the elevated levels of body weight, and ovarian weight and diameter. Numerous researchers have previously shown that CAB can be used to attenuate OHSS clinical symptoms (12, 13, 16, 19, 25, 28, 29). According to our findings, induction of OHSS resulted in several histological changes in the ovarian tissue, which included increased numbers of atretic follicles, increased diameter and number of the corpora lutea, decreased follicular reserve count, and increased numbers of blood vessels in the corpora lutea and ovarian stroma. These changes have been previously reported by many researchers and support our findings (30, 31). The results of the current study showed that administration of CAB or BET significantly attenuated the histological changes induced by OHSS, which supported the results of previous researches. We observed significantly increased VEGF and COX-2 protein expressions in ovaries of the OHSS group compared to the control group (25, 32). Treatment of OHSS group with CAB or BET significantly downregulated the increased VEGF and COX-2 protein expressions. These findings agreed with previous studies (19, 25, 33) that reported significant increases in serum levels of E2 and P4 in OHSS animals. Various studies have shown that CAB administration to an experimental OHSS model reduced the increased levels of E2 and P4 (7, 12, 13, 19). Although CAB administration reduces OHSS complications and
is a definitive drug to prevent OHSS, it does have side effects (headache, peripheral oedema, shortness of breath, bleeding from the nose), contraindications (breastfeeding, cardiovascular problems, pulmonary and respiratory problems, high blood pressure), and drug interactions (vascular contraceptives, oral contraceptive pills, blood pressure-lowering drugs) (11, 14).

BET is a natural compound that is released in biological systems for protection against environmental osmotic and oxidative stressors (34, 35). BET appears to decrease VEGF protein expression and suppress new vascularisation in the ovaries of rats (17, 36-38). To the best of our knowledge, there are no reported results that show the benefits of treatment with BET compared with CAB in reducing the side effects of OHSS. Our results indicate that BET reduced VEGF and COX-2 protein expressions along with attenuation of vascularisation in cortical tissue and ovarian medulla of an OHSS animal model. In the BET-treated group, the increased body weight, and microscopic and macroscopic ovarian changes from OHSS were reduced, and this was similar to CAB, as an efficient drug for attenuation of OHSS complications. In comparison between the two treated groups, BET group had evidence of increased improvements in the effects on all molecular and histological changes compared with the CAB group.

Conclusion

Our results show a wealth of evidence to support the effectiveness of BET in reducing OHSS complications. We noted that BET administration reduced the vasodilator protein (VEGF and COX-2) expressions, improved ovarian steroidogenesis, and mitigated the histological alterations associated with OHSS. Further research is needed to optimise BET dosages and administration route. Additionally, ongoing monitoring and long-term studies are crucial to ensure the safety and efficacy of this treatment in OHSS patients.

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Authors’ Contributions

Z.B.; Is the guarantor of this work, Study design, Data analysis, Wrote the original draft, and Full access to the study data. M.R.A.; Experimental work and Study design. J.J.; Assisted with dosage design, Study design, and Data collection. M.R.T.; Performed the biochemistry, Molecular studies, Study design, and Edited the final manuscript. All authors read and approved the final manuscript.

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BET and OHSS


