

Estrogen provides neuroprotection against brain edema and blood brain barrier disruption through both estrogen receptors α and β following traumatic brain injury

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ARTICLE INFO	ABSTRACT
<p>Article type: Original article</p> <p>Article history: Received: Apr 13, 2014 Accepted: Aug 3, 2014</p> <p>Keywords: Blood-brain-barrier Brain edema ERα antagonist ERβ antagonist ICI182780 Traumatic brain injury</p>	<p>Objective(s): Estrogen (E₂) has neuroprotective effects on blood-brain-barrier (BBB) after traumatic brain injury (TBI). In order to investigate the roles of estrogen receptors (ERs) in these effects, ER-α antagonist (MPP) and, ER-β antagonist (PHTPP), or non-selective estrogen receptors antagonist (ICI 182780) were administered.</p> <p>Materials and Methods: Ovariectomized rats were divided into 10 groups, as follows: Sham, TBI, E₂, oil, MPP+E₂, PHTPP+E₂, MPP+PHTPP+E₂, ICI+E₂, MPP, and DMSO. E₂ (33.3 μg/Kg) or oil were administered 30 min after TBI. 1 dose (150 μg/Kg) of each of MPP, PHTPP, and (4 mg/kg) ICI182780 was injected two times, 24 hr apart, before TBI and estrogen treatment. BBB disruption (Evans blue content) and brain edema (brain water content) evaluated 5 hr and 24 hr after the TBI were evaluated, respectively.</p> <p>Results: The results showed that E₂ reduced brain edema after TBI compared to vehicle ($P < 0.01$). The brain edema in the MPP+E₂ and PHTPP+E₂ groups decreased compared to the vehicle ($P < 0.001$). There was no significant difference in MPP+PHTPP+E₂ and ICI+E₂ compared to TBI. This parameter in MPP was similar to vehicle. Evans blue content in E₂ group was lower than vehicle ($P < 0.05$). The inhibitory effect of E₂ on Evans blue was not reduced by MPP+E₂ and PHTPP+E₂ groups, but decreased by treatment with MPP+PHTPP or ICI. MPP had no effect on Evans blue content.</p> <p>Conclusion: A combined administration of MPP and PHTPP or ICI inhibited the E₂-induced decrease in brain edema and BBB disruption; this may suggest that these effects were mediated via both receptors.</p>

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Introduction

Traumatic brain injury (TBI) is a complex neurodegenerative disease, involving many cellular and molecular pathways, including inflammation (1). Paradoxically, the inflammatory response can lead to both aggravation and amelioration of the resulting tissue damage, depending on the tissue involved and the type of injury evoked (2). After traumatic brain injury (TBI), disruption of blood-brain-barrier (BBB) will lead to the release of neutrophils, lymphocytes and monocytes from blood to central nervous system and cause inflammatory responses (3).

Estrogen (E₂) is a cholesterol-derived hormone that is often known as a reproduction regulator and can play a protective role in different kinds of brain injuries; it has been revealed that the treatment by estradiol following ischemia, will decrease cortical injuries (4). E₂ regulates the selective permeability (5) of the BBB in several models of injury, including systemic inflammation, neurotrauma, and ischemia

(6). Previous study in our laboratory showed that the administration of estrogen and progesterone, alone and in combination after TBI decreases brain edema, blood-brain-barrier permeability, and intracranial pressure (7).

Many of E₂ functions are carried via the classic receptors, estrogen receptor alpha and beta. Estrogen receptors (ERs) are involved in signaling pathways of classical and non-classical pathways (4). Both receptors express throughout the CNS, including microglia, astrocytes, and neurons, and also in many immune system cells such as macrophages/monocytes, T and B cells and dendrites are expressed (8). It has been reported that ER α expression in ischemic brain after stroke greatly increases (9). Beta receptor agonist in animal models of arthritis and inflammatory bowel disease, has anti-inflammatory effects and suppress gene expression of pro-inflammation. Both receptors ER α and ER β are involved in cell proliferation induced by

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estradiol after ischemia (10). After brain ischemia, the expression of ER α in injured cortex increases rapidly (11). Which of these two receptors alpha and beta-estradiol-mediated control inflammation, depends on the inflamed tissue (12).

In agreement with previous studies, it was shown that estrogen reduces brain edema and BBB permeability after trauma (7, 13), although the roles of estrogen receptors ER α and ER β in the neuroprotective actions of estrogen on TBI remain poorly understood. The goal of the present study was to use these ER-selective and non-selective antagonists (MPP, PHTPP and ICI 182780) to evaluate the relative necessity of ER α - and ER β -activation in mediating the neuroprotective effect of estrogen following diffuse TBI in OVX rats.

Materials and Methods

Animals

This study was conducted in accordance with the guidelines for the animal experimental protocols of Kerman University of Medical Sciences. The protocol was approved by the ethics committee (no. KNRC/90-40) of this University, in accordance with the internationally accepted principles for laboratory animal use and care, as found in the European Community guidelines (EEC Directive of 1986; 86/609/EEC) or US guidelines (NIH publication #85-23, revised in 1985). Adult female Albino N Mary rats (weighing 200-250 g) were housed in an air-conditioned room at 22-25°C, with a 12 hr light: 12 hr dark cycle and free access to food and water.

Ovariectomy surgery

In the first step, animals were anesthetized by injection of 60 mg/kg thiopental (IP), sub abdominal area was shaved and an incision of 1 cm was made. Afterwards, skin, fascia, and abdominal muscles were opened. Both fallopian tubes and vascular base of the ovaries were ligated in the proximal parts and then cut from the distal parts. Abdominal cavity was irrigated by 2 ml of normal saline. The incision was closed with a 2-0 silk surgical suture. To avoid interference due to the estrus cycle, all experimental animals were ovariectomized (OVX) 2 weeks before the experiments (14).

Experimental protocols

Before induction of TBI, ovariectomized rats were randomly divided into 10 groups, as follows: sham (animals that underwent ovariectomy 2 weeks before the start of the experiment and underwent false brain trauma under anesthesia but did not receive hormones or vehicle), TBI (brain injury was induced 2 weeks after ovariectomy), E₂, MPP+E₂, PHTPP+E₂, MPP+PHTPP+E₂, MPP, ICI+E₂, oil (sesame oil, which was used as estrogen solvent) and DMSO (dimethyl sulfoxide, which was used as antagonists solvent). (n=6 in each group).

E₂(33.3 μ g/Kg) or oil were administered (IP) 30 min after TBI (15, 16).

MPP(1,3-Bis(4-hydroxyphenyl)-4-methyl-5-[4-(2-piperidinyloxy)phenol]-1H-pyrazole hydrochloride), ER α antagonist, and PHTPP(4-[2-Phenyl-5, 7-bis(trifluoromethyl) pyrazolo [1,5-a] pyrimidine-3-yl] phenol), ER β antagonist, or DMSO were used as follows: 1 dose (150 μ g/Kg) of each of MPP and, PHTPP was injected two times, 24 hr apart, before TBI and oestrogen treatment (17). Then treated with E₂ (33.3 μ g/Kg) 30 min after TBI (13). One group received only an injection of MPP(150 μ g/Kg) 30 min after TBI (17). 4 mg/kg ICI182780 was injected similar to the other two antagonists (18).

Induction of diffuse traumatic brain injury (TBI)

All animals were intubated before TBI. The TBI was induced by the Marmarou method (16), using a TBI induction device made by the Department of Physiology, Kerman University of Medical Sciences. The protocol was as follows: a weight of 300 g was dropped from a 2 m height onto the head of the anesthetized rat while a metal disc (stainless steel, 10 mm in diameter, 3 mm thick) was attached to the animal's skull. After induction of the trauma, the rats were immediately connected to a respiratory pump until spontaneous breathing had been restored.

Determination of brain edema

The brain edema of each animal was assessed by measuring brain water content 24 hr after TBI. Anesthetized animals were sacrificed by cervical dislocation; the brain was removed, and brain samples were placed in pre-weighed glass vials and weighed (wet weight). The lids were removed and the vials placed in an incubator (Mettler, Germany) at 60°C for 72 hr, and then reweighed (dry weight). The percentage of water in each sample was then calculated using a formula published previously (16, 19): $(100 * [(wet\ weight - dry\ weight) / wet\ weight])$.

Determination of blood-brain barrier (BBB) disruption

The degree of BBB disruption was assessed by measuring Evans blue dye leakage (16). Briefly, Evans blue dye was dissolved in 0.01 mol/l PBS at a concentration of 2%, then the dye (2 ml/kg IV) was injected into the tail vein 4 hr after TBI, as a BBB permeability tracer. The rats were then deeply anesthetized and transcardiac perfused with 200 ml heparinized saline through the left ventricle to remove the intravascular dye. The brains were removed, then dissected, weighed, and stored at -80°C for quantitative measurement. Brain samples were homogenized in 1 ml of 0.1 mol/l PBS, and 0.7 ml of 100% (w/v) trichloroacetic acid was added, and centrifuged. After centrifugation for 30 min at 1000 g, the absorbance of Evans blue in supernatant

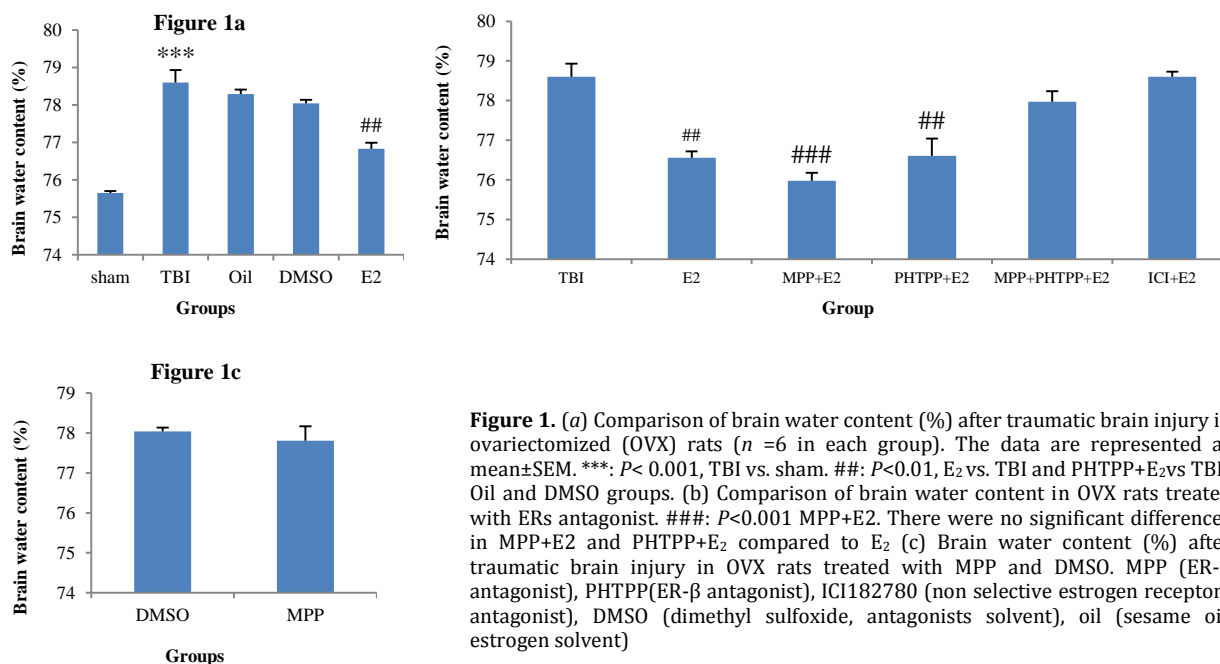


Figure 1. (a) Comparison of brain water content (%) after traumatic brain injury in ovariectomized (OVX) rats ($n = 6$ in each group). The data are represented as mean \pm SEM. ***: $P < 0.001$, TBI vs. sham. #: $P < 0.01$, E₂ vs. TBI and PHTPP+E₂ vs TBI, Oil and DMSO groups. (b) Comparison of brain water content in OVX rats treated with ERs antagonist. ###: $P < 0.001$ MPP+E₂. There were no significant differences in MPP+E₂ and PHTPP+E₂ compared to E₂ (c) Brain water content (%) after traumatic brain injury in OVX rats treated with MPP and DMSO. MPP (ER- α antagonist), PHTPP (ER- β antagonist), ICI182780 (non selective estrogen receptors antagonist), DMSO (dimethyl sulfoxide, antagonists solvent), oil (sesame oil, estrogen solvent)

was measured at 610 nm using a spectrophotometer (UV/VIS, Spectrometer, UK). The amount of extravasated Evans blue dye was quantified as $\mu\text{g/g}$ brain tissue.

Statistical analysis

Quantitative data were expressed as mean \pm SEM. The data were analyzed by parametric analysis of variance (ANOVA) or independent t test. Fisher's LSD was employed for the post-hoc analysis. The criterion for statistical significance was sign at $P < 0.05$.

Results

Brain edema

Changes in the brain water content of the ovariectomized rats are shown in Figure 1. Figure 1a shows that the brain water content in the TBI ($78.6 \pm 0.33\%$) group was significantly higher than in sham ($75.65 \pm 0.001\%$, $P < 0.001$) group and brain water content in the group treated with E₂ ($76.76 \pm 0.17\%$) was significantly lower than those in the oil ($78.32 \pm 0.15\%$) and DMSO ($78.4 \pm 0.22\%$) groups, ($P < 0.01$). Figure 1b shows the effects of administration of estrogen receptors antagonists. In the presence of MPP and PHTPP, brain water content decreased and was $74.86 \pm 0.54\%$ in MPP+E₂ group and $75.41 \pm 0.88\%$ in PHTPP+E₂ group. There was no significant difference in MPP+PHTPP+E₂ and ICI+E₂ compared to TBI. Figure 1c shows that the brain water content in the group treated with MPP ($78.06 \pm 0.32\%$) has no significant difference with the water content in DMSO group.

Extravasation of Evans blue content

The effects of estrogen and ERs antagonists on BBB disruption after TBI are shown in Figure 2.

Figure 2a shows that the Evans blue content in the TBI ($37.18 \pm 0.4 \mu\text{g/g}$ tissue) group was significantly higher than in the sham ($33.25 \pm 0.05 \mu\text{g/g}$ tissue, $P < 0.05$) group and the Evans blue content in E₂ ($33.5 \pm 0.65 \mu\text{g/g}$ tissue) was lower than oil ($37.67 \pm 0.44 \mu\text{g/g}$ tissue), TBI, and DMSO ($37.34 \pm 0.22 \mu\text{g/g}$ tissue) groups. $P < 0.05$. Figure 2b shows the effect of administration of estrogen receptors antagonist. In the presence of MPP and PHTPP, Evans blue content decreased and was $30.78 \pm 0.31 \mu\text{g/g}$ tissue in MPP+E₂ and $31.4 \pm 0.38 \mu\text{g/g}$ tissue in PHTPP+E₂ groups ($P < 0.001$). There were no significant differences in MPP+PHTPP+E₂ & ICI+E₂ compared to TBI. Figure 2c shows that the Evans blue content in the groups treated with MPP (36.46 ± 0.68) has no significant difference with the Evans blue in the E₂ group.

Discussion

Previous studies have reported, the neuroprotective effects of estrogen on brain edema and blood-brain barrier permeability (7, 13). In the present study, the roles of alpha and beta estrogen receptors in the development of these effects were investigated. The main findings of this research are as follows: 1-estrogen exerts its anti-edema effects through both receptors, so that any differences between the results of combined use of selective antagonists and a non-selective antagonist do not exist. 2-BBB protection after TBI by estrogen is mediated by both receptors, which is similar to the results of the anti-edema effect of estrogen. 3-The estrogen receptor alpha antagonist (MPP) has no agonistic effect on its receptor.

TBI caused an increase in brain water content and BBB permeability, these increases were inhibited by estrogen compared to vehicle (oil).

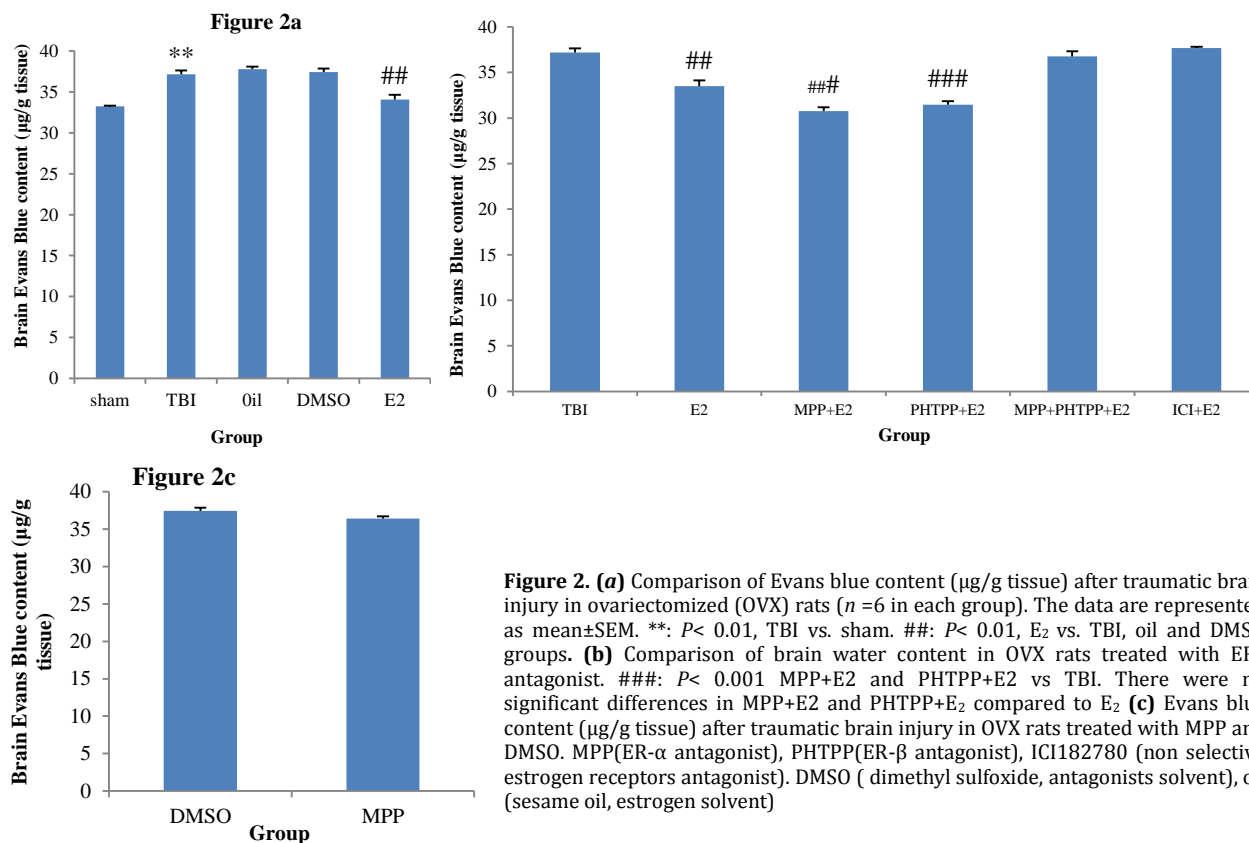


Figure 2. (a) Comparison of Evans blue content ($\mu\text{g/g}$ tissue) after traumatic brain injury in ovariectomized (OVX) rats ($n=6$ in each group). The data are represented as mean \pm SEM. **: $P < 0.01$, TBI vs. sham. ##: $P < 0.01$, E₂ vs. TBI, oil and DMSO groups. (b) Comparison of brain water content in OVX rats treated with ERs antagonist. ###: $P < 0.001$ MPP+E₂ and PHTPP+E₂ vs TBI. There were no significant differences in MPP+E₂ and PHTPP+E₂ compared to E₂. (c) Evans blue content ($\mu\text{g/g}$ tissue) after traumatic brain injury in OVX rats treated with MPP and DMSO. MPP (ER- α antagonist), PHTPP (ER- β antagonist), ICI182780 (non selective estrogen receptors antagonist). DMSO (dimethyl sulfoxide, antagonists solvent), oil (sesame oil, estrogen solvent)

There have been a number of reports supporting a neuroprotective role for estrogen after various insults to the CNS. Ovariectomy increases ischemic brain damage in females, and estradiol exerts neuroprotective and anti-inflammatory action against the ischemic brain when administered immediately upon ovariectomy (20). E₂ regulates the selective permeability of the BBB in several models of injury (5). The BBB permeability decreased in E₂-treated mice (4). Recently, our research showed that estrogen loss facilitates, whereas administration of a different dose of estradiol could prevent edema formation (7) and destruction of the BBB after TBI (13). A significant reduction in the brain water content after TBI was induced by estrogen; there is no difference between estrogen and solo administration of antagonist groups. This suggested that neither selective ER α antagonist (MPP) nor ER β antagonist (PHTPP) can abolish these effects of estrogen.

On the other hand, administration of the two antagonists together (MPP + PHTPP) or the use of non-selective estrogen receptor antagonist (ICI182780) eliminates the inhibitory effect of estrogen on brain edema and BBB permeability, this means that, the effects of estrogen were not inhibited by the administration of antagonists alone, but when these antagonists were administered in combination, they have an inhibitory effect. It is suggested that, probably both receptors are involved in the

development of neuroprotective effects of estrogen. We found that more than one receptor is activated by estrogen at the same time, perhaps activation of both receptors at the same time has reciprocal modulatory effects. Whenever the effect of estrogen was observed during receptor inhibition, it was caused by activation of the other receptor at the same time. To the best of our knowledge, the roles of neither ER α antagonist nor ER β antagonist in TBI have been reported thus far. Because this study is the first to use a selective ER α and ER β antagonist to investigate the effects on TBI. However, the effects of these receptors or antagonists in other non-TBI studies, and other estrogen-mediated effects have been demonstrated, including: The receptors ER α and ER β are essential for E₂-mediated regulation of BBB permeability, anti-inflammatory properties, and regulation of neutrophil recruitment into the brain (4). Both ER receptors are involved in the control of inflammation by estradiol (21), neuromodulation, and neuroprotection processes after brain injury (22). The affinity of MPP to ER α is only 11% of that of E₂ (23). ER antagonists (MPP and ER β antagonist (R,R)-THC) treatments alone did not significantly change NPY release levels (24). MPP failed to attenuate the anorexigenic effects of estradiol and an ER α agonist in OVX rats (25). MPP did attenuate the estrous-related decrease in food intake in cycling rats (25). MPP shows no activation of either receptor subtype (23). 17 β estradiol protects dopamine neurons

from injury, an effect that was blocked by ICI 182,780 (26). ICI 182,780 after ischemia abolished estrogen protection (27). Estrogen neuroprotection was related to microglial activation of estrogen receptors (ERs), this protective effect was overridden by pretreatment with ICI 182,780, the specific ER α , only partially blocked the effects of estrogen (6).

An alternate mechanism which could explain the differential effects of ICI and MPP, is an involvement of non-genomic/non-classical signaling mechanisms activated by anti-estrogen-occupied ER. Indeed, ICI-mediated Erk1/2 (extracellular signal-regulated kinases) activation appears to be critical for modulation in immature cerebellar neurons (28). Some studies suggest that MPP acts as an ER α antagonist following *in vitro* applications, but exerts mixed ER α agonist/antagonist actions following *in vivo* applications (29). It is possible that the MPP concentration in this experiment was too low to completely compete with E2 for binding to ER α , because it is reported that MPP is needed in high excess to fully antagonize E2 (30). There are several probable mechanisms by which ER α and ER β attenuate BBB permeability and brain edema after TBI that result from the inflammatory response; up regulation of defense-related genes, IL-6, MMP-9, and some of phagocytic receptors (31), protection of cortical cells against oxidative glutamate toxicity (32), neuroprotection of estrogen against activated microglia (33) and inhibition of the cytoplasmic transport of NF- κ B (nuclear factor kappa-light-chain-enhancer of activated B cells) (34).

There is, however, some controversy. MPP reduced the E2 effect on protein vitellogenin (35). MPP and PTHPP inhibited DNA synthesis, oxygen species (ROS) formation induced by estrogens according to their receptors affinity (36). Experiments with PHTPP, indicated that E2-mediated neuroprotection against NMDA toxicity was mediated via ER β receptor (37). Many studies have reported a primary role for ER α as a regulator of the anti-inflammatory properties of estradiol, including the anti-inflammatory and neuroprotective activity of estradiol (38), neuroprotection after ischemia and inhibition of microglial activation (39). Studies involving transgenic mice with targeted deletion of either ER α or ER β indicate that ER α , but not ER β , is required for the E2-mediated suppression of BBB permeability (8). In addition, a growing number of studies have reported anti-inflammatory roles for ER β , such as neuroprotection after ischemia and, inhibition of microglia activity (40). In mammals, the actions of MPP vary, depending on the species, physiological conditions, dosing method, and tissue (17, 41-43).

Since currently, there is no evidence that PHTPP, exerts the mixed agonist/antagonist properties that are often seen for MPP (29), in another part of the present study, MPP was used to replace estrogen

after TBI, it was observed this compound did not reduce brain edema, and BBB permeability. In other words, MPP has no agonist effect on ER α . This finding is consistent with other studies that have reported that MPP counteracted the effects of PPT (estrogen receptor agonist) but had little effect on its own (44). On the other hand, it has also been reported that estrogen receptor/(ER α), even in absence of estrogen (E2), plays a critical role in lactotroph homeostasis (45). *In vivo* tests of MPP's effects on food intake revealed estrogenic activity, suggesting that MPP functions as a SERM (Selective estrogen receptor modulator), rather than as an ER α antagonist (46).

Conclusion

The present study showed that the administration of PPT or PHTPP alone will not lead to reduced brain edema, and prevention of BBB disruption following TBI, but a combined administration of MPP and PHTPP or ICI inhibited the E2-induced decrease in brain edema and BBB disruption, this may suggest that estrogen neuroprotection was mediated via both ER α and ER β . Both of them can play roles in classic and non-classic pathways. Further study is needed and intracellular mechanism activation in the presence of E2 should be evaluated. These data, makes them exciting therapeutic targets for further study, in addition, perhaps it is essential that non-genomic mechanisms are also investigated.

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