Expression of HSP70 in cerebral ischemia and neuroprotervative action of hypothermia and ketoprofen

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ABSTRACT

Heat shock proteins (HSPs) are molecular chaperones that bind to other proteins to shepherd them across membranes and direct them to specific locations within a cell. Several injurious stimuli can induce Hsp70 expression, including ischemia. This study aimed to investigate the pattern of expression of protein (immunohistochemistry) and gene (real-time PCR) Hsp70 in experimental focal cerebral ischemia in rats by occlusion of the middle cerebral artery for 1 hour and the role of neuroprotection with hypothermia (H) and ketoprofen (K). The infarct volume was measured using morphometric analysis defined by triphenyl tetrazolium chloride. It was observed increases in the protein (p=0.0001) and gene (p=0.0001) Hsp70 receptor in the ischemic areas that were reduced by H (protein and gene: p<0.05), K (protein: p<0.001), and H+K (protein: p<0.01 and gene: p<0.05). The Hsp70 increases in the ischemic area suggests that the Hsp70-mediated neuroexcitotoxicity plays an important role in cell death and that the neuroprotective effect of both, H and K are directly involved with the Hsp70.

Key words: Hsp70 expression, cerebral ischemia, neuroprotection, hypothermia, ketoprofen.

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Focal cerebral ischemia produces changes in gene transcription and translation that are related to the extent of blood flow reduction. While total protein synthesis is generally reduced with ischemia, some genes, including the c-fos immediate ear-
ly gene and the 70 kD heat shock stress gene Hsp70, are induced with ischemia.1,3

The heat shock protein (HSP) family is induced in the brain by several insults including excitotoxin exposure, trauma, elevated body temperature and cerebral ischemia.4-7 The Hsp70 expression is elevated mainly in the area of ischemic penumbra8-10 and detection of the Hsp70 gene exacerbate apoptosis after focal cerebral ischemia in mice.11 Olsson et al.12 suggested that the increased expression of the Hsp70 is an indicator of cellular stress. Recent work has shown that HSP proteins modulate inflammatory response either by potentiating13 or inhibiting them14 and suggest that Hsp70 brain protection against ischemia is done through an anti-inflammatory mechanism.

Several pharmacological procedures have been used in an attempt to reduce the area involved in the ischemic process. Among them, hypothermia15,16 and anti-inflammatory drugs17-19 and hypothermia in combination with anti-inflammatory drugs20, has proved to be effective in providing protection after cerebral ischemia. Anti-inflammatory agents are used to attenuate the inflammatory response triggered by the ischemic phenomenon, with emphasis on ketoprofen21-23, a non steroid agent with a very potent analgesic and anti-inflammatory action, acting as a possible antagonist of the N-methyl-d-aspartate receptors (NMDARs)24. Its neuroprotective effect was attributed by Asanuma et al.25 to decrease the antagonism of acetylcholine receptors caused by transient ischemia in gerbil. Experimental studies have also demonstrated the protective action of ketoprofen against ischemia in neurons of the retina and brain tissues in transient focal ischemia26.

This study aimed to investigate the pattern of expression of protein and gene Hsp70, in an experimental model of transient focal cerebral ischemia without reperfusion and the possible neuroprotective effect of hypothermia and ketoprofen.

**METHOD**

One hundred and twenty adult male rats (*Rattus norvegicus*) weighing 280-310 g were used and the animals were randomly divided into 3 experimental groups: control (C), 20 animals sacrificed without being submitted to the surgical procedure; sham (S), 20 control animals submitted to complete simulation of the surgical procedure but without obstruction of the middle cerebral artery (MCA) and then sacrificed, and ischemic (I), 80 animals submitted to focal ischemia by occlusion of the MCA for 1 hour and then sacrificed. This group was subdivided into 4 subgroups of 20 animals each: ischemic subgroup (Sg I), ischemic subgroup submitted to hypothermia (Sg IH), ischemic subgroup treated with ketoprofen (Sg IK), and ischemic subgroup submitted to hypothermia and treated with ketoprofen (Sg IHK). Ten animals from each experimental group, for a total of 60 animals, were used for the delimitation of the ischemic area using a morphometric method for controlling the efficacy and the extension of the ischemia. The experiments were carried out according to the Ethical Principles for Experimental Animals (COBAO) and the study was approved by the Animal Experimentation Committee (CETEA) of the Faculty of Medicine of Ribeirão Preto, University of São Paulo (Protocol nº 0347/2005).

**Induction of cerebral ischemia**

All animals were partially anesthetized by halothane inhalation and intubated with an orotracheal cannula. At two times during the ischemic period, arterial blood samples were collected for the determination of glycemia, paco2, pao2 and pH. The animals in the Sg IH subgroup were submitted to moderate intra-ischemic hypothermia (32-34°C) by anesthesia itself and by air conditioning the room, starting 15 minutes before the surgical procedure to induce ischemia. The animals in the Sg IK subgroup received an intravenous injection of ketoprofen at the dose of 10 mg/kg body weight 15 minutes before the induction of ischemia, and the animals in the Sg IHK subgroup were submitted to intra-ischemic hypothermia and similarly injected with ketoprofen. MCA occlusion was carried out through the external carotid artery which was ligated cranially and sectioned for the retrograde introduction of a 2.5 cm long obstructive 4-0 mononylon suture with one end thickened with silicone over an extension of 5 mm27. The suture was introduced until to reach the common carotid artery and then cranially progressed through the internal carotid artery until to reach and to obstruct the MCA.

**Morphometric analysis of infarct volume**

Infarct volumes were measured by morphometric analysis of infarct areas that were defined by 2,3,5-triphenyl tetrazolium (TTC). The brains of the animals were removed and cut into coronal sections using a rat brain matrixe which were placed in 10% buffered formalin solution. Cross-sectional area of the TTC-stained region for each brain slice was determined by analysis of infarct areas that were defined by 2,3,5-triphyl tetrazolium (TTC). The brains of the animals were removed and cut into coronal sections using a rat brain matrixe which were placed in 10% buffered formalin solution. Cross-sectional area of the TTC-stained region for each brain slice was determined by analysis of infarct areas that were defined by 2,3,5-triphyl tetrazolium (TTC). The brains of the animals were removed and cut into coronal sections using a rat brain matrixe which were placed in 10% buffered formalin solution. 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and Pierce solution to block endogenous peroxidase and biotin, respectively. Sections were subsequently incubated with primary antibody against Hsp70 (1:200 dilution - CHEM - AB9864 - Chemicon*) and with biotin-conjugated secondary anti-rabbit antibody (1:1000; Vector Laboratories Inc., Burlingame, CA) and streptavidin-conjugated peroxidase (Vecstatin Abc kit, Vector Laboratories Inc.). Color was developed by the addition of DAB (Sigma Chemical, St. Louis, MO). To evaluate the background reaction, procedures were also performed in sections incubated only with the secondary antibodies (indirect technique) or in the absence antibodies (direct technique). For the analysis of protein expression of Hsp70 a field in an increase of 400× in the area where there was greater concentration of positive cells or marked (areas of “hot spots”) was selected in each layer of immunohistochemical reaction. The number with positive staining for Hsp70 was measured by using a camera (Axio Cam, Zeiss, Germany) and the software Axiovision 4.6 (Zeiss, Germany).

The method used for counting the cells positive to antibody study was to quantify “in crosses” of cytoplasmic immunoreactivity of cells of nervous tissue. The fields chosen in the slides were evaluated as the percentage of cells marked and graded from 0 (zero): no mark, up to 25%: 1 + (low expression), 25% to 50%: 2 ++ (intermediate expression); 50% to 75%: 3 +++ (high expression) and over 75% of cells marked: 4 +++++ (high expression).

RNA extraction and cDNA synthesis
After removal of the brain, the cortex of the left cerebral hemisphere was isolated and a sample measuring 7 mm in diameter, was obtained by biopsy from the central ischemic region, corresponding to the irrigation zone of the MCA. The samples were placed in cryotubes and stored in liquid nitrogen at -196°C until the time for RNA extraction. For this procedure, 250 μl PBS + 750 μl TRIZOL were added to the brain tissue samples which were then lysed in a Polytron homogenizer for about 3 minutes. Next, total RNA was extracted by the TRIZOL method (Invitrogen, Carlsbad, CA, USA) according to manufacturer instructions. Complementary DNA (cDNA) was constructed by reverse transcription using the enzyme Superscript II and the DNA obtained was amplified by real-time quantitative PCR (RQ-PCR).

Analysis of gene expression patterns by RQ-PCR
For the quantitative analysis of the gene under study, Hsp70 (Rn 01525984), we used the commercially available system TaqMan Assay-on-demand, which consists of oligonucleotides and probes (Applied Biosystems, Foster City, CA, USA). Reverse transcription was performed using 4 μg total RNA for each sample in 40 μL of the total reaction mixture, with the addition of oligo (dT)20 (Cat. No. 18418-020, Invitrogen) and Superscript II reverse transcriptase (Cat. No. 18064-022, Invitrogen), according to manufacturer instructions. The cDNA obtained was diluted 1:10 and 2.5 μL was used for each 15 μL of the RQ-PCR mixture using the TaqMan Master Mix (Applied Biosystems).

All reactions were carried out in duplicate and analyzed with the 7500 Sequence Detection System apparatus (Applied Biosystems). Data were analyzed using the ABI-7500 SDS software. The maximum standard deviation between duplicates was 10%. The total RNA absorbed was normalized on the basis of the Ct value for the GAPDH gene (Rn_99999916). The variation of expression among samples was calculated by the 2^-ΔΔCt method, with the mean delta Ct value for a group of 20 samples from control rats being used as a calibrator.

Statistical analysis
The expression of the gene and protein under study showed normal distribution in the various groups. Data concerning the mean gene and protein expression in the various groups were analyzed statistically by one-way ANOVA followed by the Bonferroni post-test using the GraphPad Prism software, version 4.00 for Windows, (GraphPad Software, San Diego, CA, USA). The level of significance was set at p<0.05 for two-tailed tests.

RESULTS
Rats subjected to a 60 minutes period by occlusion of the MCA showed after morphometric analysis that the infarct area corresponding to the territory of irrigation of the MCA was clearly observed in the animals of the ischemic group.

The analysis of the percentage of the volume of ischemic area in relation to the normal hemisphere among the animals of the ischemic group (SgI, SgIK and SgIHK) revealed a smaller area of infarction in relation to animals of the ischemic subgroup (Sgl) (p<0.0001, one-way ANOVA, p=0.0001, Bonferroni post-test, and similar areas to that observed in animals of the control group.

The analysis of protein expression of Hsp70 by immunohistochemistry (Fig 1) showed that it was intermediate (++) for all animals in subgroups Sgl, SglH and SglHK, and that average percentage of positive cells were 40.96%; 36.73% and 36.20%, respectively. The expression was also intermediate (++) for all animals of the subgroup SglK, with low intensity (+) only in animal 2 and with the average percentage of positive cells of 28.61%.

Fig 2 shows mean and standard deviation of the percentage of positive cells to Hsp70 antibody in the ischemic group. There was a statistically significant difference (p=0.0001 one-way ANOVA) between the subgroups
SgIxSgIH (p<0.05), SgIxSgIK (p<0.001), SgIxSgIHK (p<0.01), Bonferroni post test.

Fig 3 illustrates the mean expression of the Hsp70 gene in the experimental groups. The expression in the animals of the ischemic group was higher when compared with control and sham (p=0.0001, one-way ANOVA, p<0.05, Bonferroni post-test), and it was lower in the animals of the subgroups SgIH and SgIHK compared to control, sham and ischemic (SgI) animals (p=0.0001, one-way ANOVA, p<0.05, Bonferroni post-test).

DISCUSSION

Cerebral ischemia is one of the devastating neuropathological disorders known to humans. Though several putative molecules have been investigated for pharmacological intervention for preventing or reducing cerebral ischemia, no successful therapy has been achieved24.

The model of ischemia induction by occlusion of the MCA with an obstructive suture has been used by several investigators and has been shown to produce constant and reproducible focal injuries19,22. In our study, morphological measure of the infarct volume was randomly performed in 10 animals of each experimental group for controlling the occurrence of ischemia. All animals submitted to ischemia showed some evidence of infarct in the ischemic hemisphere. The sham group, consisting of animals subjected to complete simulation of the surgical procedure, but without obstruction of the MCA, was used in order to demonstrate that the surgical procedure that these animals are subjected to access to the MCA does not affect the results obtained and therefore different from control animals, which does not undergo any procedure before the sacrifice.

Modulation of inflammatory response by Hsp70 is controversial: some suggest potentialization by this protein12 and others inhibition13,14. Zheng et al14, suggest that overexpression of Hsp70 not only protect brain against ischemia but also appear to do via an anti-inflammatory mechanism.

As several pathways leading to cell death are activated after an ischemic insult, effective neuroprotection might require a combination of drugs that target distinct pathways during the evolution of ischemic injury. This has led to a number of papers studying combinations of pharmacological agents25-27 and multiple mechanisms for hypothermia-induced neuroprotection have already been identified28-30.

Using a global model of cerebral ischemia in mice, Olsson et al6 noted a change in Hsp70 expression in the
CA1-CA3 regions of the hippocampus. When coupled with hypothermia (33°C) there was a decrease in the expression of Hsp70 and consequent decrease in cellular damage. The authors suggest that increased expression of Hsp70 in cerebral ischemia is an indicator of cellular stress. Our results corroborated the data of literature that decrease of Hsp70 lead to an neuroprotector effect and this effect also was observed when this protein with hypothermia and ketoprofen alone as ketoprofen plus hypothermia. More studies with this drug are necessary for better understanding participation of the Hsp70 in the inflammatory response and especially its actions as anti-inflammatory agent in ischemic brain.

In conclusion, the increased expression in the HSP70 after experimental ischemia suggests its involvement with cerebral ischemia. The neuroprotective agents associated with ischemia evaluated in this study (hypothermia, ketoprofen and hypothermia+ketoprofen) proved to be effective. This was suggested by the reduction of both gene and protein expression of HSP70. The association between hypothermia+ketoprofen do not proved to be more effective then the isolated action of these two neuroprotective agents.

REFERENCES