BASIC RESEARCH

MEMANTINE PREVENTS CARDIOMYOCYTES NUCLEAR SIZE REDUCTION IN THE LEFT VENTRICLE OF RATS EXPOSED TO COLD STRESS

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doi: 10.1590/S1807-59322009000900014

Meneghini A, Celso Ferreira, Luiz Carlos de Abreu, Vitor E. Valenti, Marcelo Ferreira, Celso F. Filho, Neif Murad. Memantine prevents nuclear size reduction in cardiomyocytes in the left ventricle of rats exposed to cold stress. Clinics. 2009;64(9):921-6.

OBJECTIVES: Memantine is an N-methyl-d-aspartate (NMDA) glutamate receptor antagonist used to treat Alzheimer’s disease. Previous studies have suggested that receptor blockers act as neuroprotective agents; however, no study has specifically investigated the impact that these drugs have on the heart. We sought to evaluate the effects of memantine on nuclear size reduction in cardiac cells exposed to cold stress.

METHOD: We used male EPM-Wistar rats (n=40) divided into 4 groups: 1) Matched control (CON); 2) Memantine-treated rats (MEM); 3) Rats undergoing induced hypothermia (IH) and 4) Rats undergoing induced hypothermia that were also treated with memantine (IHM). Animals in the MEM and IHM groups were treated by oral gavage administration of 20 mg/kg/day memantine over an eight-day period. Animals in the IH and IHM groups were submitted to 4 hours of hypothermia in a controlled environment with a temperature of -8ºC on the last day of the study.

RESULTS: The MEM group had the largest cardiomyocyte nuclear size (151 ± 3.5 μm³ vs. CON: 142 ± 2.3 μm³; p<0.05), while the IH group had the smallest mean value of nuclear size. The nuclear size of the IHM group was preserved (125 ± 2.9 μm³) compared to the IH group (108 ± 1.7 μm³; p<0.05).

CONCLUSION: Memantine prevented the nuclear size reduction of cardiomyocytes in rats exposed to cold stress.

KEYWORDS: Memantine; Cardiac myocytes; Cardiotonic agents; Cell nucleus structures; Heart.

INTRODUCTION

Reduced body temperature induced by exposure to a cold surrounding environment is considered a physiological stressor.1-3 Neurogenic lesions related to myocardial hypertrophy and changes in myocardial tissue metabolism4 can be also caused by the reaction of cardiovascular tissue to cold stress, and these lesions are hypothesized to be a marker for subsequent cardiovascular disease as well as a predictor of hypertension.5 Recently, this model was used to investigate several aspects of cardiac injury.6 Interestingly, a previous study indicated that cold stress acutely induces a reduction in the nuclear size (hence atrophy) of cardiomyocytes in rats,7 which is in contrast to the finding that is observed in myocardial hypertrophy. This reduction in nuclear size has been hypothesized to be a marker of ischemia during periods of physiological stress.7

It has been well established that glutamate excitotoxicity triggers neurodegeneration in patients with medical conditions that can lead to acute brain injuries such as stroke, status epilepticus or head trauma. Drugs that block N-methyl-d-aspartate (NMDA) glutamate receptors have been shown to be neuroprotective in animal models of these medical conditions.8,9 Memantine is a non-competitive antagonist of NMDA receptors and is currently used to treat patients

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Received for publication on May 18, 2009
Accepted for publication on June 24, 2009
with moderate to severe Alzheimer’s disease to improve cognition.\textsuperscript{10,12} Its mechanism of action is thought to be related to its effects on Ca\textsuperscript{2+} homeostasis. An increase in cytosolic Calcium (Ca\textsuperscript{2+}) is associated with functional impairment of many organelle and is also strongly associated with apoptosis.\textsuperscript{13} Previous studies have provided evidence that memantine is able to prevent ischemic injuries in the retina and in neurons exposed to different aggressive agents (that lead to increases in cytosolic Ca\textsuperscript{2+}) due its effects on Ca\textsuperscript{2+}.\textsuperscript{14,15}

Although various studies have suggested that NMDA receptor antagonists have neuroprotective properties, no previous study has specifically investigated the influence of this class of drugs may have on the heart. Previous studies, however, have reported an association between memantine and the prevention of ischemia under conditions of cellular stress.\textsuperscript{14,16} Therefore, we sought to evaluate the effects of memantine on the nuclear size of cardiomyocytes in the left ventricle of rats exposed to cold stress.

**METHODS**

**Animals**

Experiments were conducted on forty adult male Wistar rats that weighed 200-250 g. The ambient temperature of the environment in which the rats were housed was 22\degree C, with the humidity at \textasciitilde 60\%. The rats were kept on a 12-hour light/12-hour dark cycle. Animals had free access to food and water. Animals were randomized into four groups: 1) the control group (CON, n=10), in which rats were treated with gavage administration of 1 mL of water for eight consecutive days; 2) the memantine group (MEM, n=10), in which rats were treated with gavage administration of 1 mL containing 20 mg/kg of memantine for eight consecutive days; 3) the induced hypothermia group (IH, n=10), in which rats were treated with gavage administration of 1 mL of water for eight consecutive days and exposed to a \textasciitilde 8\degree C environment for four hours on the last (8\textsuperscript{th}) day of the study and; 4) the memantine plus induced hypothermia group (IHM, n=10), in which rats were treated with gavage administration of water (1 mL) containing 20 mg/kg memantine for eight consecutive days and exposed to a \textasciitilde 8\degree C environment for four hours on the last (8\textsuperscript{th}) day of the study. All experiments were performed in accordance with the ethical guidelines of the National Institutes of Health Guide for the Care and Use of Laboratory Animals and were approved by the Ethical Committee for research at our University (number 003/08).

**Induced Hypothermia Procedure**

Rats were exposed to cold stress, which was induced by placing them in wire mesh cages in an open refrigerated compartment at \textasciitilde 8\degree C for 4 hours. Rats were exposed to this environment only once and their behavior was observed throughout the stress experiment.\textsuperscript{7} Rats’ body temperatures were monitored and maintained near 37\degree C. No rats died during the induced hypothermia procedure.

**Histological Examination**

To verify if the exposure to cold stress was sufficient to cause stress responses characterized by lipid and glycogen depletion in adrenal gland and liver, we evaluated lipid depletion in adrenal gland cortical cells and glycogen depletion in hepatocytes. After administration of an adequate level of ether anesthesia, we examined the rats’ tail tone and response to external stimuli before and during surgical procedure through the evaluation of vibrissa movements. All animals were then submitted to laparotomy. Two pieces of the left lobe of the liver and right adrenal gland were removed for microscopy investigation. These specimens were cut into small pieces (1 mm\textsuperscript{3}), post-fixed in 1% OsO\textsubscript{4} solution for 2 hours, dehydrated and embedded in araldite. Silver or gray thin sections (60-90 nm) were produced using a Porter-Blum MT-B ultramicrotome. The sections were then mounted on copper silver grids with 200 patches and stained with uranyl acetate and lead citrate. We presented data regarding lipid depletion in adrenal gland cortical cells and glycogen depletion in hepatocytes, respectively, as staining intensity levels (+=small intensity; ++++=high intensity).\textsuperscript{6}

**Nuclear Volume Measurement**

All animals were also submitted to thoracotomy. The thorax of each rat was opened and the left ventricle was exposed and removed. Fragments of heart material were fixed in Bouin’s solution, mounted and paraffinized. Sections measuring 10\mu m were stained with hematoxylin-eosin. In order to estimate the nuclear size of each cell, karyometry was used according to the same principles described in a previous publication.\textsuperscript{17} Morphometric evaluation was performed using the Quantimet Color Option (Leica, Cambridge) image analysis system. Measurements of cardiomyocyte nuclear parameters were performed exclusively on clearly visible longitudinal sections of muscle fibers in which cardiomyocyte nuclei had a clear outline. To calculate nuclear volume, we used the following equation proposed by Salvatore:\textsuperscript{17}

\[
V = \frac{A^2 \times B}{1.91}
\]

\(V\) = volume; \(A\) = smaller axial measure; \(B\) = bigger axial measure; \(1.91\) = constant.
Using a low power field diameter of 1800 μm, outlines of 150 nuclei were drawn for each sample by camera lucida. Arithmetical means of the diameters of nuclei were grouped into frequency classes. The samples were examined by three independent investigators using standardized criteria.\textsuperscript{7}

**Statistical Analysis**

The results are reported as means ± standard deviations. In order to examine cell volume data, the Kruskal-Wallis and Tukey post-hoc tests were applied to allow comparisons between independent groups. The concordance of the measurements performed by the three individual investigators was evaluated and analyzed using Bartko’s intra-class correlation coefficient according to the Fleiss guidelines.\textsuperscript{17} The significance level was set at R>0.75 or p<0.05.

**Bartko’s test formula**

\[
R = \frac{N(PMS - EMS)}{N(PMS) + (K-1)(RMS) + (N-1)(K-1)EMS}
\]

- R: Bartko’s correlation index
- PMS: Patients Mean Square
- RMS: Researcher Mean Square
- EMS: Error Mean Square
- N: Number of events
- K: Number of investigators

**RESULTS**

We noted the following responses in rats after cold stress exposure: hair bristling, paw edema, tremor, tail stretching and tachypnea. There were no response differences noted between the memantine-treated group and the untreated group. Body weights were not statistically different among the groups (IH group: 306.2±20.3 grams; CON group: 308.6±29.7 grams; MEM group: 265.8±29.4 grams; IHM group: 254.6±21.9 grams).

To validate whether or not exposure to cold stress was sufficient to cause stress response, we examined lipid depletion in adrenal gland cells (Table 1) and glycogen depletion in hepatocytes (Table 2) of rats from each group.

The IH group had highest rate of lipid depletion in adrenal gland cells (Table 1) and glycogen depletion in hepatocytes (Table 2) of rats from each group. Body weights were not statistically different among the groups (IH group: 306.2±20.3 grams; CON group: 308.6±29.7 grams; MEM group: 265.8±29.4 grams; IHM group: 254.6±21.9 grams).

The IH group had highest rate of lipid depletion in adrenal gland cells (p<0.05) and the highest rate of glycogen depletion in hepatocytes (p<0.05), which supports our hypothesis that exposure to -8ºC for four hours was effective at inducing physiological stress.

The rats in the IH group had a smaller cardiomyocyte nuclear size (Figure 1) compared to the other three groups (p<0.05). Our analysis revealed that the rats in the IHM group had a larger nuclear size than rats in the IH group, likely because the memantine treatment provided protection on the ultrastructural level (p<0.05).

We compared the groups on the basis of nuclear size (Figure 1). The nuclear size of the IH group (Figure 2A) was significantly decreased as compared to the control group (Figure 2B) (p<0.05). Furthermore, rats in the IHM group (Figure 2C) had a 76% reduction in their cardiomyocyte nuclear size as compared to rats stressing the MEM group (Figure 2D) (p<0.05). Data variance analysis using Bartko’s correlation index yielded results ranging from 0.44-0.96 in all experimental groups, thus validating our methodology. Furthermore, when we performed variation analysis

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**Table 1 - Lipid depletion intensity in adrenal gland cortical cells stained with Sudan IV* (+=small intensity; ++++=high intensity)**

| Animal | Cold Stress | Control | Memantine | Memantine + Cold Stress |
|--------|-------------|---------|-----------|-------------------------|
| 1      | ++          | ++++    | +++       | +                       |
| 2      | ++          | +++     | +++       | +                       |
| 3      | +           | +++     | +++       | +                       |
| 4      | ++          | +++     | +++       | +                       |
| 5      | +++         | ++++    | +++       | +                       |
| 6      | +           | +++     | +++       | +                       |
| 7      | +           | ++     | +++       | +                       |
| 8      | +           | +++   | ++        | +                       |
| 9      | +           | ++++  | +++       | +                       |
| 10     | +           | ++++  | +++       | +                       |

* + = Low intensity (more lipid depletion); ++++ = High intensity (less lipid depletion)

**Table 2 - Glycogen depletion intensity in hepatocytes blushed with PAS method* (+=small intensity; ++++=high intensity)**

| Animal | Cold Stress | Control | Memantine | Memantine + Cold Stress |
|--------|-------------|---------|-----------|-------------------------|
| 1      | ++          | ++++    | +++       | +                       |
| 2      | ++          | +++     | +++       | +                       |
| 3      | +++         | ++++    | +++       | +                       |
| 4      | ++          | +++     | +++       | +                       |
| 5      | +++         | ++++    | +++       | +                       |
| 6      | +           | ++++    | +++       | +                       |
| 7      | +           | ++     | ++++      | +                       |
| 8      | +           | ++     | ++++      | +                       |
| 9      | +           | +++   | ++++      | +                       |
| 10     | +           | ++++  | ++++      | +                       |

* + = Low intensity (more glycogen depletion); ++++ = High intensity (less glycogen depletion)
examining group mean differences, we found a statistically significant difference between groups ($p<0.05$).

**DISCUSSION**

We sought to evaluate the effects of memantine administration on the heart tissue of rats exposed to cold stress. As demonstrated in a previous study, we found that exposure to cold stress for four hours at -8ºC decreased cardiomyocyte nuclear size in rats. We also verified that rats in the group treated with memantine had the largest mean cardiomyocyte nuclear size of all groups in this study. Moreover, rats treated with memantine and exposed to cold stress were found to have a larger cardiomyocyte nuclear size than rats that were exposed to cold stress but not treated with memantine.

Our investigation demonstrated that induced hypothermia causes cardiomyocyte nuclear volume reduction. According to prior research, physiological stressors such as exercise, fasting and cold are known triggers of different physiological pathways through mechanisms that are not yet completely understood. In a study performed by Matsuzuka et al., it was observed that the cellular stress caused by hypothermic exposure exerted a lethal effect on hamsters with cardiomyopathy and therefore suggested that cold stress leads to, or increases cardiovascular dysfunction in animals. Previous studies have shown that cold stress triggers activation of a mitochondria-dependent signaling pathway in cardiomyocytes, which is an important finding, given that mitochondrial stress either leads to cellular adaptation and survival or induces apoptotic signaling in various types of cells. Thus, the increase in metabolic load and reduction in ATP production caused by stress cause a cytosolic Ca$^{2+}$ imbalance that leads to hypoxia and irreversible cellular injury. Our histological analysis, which showed an increased rate of lipid depletion in the adrenal gland and a higher rate of glycogen depletion in liver cells, supports these recent findings.

We observed that rats in the IHM group had a larger nuclear size than rats in the IH group. However, we cannot confirm the effects that memantine administration has on cardiovascular parameters identified in prior studies because we did not examine hemodynamic determinants. Several prior studies have described the association between memantine and the prevention of ischemia. Osborne et al. examined whether or not memantine could slow down the changes observed in rat retina following ischemia and reperfusion. When memantine (5 mg/kg) was given at the onset of ischemia, it prevented ischemia/reperfusion injuries to varying degrees. However, when memantine was given at the onset of the reperfusion, this was not observed. The authors therefore suggested that a single injection of memantine given intraperitoneally or intravitreally protects the retina from a subsequent ischemic insult. D’Amico et al. observed that the incidence of ventricular tachycardia, ventricular fibrillation and mortality induced by reperfusion was greatly reduced in animals treated with intravenous memantine (1.5 mg/kg) injected five minutes prior to coronary artery occlusion ($p<0.01$). They also indicated that reperfusion-induced arrhythmias, but not ischemia-induced arrhythmias, were preventable by administration of both NMDA and non-NMDA ionotopic excitatory amino acid receptor antagonists.

In our study, Bartko’s intra-class correlation coefficient revealed that there was significant concordance among the investigators. The mean indexes of intra-class correlation

**Figure 1** - The nuclear size of cardiomyocytes in the control (CON), memantine-treated (MEM), induced hypothermic/memantine-treated (IHM) and induced hypothermic (IH) groups. *$p<0.05$ for IH as compared to the other groups

**Figure 2** - In the IH group (A), cardiomyocyte nuclei were found to be smaller and have a lower volume than the nuclei in other groups. Cardiomyocytes in the CON group (B) had normal cardiomyocyte nuclear size without evidence of any type of cellular injury. In the IHM group (C) decreased cardiomyocyte nuclear volume was observed, but the nuclei were larger than those of cardiomyocytes in the IH group. In the MEM group, (D) we observed increased cardiomyocyte nuclear size. The nuclei were normal in appearance without evidence any type of cellular injury (HE – Amplification 400 x)
Memantine role in heart tissue

Herrero et al. found that memantine did not significantly affect blood pressure.

To our knowledge, there has been no prior study that has examined the association between memantine administration and alterations in cardiomyocyte nuclear size. We found that memantine was an effective Ca\(^{2+}\) channel blocker in cardiac myocytes, which is evidenced by the preservation of nuclear size that occurred in this rat model of cardiac injury. In addition, an unexpected observation made in this study was that the group treated with memantine had the largest nuclear size, which supports the idea that memantine may also act as a cardioprotective drug. This is the first study to demonstrate that this drug preserves the nuclear size of cardiomyocytes in animals that have been exposed to cold stress. We believe that these data are important in developing future therapies to preserve cardiomyocyte structure under conditions of cold stress.

In summary, our results indicate that memantine prevents cardiomyocyte nuclear size reduction in the left ventricles of rats exposed to cold stress.

**ACKNOWLEDGMENT**

We thank Mr. Brandon Saltzgiver for his critical evaluation of English Grammar.

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(R) ranged between 0.44 and 0.96 for each animal studied. Statistically significant differences were demonstrated between all groups, and the results were complemented by a group mean difference analysis. Therefore, based on our statistical analyses, we were able to validate the reliability of this method for assessing differences among groups.

The effect that memantine was found to have on the cardiac cells in our study may lead to new perspectives regarding its role in the cardiovascular system. Recent studies have demonstrated the association between memantine and various cardiovascular parameters. Lapchak et al.

ACKNOWLEDGMENT

We thank Mr. Brandon Saltzgiver for his critical evaluation of English Grammar.
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