Zeolite pretreatment accomplishes partial brain radioprotective role by reducing iron and oxidative / nitrosative stress in rats

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Abstract
The aim of our study was to test the effect of subacutely applied micronized zeolite [micronized clinoptilolite (MZC)] on brain status of iron (Fe), reactive oxygen and nitrogen species (ROS, RNS), and radioprotective role against brain oxidative/nitrosative stress (OS/NS) initiated by single ionizing radiation of 2 or 10 Gray (Gy). Wistar rats on normal (n=18) and 5% MZC supplemented diet (n=18), during 4 weeks, were internally subdivided into 3 subgroups (6 rats in each subgroup), with one of subgroup remaining as a control, and the other two subjected to single ionizing radiation of 2 Gy or 10 Gy. Thus, we had groups on normal diet: C – controls, 2Gy and 10Gy; and on 5% MZC supplemented diet: MZC, MZC+2Gy and MZC+10Gy. Concentrations of nitrates (a final RNS metabolite) and superoxide anion radical (O2−) (an initial ROS) were measured in homogenates of selective vulnerable brain regions (cerebellum, hippocampus and forebrain cortex), while Fe was determined in whole brain of rats. Results documented a significant drop of Fe in MZC and MZC+2Gy/10Gy groups; decrease of O2− and nitrate in MZC group; almost equal drop of O2− in 2Gy and MZC+2Gy groups; and nitrate increase in 10Gy and MZC+10Gy groups. We confirmed that subacute MZC pretreatment contributes to partially accomplished brain radioprotective effect in rats exposed to single radiation dose of 2 Gy and 10 Gy, probably due to reduced OS/NS and Fe.

Key words: brain, iron, ionizing radiation, nitrosative stress, oxidative stress, zeolite (MZC)

INTRODUCTION
Zeolite, a natural clinoptilolite, is a strong non-selective adsorbent and/or ion exchange agent for different compounds, including metals, mycotoxins, some strains of bacteria, gases, essential nutrients, etc. [1–3]. Adsorptive potential of zeolite grows with the decrease of particle size, which is why we used micronized clinoptilolite (MZC).

Taken orally, zeolite remains intact within the alimentary tract. It is used worldwide as a supplement for animal feed and in human medicine [4] . However, there is plenty of evidence (pro and contra) regarding daily intake of zeolite in humans [5].

In respect to its binding properties, zeolite reduces bioavailability of essential transition metals (iron (Fe), copper (Cu), manganese (Mn), etc., which contribute to free radical production through Fenton like reactions). Also, zeolite absorbs gases released during food fermentation within the alimentary tract (oxygen (O2) and nitric oxide (NO), which altogether indicates the certainty of its antioxidative role [6]. Homeostasis of essential metals is particularly important for regular development and physiological functioning of living organisms [7]. Iron is the most abundant transition element in humans with ascertained physiological importance. Nevertheless, Fe contributes to oxidative stress (OS) development, via Fenton reaction, where it reacts with hydrogen peroxide (H2O2) to form the potent hydroxyl radical (HO•) [8]. Reactive oxygen and nitrogen species (ROS, RNS) affect all classes of biomolecules (proteins, lipids and DNA), also, they alter cell signaling pathways, gene expression deplete energy and eventually, lead cells into death by apoptosis [9].
It is well known that ionizing radiation induces OS in living organisms [10]. We supposed that nitrosative stress (NS) will develop in irradiated rats as well, based on the fact that blood vessel endothelium produces NO which spontaneously reacts with superoxide anion radical (O$_2^{-}$·) and produces harmful RNS, peroxynitrite anion (ONOO$^-$). The final metabolic products of entirely generated RNS are nitrates (NO$_3^-$) [9]. Additionally, we assumed that reduced bioavailability of Fe, NO and O$_2$ from gastrointestinal tract (GIT) during prolonged MZC intake will result in systemic OS and NS decrease, including brain. Since previous studies confirmed that ionizing radiation causes increase of OS and NS [11, 12], we were tempted to check if the prolonged MZC pretreatment will have a radioprotective role in the brain, by analyzing certain OS/NS parameters in rats.

The use of zeolite as a food supplement is based on its detoxification capabilities due to ability to bind various toxins within the GIT [13–15]. Furthermore, zeolite improves some physiological occurrences in treated animals, such as growth improvement, reproductive performance, boosts immune system, etc. [16–18].

Accordingly, the aim of our study was to test if oral pretreatment with 5% MZC, for 4 weeks, will make any influence on brain status of Fe, ROS and RNS and additionally, will radioprotective role against brain OS/NS initiated by single ionizing radiation doses, 2Gray (Gy) or 10Gy, will be attained.

MATERIALS AND METHODS

Experimental animals

Adult male Wistar rats (weighing 220–250g) were kept under standardized housing conditions (temperature 23±2°C, lighting 12:12 light:dark, light on from 8:00 to 20:00h) with free access to tap water and a custom pellet rat diet. Suspension of MZC was administered daily by gavage. The experimental protocol followed the “Guide for the Care and Use of Laboratory Animals” [19] and was approved by Ethical Committee for Experimental Animals, „Vinča” Institute (No. 6/12).

Experimental design

Wistar rats on normal diet were randomly subdivided into three groups (n=6): C – control (not treated) and 2Gy and 10Gy groups – rats subjected to a single dose of radiation of 2Gy or 10Gy, respectively; and accordingly, rats on 5% MZC supplemented diet covered three groups (n=6): MZC, MZC+2Gy and MZC+10Gy groups. The MZC amount was calculated in respect to the quantity of ingested food and rat body mass. The suspension of 0.85–1g of MZC/day (corresponds to 5% of 17–20 g of custom pellet/day) was administered orally, by gavage, during four weeks [5]. The animals gained 133.85 ± 24.7g body weight after 4 weeks. No statistically differences were observed between the C and MZC groups.

Gamma irradiation of rats was performed in the Laboratory of Radiation Chemistry and Physics, Vinča Institute of Nuclear Sciences, using 60Co gamma source designed for radiobiological and radiation chemistry experiments. The animals were confined in custom made individual cages, made of wire, sideways positioned to the irradiation source and then subjected to the whole body irradiation at a dose rate of 0.1676Gy/min for 12 minutes (corresponds to a total sublethal dose of 2Gy) or 60 minutes (corresponds to a total lowered lethal dose of 10Gy) [20]. Five days after the irradiation, the animals were anesthetized with a 50 mg of sodium pentobarbital/kg injection before being sacrificed (by decapitation), when brains were removed immediately and stored at -80 °C, until analyzing. The impact of the administered anesthesia on OS and NS parameters measured in brain of rats was not achieved (the results of intact and anesthetized animals were almost identical, therefore, were not presented).

Reagents

All reagents and chemicals were of analytical grade or higher purity. In this study we used ethylenediaminetetraacetate acid – EDTA, nitroblue tetrazolium – NBT, nitric acid – HNO$_3$ (65%) (Fisher Chemical, UK), perchloric acid – HClO$_4$ (65%), hydrochloric acid – HCl, sodium hydroxide – NaOH, sulfanilic acid, N – (1-naphthyl) amine (Sigma–Aldrich, USA), sodium nitrate – NaNO$_3$, sodium phosphate – NaHPO$_4$ (Merck, Germany), Fe standard solution (AccuStandard, USA), sodium nitrate – NaNO$_3$, sodium phosphate – NaHPO$_4$, glycerol (Merck, Germany), Fe standard solution (AccuStandard, USA), sodium pentobarbital Vetanarcol (0.162 g/mL) (Werff – Chemie, Austria), saline solution (0.9% w/v) and de-ionized water (Hospital Pharmacy Military Medical Academy, Belgrade, Serbia).

Analysis of iron

Brain tissue samples (around 1g) were mineralized in a mixture of concentrated HNO$_3$ and HClO$_4$ (4:1, v/v), heated at temperature between 250 °C and 300 °C till dryness and diluted up to 10mL with 0.1M HNO$_3$. The concentrations of Fe were analyzed by inductive coupled atomic absorption spectrometry (IC-AAS, Analyst 200, PerkinElmer), using air-acetylene flame. Standard solutions of Fe were prepared according to the Perkin-Elmer Pure Atomic Spectroscopy Standards guidelines (NIST traceable CRM, Perkin-Elmer Corporation, USA and Merck – Germany). The absorption wavelength was 305.91 nm.

Superoxide anion radical measurement

Quantification of O$_2^{-}$· was based on nitroblue tetrazolium (NBT) reduction by O$_2^{-}$· to yellow colored monofor- mazan, which absorbance was measured at 550 nm [21].

Nitrate measurement

After deproteinization, concentration of nitrates (NO$_3^-$, the final metabolic product of all RNS) was calculated...
as the difference between two sets of measurements with Griess reagent (1.5 % sulfanilamide in 1 mol HCl plus 0.15 % N-(1-naphthyl) ethylenediamine dihydrochloride in distilled water), before (only NO$_3^-$ was measured) and after the addition of cadmium, which reduces NO$_3^-$ into NO$_2^-$ (NO$_2^-$ + NO$_3^-$ were measured) [22]. The measurements were performed spectrophotometrically at 492 nm. The results were expressed as nmol NO$_3^-$/mg protein.

**Protein determination**

The Lowry method was used for protein measurement in rat vulnerable brain regions (VBR) homogenates [23].

**Statistical analysis**

One-way ANOVA and post-hoc Dunnett’s C tests were used (software GraphPad Prism, version 5.01) for statistical data analysis. Values were presented as average ± SD. Differences were considered statistically significant for p<0.05.

**RESULTS**

Decrease of brain Fe concentrations was observed in all experimental groups: MZC, 2Gy and MZC+10Gy (p<0.01); 10Gy (p<0.05) and MZC+2Gy (p<0.001), compared to the controls. Also, lower Fe was found in MZC+2Gy and MZC+10Gy (p<0.05) groups than in corresponding irradiated groups, 2Gy and 10Gy (Figure 1).

![Graph 1](image1.png)

**Figure 1.** Brain iron in the experimental groups. Brain Fe (µg Fe/mg wet tissue) concentrations are presented as means ± SD, for 10 animals/group. Differences were considered statistically significant at: *p<0.05, **p<0.01 and ***p<0.001 (compared to control, # compared to MZC group). One-way ANOVA, Dunnett’s C test were used for statistical analysis.

Significant increase of NO$_3^-$ was reached in the experimental groups, as follows: 10Gy group (in: cerebellum (p<0.05), hippocampus (p<0.001) and cortex (p<0.001)); MZC+10Gy (in: hippocampus (p<0.05) and cortex (p<0.001)); and 2Gy (in: hippocampus (p<0.01)). Contrary, NO$_3^-$ decline was documented in: MZC group (in: cerebellum and hippocampus (p<0.01) and cortex (p<0.001)); 2Gy (in: cerebellum (p<0.001) and cortex (p<0.01)); and MZC+2Gy group (in: cerebellum and cortex (p<0.001)). In MZC+2Gy group, NO$_3^-$ was lower in hippocampus (p<0.01) and cortex (p<0.05) compared to 2Gy group (Figure 3).

![Graph 3](image3.png)

**Figure 3.** Nitrates in selective vulnerable brain regions of rats exposed to zeolite. Nitrates (NO$_3^-$: nmol NO$_3^-$/mg proteins), the final metabolic product of RNS, were measured in VBRs: cerebellum (Cer), hippocampus (Hipp) and cortex (Cx). Values are presented as means ± SD, for 10 animals/group. Differences were considered statistically significant at: *p<0.05, **p<0.01 and ***p<0.001 (compared to control, # compared to MZC group). One-way ANOVA, Dunnett’s C test were used for statistical analysis.
DISCUSSION

Zeolite reversibly binds gases, such as O\(_2\), and NO, as well as other nutrients, including metals (i.e. Fe) within GIT [24]. Reduced bioavailability of Fe, ROS and RNS in rats on 5% MZC supplemented diet during four-week diet regimen imposed a systemic effect, documented by the decreased brain levels of the aforementioned substances. Moreover, according to our results, MZC imposed a partial radioprotective role against brain OS/NS, induced by lower doses of ionizing irradiation (2Gy) in rats (Figures 1–3).

We confirmed positive association between reduced brain ROS and RNS and lower brain Fe concentrations in MZC treated rats. These occurrences confirmed that reduced bioavailability of Fe, O\(_2\), and NO is followed by a systemic drop of Fe, ROS and RNS.

The brain has high levels of O\(_2\), transitional metals and polyunsaturated fatty acids and therefore is particularly susceptible to OS [25]. Excessive amounts of ROS and RNS cause OS in all neurons, yet, the vulnerability of neurons to OS varies from one brain region to another, as well as within the same brain region. Although the hippocampal regions CA1 and CA3 are next to each other and are composed of morphologically similar pyramidal neurons, there are differences in their sensitivity to OS. When exposed to pro-oxidative substances, the neurons in the CA1 region suffer massive cell death while those in CA3 mostly survive [26, 27]. Also, significant differences in vulnerability to OS of cerebellar and cortical neurons are documented. Exposed to some OS-inducing agents, such as paraquat, or conditions involving OS, such as ischemia and re-oxygenation, extensive death of neurons in the cerebellum, but not in the cerebral cortex was observed [25].

Overproduction of ROS, including O\(_2\)^•-, H\(_2\)O\(_2\), HO• and others, can be triggered by different stimuli, including metabolism of endogenous exogenous compounds in the presence of O\(_2\). Fenton-like reactions, ionizing radiation, etc. Oxidative stress development assumes time and spatial spreading of free radical chain reactions in living organisms. In the Fenton reaction, a low-valent transition metal (such as Fe\(^{2+}\)) reacts with H\(_2\)O\(_2\) and produces HO•, which instantly oxidatively damages various classes of essential biomolecules such as lipids, proteins and nucleic acids [28, 29]. According to our results, reduced brain Fe level is associated with lower concentrations of O\(_2\)^•- (Figures 1, 2).

Keeping the homeostasis of NO sustained throughout all body organ systems is immensely important, considering its regulatory role in many physiological processes. However, healthy blood vessel endothelium is defined by the ability to produce NO [30]. Spontaneous reactions between NO radical (NO\(^-\)) and O\(_2\)^•- results in ONOO• production. Peroxynitrite is an extremely harmful molecule, blamed for atherosclerotic and other NS associated diseases [31].

In relation to this, MZC capability to reduce OS, as well NS in the body, including brain, was anticipated (Figures 2, 3). Decreased concentrations of O\(_2\)^•- and NO\(_3\)^• in all examined brain regions of rats on MZC supplemented diet can be explained by the ability of zeolites to bind transition metals (including metals participating in Fenton reaction, such as Fe) in the alimentary tract and reduce their availability to other organs. Very few studies on zeolite pointed at its indirect, antioxidant effect, confirmed by suppressed lipid peroxidation [5, 32].

However, ionizing radiation can alter molecules within the cells, both directly and indirectly, affecting cell viability. Radiation energy absorbed by tissues and fluids leads to radiolysis of water and biomolecules, resulting in consequent OS and NS development [28, 33, 34].

Human population can be exposed to the dose range of 1–10Gy during radiation therapy treatment or as the result of radiation accidents or nuclear/radiological terrorism [35].

According to our results, a higher irradiation dose caused large elevation of RNS in all examined brain regions of treated rats, while at lower doses elevation of RNS was noticed only in hippocampus. This is consistent with the results of previous studies that confirm that hippocampus is one of the most sensitive brain regions to OS/NS. Radioprotective effect of MZC against brain NS induced by ionizing radiation was accomplished only at lower dose of 2Gy in hippocampus and cortex, but not in cerebellum (Figure 3). This is in line with the results from a recent study indicating that lower doses of irradiation increased blood-brain barrier (BBB) permeability, decreased blood flow and content of antioxidants in the cerebellum, more than in other brain regions, causing OS [36].

Significantly enlarged brain NS was obtained after irradiation with 10Gy regardless of the previous dietary regimen, normal or MZC supplemented (Figure 3). This suggests that the antioxidant potential of MZC is not sufficient to reduce brain damage caused by high content of ONOO• produced by the higher irradiation dose.

Herein, we demonstrated that the most resistant VBR structure to OS, initiated by 2Gy irradiation was hippocampus (no change of O\(_2\)^•- level was noticed), while the most sensitive was cortex (the highest O\(_2\)^•- concentration was observed). This can be explained by the different concentrations of Cu, zinc (Zn) and Mn within these brain regions. The highest concentrations of these metals were documented in hippocampus [37, 38]. These metals are cofactors of most important enzymatic antioxidants superoxide dismutases (SODs – CuZnSOD and MnSOD) which catalyze dismutation of O\(_2\)^•- to H\(_2\)O\(_2\). Therefore we can conclude that after exposure to 2Gy irradiation concentration of O\(_2\)^•- in hippocampus is lower than in other brain regions due to higher total SOD activity in this brain section.

After radiation of 10Gy, no change in the O\(_2\)^•- concentration was observed, contrary to the increased NO\(_3\)^• concentration in all tested brain regions of the ex-
posed rats. We assumed that $O_2^-$ spontaneously reacts with NO$^+$ producing more potent ONOO$^-$. Regarding NS, cerebellum was proven to be the most resistant VBR region against NS, initiated by 2Gy irradiation (an intense NO$^+$ drop was documented), while hippocampus was the most sensitive. The NS respond of all VBRs was similar at 10Gy (Figures 2, 3). Nitric oxide in the brain is synthesized primarily by neuronal nitric oxide synthase (nNOS). However, in the conditions of exposure to various OS-inducing agents activation of inducible nitric oxide synthase (iNOS) can lead to overproduction of NO. Supraphysiological levels of NO cause apoptosis with resultant decreased regional neuronal function. The sensitivity of iNOS to radiation is probably the highest in the hippocampus compared to other brain structures [39].

CONCLUSION
Herein, we confirmed that decreased bioavailability of Fe, NO and O$_2$ by subacute intake of MZC resulted in systemic drop of Fe, ROS and RNS, judging by brain Fe, ROS and RNS levels in the exposed rats. Zeolite achieved a partial antioxidative effect initiated by a lower dose of ionizing radiation and limited antioxidative effect initiated by a higher irradiation dose in the exposed rats.

Also, the obtained results suggest that there are differences in the sensitivity to radiation between the individual brain structures evaluated by OS/NS development that can be explained by variations in cerebral-vascular permeability, content of antioxidant enzymes and transition metals.

CONFLICT OF INTEREST STATEMENT
The authors disclosed any financial or personal relationships that could inappropriately influence or bias the content of the paper.

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