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Antioxidative, membrane protective and antiapoptotic effects of melatonin, *in silico* study of physico-chemical profile and efficiency of nanoliposome delivery compared to betaine

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The mechanisms through which melatonin exerts its inhibitory effect on cell death remains insufficiently clarified. The subject of the present study is the evaluation of the hepatoprotective effects of melatonin on the inhibition of apoptotic and oxidative processes and activation of survival pathways, in comparison with betaine, in a primary culture of hepatocytes that have undergone Fas-ligand apoptosis. Melatonin exerted a protective effect on membrane bilayer stability through the inhibition of phosphatidylserine externalization, Bax expression and xanthine oxidase catalyzed free-radical liberation. It also reduced liberation of cellular and membrane-bound enzymes and endonuclease catalyzed DNA fragmentation. Betaine hydrochloride did not exert these effects, when administered alone or co-incubated with anti-Fas antibodies. The *in silico* Molinspiration tool, which was employed to calculate melatonin and betaine physico-chemical properties and membrane interaction indicated that melatonin may easily cross and interact with biological membranes and maintain membrane phospholipid structural topography. It was documented to occur *via* non-receptor mechanisms in more than 75%; this may clarify melatonin as suitable for nanoliposome-based delivery, where it was also able to successfully counteract induced oxidative stress. This could not be considered for betaine hydrochloride. The use of lipophilic compounds like melatonin, encapsulated in nanoliposomes, could therefore be a preferable tool in the successful membrane-preservation therapy of liver apoptosis, rather than the use of hydrophilic compounds, like betaine hydrochloride.

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Introduction

Apoptosis mediated by the Fas receptor (CD95) has been implicated in a growing number of liver diseases including viral hepatitis, alcoholic, cholestatic, autoimmune and Wilson's disease.¹ In the extrinsic apoptotic pathway, initiation of apoptosis in the liver occurs presumably through Fas death receptors, located on the cell membrane.² Hepatocytes constitutively express Fas receptors and have been shown to be sensitive to the Fas death signal, both, *in vitro* and *in vivo*.³ The anti-Fas monoclonal antibodies were shown to trigger apoptotic cell death *in vivo* and *in vitro*⁴ and our previous report documented Fas-ligand apoptosis as a useful model in evaluating

potentially hepatoprotective substances.⁵ The binding of specific ligands to Fas receptors causes a cascade of signaling effects following the activation of executor caspase and endonuclease enzymes.^{1,2} Cells undergoing Fas-mediated apoptosis such as hepatocytes and T lymphocytes show PS externalization and subsequent clearance by phagocytes before lysis, or inflammatory reaction.⁶ In an intrinsic apoptotic pathway the Bcl-2 family of proteins are potent regulators, where Bcl-2 and Bcl-xL (group I) possess antiapoptotic activity, whereas members of group II (Bax and Bak) and group III (Bid) promote cell death.⁷ Alteration of membrane structural stability is the mechanism responsible for the translocation of Bax to the external mitochondrial membrane leaflet, when apoptosis is induced.⁸

Melatonin (*N*-acetyl-5-methoxytryptamine) is a compound derived from L-tryptophan.⁷ It is a lipophilic molecule, which to some extent may freely cross cell membranes and enter cells. Moreover, it can also bind to plasma membrane melatonin receptors (MT₁ and MT₂), members of the seven transmembrane G protein-coupled receptors, initiating downstream cell signaling.^{9,10} It has been documented that melatonin exerts

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a broad range of physiological⁹ and protective antioxidant effects, preventing the peroxidation of membrane lipids and protecting the proteins and DNA from oxidative damage induced by hydroxyl radical ($\cdot\text{OH}$), peroxynitrite anion (ONOO^-), hydrogen peroxide (H_2O_2), superoxide anion radical ($\text{O}_2^{\cdot-}$) or singlet oxygen ($^1\text{O}_2$), produced in different *in vitro* and *in vivo* conditions.^{11,12} Our previous results pointed out melatonin as protector against microwave irradiation.^{13–15} Like an electron donor, melatonin has been shown to increase the efficiency of the electron transport chain and, as a consequence, to reduce electron leakage and the generation of free-radicals.¹⁶ Beside direct free-radical scavenging properties, it also has been shown to have indirect antioxidative effects, through the up-regulation of several antioxidative enzymes. When compared to vitamin E, melatonin was found to be more effective in limiting lipid membrane destruction,^{12,17} serving as naturally mitochondria-targeted protective molecule.¹⁸ Metabolites of melatonin such as major hepatic metabolite 6-hydroxymelatonin as well as N^1 -acetyl- N^2 -formyl-5-methoxykynuramine and N^1 -acetyl-5-methoxykynuramine, have been shown to detoxify radicals themselves. This effect, independent of the species and the type of injury and extent of organ damage, is especially true for the liver and a variety of its pathophysiological conditions.¹⁹ Furthermore, there are data documenting the influence of melatonin on Bcl-2/Bax levels in rat peripheral blood lymphocytes²⁰ and pancreatic cancer cell line SW-1990.²¹ It was recently demonstrated that melatonin may be involved in epigenetic DNA reprogramming, decreasing the levels of unmethylated DNA promoter regions.²² It has been further proven that in liver disorders with a dysfunction of the apoptotic program, like ischemia/reperfusion injury,²³ viral hepatitis,²⁴ acetaminophen-induced acute liver failure²⁵ or cadmium-induced hepatotoxicity,⁹ melatonin may enhance regenerative response.

Betaine is *N*-trimethylated amino acid glycine, a quaternary ammonium compound, which exists in zwitterionic form at neutral pH. As a positively-charged amine, it requires a protein-mediated mechanism to effectively pass the membrane-lipid barrier *via* lipid-associated betaine-binding protein (BetP).²⁶ Because of its three chemically-reactive methyl groups, it has a role as a donor of methyl group in many cell pathways, such as synthesis of phospholipids, proteins and in stabilization of DNA structure through the methylation.²⁷ It may enhance the secretion of VLDL *via* methylation of membrane phospholipids increasing phosphatidylcholine content.²⁸ Betaine may be a promising therapeutic agent in relieving defects associated with alcoholic liver disease.²⁹

The subject of the present study was the hepatoprotective effects of melatonin mediation by the inhibition of apoptotic and oxidative processes and activation of survival pathways, in comparison with betaine in the primary culture of mouse hepatocytes which had undergone Fas-ligand apoptosis. The effects of melatonin and betaine on the integrity of the membrane, phospholipid asymmetry and PS (phosphatidylserine) externalization in relation to certain biomarkers of oxidative stress and apoptosis were evaluated, as well as cellular and membrane enzymes liberation, mitochondrial Bcl-2/Bax distribution and DNA degradation. Since a nanoliposome-based drug

delivery system could improve the pharmacokinetic properties of certain drugs, the efficacy of melatonin and betaine hydrochloride to counteract oxidative stress in encapsulated nanoliposomes was employed. Additionally, an *in silico* study by using Molinspiration³⁰ and ACD/I-Lab software³¹ was performed in order to gain a better insight into the physico-chemical properties of melatonin and betaine as well as to explain obtained biological effects.

Materials and methods

Chemicals used

All chemicals were of a pro analysis degree of purity. Anti-Fas antibodies (Jo CD95) were obtained from BD Biosciences. The cell culture medium (RPMI 1640), fetal calf serum (FCS), collagenase and betaine hydrochloride were purchased from Sigma Aldrich (USA), melatonin was purchased from ICN (USA).

Ethical statement

All experimental procedures were performed in compliance with the Ethical Committee guidelines and rules for the protection of experimental animals according to the Animal Welfare Rules Republic of Serbia. The approval for the animal experiment was obtained from the Animal Welfare Committee No. 01-5518-6 (University Nis Medical Faculty) after submitting the required form for approval.

Isolation of primary mouse hepatocyte culture

The experimental study was conducted on inbred, male, three-month-old, BALB mice (22–25 g weight) which were killed under ketalar anesthesia, and whose livers were removed according to the method explained in our previous article.⁵ The hepatocytes were isolated from the liver tissue by use of the collagenase technique (1 mg collagenase per ml RPMI 1640 medium). Isolation of hepatocytes was monitored by microscope and collagenase activity was inhibited by 10% FCS/RPMI 1640 medium. The digested tissue was sieved and low-speed centrifuged (600 g at 4 °C for 15 min), the cells were collected, dissolved in 0.9% NaCl and centrifuged again (1000g for 10 min at 4 °C). Supernatant was drained and the cells were dissolved in RPMI 1640 medium with 10% FCS and penicillin/streptomycin (cell final concentration was 10^5 mm^{-3}). The cells were allocated into six experimental groups, each containing six samples in a following way: the first group of hepatocytes was treated with anti-Fas antibodies in final concentration of 100 ng ml^{-1} ; the second group was treated with anti-Fas antibodies (in the same concentration) and 1 mM melatonin; the third group was treated with anti-Fas antibodies (in the same concentration) and 1 mM betaine hydrochloride; the fourth group was treated with melatonin only (the same concentration); the fifth group was treated with betaine hydrochloride only (in the same concentration); the sixth group was a control with standard culture conditions (RPMI 1640 medium, 10% FCS and penicillin/streptomycin). The cells were grown for 24 h, afterwards they were centrifuged (1000g for 10 min at 4 °C), the



supernatant was collected and the cells were resuspended in 1.2 ml of a physiological saline solution.

Detection of PS membrane externalization, oxidative stress generation, plasma and nuclear membrane integrity and nuclear fragmentation of isolated hepatocyte culture

In order to detect PS on the membrane external leaflet, Annexin V was used. To the amount of 500 μ l of cells suspension, the Annexin V-FITC (Santa Cruz Biotechnology, USA) solution was added and fluorescence was detected on a flow-cytometer (Epics-XL Coulter, Germany). Nuclear membrane permeability (usually seen in dead cells) was detected adding the equal aliquot of propidium iodide (PI) (Santa Cruz Biotechnology, USA) to cells suspension. Viable cells do not normally bind these chemicals. Annexin V positive, PI negative cells represent early apoptosis events. Both, Annexin V and PI positive cells document late apoptosis. If the cells do not bind Annexin V, but bind PI, they are in the necrotic phase. In this way, the following possibilities are: Annexin V–/PI– (healthy cells); Annexin V+/PI– (early apoptosis); Annexin V+/PI+ (late apoptosis); Annexin V–/PI+ (necrosis).

Xanthine oxidase (XO) activity was evaluated quantifying the formation of uric acid by using method modified and explained in our previous report³² with the calculated molar absorption coefficient of uric acid ($1.22 \times 10^4 \text{ l mol}^{-1} \text{ cm}^{-1}$). In brief, the reaction mixture containing hepatocytes, treated with anti-Fas antibodies, melatonin and/or betaine, and Tris/HCl buffer was preincubated at 37 °C. The reaction was started by adding xanthine and, after incubation, stopped by adding perchloric acid. The enzyme activity was determined spectrophotometrically by measuring the absorption at 293 nm associated with uric acid formation. Xanthine oxidase activity was expressed as U g^{-1} protein.

Generation of malondialdehyde (MDA) was used as an index of lipid peroxidation intensity in isolated hepatocytes, evaluated spectrophotometrically, according to the thiobarbituric acid (TBA) assay.³³ The MDA level was expressed as nmol (MDA)/mg (proteins).

For detection of plasma membrane integrity and cell damage, the liberation of hepatic plasma membrane alkaline phosphatase (ALP) and gamma-glutamyl transferase (γ -GT), and cellular enzymes alanine aminotransferase (ALT) and aspartate aminotransferase (AST) were all measured in collected supernatant using the Dade Behring RxL Max automatic analyzer. The activity of enzymes, evaluated quantifying the liberation in collected supernatant, was expressed as U l^{-1} .

The activity of alkaline DNase was measured spectrophotometrically at 260 nm by the method of Bartholeyns *et al.*,³⁴ modified in our laboratory for cell culture systems.⁵ The method is based on liberated acid soluble nucleotide determination, after precipitation, at pH 7.6 obtained by adding Tris/HCl buffer and in the presence of Mg^{2+} ions as activators, using corresponding standard, commercial DNase I (Sigma Aldrich, USA). The enzyme activity was expressed as U g^{-1} protein.

Detection of Bcl-2 and Bax expression

For determining the levels of Bcl-2 and Bax proteins, hepatocytes were isolated and treated as it was described in the

previous section (Isolation of primary hepatocyte culture). Obtained cells were further processed according to the protocol by Kocic *et al.* 2011.³⁵ Briefly, the hepatocytes, resuspended in U-bottom 96-well culture plates, were fixed using 70% methanol and permeabilized with 0.1% Triton PBS. Specific antibodies were used: for Bcl-2 (N-19: sc-492 Santa Cruz Biotechnology, USA) and for Bax (P-19: sc-526 Santa Cruz Biotechnology, USA). The cells were incubated with the primary anti-Bcl-2 and anti-Bax, washed three times, and incubated with the FITC-conjugated secondary antibodies. The excess of antibodies following staining was washed adding PBS (phosphate buffer saline) to each well. The mean fluorescence intensity (MFI; logarithmic scale) of cell populations was obtained following subtraction of blank values obtained by the treatment with secondary antibodies only.

Nanoliposome encapsulation with melatonin or betaine hydrochloride and detection of susceptibility to Fe^{2+} and H_2O_2 induced oxidative stress

In order to detect percentage of non-receptor phospholipid membrane melatonin uptake, phospholipid nanoparticles in the form of nanospheres, purchased from Nattermann Phospholipids (Germany) were dissolved as a 10% solution and one group was encapsulated by melatonin or by betaine hydrochloride at a concentration of 1 mM. Encapsulated nanoparticles were isolated by centrifugation at 6500g for 30 min at 4 °C according to the method of Gortzi *et al.*³⁶ In obtained supernatant melatonin concentration was estimated by using HPLC method³⁷ and betaine hydrochloride by method of Storey and Jones³⁸ and calculated as the encapsulation efficiency (%), according to the corresponding standard of melatonin or betaine hydrochloride (1 mM) dissolved in non-encapsulated liposome supernatant.

Sedimented nanoparticles were suspended in water and both encapsulated and native nanoparticles were exposed to oxidative stress by incubation with H_2O_2 or FeSO_4 in a reaction mixture at a concentration of 1 mM at 37 °C for another 30 min. The intensity of lipid peroxidation was estimated by the reaction of terminal lipid peroxidation product MDA with TBA, and measured spectrophotometrically at the wavelength of 532 nm, according to the method of Buege and Aust.³³ The MDA level was expressed as nmol (MDA)/ml (nanoliposome mixture).

In silico study of physico-chemical properties of melatonin and betaine

The physico-chemical properties of melatonin and betaine, such as lipophilicity (octanol–water partition coefficient and polar surface area), molecular size (molecular weight and volume), conformational flexibility (number of rotatable bonds) and number of present hydrogen-donors and acceptors, were calculated by using *in silico* Molinspiration tool.³⁰ With physico-chemical characteristics taken into account, some other important features, relating to the behavior and interaction of compounds to the membrane, were predicted. ACD/I-Lab software³¹ was used in order to estimate the membrane permeability of both tested compounds through the quantified



percentage represented information about their absorption rate and bioavailability predicted on the basis of their chemical structure.

Statistical analysis

The results are presented as mean \pm SD. The statistical significance was evaluated using ANOVA test. A p -value of 0.05 was considered statistically significant.

Results

Detection of PS membrane externalization, oxidative stress generation, plasma and nuclear membrane integrity and nuclear fragmentation of isolated hepatocyte culture

Fig. 1 illustrates PS membrane externalization of mouse hepatocytes assayed by Annexin V/PI tests. The treatment of hepatocytes with anti-Fas antibodies significantly increased the number of Annexin V+ hepatocytes, where PS was on the external membrane leaflet. Melatonin greatly preserved the number of Annexin V-/PI- cells, when was co-incubated with anti-Fas antibodies, compared to only anti-Fas antibodies treated hepatocytes. Contrary to melatonin, betaine hydrochloride was able to achieve protective effect neither alone nor in co-incubation with anti-Fas antibodies. According to obtained results, it is supposed that in the early phase of apoptosis melatonin exerts antioxidative effect in order to reduce the generation of free-radicals and subsequent oxidation and externalization of PS. Only antioxidative effect was shown as non sufficient for maintaining the membrane stability. A more effective protective effect of melatonin was exerted in the late apoptotic conditions, when antioxidative effects were manifested in a synergy with antiapoptotic effects latter discussed. Betaine exerted even stronger effects on nuclear membrane disintegration, regarding the quantity of Annexin V+/PI+ cells. Melatonin combined with anti-Fas antibodies significantly decreased the quantity of late apoptotic Annexin V+/PI+ cells. Betaine hydrochloride alone increased the level of Annexin V+/PI+ cells compared to control hepatocytes. There were no Annexin V-/PI+ cells which were considered necrotic.

The obtained results provided evidence that melatonin exerts hepatoprotective effects especially in the late phases of

apoptosis, by maintaining plasma and nuclear membrane integrity. Betaine hydrochloride would not be considered as protector of membrane stability, it exerted strong effect on nuclear membrane disintegration.

As shown in Fig. 2, exposure to anti-Fas antibodies and their combination with betaine hydrochloride induced an increase in lipid peroxidation (LP), when compared to the control hepatocytes. Decrease in LP was evidenced in the combined treatment with anti-Fas antibodies and melatonin in comparison with only anti-Fas antibodies-treated hepatocytes or the combination of anti-Fas antibodies and betaine-treated hepatocytes. Melatonin, given alone, decreased the intensity of LP in comparison with control hepatocytes. Betaine hydrochloride given alone was without significant effect. Melatonin as antioxidant prevented the peroxidation of membrane lipids in mouse hepatocytes and maintained membrane stability, while betaine hydrochloride could not be considered as hepatoprotective agent.

As depicted in Fig. 3, exposure to anti-Fas antibodies induced an increase in xanthine oxidase (XO) activity, when compared to the control hepatocytes. Significant decrease in XO

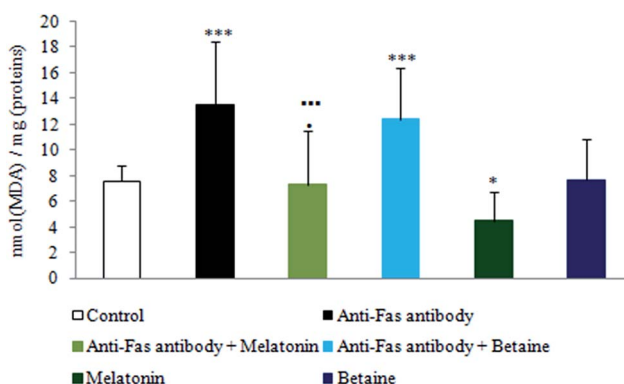


Fig. 2 The intensity of lipid peroxidation in hepatocytes treated with anti-Fas antibodies, melatonin and betaine. Generation of malondialdehyde (MDA) used as an index of lipid peroxidation intensity was evaluated spectrophotometrically and expressed as nmol (MDA)/mg (proteins). *** $p < 0.001$ vs. control; * $p < 0.05$ vs. control; ■■■ $p < 0.001$ anti-Fas antibody vs. anti-Fas antibody + melatonin; ●● $p < 0.01$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine.

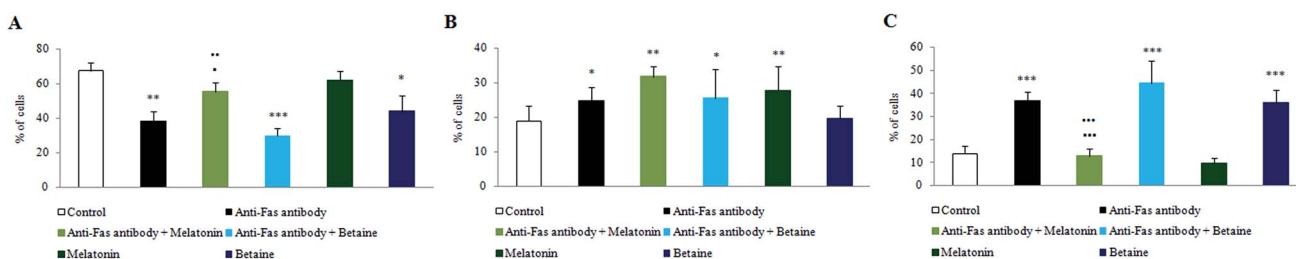


Fig. 1 PS membrane externalization and changes in nuclear membrane integrity of mouse hepatocytes treated with anti-Fas antibodies, melatonin and betaine, assayed by Annexin V/PI tests. The represented stages are: (A) Annexin V-/PI- (healthy cells); (B) Annexin V+/PI- (early apoptosis); (C) Annexin V+/PI+ (late apoptosis). *** $p < 0.001$ vs. control; ** $p < 0.01$ vs. control; * $p < 0.05$ vs. control; ■■■ $p < 0.001$ anti-Fas antibody vs. anti-Fas antibody + melatonin; ■ $p < 0.05$ anti-Fas antibody vs. anti-Fas antibody + melatonin; ●●● $p < 0.001$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine; ●● $p < 0.01$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine.



activity was evidenced in the combined treatment with anti-Fas antibodies and melatonin in comparison with only anti-Fas antibodies treated hepatocytes. Melatonin, given alone, decreased the activity of this enzyme in the comparison with

control hepatocytes. Betaine hydrochloride given alone was without significant effect. In that way, melatonin decreased the generation of free-radicals by the XO catalyzed reactions, what represents one more mechanism of its indirect antioxidative effect. Betaine was without significant effect on the activity of this enzyme.

The exposure to anti-Fas antibodies, specifically their combination with betaine hydrochloride, induced an increase in DNase activity when compared to the control hepatocytes. Melatonin, given alone, decreased the DNase activity in the comparison with control hepatocytes. Betaine hydrochloride given alone was without significant effect (Fig. 4). Besides the protection of DNA from oxidative damage, melatonin may protect it by the inhibition of DNase activity as well. Destabilization of DNA potentially may be provoked by betaine.

The integrity of hepatocytes and their damage was assessed by the levels of liberated enzymes due to cell damage. The hepatocytes treated with anti-Fas antibodies were evidenced by significant increase ($p < 0.001$) in the liberation of AST, ALT, ALP and γ -GT when compared to control cell supernatant (Table 1). However, melatonin exhibited hepatoprotective effects and attenuated the elevated activity of liberated hepatocyte enzymes, with significant effectiveness when was combined with anti-Fas antibodies in comparison to alone anti-Fas antibody treatment ($p < 0.001$). Betaine exerted a less effective hepatoprotective effect.

Detection of Bcl-2 and Bax expression

Fig. 5 shows Bcl-2 and Bax protein expression on the mitochondrial membrane under the given experimental conditions. The treatment with anti-Fas antibodies led to inverse relation between Bcl-2 and Bax protein compared to control hepatocytes, reducing the values of Bcl-2 and enhancing Bax membrane expression. Melatonin almost fully preserved mitochondrial Bcl-2/Bax expression and distribution, while betaine hydrochloride was without significant effect under the co-incubation conditions with anti-Fas antibodies, compared to anti-Fas antibodies treated hepatocytes. Melatonin, given alone, significantly attenuated Bax expression in the cell culture, compared to control cells, while betaine hydrochloride alone significantly decreased Bcl-2 expression, compared to the control hepatocytes. Thus, melatonin was

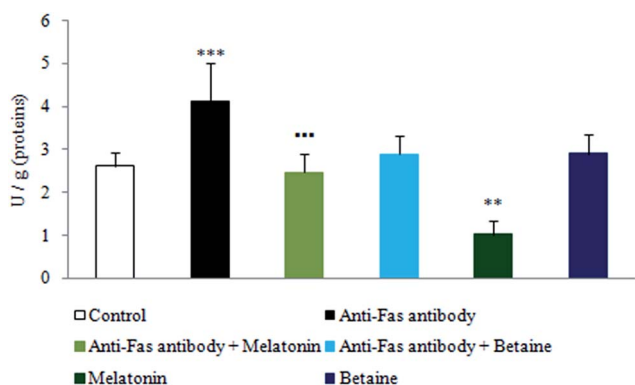


Fig. 3 Xanthine oxidase activity in hepatocytes treated with anti-Fas antibodies, melatonin and betaine. The enzyme activity, evaluated spectrophotometrically quantifying the formation of uric acid, was expressed as $U\ g^{-1}\ protein$. *** $p < 0.001$ vs. control; ** $p < 0.01$ vs. control; ■■■ $p < 0.001$ anti-Fas antibody vs. anti-Fas antibody + melatonin.

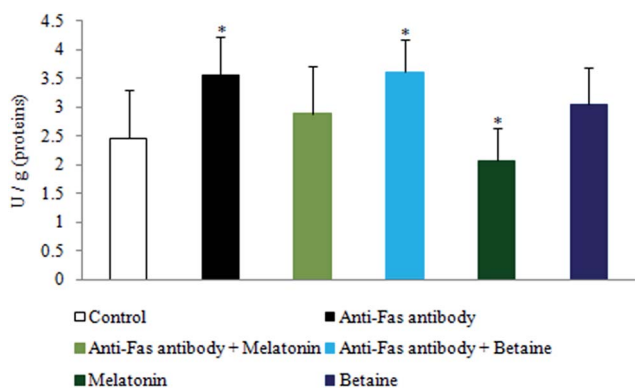


Fig. 4 DNase activity in hepatocytes treated with anti-Fas antibodies, melatonin and betaine. The activity of alkaline DNase was spectrophotometrically measured by the liberation of acid-soluble nucleotides and expressed as $U\ g^{-1}\ protein$. * $p < 0.05$ vs. control.

Table 1 The integrity of hepatocytes treated with anti-Fas antibodies, melatonin and betaine assessed by the levels of liberated enzymes^a

Experimental groups of hepatocytes/enzymes ($U\ l^{-1}$)	AST	ALT	γ -GT	ALP
Control	49.50 \pm 12.32	247.25 \pm 42.12	5.65 \pm 0.62	12.75 \pm 3.76
Anti-Fas antibody	166.15 \pm 23.45*** ■■■	616.68 \pm 83.85*** ■■■	7.52 \pm 1.95*** ■	27.85 \pm 5.23*** ■■■
Anti-Fas antibody + melatonin	63.50 \pm 18.24●	292.75 \pm 48.65●●●	5.50 \pm 1.04	10.75 \pm 2.99
Anti-Fas + betaine	98.00 \pm 12.54***	459.12 \pm 62.34***	7.00 \pm 1.34*** ●	13.60 \pm 3.87
Melatonin	50.25 \pm 12.22	329.75 \pm 42.81	3.25 \pm 1.02***	8.75 \pm 2.97***
Betaine	86.50 \pm 33.34***	324.71 \pm 53.13	6.50 \pm 2.95	10.50 \pm 3.19

^a The activities of liberated aspartate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphatase (ALP) and gamma-glutamyl transferase (γ -GT) were measured in collected supernatant and expressed as $U\ l^{-1}$. *** $p < 0.001$ vs. control; ■■■ $p < 0.001$ anti-Fas antibody vs. anti-Fas antibody + melatonin; ■ $p < 0.05$ anti-Fas antibody vs. anti-Fas antibody + betaine; ●●● $p < 0.001$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine; ● $p < 0.05$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine.



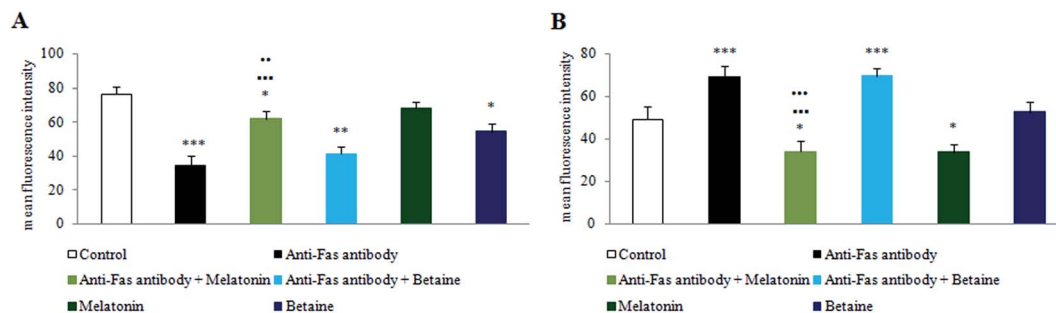


Fig. 5 Bcl-2 (A) and Bax (B) expression in hepatocytes treated with anti-Fas antibodies, melatonin and betaine. The mean fluorescence intensity of cells indicated for both, up and down regulation of populations for compounds and antibodies used, compared to intact hepatocytes. *** $p < 0.001$ vs. control; ** $p < 0.01$ vs. control; * $p < 0.05$ vs. control; ■■■ $p < 0.001$ anti-Fas antibody vs. anti-Fas antibody + melatonin; ●●● $p < 0.001$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine; ●● $p < 0.01$ anti-Fas antibody + melatonin vs. anti-Fas antibody + betaine.

shown to increase the expression of Bcl-2 and to decrease the expression of Bax protein in mouse hepatocytes, indicating decreased susceptibility of these cells to apoptosis and hepatic damage. Betaine hydrochloride exerted the opposite effect.

Melatonin and betaine hydrochloride nanoliposome encapsulation and detection of susceptibility to Fe^{2+} and H_2O_2 induced oxidative stress

Results obtained by using nanoliposomes as a possible artificial membrane-like phospholipid system and potential drug

delivery system suggest that melatonin as a lipophilic molecule may freely cross external cell membrane and be localized within the lipid nanospheres in about $74.48 \pm 1.04\%$, while the encapsulation efficiency of betaine hydrochloride was $51.40 \pm 3.99\%$. In this way melatonin may additionally contribute to the intracellular membrane structure stability and counteract oxidative stress-induced membrane damage.

Fig. 6 shows the intensity of H_2O_2 and FeSO_4 induced lipid peroxidation measured *via* thiobarbituric acid reactive substances (TBARS) production, regardless of the absence or presence of encapsulated melatonin or betaine hydrochloride in phospholipid liposome nanospheres. According to the results of the encapsulation of nanoliposomes with betaine hydrochloride, it would not be considered as a successful drug-delivery system. Fe^{2+} -dependent TBARS (MDA) production in nanoliposomes was markedly enhanced compared to H_2O_2 , significantly suppressed after melatonin encapsulation, but not after betaine hydrochloride encapsulation.

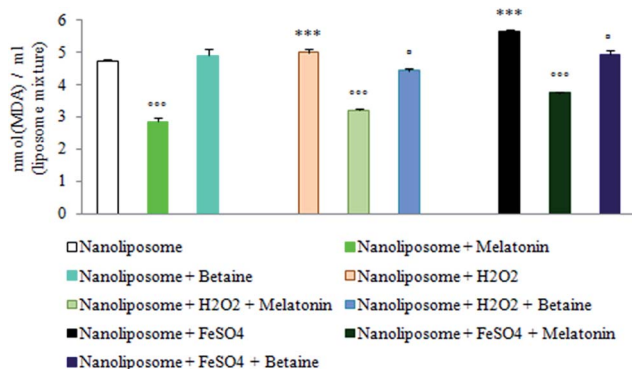


Fig. 6 Capability of melatonin- or betaine-encapsulated nanoliposome spheres to counteract H_2O_2 and FeSO_4 induced oxidative stress. The intensity of lipid peroxidation was estimated by the reaction of MDA with TBA, and measured spectrophotometrically. The MDA level was expressed as nmol (MDA)/ml (nanoliposome mixture). *** $p < 0.001$ vs. nanoliposome; *** $p < 0.001$ melatonin-encapsulated nanoliposome vs. nanoliposome; * $p < 0.05$ betaine-encapsulated nanoliposome vs. nanoliposome.

In silico study of calculated molecular physico-chemical properties of melatonin and betaine

Table 2 represents molecular physico-chemical properties of melatonin and betaine calculated by using Molinspiration tool.³⁰ Lipinski and coworkers established an effective methodology for estimation of potential drug solubility and permeability based on the calculation of molecular weight, octanol/water partition coefficient, number of H-bond donors and acceptors. Based on the analysis of large number of drugs, a cut-off for each of the four physico-chemical properties was set into the "Rule of five": poor absorption or permeation are more likely to occur when the molecule has molecular weight more

Table 2 Molecular physico-chemical properties of melatonin and betaine calculated by using *in silico* Molinspiration tool³⁰

Compound	$m_i \log P^a$	TPSA ^b	N_{atoms}^c	MW ^d	N_{ON}^e	N_{OHNH}^f	$N_{\text{viol.}}^g$	$N_{\text{roth.}}^h$	Vol ⁱ
Melatonin	1.45	54.12	17	232.28	4	2	0	4	220.12
Betaine	-5.41	40.13	8	117.15	3	0	0	2	119.60

^a Octanol-water partition coefficient, calculated by the methodology developed by Molinspiration. ^b Polar surface area. ^c Number of nonhydrogen atoms. ^d Molecular weight. ^e Number of hydrogen-bond acceptors (O and N atoms). ^f Number of hydrogen-bond donors (OH and NH groups). ^g Number of "Rule of five" violations. ^h Number of rotatable bonds. ⁱ Molecular volume.



than 500, log *P* over 5, and contains more than 5 H-bond donors or 10 H-bond acceptors. More than one violation of the rule is the critical limit for acceptable drug-likeness.³⁹ Summarizing the calculated physico-chemical properties of melatonin and betaine, it was concluded that both compounds obey the Lipinski "Rule of five" and meet all criteria for good solubility and permeability. Based on the physico-chemical determinants, absorption rate and oral bioavailability were predicted in order to gain a better insight into membrane permeability for melatonin and betaine. According to the results obtained by using ACD/I-Lab software³¹ betaine was predicted to have poor passive absorption of 34% (paracellular route), while melatonin was shown to have complete passive absorption of 100% (transcellular route).

Discussion

Membrane phospholipids represent a specific sensing system for external "death signals" or for intracellular damage, undergoing to perturbations of the membrane lipid bilayer in a way that phosphatidylserine (PS) is redistributed from the inner to the outer leaflet of the plasma membrane. The appearance of anionic PS at the cell surface is the most remarkable feature in apoptosis because of the abundance of this phospholipid, its negative charge and the ability to change interactions with other lipids that probably disrupts lipid rafts.⁴⁰ There is accumulating evidence showing that the generation of reactive oxygen species and subsequent oxidative stress associated with the initiation of apoptosis leads to selective oxidation of phospholipids such as PS in the cytoplasmic layer of membrane, followed by egress of oxidized PS (oxPS) to the surface of the apoptotic cells, effective recognition and cell clearance by phagocytes.^{41,42}

It has been shown that Fas-mediated apoptosis induces caspase-3 activation, chromatin condensation, PS externalization, generation of reactive oxygen species, intracellular glutathione depletion, disruption of mitochondrial transmembrane potential, release of cytochrome c (cyt c) from mitochondria, followed by the increased presence of oxPS within the cells and on the cell surface.⁴² Mitochondria play an essential role in production of oxPS and its subsequent transmigration to the cell surface during extrinsic apoptosis.⁴³ Serving as a naturally mitochondria-targeted protective molecule,¹⁸ melatonin regulates membrane stability.⁷ It is considered that a potential mechanism underlying selective oxidation of PS by H₂O₂ during apoptosis relies on the specific interactions between positively charged cyt c and negatively charged PS. Formed oxPS (PSOOH) may function as a substrate for cyt c peroxidase activity and further propagate lipid peroxidation. Therefore, the departure of cyt c from mitochondria is required for production of oxPS as well as its externalization. It has been reported that oxPS facilitates translocation of both PS and oxPS molecules to the cell surface.⁴¹ By the decrease of the Bax expression reported in our study, melatonin may inhibit the release of cyt c from mitochondria and, additionally, in this way, besides its antioxidative activity, reduce PS externalization.

Moreover, it has been shown that the inhibition of caspase 8 reduces mitochondrial cyt c release, the amount of oxPS and caspase 3 activation in Fas-induced apoptosis.^{1,44} Suppression of caspase-8 activity in Fas-mediated apoptotic pathways provides a new insight into the cytoprotective effect of melatonin.⁴⁴ It has been found that melatonin abrogates the caspase-8 activity in rabbit liver modulating TNF-mediated apoptotic pathway.⁴⁵ On the other hand, cyt c release induced by the proapoptotic Bcl-2 family member Bid inserted in the outer mitochondrial membrane can be inhibited by Bcl-2 overexpression. The overexpression of Bcl-2 protects against Fas-mediated apoptosis.¹ By the increase of Bcl-2 expression reported in this study, melatonin may inhibit mitochondrial release of cyt c, as well. The apoptotic-induced production, and externalization of oxPS has been detected in patients with alcoholic liver disease.⁴⁶

In the study on prevention of bile acid-induced apoptosis by betaine in rat hepatocytes *in vitro* and *in vivo*, it has been shown that betaine affected neither the bile acid-induced membrane trafficking of CD95 and tumor necrosis factor receptor nor recruitment of Fas-associated death domain and caspase 8 to the CD95 receptor.⁴⁷ By using a model of Fas ligand-induced apoptosis, we have obtained similar results.

Understanding the interaction and positioning of melatonin in biological membranes is an essential step towards decoding the mechanism of its influence on the membrane integrity, its protective abilities⁴⁸ and potential delivery of nanoliposome. Results of the evaluation of non-receptor phospholipid membrane melatonin uptake indicated that more than 74% of melatonin was encapsulated into lipid nanospheres, where it was able to counteract oxidative stress-induced membrane damage (Fig. 6). Since lipid peroxidation of cell membrane phospholipids caused by free-radicals has been proposed to be a major mechanism for the onset of a number of pathological events including apoptosis, the obtained result may suggest that membrane localization of melatonin may serve as the first line of defense against cell damage and apoptosis, which would not be considered in the case of betaine hydrochloride.

These results are in accordance with other literature data.^{47–51} Assuming that melatonin is located in the proximity of the polar heads of membrane phospholipids, molecules should exaggerate its activity against free-radicals coming from the aqueous compartment⁵⁰ protecting phospholipids from oxidation. Understanding the organization and interactions of melatonin in membranes may shed light into its antioxidative and cytoprotective properties,⁴⁸ and help develop a structural basis for construction of encapsulated nanoliposomes as suitable drug delivery system. By virtue of its ability to protect lipids from free-radical damage, melatonin is remarkably beneficial in preserving the morphological and functional integrity of cell membranes, in maintaining them at optimal fluidity and preventing them from becoming rigid. This contributes significantly to the function of proteins (receptors, channels, pores, *etc.*) in the cell membranes and helps in preserving their functional stability.¹⁷

The change in the Bax/Bcl-2 ratio may cause mitochondrial damage mediated by the breakdown of the mitochondrial transmembrane potential and the release of cyt c into the



cytoplasm along with subsequent caspase activation in the final apoptotic phenotype. The relative expression of antiapoptotic Bcl-2 is the key deciding point in regard to cell fate.²⁰ Oxidative stress is considered to be an important mechanism in the apoptosis of hepatocytes and the resulting liver damage. Melatonin has been documented to play a direct role in mitochondrial homeostasis by scavenging reactive oxygen species (ROS) and reactive nitrogen species (RNS), improving mitochondrial membrane integrity.⁵² It has been shown that melatonin may significantly decrease the Bax/Bcl-2 ratio in irradiation-induced apoptosis in rats' peripheral blood lymphocytes and liver.^{20,53} Some BH3-only proteins such as Bim may directly bind to and activate Bax, which leads to its oligomerisation, the permeabilization of the mitochondrial outer membrane, cyt c release and the activation of caspase cascade.^{54,55} Bim phosphorylation by the extracellular signal regulated kinase 1/2 (ERK1/2) promotes its degradation through the proteasome-ubiquitination system.⁵⁵ Contrary to Bim, the phosphorylation of Bcl-2 mediated by ERK2 might be of substantial importance for Bcl-2 stability and cell survival. Bcl-2 phosphorylation lends it its stability and rescues it from degradation and eventual cell death through the proteasome-ubiquitination system.⁵⁶ The activity of ERK was shown to be significantly stimulated by melatonin, suggesting that the mitogen-activated protein kinase (MAPK)/ERK pathway is involved in constitutive and stress-induced signaling of melatonin to promote mitochondrial protection and cell survival.⁵⁷ Moreover, it has been shown that the activation of the ERK1/2 pathway confers hepatocyte resistance to death.⁵⁸ It has been shown that melatonin may sustain the activation of Akt.^{59–62} In that way, by the activation of Akt, melatonin may inhibit Bax proapoptotic actions. It has been documented that Akt protects mouse hepatocytes from TNF- α and Fas-mediated apoptosis.⁶³ Therefore, this is one more possible mechanism of melatonin hepatoprotection mediated by the activation of Akt. Furthermore, MT₁/MT₂ signal transduction seems to be responsible for Bcl-2 re-localization to mitochondria, suggesting that Bax activation may in fact be antagonized by Bcl-2 at the mitochondrial level.⁶⁴ Bax translocation to the mitochondria and the antiapoptotic effect of melatonin have been shown to be dependent on the presence of Bcl-2 re-localized in the mitochondria.⁶⁵

Changes in the mitochondrial membrane can trigger further DNA cleavage, including nuclear membrane disintegration.² Melatonin was observed to be a protector of nuclear DNA fragmentation. The possible mechanisms would be free-radicals scavenging property, significant inhibition of free-radicals' formation by xanthine oxidase (Fig. 3) and the inhibition of DNase activity (Fig. 4). Obtained results are in accordance with those in our previous study about the protective effect of melatonin on rats exposed to microwave irradiation.^{13–15} In the present study, it was shown that exposure to betaine in Fas-mediated apoptotic conditions may induce an increase in DNase activity in treated hepatocytes. This result is in accordance with known literature data which claim that betaine was proven to considerably destabilize DNA.⁶⁶

Liver biomarkers (AST, ALT, ALP and γ -GT) are important criteria for the evaluation of the structural integrity of

hepatocytes and hepatotoxicity. The liberated amounts of enzymes indicate the severity of hepatocyte damage. In the present study, melatonin exhibited hepatoprotective effects by preserving liberation of hepatocyte enzymes, with significant effectiveness when it was combined with anti-Fas antibodies in comparison to anti-Fas antibody treatment alone. These results are in accordance with the literature data.^{67,68} Betaine combined with anti-Fas antibodies show less effectiveness in preserving membrane stability and liberation of hepatocyte enzymes. In the evaluation of the prevention of bile acid-induced apoptosis by betaine in rat hepatocytes *in vitro* and *in vivo* it has been shown that betaine may decrease the activity of these liver enzymes, but only to a low degree of effectiveness, when it was supplemented after bile duct ligation.⁴⁷ Our results documented that betaine hydrochloride could not be considered as a protector against membrane PS externalization nor Bcl-2/Bax expression and redistribution, lipid peroxidation, generation of free-radicals in reactions catalyzed by xanthine oxidase and DNA fragmentation during apoptosis, when it was co-incubated with anti-Fas antibodies or given alone. Obtained results may be interpreted in terms of betaine structure and its transport protein dependence on surrounding membrane phospholipids. The membrane receptors and transport proteins function in a fluid lipid bilayer environment, which is crucially involved in their proper organization and function. The activity and regulation of lipid-associated BetP is strongly dependent on the presence of negatively charged lipids around it.²⁶ These lipids may contribute to inactivation or activation of BetP *via* specific electrostatically favorable interactions. Modulation of transport protein activity by external stimuli may be an important aspect of cellular stress, including apoptosis. The influence of the membrane surface charge has shown that the increase of strongly negatively charged phospholipids may shift the threshold of activation or may lead to inactive BetP conformation.⁶⁹

The lipophilicity of compounds is a common property used to estimate the membrane permeability of a molecule. It is often measured as the log of the partition coefficient between *n*-octanol and water. The partition coefficient values (Table 2), calculated by using the Molinspiration tool,³⁰ indicated that betaine is more hydrophilic than melatonin, which means that it may show lower membrane permeation, absorption rate and bioavailability. Regarding hydrogen binding capacity, both melatonin and betaine possess less than 5 H-bond donors and acceptors. The conformational flexibility of the molecules described by the number of rotatable bonds is important factor, among others, for the assessment of the binding efficiency of receptors and channels. The satisfactory number of rotatable bonds is 10 or fewer, and both of the tested compounds meet these criteria. In addition to the number of rotatable bonds, Veber and coworkers found that the polar surface area can be used as a good indication of permeation. The membrane permeation rate is recognized as a common requirement for bioavailability. The higher hydrophilicity makes the interaction and uptake over a lipid bilayer difficult. The suggested Veber rules are: acceptable oral bioavailability is more likely to occur when the molecule has 10 or fewer rotatable bonds, and polar surface area equal to or less than 140 Å² and 12 or fewer



H-donors and acceptors. Veber and coworkers therefore also raise the issue of molecular weight as a proper descriptor for permeation and absorption measurement, as it might just be positively correlated with more precise properties like the rotatable hydrogen bonds count and polar surface area.⁷⁰ It was concluded that both melatonin and betaine obey the Veber rules. The data obtained by ACD/I-Lab³¹ indicate that the oral bioavailability of melatonin is more than 70%, and of betaine less than 30%. These values are in accordance with criteria of Egan rule, relating to the assessment of potential drug oral bioavailability based on the calculation of the octanol/water partition coefficient and polar surface area.⁷¹ All of the above may indicate that the cell membrane permeation of melatonin is higher than the permeation of betaine.

In conclusion, increased formation of reactive oxygen species and lipid peroxidation in anti-Fas-antibody induced apoptosis was reduced by melatonin. In this way, melatonin may contribute to mitochondrial membrane stability and integrity and to nuclear membrane stability. The above mentioned effects are supported by data on the physico-chemical characteristics of both tested compounds obtained using *in silico* studies. Considering the importance of apoptosis in a variety of liver diseases, the evidence and clarified mechanisms of melatonin actions in the apoptotic and oxidative processes, knowing its physico-chemical properties, represent important prerequisites for the formulation of future non-toxic nanoliposome-mediated therapy for clinical practice. In order to provide recommendations for successful treatment of hepatic apoptosis, it could therefore be preferable the use of lipophilic compounds, like melatonin, rather than more hydrophilic ones like betaine.

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