Neuroprotective and MAOB inhibitory effects of a series of caffeine-8-thioglycolic acid amides

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The effects of new derivatives of caffeine-8-thioglycolic acid (100 μM) on isolated rat brain synaptosomes, human neuroblastoma cell line SH-SY5Y and human recombinant MAOB enzyme (hMAOB) (1 μM) were evaluated. Most of the compounds, administered alone, didn’t show statistically significant neurotoxic effects on SH-SY5Y, when compared to the control (non-treated cells). Of all studied structures JTA-2Ox, JTA-11, JTA-12 and JTA-13 decreased cell viability. In combination with 6-hydroxydopamine (6-OHDA) (100 μM), only JTA-1 and JTA-2 revealed neuroprotective effects, stronger than those of caffeine. All compounds administered alone revealed, neurotoxic effects on synaptosomes, as compared to non-treated synaptosomes. JTA-1, JTA-2 and JTA-3 showed lowest neurotoxic effects and were investigated in a model of 6-OHDA-induced oxidative stress. In this model of neurotoxicity, only JTA-1 and JTA-2 showed statistically significant neuroprotective effect, by preserving the synaptosomal viability and the level of reduced glutathione. Inhibition of hMAOB, was revealed by JTA-1 and JTA-2. They inhibited the enzyme by 23% and 25% respectively, thus approaching the selegiline activity, which was 42%. The possible mechanisms of neuroprotection of JTA-1 and JTA-2 might be a result from the inhibition of hMAOB, which catalyze the production of neurotoxic p-quinone from 6-OHDA.

Keywords: Neuroprotection. Caffeine. Synaptosomes. SH-SY5Y. hMAOB.

INTRODUCTION

Neurodegenerative diseases are characterized by loss of neurons in certain regions of the brain. They are incurable and debilitating conditions that result in progressive degeneration and/or death of nerve cells. This causes problems with movement (ataxias), or mental functioning (dementias).

For a couple of decades, significant ageing of the population has been observed worldwide and the prevalence of such disorders is growing continuously. At present, therapeutic strategies for treatment of neurodegenerative diseases are only symptomatic and none of them is able to terminate or delay the disease progression.

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Parkinson’s disease (PD) is one of the most common chronic neurodegenerative diseases following Alzheimer’s disease (Tieu, 2011). Approximately 630,000 people in US were diagnosed with PD only in 2010 (Kowal et al., 2013). The symptoms of the disease were described by James Parkinson in his monograph “An Essay on the Shaking Palsy” for the first time in 1817. Parkinsonism is a syndrome characterized by resting trepidation, rigidity, delayed or absent conscious movements and postural instability. So far, there has no cure been found and the causes of the disease remain not fully understood, with less than 10% of the cases being directly associated with monogenic mutations (Tieu, 2011).

PD is a progressive disease caused by damage of the central nervous system nigrostriatal neuronal pathways. The pathophysiological distinguishing features of PD are loss of dopaminergic neurons in substantia nigra pars compacta and the presence of cytoplasmic protein aggregates called Lewy bodies. When degeneration of these neurons reaches a threshold of about 60% and
striatal dopamine decreases by about 80%, symptoms of PD appear (Dauer, Przedborski, 2003).

The current treatment of PD is generally focused in compensating dopamine deficiency, which is ineffective in delaying progression. Dopamine replacement therapy is associated with many side effects, especially dyskinesia. Recent progress in understanding of molecular mechanisms involved in PD has revealed new perspectives for PD therapy. In these strategies, caffeine and xanthine derivatives are promising scaffolds for design of novel structures with valuable neuroprotective properties.

Although pathogenesis of PD remains unclear, there is an increasing evidence for the involvement of mitochondrial function, oxidative damage and inflammation in the development of the neurodegenerative process (Dauer, Przedborski, 2003; Bagga, Patel, 2016).

Caffeine (1,3,7-trimethylxanthine) is a natural alkaloid and similarly toxanthines, act as non-selective adenosine A₃ receptors antagonist. Caffeine easily crosses the blood-brain barrier and exhibits a variety of behavioral effects, including sharpened alertness, reduced fatigue and increased motor activity (Smith, 2002) by inhibiting the suppression of neuronal function (Garrett, Griffiths, 1997).

Several clinical trials and meta-analyses have showed that caffeine consumption is associated with a decreased risk of developing PD and dementia (Ross, Petrovitch, 2001; Ascherio et al., 2001; Costa et al., 2010; Altmann, Lang, Postuma, 2011; Palacios et al., 2012; Liu et al., 2012; Kolahdouzan, Hamadeh, 2017).

Other studies report similar results and the evidence for the potential beneficial effects of caffeine and xanthine derivatives administration is growing.

An investigation on the effect of pre-treatment with caffeine on the expression of genes in MPTP-lesioned mouse striatum, has been performed by Singh and colleagues in 2010. Several genes responsible for apoptotic cell death, oxidative stress, cell cycle regulation, protein modification and mitochondrial dysfunction were examined. The results revealed significant reduction of damaging events after caffeine treatment (Singh et al., 2010).

Sonsalla and co-workers found out in their study that caffeine administration upon the onset or throughout the course of ongoing neurodegeneration, reduced the loss of dopaminergic neurons in substantia nigra and might stop or slow down the degeneration process (Sonsalla et al., 2012).

In their paper, Kaster and co-workers (2015) revealed the relation of caffeine and istradefylline with mood and memory alterations provoked by chronic stress. The results show that chronic stress induces behavioral and synaptic alterations and xanthine based structures, such as caffeine and istradefylline, prevented those alterations (Kaster et al., 2015).

A lot of efforts have been made to uncover the therapeutic potential of xanthine structures as neuroprotective agents over the last decades. The substituted xanthine molecule is used as a scaffold for the synthesis of new compounds with protective effects in neurodegenerative diseases. The use of the xanthines has been proposed as a non-dopaminergic strategy for neuroprotection in Parkinson’s disease and the mechanisms of protection have been associated with antagonism of adenosine A₂A receptors and monoamine oxidase type B (MAOB) inhibition (Palacios et al., 2012). Adenosine A₂A receptor antagonism leads to reduced inflammation through decreased proliferation and activity of the astroglial cells in the midbrain (Bagga, Patel, 2016; Madeira et al., 2017). On the other hand, the neuroprotective effects of MAOB inhibitors can be explained in part, taking into account the metabolic products generated by the action of MAOB on monoamines. In the catalytic reaction of MAOB, aldehyde product and hydrogen peroxide are produced for each mole of oxidized monoamine substrate (Petzer et al., 2009). These metabolic products may be neurotoxic, if they are not rapidly inactivated by antioxidant systems, whether enzymatic or non-enzymatic.

Additionally, some data indicate a possible dual target activity of methylxanthine derivatives with pronounced neuroprotective effect, which is expressed in a potent and selective antagonism on adenosine A₂A receptors coupled with MAOB inhibition (Petzer et al., 2009).

Based on this data, a series of synthesized caffeine-8-thioglycolic acid amides was evaluated for possible neuroprotective effects as MAOB inhibitors. The neurotoxicity of the compounds on SH-SY5Y neuroblastoma cells was also investigated.

**MATERIAL AND METHODS**

**Synthesis of caffeine-8-thioglycolic acid (KTG) derivatives (JTA1 to JTA 13)**

A modified method of Persch and Beyerle (1957) was used to synthesize caffeine-8-thioglycolic acid (Persch, Beyerle, 1957). The reaction was carried out in 60% ethanol and the yield was 92%. Methyl caffeine-8-thioglycolate was prepared by the method used for synthesizing ethyl esters of 7-alkylxanthinyl-8-thioglycolic acids. The reaction was carried out in anhydrous methanol in the presence of a catalytic amount
of sulfuric or phosphoric acid and the yield was 95%. After aminolysis of methyl caffeine-8-thioglycolate, a series of amide derivatives were obtained (Mitkov et al., 2007). The structures of the synthesized amide derivatives were proven by IR, ¹H and ¹³C-NMR spectral data. The purity of the compounds was confirmed by the corresponding TLC characteristics, melting points and elemental analyses.

**Chemicals**

The chemicals used in the experiments:
- 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), Roswell Park Memorial Institute (RPMI) 1640 Medium, Fetal Bovine Serum 10%, L-glutamine, 6-OHDA, Percoll, 4-(2-hydroxyethyl)-1-piperazinenuenesulfonic acid (HEPES), sucrose, 2-thiobarbituric acid (4,6-dihydroxypyrimidine-2-thiol) and Tyramine hydrochloride were obtained from Sigma Aldrich, Germany;
- NaCl, KCl, CaCl₂.2H₂O, MgCl₂, NaHPO₄, D-glucose, trichloroacetic acid, 2,2’-dinitro-5,5’-dithiobisbenzoic acid (DTNB) and selegiline were obtained from Merck, Germany;
- Amplex UltrRed Kit was obtained from Invitrogen, USA.

**Cultivation and incubation of human neuroblastoma cell line SH-SY5Y**

The SH-SY5Y human neuroblastoma cell line was received from Sigma Aldrich (ECACC cell lines). The cultivation process was carried out with nutritional media RPMI-1640, supplemented with fetal bovine serum 10% and L-glutamine. Sub-cultivation process was carried twice a week. When about 80% confluence was reached the cells were seeded in 96-well plate, and placed for 24 h in an incubator under standard conditions (37 °C, 5% CO₂) for precipitation and adhesion (seeding cell density 3.10⁴ per well). The cells were incubated with solutions of the test substances in nutrient medium (final concentration 100 µM) (Timson, 1970) followed by 24 hours incubation.

**Model of neurotoxicity on SH-SY5Y cell line**

The neurotoxicity experiments were performed using 100 µM 6-OHDA and the test substance for 24 hours. In our preliminary experiments, we investigated 5 different concentrations of the toxic agent: 50 µM, 100 µM, 150 µM, 200 µM and 250 µM and as a result we found that concentration 100 µM of 6-OHDA lead to 50% cell death and was convenient for the experiments.

**MTT test: vitality assessment method**

The tetrazole salt, MTT, is reduced by the action of mitochondrial dehydrogenases to water insoluble violet crystals formazan. The culture medium was aspirated and added 100 µL of a solution of freshly prepared MTT (0.5 mg/mL) in culture medium and 3 hours incubation was carried out. The liquid from the wells was aspirated and added 100 µL of dimethyl sulfoxide (DMSO) and maintained 10 minutes with shaking at dark. Absorbance was measured at 570 nm and 690 nm. The amount of formed formazan is proportional to viability (Mosmann, 1983).

**Animals**

Male Wistar rats (body weight 200–250 g) were used. The rats were housed in plexiglass cages (3 per cage) in a 12/12 light/dark cycle, under standard laboratory conditions (ambient temperature 20 ± 2 °C and humidity 72 ± 4%) with free access to water and standard pelleted rat food 53-3, produced according ISO 9001:2008. The animals were purchased from the National Breeding Center, Sofia, Bulgaria. Seven days acclimatization was allowed before the commencement of the study and a veterinary physician monitored the health of the animals regularly. Vivarium (certificate of registration of farm No. 0072/01.08.2007) was inspected by the Bulgarian Drug Agency in order to check the husbandry conditions (No. A-11-1081/03.11.2011). All performed procedures were approved by the Institutional Animal Care Committee and the principles stated in the European Convention for the Protection of Vertebrate Animals used for Experimental and other Scientific Purposes (ETS 123) (Council of Europe, 1991) were strictly followed throughout the experiment.

**Isolation and incubation of synaptosomes**

The synaptosomes were prepared by multiple, subcellular fractionation using a Percoll gradient (Taupin et al., 1994). Two types of buffers were prepared – Buffer A: HEPES 5 mM and Sucrose 0.32 M; Buffer B: NaCl 290 mM, MgCl₂,2H₂O, 0.95 mM, KCl 10 mM, CaCl₂,2H₂O, 2.4 mM, NaH₂PO₄, 2.1 mM, HEPES 44 mM, D-Glucose 13 mM.

Brain homogenates were prepared with buffer A and centrifuged at 1000 x g for 10 minutes at 4 °C. After centrifugation, the supernatants were removed and centrifuged again under the above-mentioned conditions. Supernatants were then taken and subjected to three times centrifugation at 10 000 x g for 20 minutes at 4 °C. The last two centrifuges were for purification of the synaptosomes.
Isolation of synaptosomes was performed with the help of a colloidal silicon solution (Percoll).

1. Preparation of 90% stock solution of Percoll.
2. Preparation of Percoll solutions with two concentrations 16% and 10%. 4 mL of Percoll 16% and 10%, were added in the tubes.
3. 90% Percoll (7.5% Percoll) was added to the sediment from the last centrifugation. The tubes were centrifuged for 20 minutes at 15 000 x g at 4 °C.

After centrifugation, three layers were formed in the tubes. The lower layer contained mitochondria, the upper layer - lipids and the middle layer (between 16% and 10% Percoll) - synaptosomes. The middle layer of each tube was harvested and buffer B with glucose was added. The mixture was centrifuged at 10 000 x g for 20 minutes at 4 °C. After centrifugation, the sediment with the synaptosomes was mixed with buffer B with glucose.

The synaptosomes and the cells were incubated with 100 µM of the test substances (Timson, 1970).

After the incubation, MTT-test was performed to determine synaptosomal vitality by method described by Mungarro-Menchaca et al. (2002).

**Determination of reduced glutathione (GSH)**

The level of reduced glutathione was determined by measuring the non-protein SH-groups after precipitation of the proteins with trichloroacetic acid.

After the incubation, synaptosomes were centrifuged at 400 x g for 3 minutes. The sediment was treated with 5% trichloroacetic acid and left for 10 minutes on ice. Samples were centrifuged at 8 000 x g for 10 minutes (2 °C). The supernatant was removed to determine the level of GSH and was stored at -20°C. Immediately before the measurement, the samples were neutralized with 5 N NaOH.

The presence of thiols in the supernatant was determined using Elman reagent. The resulting yellow color was measured spectrophotometrically (λ = 412 nm) (Robyt, Ackerman, Chittenden, 1971).

**Model of 6-OHDA-induced neurotoxicity**

This in vitro model resembles the neurodegenerative processes occurring in PD. Dopamine metabolism and oxidation lead to the formation of reactive oxygen species (ROS) and reactive quinones. They induce dopamine neurotoxicity and neurodegeneration (Stokes et al., 2002). The synaptosomes were incubated with 150 µM 6-OHDA and the test substance for 1 hour. After centrifugation, on a microfuge, the probes samples were centrifuged for 1 minute at 15 000 x g. The pellet was mixed with Buffer B and centrifuged again for 1 minute at 15 000 x g. The precipitate containing the synaptosomal fraction was mixed with buffer B with glucose.

**Measurement of Monoamine oxidase B activity**

Monoamine oxidase activity assay of recombinant human MAOB was performed using a fluorometric method with Ampex UltraRed reagent (Bautista-Aguilera et al., 2014) with small modifications. Tyramine hydrochloride was used as substrate.

**Statistical methods**

Statistical analysis was performed using statistical programme “MEDCALC”. Results are expressed as mean ± SEM for 6 experiments. The significance of the data was assessed using the non-parametric Mann-Whitney test (synaptosomes). Values of p ≤ 0.05; p ≤ 0.01 and p ≤ 0.001 were considered statistically significant. The cell survival data were normalized as percentage of the control (set as 100% viability). Statistical evaluation was performed using the GraphPad Prism 5.0 software.

**RESULTS AND DISCUSSION**

The general procedure for the preparation of the desired amides is described elsewhere (Mitkov et al., 2007) and shown in Figure 1.

![FIGURE 1 - Synthesis of caffeine-8-thioglycolic acid (KTG) amides.](image-url)
Neuroprotective and MAOB inhibitory effects of a series of caffeine-8-thioglycolic acid amides

Effects of newly synthesized caffeine-8-thioglycolic acid (KTG) derivatives on neuroblastoma cell line SH-SY5Y

The SH-SY5Y neuroblastoma cells are often used as in vitro models of neuronal function and differentiation. They are adrenergic in phenotype but also express dopaminergic markers and therefore are appropriate to study Parkinson’s disease.

Caffeine and caffeine-8-thioglycolic acid (KTG), administered alone, didn’t reveal statistically significant neurotoxic effects, compared to the control (non-treated cells). Only JTA-2Ox, JTA-11, JTA-12 and JTA-13 decreased significantly the cell viability by 41%, 25%, 37% and 39%, respectively (Figure 2).

Some data suggest the presence of free radical scavenging activity of caffeine and other methylxanthines (Shi, Dalal, Jain, 1991). Thus, it was of interest to determine the effect of the less toxic caffeine amides JTA-1 – JTA-10 on a model of 6-OHDA-induced oxidative stress in SH-SY5Y cell line.

When administered alone, 6-OHDA decreased the cell viability with 48%, compared to the control (non-treated cells). After the administration of the evaluated structures on this model, only JTA-1 and JTA-2 revealed good statistically significant neuroprotective effects, better than those of caffeine-8-thioglycolic acid (KTG) and caffeine. JTA-1 preserved cell viability by 56% and JTA-2 by 54%, while caffeine-8-thioglycolic acid and caffeine – by 31%, compared to the toxic agent (Figure 3).

**FIGURE 2** - Effects of caffeine-8-thioglycolic acid (KTG) derivatives, administered alone, on SH-SY5Y cell viability, measured by MTT-test. After 24 h of treatment with test substances cell viability is expressed as a percentage of a control (non-treated cells). Mean ± SEM (n = 6). ** P < 0.01; *** P < 0.001 – vs control.

**FIGURE 3** - Effects of JTA-1 to JTA-10, in conditions of 6-OHDA-induced oxidative stress, on SH-SY5Y cell viability, measured by MTT-test. After 24 h of treatment with test substances and toxic agent, cell viability is expressed as a percentage of a control (non-treated cells). Mean ± SEM (n = 6). *** P < 0.001 – vs control; ++ P < 0.01; +++ P < 0.001 – vs 6-OHDA.
Effects of newly synthesized caffeine-8-thioglycolic acid (KTG) derivatives on isolated rat brain synaptosomes

Along with the other brain models (brain slices, primary neuronal cultures), the isolated nerve terminals – synaptosomes, are an important tool for investigation of the synaptic function in the brain on molecular level (Evans, 2015).

Using this useful model system for assessment of the neuroprotective properties of the studied amides we evaluated their effect on isolated rat brain synaptosomes. The necessary rat brain synaptosomes were prepared by using Percoll reagent. All the tested derivatives, along with the initial caffeine and KTG expressed statistically significant neurotoxic effects on synaptosomal viability and level of GSH, compared to the control (non-treated synaptosomes).

Only compounds JTA-1, JTA-2 and JTA-3 presented lower neurotoxic effects on isolated rat brain synaptosomes (Figures 4 and 5). These 3 compounds were further evaluated for possible neuroprotective effects on a model of 6-OHDA-induced oxidative stress.

Based on data showing that caffeine prevents lipid peroxidation (Devasagayam et al., 1996) we determined the effect of the less toxic amide derivatives on 6-OHDA induced oxidative stress.

The treatment of isolated rat brain synaptosomes with 6-OHDA is a reliable and commonly used in vitro model for the investigation of processes, which play role in the neurodegenerative disease, including Parkinson’s and Alzheimer’s disease. The mechanism of 6-OHDA neurotoxicity includes the formation of ROS and reactive metabolites, as a result of its metabolism in mitochondria of the neuronal cells (Stokes et al., 2002).

The mechanism of destruction of the nerve terminals is thought involvement of oxidation of 6-OHDA to a p-quinone and production of a free radical or of superoxide anion. The species intermediate reacts covalently with the nerve terminal and permanently inactivates it (Timbrell, 2003).

When administered alone, 6-OHDA decreased synaptosomal viability and GSH level by 55% and 50%, respectively, compared to the non-treated synaptosomes, while both, JTA-1 and JTA-2 revealed stronger than JTA-3 statistically significant neuroprotective effects compared to 6-OHDA by preserving the synaptosomal viability and level of GSH to higher extent (Figure 6).

JTA-1 preserved synaptosomal viability and level of GSH by 56% and 44%, respectively, compared to toxic agent, while JTA-2 preserved synaptosomal viability and level of GSH by 58% and 46%, respectively, compared to toxic agent. The neuroprotective effects of JTA-1 and JTA-2 were similar to those of caffeine and KTG on this model of oxidative stress.

The third non-toxic amide JTA-3 - preserved the synaptosomal viability and level of GSH by 24% and 20%, respectively, compared to the toxic agent, which demonstrates its lower effect than those of caffeine and KTG.

Effects of 14 newly synthesized derivatives of caffeine-8-thioglycolic acid (KTG) and selegiline on the activity of hMAOB enzyme

MAOB is involved in the neurodegenerative process associated with aging, and is known to participate in neurodegenerative diseases including Parkinson’s and Alzheimer’s disease. MAOB inhibitors could prevent dopaminergic neuron degeneration (Timbrell, 2003) and reduce parkinsonian symptoms (Löhle, Storch, 2012). MAOB inhibitors could decrease metabolism of dopamine

**FIGURE 4** - Effects of caffeine-8-thioglycolic acid (KTG) derivatives, administered alone, on synaptosomal viability, measured by MTT-test. After 1 h of treatment with test substances synaptosomal viability is expressed as a percentage of a control (non-treated synaptosomes). Mean ± SEM (n = 6). **P < 0.01; ***P < 0.001 – vs control.
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selectively and balance its concentration in synaptic cleft in a relatively stable state (Timbrell, 2003).

Based on the information on dual-target-acting agents developed in the last years as neuroprotective molecules, we assessed newly synthesized caffeine-8-thioglycolic acid amides JTA-1-JTA-13 for potential MAOB inhibitory activity.

The results revealed that most of the examined compounds administered alone (at concentration 1 μM) didn’t have statistically significant inhibitory effects on the activity of hMAOB, if compared to the control (pure hMAOB).

Only two of the compounds: JTA-1 and JTA-2 showed significantly good inhibitory activity on hMAOB, closer to Selegiline’s activity. JTA-1 inhibited hMAOB by 23% and JTA-2 – by 25%, compared to pure hMAOB (Figure 7).

Selegiline (at concentration 1 μM) inhibited hMAOB by 42%.

This study, lead us to the following conclusions:

- In SH-SY5Y cell line, most of the derivatives, administered alone, didn’t show statistically significant neurotoxic effects, compared to the control (non-treated cells). In conditions of 6-OHDA-induced oxidative stress, only JTA-1 and JTA-2 revealed good neuroprotection which was better than that exerted by caffeine-8-thioglycolic acid and caffeine.

On rat brain synaptosomes, administered alone, all compounds revealed statistically significant neurotoxic effects compared to the control (non-treated synaptosomes). JTA-1, JTA-2 and JTA-3 showed lowest neurotoxic effects and were investigated in a model of 6-OHDA-induced oxidative stress. Under the conditions of 6-OHDA neurotoxicity, only JTA-1 and JTA-2, with proven antihypoxic effects, showed significant neuroprotective effects on brain synaptosomes.
FIGURE 7 - Effects of caffeine-8-thioglycolic acid (KTG) derivatives and selegiline (at concentration 1 μM), on hMAOB enzyme. Mean ± SEM (n = 6). ** P < 0.01; *** P < 0.001 vs control (pure hMAOB).

- Statistically significant inhibitory activity on hMAOB, was revealed by JTA-1 and JTA-2. They inhibited the enzyme by 23% and 25%, respectively, which was close to selegiline’s effect, which inhibited hMAOB by 42%.

- The possible mechanisms of JTA-1 and JTA-2 neuroprotection on 6-OHDA-induced toxicity model might be due to the inhibition of hMAOB, which catalyze the production of neurotoxic p-quinone from 6-OHDA.

- The obtained results highlight JTA-1 and JTA-2 as the most promising compounds for further consideration as a structural base for design of new neuroprotective agents.

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