

## Synthesis and antioxidant activity of phenylcarbamic acid derivatives acting on the cardiovascular system

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### Introduction

Reactive oxygen species are formed during many metabolic reactions—metabolism of nutrients, biosynthesis of certain substances, electron transport in the mitochondria, for instance phagocytosis and inflammatory processes, trauma and physical exercise. To get rid of reactive oxygen species, the human body uses various antioxidant systems. Antioxidants used<sup>1)</sup> are hydrophilic, i.e., water soluble, which are greater in the extracellular fluid (vitamin C, uric acid, selenium, bioflavonoids, and i.) and the lipophilic ones, i.e., fat-soluble, which act mainly on the membranes within cells (vitamin E, beta-carotene, ubiquinone Q10, etc.)<sup>1)</sup>. Antioxidant activity is defined as the ability of a particular compound or a mixture of compounds to inhibit oxidative degradation. One should distinguish two concepts, namely antioxidant capacity and activity. Antioxidant capacity provides information on the duration of the antioxidant effect, whereas the activity is characterized by an initial process dynamics during the antioxidant at a specific concentration of an antioxidant<sup>2)</sup>.

The  $\beta$ -adrenergic receptors antagonists ( $\beta$ -ARAs) have been primarily used as drugs in the treatment of cardiovascular diseases such as ischaemic heart disease, *angina pectoris*, arrhythmia, hypertension, cardiomyopathy, or as prevention after myocardial infarction<sup>3)</sup>. In addition, more recently they have been accepted in the heart failure therapy. Part of the beneficial cardiovascular effects shown by such a group of compounds has already been associated with the antioxidant properties that some of them have possessed – carvedilol, atenolol, carteolol or metipranolol<sup>4)</sup>.

### Experimental methods

Compounds under the study:

Alkylesters of 4{2-hydroxy-3-[(4-pyridine-2-yl)pipezine-yl]-propoxy}phenylcarbamic acid monochloride.

### ABTS method

**Chemicals:** Potassium dihydrogen phosphate (CentralChem, SR), potassium persulfate (Sigma-Aldrich, USA), 2,2'-azinobis-(3-ethylbenzothiazoline-6-sulfonic acid), (ABTS, Sigma-Aldrich, USA), ethanol (CentralChem, SR), distilled water. **Method used:** Into a 10 ml volumetric flask an amount of 0.038 g of ABTS radical was added and was made up with distilled water. Into another 10 ml flask, 0.007 g of potassium dihydrogen phosphate and water were added as well. Prepared solutions were mixed in a ratio of 1 : 1 to give a blue-green colour, and left to stand for 24 h in the dark at room temperature. After 24 h, a volume of 1.1 ml of concentrated radical was transferred to a 50 ml volumetric flask and ethanol was added up to the mark. To the volume of 2 ml of ABTS solution, 100  $\mu$ l of the sample ( $c = 10^{-4}$  mol/l) was added. After 5 minutes, the absorbance of each solution was measured at  $\lambda = 734$  nm. As the reference solution, a reaction system without the sample was used. Ascorbic acid was applied as the control solution<sup>5)</sup>.

**Evaluation of results:** Presence of active substances with antioxidant agent led to discolouring of the solution, which was measured spectrophotometrically. Antioxidant activity (%) =  $100 \times (A_p - A)/A_p$ , where  $A_p$  is the absorbance of the reference solution,  $A$  the absorbance of the samples DPPH method.

**Chemicals:** 1,1-Diphenyl-2-(2,4,6-trinitrophenyl)hydrazyl (DPPH, Sigma-Aldrich, USA), methanol (CentralChem, SR).

**Method used:** DPPH ( $m = 1$  mg) was dissolved in methanol and the solvent was made up to a 50 ml volumetric flask. To the 200  $\mu$ l of evaluated compounds ( $c = 10^{-4}$  mol/l) was added 1.8 ml of DPPH radical. The activity of the samples was evaluated after 30 min by measuring the absorbance of each solution at  $\lambda = 517$  nm. As referenced solution was considered reaction system without the sample. Ascorbic acid was applied as a control solution<sup>5)</sup>.

**Evaluation of results:** After DPPH radical solution reduction, the colour changed from the original purple to yellow. The colour change was measured spectrophotometrically according to the equation given below.

Antioxidant activity (%) =  $100 \times (A_p - A)/A_p$ , where  $A_p$  is the absorbance of the reference solution,  $A$  the absorbance of the samples.

### FRAP method

**Chemicals:** Sodium acetate (CentralChem, SR), 35%

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solution of hydrochloric acid (CentralChem, SR), ferric chloride (CentralChem, SR), 2,4,6-tris-(2-pyridyl)-s-triazine (TPTZ, Sigma-Aldrich, USA).

**Used method:** Acetate buffer was prepared by adding 3.1 g of sodium acetate and 16 ml of concentrated acetic acid to a 1000 ml volumetric flask and made up with distilled water. TPTZ ( $m = 0.1562$  g) and 0.177 ml of a 35% HCl solution were made up with distilled water into a 50 ml volumetric flask.  $\text{FeCl}_3$  ( $m = 0.2703$  g) was dissolved in a 50 ml volumetric flask and made up to the mark with distilled water. FRAP solution was made by mixing 25 ml of acetate buffer, 2.5 ml of TPTZ and 2.5 ml of  $\text{FeCl}_3$  solution. To 100  $\mu\text{l}$  of a compound solution ( $c = 10^{-4}$  mol/l) 3 ml of FRAP solution were added. Absorbance was measured after 5 min at  $\lambda = 593$  nm, using an ascorbic acid solution as the control solution<sup>9</sup>.

**Evaluation of results:** The results were evaluated as the analogical amount of ascorbic acid ( $\mu\text{g/ml}$ ) =  $A/0,134$ , where  $A$  is the absorbance of compounds.

## Results and discussion

In the area of chemical analysis and biological evaluation of drugs, many methods have been developed to determine the total antioxidant activity of the sample (Total Antioxidant Activity – TAA). The purpose of these methods is to characterize compounds in such conditions close to the physiological environment of the antioxidant<sup>7</sup>.

There is an effort to interpret antioxidant activity of the sample in a way that would be methodical, material and instrumental accessible and usable for numerous serial analyses. Their results should correspond to the biological value of the test substance. Chemical methods (TEAC, FRAP, DPPH, ABTS, ORAC or DCI) are based on the use of agents providing the oxygen free radical colour products which prevent the formation of antioxidants contained in the sample<sup>8</sup>.

The capability of inspected compounds to act as potential antioxidants was evaluated following the given methods (DPPH, ABTS and FRAP), respectively. The results obtained could be influenced by some physico-chemical aspects: lipohydrophilic properties of the molecules under the study, by isosteric replacement of ether bond for carbamoyloxy moiety, and by electrostatic and steric effects of pyridin-2-yl fragment<sup>9</sup> (Table 1).

In terms of the chemical structures of tested compounds  $4a_1$ – $4d_1$ , the elongation of the alkyl side chain has not led

to more promising antioxidative substances in all performed methods. Following the DPPH method in the compound  $4b_1$  could be regarded as the most potent substance. On the other hand, results of ABTS testing have shown a connection between elongation of the alkyl side chain and activities of analyzed substances. The highest potential has been estimated for compound  $4d_1$ . Antioxidant properties of aryloxyaminopropanols  $4a_1$ ,  $4b_1$ ,  $4c_1$ ,  $4d_1$  determined by the FRAP method have not characterized their antioxidant activity because the method only describes the reduction potential of antioxidants and has not defined the ability to scavenge free radicals. Therefore the antioxidant activity of the inspected substances by the DPPH and ABTS method could not be compared to the antioxidant activity of the FRAP method. Nevertheless, any connection between the length of the alkyl side chain and the activity has not been revealed. The highest activity has been shown for the molecule  $4c_1$ , the lowest one for compound  $4a_1$ . For more convenient interpretation of the results, the outgoings of the analyzed substance have been compared to those of  $\beta$ -blockers in clinical use (carvedilol, atenolol) with antioxidative features and two strong antioxidants (Trolox, ascorbic acid).

It has been revealed that the antioxidant activity of atenolol and carvedilol determined by the DPPH method was notably lower than that of any other substance under study. Results from the ABTS study have pointed a similar antioxidant ability of carvedilol as compared to tested compounds. Atenol has reached the capability similar to the lowest active substance  $4a_1$  only. Results from the FRAP determination have shown a lower redox potential of both analyzed  $\beta$ -blockers. On the other hand, trolox and ascorbic acid have shown a strong antioxidative potential in all performed methods.

## Conclusion

Considering the important role of antioxidants in biological systems, series of original  $\beta$ -adrenergic receptor antagonists have been evaluated *in vitro* for their ability to reduce the DPPH, ABTS and FRAP radicals as well. The results might contribute to their cardiovascular therapeutic benefits.

Focusing on performed *in vitro* experiments, it has been suggested that a presence of the carbamoyloxy moiety could be regarded as favourable structural modification. However, relatively high lipophilicity of all the

Table 1. The potential of the compounds  $4a_1$ – $4d_1$  to reduce the DPPH, ABTS and FRAP radicals

Entry	$R_1$	Molecular weight	%DPPH	%ABTS	FRAP ( $\mu\text{g/ml}$ )
$4a_1$	$\text{CH}_3$	422.91	54.41	30.46	1.00
$4b_1$	$\text{C}_2\text{H}_5$	436.94	64.26	32.39	1.12
$4c_1$	$\text{C}_3\text{H}_7$	450.97	61.28	39.86	1.05
$4d_1$	$\text{C}_4\text{H}_9$	464.99	63.28	47.58	0.99
Carvedilol	–	406.74	14.81	26.05	0.91
Atenolol	–	266.34	14.92	39.47	0.91
Trolox	–	250.29	97.31	98.11	12.04
ascorbic acid	–	176.12	93.25	96.67	11.95

compounds under study might not be considered an explicit guarantee of their antioxidant potential. No significant relationships between elongation of the alkyl side chain and antioxidant activity have been revealed.

Potential therapeutical benefits based on current *in vitro* assays results should be the motivation for a closer study of other *in vitro* or *in vivo* systems and processes, in search of new cardiovascular therapeutics.

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**Conflicts of interest:** none.

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