

PŮVODNÍ PRÁCE

Synthesis, identification and physicochemical properties of novel potential ultrashort-acting beta-adrenergic blockers

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SUMMARY

Synthesis, identification and physicochemical properties of novel potential ultrashort-acting beta-adrenergic blockers

The paper deals with the synthesis, structure confirmation, and study of physicochemical properties of some aryloxyaminopropanol derivatives, compounds of 2-hydroxy-3-[4-(2-fluorophenyl)-piperazine-1-yl]-propyl-4-[(alkoxycarbonyl)amino]benzoates, their salts with hydrochloric acid, with one to four carbon atoms in the alkoxy group of the carbamoyl group. Their structure was confirmed by elemental analysis, IR, UV and mass spectra. The melting point, solubility, surface activity, dissociation constant, and some lipophilicity parameters, i.e. partition coefficient P , R_M values from reversed phase thin-layer chromatography, and retention factor k obtained from HPLC of the substances under study were determined.

Key words: ultrashort-acting beta-blockers – synthesis – physicochemical properties

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SÚHRN

Syntéza, identifikácia a fyzikálno-chemické vlastnosti nových potenciálnych ultrakrátkopôsobiacich beta-adrenergických blokátorov

Práca sa venuje syntéze, overeniu štruktúry a štúdiu fyzikálno-chemických vlastností derivátov aryloxyaminopropanolu, 2-hydroxy-3-[4-(2-fluórfenyl)-piperazín-1-yl]-propyl-4-[(alkoxykarbo-nyl)amino] benzoátov, ich solí s kyselinou chlorovodíkovou, s jedným až štyrmi atómami uhlíka v alkoxy skupine karbamoylovej skupiny. Štruktúra zlúčenín bola potvrdená elementárnou analýzou, IČ, UV a hmotnostnými spektrami. V rámci štúdia bola stanovená teplota topenia, rozpustnosť, povrchová aktivita, disociačná konštanta, ako aj parametre lipofility, t.j. rozdeľovací koeficient P , hodnoty R_M z reverznej chromatografie na tenkej vrstve a retenčný faktor k pomocou vysokoúčinnnej kvapalinovej chromatografie.

Kľúčové slová: ultrakrátkopôsobiace beta-blokátory – syntéza – fyzikálno-chemické vlastnosti

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Introduction

Antagonists of adrenergic β -receptors are widely used in hypertension, arrhythmias, hypertrophic cardiomyopathy, and ischemic heart disease. Adverse effects of classic beta-blockers include hypotension, bradycardia, heart failure, bronchospasm or peripheral vasoconstrictions, which can last up to several hours after intravenous application ¹⁾. Therefore, ultrashort-acting β -blockers have been developed for emergent intravenous treatment of atrial fibrillation or when control of heart rate is desirable ²⁾.

This paper focuses on the synthesis, identification, and determination of physicochemical properties of four new derivatives of β -adrenergic receptor-blocking phenoxyaminopropanols. The structure of the pattern molecule was modified by the ester group inserted between the aromatic and amine moieties of the connecting chain.

EXPERIMENTAL PART

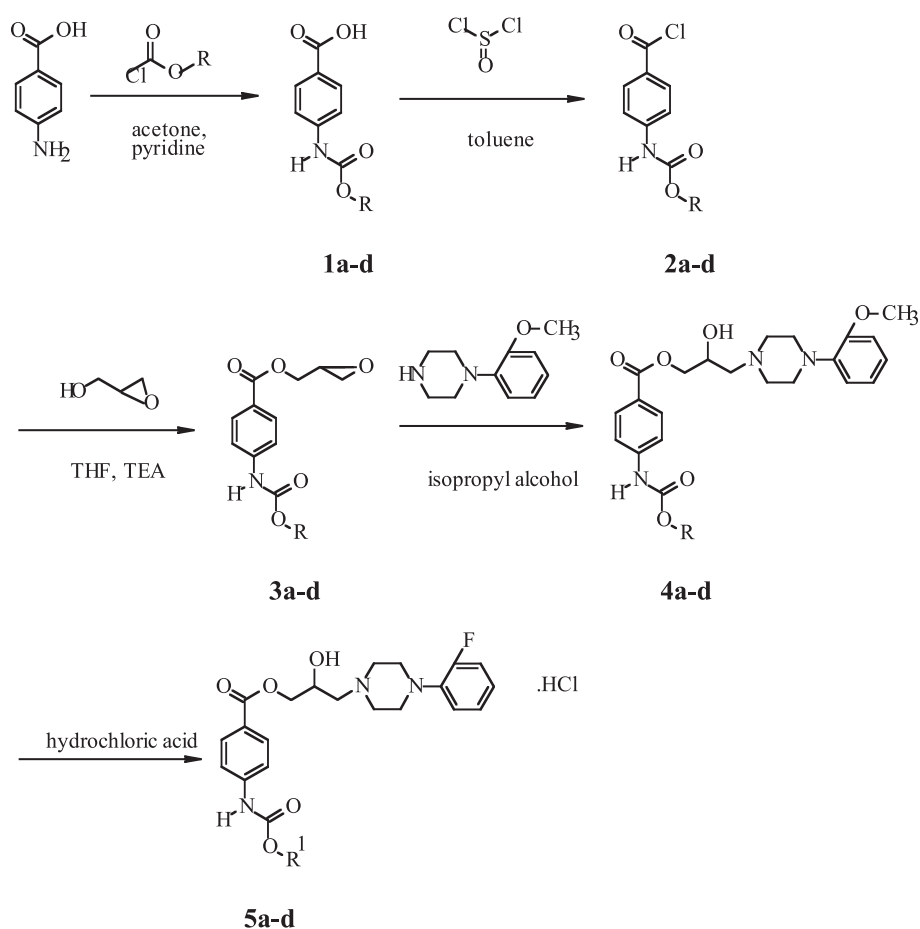
Synthesis

Synthesis of the novel derivatives of arylcarbonyloxyaminopropanols is shown in the Scheme and was carried out according to a general method first reported by Kam et al. ³⁾.

The corresponding 4-carbalkoxyaminobenzoic acids **1a-d** were prepared by the reaction of 4-aminobenzoic acid p.a. (Lachema, CR) with the appropriate alkylchloroformates. To a solution of p-aminobenzoic acid (0.1 mol.l⁻¹) and pyridine p.a. (LOBA Chemie, CR) (0.1 mol.l⁻¹) in acetone p.a. (Kulich, CR) (100 ml), alkylchloroformate (0.1 mol.l⁻¹) was added dropwise. The mixture was refluxed for 1.5 hour, evaporated to dryness and washed with water (500 ml). The crude acid was recrystallized from ethanol (96%, Lachema, CR).

To a suspension of 4-carbalkoxyaminobenzoic acid (0.1 mol.l⁻¹) in toluene p.a. (Lachema, CR) (120 ml), pure thionylchloride (Lachema, CR) (0.2 mol.l⁻¹) was added. The resulting mixture was refluxed for 2 hours and evaporated to dryness. The crude 4-carbalkoxyaminobenzoyl chloride **2a-d** recrystallized from toluene.

Glycidyl esters **3a-d** were prepared by the reaction of the acid chloride with glycidol. To a three-neck round-



Scheme 1. Synthetic route of the compounds

bottom flask, glycidol (96%, Aldrich, Germany) (0.25 mol.l⁻¹) and tetrahydrofurane *p.a.* (Lachema, CR) (50 ml) were added. The solution was cooled to -5 °C and triethylamine (> 99%, Merck-Schuchardt, Germany) (0.25 mol.l⁻¹) was added. To the resulting solution, a solution of **2a-d** (0.2 mol.l⁻¹) in tetrahydrofurane *p.a.* (Lachema, CR) (250 ml) was added dropwise at -5 to 5 °C. The mixture was stirred at room temperature for 3 hours, filtered and evaporated to dryness. The product, 2,3-epoxypropyl-4-carbalkoxyaminobenzoate – **3a-d**, was recrystallized from propan-2-ol *p.a.* (Kulich, CR).

In the last step of the synthesis, a mixture of **3a-d** (0.2 mol.l⁻¹) and 1-(2-fluorophenyl)piperazine (97%, Aldrich, Germany) (0.25 mol.l⁻¹) in isopropyl alcohol *p.a.* (Kulich, CR) (150 ml) was heated at 65 °C for 4 hours. The solvent was evaporated under reduced pressure and residuing oil was dissolved in diethylether *p.a.* (Kulich, CR). The solution of the base **4a-d** was converted to its chloride salt **5a-d** by addition of ethereal hydrochloric acid *p.a.* (35%, Central System, SR). The amine salt was collected by filtration and recrystallized from an appropriate solvent to give white crystals.

Chemical structure confirmation

Elemental analysis

Elemental analysis was made by an Elemental Analyzer EA 1108 (Carlo Erba Instruments).

Ultraviolet spectra

Electronic spectra of the prepared compounds were measured in the UV region by a spectrophotometer (8425 A Diode Array, Vectra 286/12), the solvent was methanol, concentrations 1.0 × 10⁻⁶ mol.l⁻¹ and quartz (1 cm).

Infrared spectra

IR spectra of the synthesized compounds were measured on an FTIR spectrophotometer Nicolet Model Impact 410 in the 400–4000 cm⁻¹ range.

Mass spectra

Mass spectra were recorded by a mass spectrometer Agilent 1100, LC/MSD VL Trap with a linear mechanic pump (KD Scientific, USA), the solvent was methanol, concentrations were approximately 1 mg.l⁻¹.

Thermic properties

Melting points were determined using a Kofler hotplate apparatus (HMK Franz Küstner, Germany).

Estimation of solubility and dissociation constants

Solubility

Solubility of prepared compounds was determined in distilled water, 96% ethanol (Lachema, CR), and in a chloroform medium *p.a.* (Lachema, CR) according to ⁴⁾.

Dissociation constant

The dissociation constant of prepared substances was estimated by alkalimetric titration with potentiometric

indication (pH-meter PO 811/1, Radelkis, Hungary) according to ⁵⁾.

Surface activity determination

The relative surface activity was determined by stalagmometric method according to ⁵⁾. Activity of the solution of the compounds (*c* = 1.0 × 10⁻³ mol.l⁻¹) was measured where the solvent was a mixture of H₂O-methanol *p.a.* (SPOLCHIM, SR) (ϕ_r = 3 : 2). The reference solution was the same as the solvent.

Determination of lipophilic properties

The apparent partition coefficient *P* of prepared compounds was measured by the shake-flask method in octan-1-ol (Merck, Germany)/phosphate buffer medium at pH = 7.0 according to ⁶⁾ using 0.5 ml of the organic solvent.

The chromatographic reversed-phase thin layer analysis was carried out according to ⁷⁾ on the silica gel Silufol® UV₂₅₄. *R_M* values were obtained in the mobile phase consisting of 1 mol.l⁻¹ HCl-acetone (ϕ_r = 4 : 2) after silica gel plates impregnation with 1% solution of silica oil in heptane (Lachema, CR). Chromatograms after developing and drying were detected by UV light at the wavelength 254 nm using a UV lamp TL-900 (Camag, Switzerland).

The chromatographic system for HPLC analysis for the retention factor *k* determination consisted of a pump Delta Chrom SDS (Watrex, Slovakia) with an injection valve and a UV detector Delta Chrom UVD 200 (Watrex, Slovakia). The analytical chromatography column was Sepharon SGX C₁₈ (254 × 4 mm, particle size 7 µm). The mobile phase was a mixture of 94% methanol in an amount of 500 ml and 3.4 g of natrium acetate, the pH value was adjusted with acetic acid to pH = 6.0. The flow rate of the mobile phase was 0.6 ml.min⁻¹, the injection volume was 1.10⁻² ml, the chromatograms were scanned at the wavelength 274 nm. The solution of NaNO₂ (*c* = 0.1 mol.l⁻¹) was used to determine the dead time *t₀*, and the solution of the analyzed compound in a methanol medium (*c* = 0.4 × 10⁻³ mol.l⁻¹) was used for determination of *t_R*.

RESULTS AND DISCUSSION

The paper focused on the synthesis, spectral description, and study of physicochemical properties of the new compounds.

The derivatives of 2-hydroxy-3-[4-(2-fluorophenyl)-piperazine-1-yl]-propyl-4-[(alkoxycarbonyl)amino]benzoates with one to four carbon atoms in the alkoxy group of the carbamoyl group have been synthesized by a five-step reaction according to a similar procedure, which was described in the literature ²⁾. The final compounds were isolated as the salts of hydrochloric acid (Scheme 1).

The methods of identification included elemental analysis, ultraviolet, infrared, and mass spectra. The

equation. The values of pK_a are shown in Table 4.

The relative surface activity values of the compounds increased with the alkoxy chain elongation. The values of relative surface tension were obtained as the average value from six measurements and are summarized in Table 4.

Lipophilicity is the key parameter used in quantitative structure-activity relationships (QSAR), quantitative structure-property relationships (QSPR), or quantitative structure-retention relationships (QSRR) studies, for modelling the biological and physicochemical properties of large classes of chemical compounds. In many cases, this molecular parameter strongly correlates with the biological activity of chemicals, as well as with other physicochemical properties¹⁰⁾.

The logarithm of partition coefficient between octan-1-ol and water ($\log P$) is the most widely used parameter in chemistry, pharmaceutical chemistry, and many other practical fields¹⁰⁾.

Partition coefficient was determined by the shake-flask method in octan-1-ol/phosphate medium. The $\log P$ values of the compounds increased with rising molecular weight of the substances. Degression was found in the substance with the working label 5d2. The values of $\log P$ are given in Table 4.

Logarithm of the retention factor k , like the lipophilicity parameter, was obtained by HPLC method. The results show that the $\log k$ values of

evaluated compounds increase with the increasing number of carbon atoms in the alkoxy chain of the carbamoyl group. The values of $\log k$ are summarized in Table 4.

The R_M values obtained from reversed-phase thin layer chromatography increased with the side alkyl chain elongation (Table 4).

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