

PŮVODNÍ PRÁCE

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

KEČKÉŠOVÁ, S., SEDLÁROVÁ, E., CSÖLLEI, J.¹, MOKRÝ, P.¹

Comenius University, Faculty of Pharmacy, Department of Pharmaceutical Chemistry, Bratislava

¹University of Veterinary and Sciences Pharmaceutical, Faculty of Pharmacy,

Institute of Chemical Drugs, Brno

Received 2 Juny / Accepted 26 Juny 2008

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

Čes. slov. Farm., 2008; 57, 160–164

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-[(alkoxykarbonyl)amino] benzoátov, ich soli 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

Čes. slov. Farm., 2008; 57, 160–164

Má

Address for correspondence:

PharmDr. Simona Kečkéšová

Comenius University, Faculty of Pharmacy, Department of Pharmaceutical Chemistry

Odbojárov 10, 832 32 Bratislava, SR

e-mail: skeckesova@yahoo.co.uk

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.

Synthesis

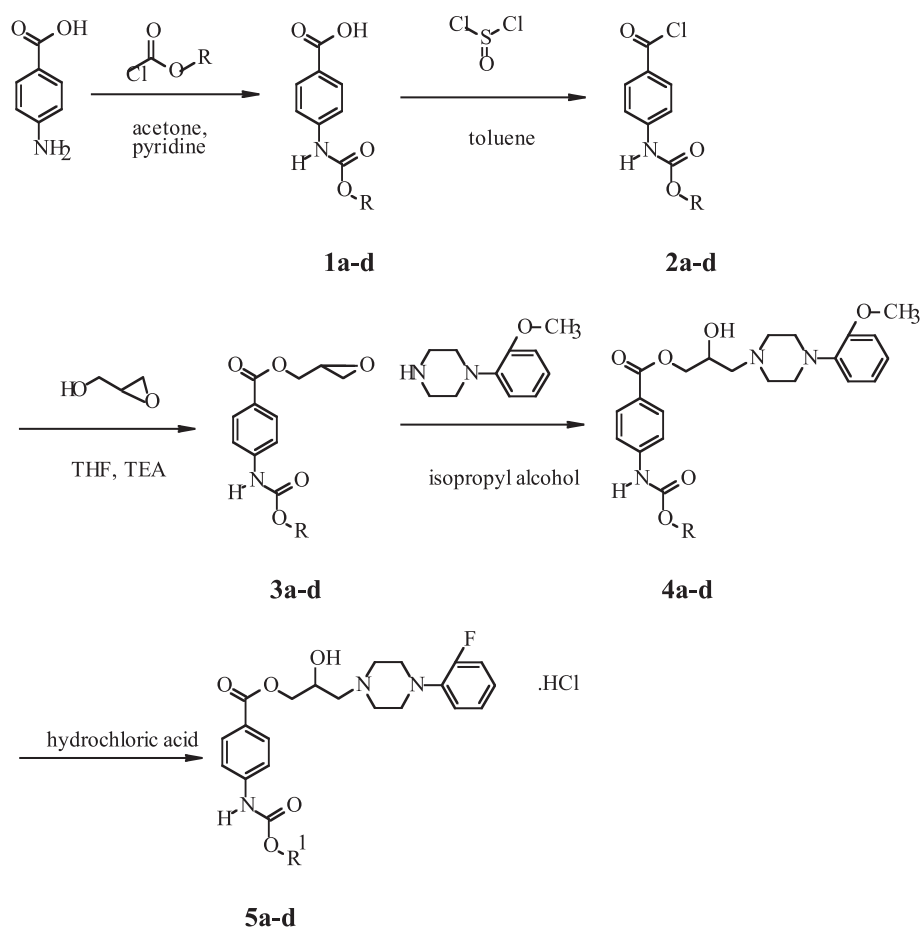
Synthesis of the novel derivatives of arylcarbonyloxyaminopropanols is shown in the Scheme and was carried out according to a general method first reported by Kam et al. ³⁾.

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-

EXPERIMENTAL PART



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 sodium 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*.

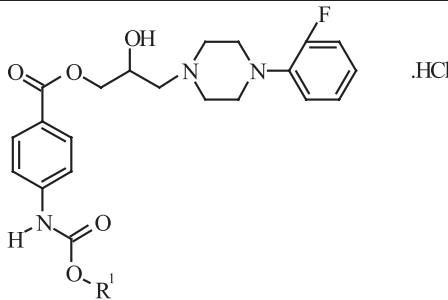
RESULTS AND DISCUSSION

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

Tab. 1. Chemical structure of studied compounds

								
Compound	R	Formula Mr	M.p. (°C)	Yield (%)	Solv. for Recrystall	% C	Analysed Found/Calc. % H	% N
5a	-CH ₃	C ₂₂ H ₂₇ O ₅ N ₃ ·FCl 467.92	188	73.1	ethanol	8.91 8.98	56.71 56.47	6.09 5.82
5b	-C ₂ H ₅	C ₂₃ H ₂₉ O ₅ N ₃ ·FCl 481.95	181–183	49.8	isopropyl alcohol	8.73 8.72	57.31 57.32	5.92 6.07
5c	-C ₃ H ₇	C ₂₄ H ₃₁ O ₅ N ₃ ·FCl 495.98	167	81.2	isopropyl alcohol	8.42 8.47	58.34 58.12	6.53 6.30
5d	-C ₄ H ₉	C ₂₅ H ₃₃ O ₅ N ₃ ·FCl 510.00	162–164	34.4	isopropyl alcohol	8.13 8.24	58.96 58.88	6.59 6.52

prepared compounds were solid, colourless and their melting points, yields, and formulas are presented in Table 1.

The structure of the substances under study was confirmed by elemental analysis, the values were in good agreement with the respective structures for carbon, hydrogen and nitrogen. The differences between the calculated and experimentally obtained data were in the range of ± 0.4 % and both data are given in Table 1.

The mass spectra of prepared compounds revealed the presence of $[M + H]^+$ ions in all cases, the value $m/z = 432.3$ for the compound 5a2, $m/z = 446.3$ for 5b2, $m/z = 460.3$ for 5c2 and $m/z = 474.3$ for 5d2.

The spectrum in the ultraviolet region showed two absorption maxima at the wavelength $\gamma = 202$ –203 nm and $\gamma = 270$ –271 nm. The values of $\log \epsilon$ of the analyzed substances are shown in Table 2.

The results of the spectral parameters of the IR region are summarized in Table 3.

The solubility of liquids and solids in water is no doubt one of the most important molecular properties that affect their biological activity. Solubility of solid

Tab. 2. UV spectra of prepared compounds

Compound	λ_{\max} (nm)	$\log \epsilon$ (m ² .mol ⁻¹)	λ_{\max} (nm)	$\log \epsilon$ (m ² .mol ⁻¹)
5a	203	4.92	270	4.79
5b	203	4.91	270	4.79
5c	202	4.89	271	4.79
5d	202	4.98	271	4.85

compounds depends on the free energy changes involved in changing the solid state to the liquid state, in addition to the interactions in the liquid phase ⁸⁾.

Despite the fact that the compounds were salts with hydrochloric acid, their solubility was practically indissoluble in water, very limited in ethanol and practically indissoluble in chloroform.

The dissociation constant presents the rate of the dissociated and undissociated forms of the drug ⁹⁾. The value of dissociation constant was obtained as the average value from three measurements and was determined according to the Henderson-Hasselbalch

Tab. 3. Parameters of IR spectra (cm⁻¹) of prepared compounds

Compound	ν N-H	ν_{ar} C-H	ν_{al} C-H	ν_{es} C=O	ν_{carb} C=O	ν_{es} C-O-C	ν_{carb} C-O-C
5a2	3239	3040	2947	1727	1704	1222	1065
5b2	3318	3074	2970	1710	1687	1231	1056
5c2	3326	3047	2972	1722	1689	1219	1058
5d2	3311	3048	2962	1723	1689	1219	1062

Tab. 4. Physicochemical parameters of substances under study

Compound	pKa	σ_{rel} (N.m ⁻¹)	$\log P$	$\log k$	Rf	RM
5a	7.48	1.257	4.23	- 0.35	0.545	- 0.078
5b	7.64	1.269	4.55	- 0.31	0.456	0.077
5c	7.35	1.368	4.88	- 0.26	0.361	0.248
5d	7.27	1.444	4.07	- 0.20	0.274	0.423

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 properties ¹⁰.

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 fields ¹⁰.

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).

This study was supported by the Comenius University grant No. UK/2/2008.

REFERENCES

1. **Bartošová, L. et al.:** *Pharmazie*, 2003; 58, 841-842.
2. **Mokřý, P. et al.:** *Pharmazie*, 2003; 58, 18-21.
3. **Kam, S.T. et al.:** *J. Med. Chem.*, 1984; 27, 1007-1016.
4. *Pharmacopea Slovaca PhS 1*, 1st Edition (in Slovak). Bratislava, Herba 1999, p. 22.
5. **Malík, I. et al.:** *Čes. Slov. Farm.*, 2005; 54, 235-239.
6. **Sedlářová, E. et al.:** *Farm. Obzor*, 2007; 76, 86-89.
7. **Malík, I. et al.:** *Farm. Obzor*, 2005; 74, 211-215.
8. **Malík, I. et al.:** *Chem. Pap.*, 2006; 60, 42-47.
9. **Sedlářová, E. et al.:** *Čes. a Slov. Farm.*, 2000; 49, 306-312.
10. **Sárbu, C. et al.:** *Talanta*, 2008; 75, 651-657.

Abstrakta z akcí ČFS v časopisu Česká a slovenská farmacie

Redakce časopisu Česká a slovenská farmacie nabízí možnost zveřejňovat limitované množství abstrakt z odborných akcí pořádaných Českou farmaceutickou společností, například sympozií, seminářů, pracovních dnů apod.

Jednotlivá abstrakta (písmo Courier New, velikost 12, řádkování 2), by neměla přesáhnout 1 rukopisnou stranu formátu A4.

Počet abstrakt předem dohodnou předsedové příslušných sekcí, které akci pořádají, případně osoby zodpovědné za akci s redakcí časopisu, která poskytne i bližší informace.

Lze zveřejnit rovněž na internetových stránkách ČFS (www.cfs-cls.cz)

Kontakt:

doc. RNDr. Pavel Komárek, PhD., vedoucí redaktor, Katedra farmaceutické technologie a kontroly léčiv IPVZ
100 05 Praha 10, Ruská 85, e-mail: komarek@ipvz.cz, tel.: 271 019 278