

PŮVODNÍ PRÁCE

Antimycobacterial activity of novel derivatives of arylcarbonyloxyaminopropanols

KEČKÉŠOVÁ S.¹, SEDLÁROVÁ E.¹, ČIŽMÁRIK J.¹, GARAJ V.¹, CSÖLLEI J.², MOKRÝ P.², ANDRIAMAINY F.¹, MALÍK I.¹, KAUSTOVÁ J.³

¹Comenius University, Faculty of Pharmacy, Department of Pharmaceutical Chemistry, Bratislava

²University of Veterinary and Pharmaceutical Sciences, Faculty of Pharmacy, Institute of Chemical Drugs, Brno

³Regional Institute of Public Health, Ostrava

Received 5 October 2009 / Accepted 4 November 2009

SUMMARY

Antimycobacterial activity of novel derivatives of arylcarbonyloxyaminopropanols

The study examined the antimicrobial activity of 2-hydroxy-3-[4-(2-fluoro, 4-fluoro, and 2-methoxyphenyl)-piperazin-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 studied and described. The series of these substances were evaluated *in vitro* against *Mycobacterium tuberculosis*, *Mycobacterium avium*, *Mycobacterium kansasii* and against clinically isolated strains of *Mycobacterium kansasii* 6 509/96. The correlation of this activity with lipophilicity (log *k'*) was used to describe the structure-antimycobacterial activity relationships (QSARs). The tests have shown that practically all the substances under study possess antituberculous activity. *In vitro* antimycobacterial activity increased with increasing lipophilicity. According to the obtained results, the substances can be considered as potential antituberculous.

Key words: antimycobacterial activity – antituberculous activity – potential antituberculous – arylcarbonyloxyaminopropanols – lipophilicity – phenylcarbamates

Čes. a slov. Farm., 2009; 58, 203–207

SÚHRN

Antimykobakteriálna aktivita nových derivátov arylkarbonyloxyaminopropanolov

V práci je opísaná antimikrobiálna aktivita 2-hydroxy-3-[4-(2-fluór, 4-fluór, a 2-metoxifyfenyl)-piperazín-1-yl]-propyl-4-[(alkoxykarbonyl)amino] benzoátov, ich soli s kyselinou chlorovodíkovou, s jedným až štyrmi atómami uhlíka v alkoxy skupine karbamoylovej skupiny. Série týchto látok boli hodnotené *in vitro* vzhľadom na *Mycobacterium tuberculosis*, *Mycobacterium avium*, *Mycobacterium kansasii* a klinicky izolovaný kmeň *Mycobacterium kansasii* 6 509/96. Korelácia aktivity s lipofilitou (log *k'*) bola použitá pre popis vzťahov štruktúra-antimykobakteriálna aktivita (QSAR). Testy ukázali, že prakticky všetky študované látky majú antituberkulotickú aktivitu. *In vitro* sa antimikrobiálna aktivita zvyšovala so zvyšujúcou sa lipofilitou. Na základe dosiahnutých výsledkov možno študované látky považovať za potenciálne antituberkulotiká.

Kľúčové slová: antimykobakteriálna aktivita – antituberkulotická aktivita – potenciálne antituberkulotiká – arylkarbonyloxyaminopropanoly – lipofilita – fenyلكarbamáty

Čes. a slov. Farm., 2009; 58, 203–207

Má

Address for correspondence:

PharmDr. Simona Kečkéšová, PhD.
Comenius University, Faculty of Pharmacy, Department of Pharmaceutical Chemistry
Odbojárov 10, 832 32 Bratislava, SR
e-mail: skeckesova@yahoo.co.uk

Introduction

In consideration of minutely increasing counts of the multiresistant strains of *Mycobacterium tuberculosis*, the design, preparation and study of new potential antituberculars with structures different from those used in clinical praxis nowadays, belongs to the topical tasks of the drug research and development.

The task is really topical because there is also another source of the threat for European inhabitants – the occasionally pathogenic mycobacterial strains, such as *Mycobacterium avium*, *Mycobacterium kansasii* and *Mycobacterium intercellulare*¹⁾.

The paper¹⁾ described the antimycobacterial activity of 2-(4-alkylpiperazin-yl)-ethyl esters of 2-heptyloxyphenylcarbamic acid. The series of these substances was evaluated *in vitro* against *Mycobacterium tuberculosis*, *Mycobacterium kansasii*, *Mycobacterium avium* and against clinically isolated strains of *Mycobacterium kansasii* 6 509/96. According to the obtained results, all the studied substances possessed antitubercular activity. The substances under study have in their chemical structure a fragment of substituted ester of phenylcarbamic acid and 4-substituted

piperazine. These two groups are present in the studied structures and participate in the formation of some common metabolites.

The aim of this paper was to verify whether the studied structures exerted an antimycobacterial effect against hereinbefore mentioned strains.

EXPERIMENTAL PART

Synthesis

All of the compounds under study were prepared according to²⁾ and published in the papers^{3, 4)}. The structure of the substances studied is given in Table 1.

Microbiology

For the evaluation of the antimycobacterial activity *in vitro*, the following strains were used: *Mycobacterium tuberculosis* CNCTC My 331/88, *Mycobacterium avium* CNCTC My 330/88,

Table 1. Chemical structure of the compounds under study

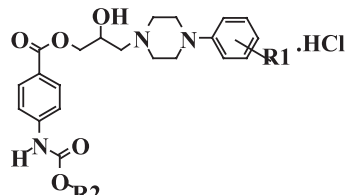
								
Compd.	R1	R2	Compd.	R1	R2	Compd.	R1	R2
5a1	4-F	-CH ₃	5a2	2-F	-CH ₃	5a3	2-OCH ₃	-CH ₃
5b1	4-F	-C ₂ H ₅	5b2	2-F	-C ₂ H ₅	5b3	2-OCH ₃	-C ₂ H ₅
5c1	4-F	-C ₃ H ₇	5c2	2-F	-C ₃ H ₇	5c3	2-OCH ₃	-C ₃ H ₇
5d1	4-F	-C ₄ H ₉	5d2	2-F	-C ₄ H ₉	5d3	2-OCH ₃	-C ₄ H ₉

Table 2. The minimum inhibitory concentration (MIC) and the logarithm of the retention factor *k* (log *k*) of the substances under study

Compd.	log <i>k</i>	MIC (μmol/l) Time of inhibition 14 d/21 d			
		<i>M. tuberculosis</i> My 331/88	<i>M. avium</i> My 330/88	<i>M. kansasii</i> My 235/80	<i>M. kansasii</i> My 6 509/96
5a1	-0.42	62.5/125	125/250	250/250	125/250
5b1	-0.38	62.5/125	62.5/250	125/250	125/250
5c1	-0.32	32/62.5	125/250	125/125	62.5/125
5d1	-0.27	32/32	32/62.5	62.5/62.5	32/62.5
5a2	-0.35	125/250	250/500	250/500	62.5/125
5b2	-0.31	125/250	125/250	125/125	125/125
5c2	-0.26	32/62.5	62.5/125	125/125	62.5/125
5d2	-0.20	32/32	62.5/125	62.5/125	32/62.5
5a3	-0.36	125/250	250/250	500/>500	125/250
5b3	-0.32	32/62.5	250/250	500/>500	62.5/125
5c3	-0.27	32/62.5	125/250	125/125	32/62.5
5d3	-0.22	16/32	125/125	62.5/125	32/32
Ison.	–	0.5/0.5	>250/>250	>250/>250	2/4

Mycobacterium kansasii CNCTC My 235/80 obtained from the Czech National Collection of Type Cultures (CNCTC), National Institute of Public Health, Prague, and the clinically isolated strain of *Mycobacterium kansasii* 6 509/96. Antimycobacterial activity was determined in Šula semisynthetic medium (SEVAC, Prague). The substances were added to the medium in a Me₂SO solution. The final concentrations of the compounds were 1–500 μmol/l. The minimum inhibitory concentration (MIC, i.e. the lowest concentration of a substance at which the inhibition of growth occurred)⁵⁾ was determined after incubation at 37 °C for 14 and 21 days. The results are summarized in Table 2. The antituberculous activity of isoniazide was used as the reference standard.

Determination of lipophilic properties

Since many years, lipophilicity is recognized as a meaningful parameter in many structure-activity and

structure-ADME relationships. It is also the single most informative and successful physicochemical property in medicinal chemistry⁶⁾.

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 a 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 of 274 nm. The solution of NaNO₂ (*c* = 0.1 mol.l⁻¹) was used to determine the dead time *t*₀, and the solution of the analyzed compound in methanol medium (*c* = 0.4 · 10⁻³ mol.l⁻¹) was used for the determination of *t*_R.

Table 3. Linear regressions (5a1–5d1 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	-2.329	0.800	0.843	0.167	4
	21 d	3.903	2.724	0.569	0.342	4
<i>avium</i> My 330/88	14 d	-2.752	0.912	0.413	0.283	4
	21 d	-3.557	1.004	0.613	0.300	4
<i>kansasii</i> My 235/80	14 d	-3.442	0.894	0.861	0.245	4
	21 d	-4.188	0.709	0.927	0.287	4
<i>kansasii</i> My 6509/96	14 d	-4.128	0.433	0.928	0.283	4
	21 d	-4.188	0.709	0.926	0.287	4

Table 4. Parabolic regressions (5a1–5d1 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	c	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	4.482	0.762	1.354	0.849	0.167	4
	21 d	-40.999	-24.370	2.016	0.676	0.342	4
<i>avium</i> My 330/88	14 d	-33.967	-26.174	-3.016	0.519	0.283	4
	21 d	-62.442	-46.615	-6.215	0.934	0.300	4
<i>kansasii</i> My 235/80	14 d	0.850	-2.856	0.993	0.861	0.245	4
	21 d	-26.584	-22.519	-2.364	0.989	0.287	4
<i>kansasii</i> My 6509/96	14 d	-25.544	-21.742	-2.519	0.989	0.283	4
	21 d	-26.584	-22.519	-2.364	0.989	0.287	4

Table 5. Linear regressions (5a2–5d2 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	-4.683	0.484	0.794	0.341	4
	21 d	-6.603	0.169	0.920	0.446	4
<i>avium</i> My 330/88	14 d	-4.048	0.882	0.834	0.287	4
	21 d	-4.048	1.182	0.834	0.287	4
<i>kansasii</i> My 235/80	14 d	-3.572	1.090	0.893	0.245	4
	21 d	-3.334	1.307	0.519	0.300	4
<i>kansasii</i> My 6509/96	14 d	-2.556	1.077	0.473	0.241	4
	21 d	-1.905	1.482	0.677	0.150	4

Table 6. Parabolic regressions (5a2–5d2 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	c	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	16.652	4.445	1.682	0.813	0.341	4
	21 d	-10.302	-12.249	-0.573	0.925	0.446	4
<i>avium</i> My 330/88	14 d	40.146	17.956	3.769	0.994	0.287	4
	21 d	40.146	17.956	4.069	0.994	0.287	4
<i>kansasii</i> My 235/80	14 d	3.504	-1.651	1.342	0.895	0.245	4
	21 d	63.358	31.393	5.863	0.882	0.300	4
<i>kansasii</i> My 6509/96	14 d	-50.448	-30.206	-2.551	0.831	0.241	4
	21 d	-28.175	-17.347	-0.545	0.965	0.150	4

Table 7. Linear regressions (5a3–5d3 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	-5.559	-0.054	0.821	0.373	4
	21 d	-5.556	0.243	0.814	0.374	4
<i>avium</i> My 330/88	14 d	-2.574	1.488	0.815	0.173	4
	21 d	-1.964	1.741	0.633	0.150	4
<i>kansasii</i> My 235/80	14 d	-7.111	0.235	0.922	0.450	4
	21 d	-5.147	0.885	0.815	0.347	4
<i>kansasii</i> My 6509/96	14 d	-4.316	0.458	0.865	0.282	4
	21 d	-6.300	0.099	0.997	0.384	4

Table 8. Parabolic regressions (5a3–5d3 derivatives)

<i>Mycobacterium</i>	Time of inhibition	a	b	c	r ²	s	n
<i>tuberculosis</i> My 331/88	14 d	36.572	15.623	2.912	0.879	0.373	4
	21 d	38.907	16.980	3.398	0.879	0.374	4
<i>avium</i> My 330/88	14 d	5.364	0.534	1.923	0.821	0.173	4
	21 d	-34.120	-21.726	-1.027	0.947	0.150	4
<i>kansasii</i> My 235/80	14 d	-23.390	-20.660	-1.662	0.938	0.450	4
	21 d	10.728	1.067	1.755	0.821	0.347	4
<i>kansasii</i> My 6509/96	14 d	41.130	19.507	3.793	0.994	0.282	4
	21 d	8.327	-1.478	0.775	0.999	0.384	4

The quantitative structure-activity relationships (QSARs)

The quantitative structure-activity relationships were determined according to two computing models of linear regression ($\log \text{MIC} = a \log k' + b$) and parabolic regression ($\log \text{MIC} = a (\log k')^2 + b (\log k') + c$). Both types of the calculation were carried out using the Microsoft Excel program. The results are summarized in Tables 3–8.

RESULTS AND DISCUSSION

The paper focused on the antimicrobial activity of the new compounds.

The logarithm of the retention factor k , like the lipophilicity parameter, was obtained by the HPLC method. The $\log k$ values of the substances under evaluation increased with the increasing number of carbon atoms in the alkoxy chain of the carbamoyl group. The results of the analysis confirm that the activity of the compounds increases with increasing lipophilicity.

The conclusions indicate that practically all of the derivatives show expected antimycobacterial activity in consideration of all of the strains studied, and some of them are more efficient than the standard isoniazide.

This study was supported by the VEGA grant No. 1/4291/07.

REFERENCES

1. Čižmárik, J., Waisser, K., Doležal, R., Kaustová, J.: Farm. Obzor, 2007; 76, 144–146.
2. Mokřý, P., Zemanová, M., Csöllei, J., Račanská, E., Tumová, I.: Pharmazie, 2003; 58, 18–21.
3. Kečkéšová, S., Sedlářová, E., Csöllei, J., Mokřý, P.: Čes. a Slov. Farm., 2008; 57, 160–164.
4. Kečkéšová, S., Sedlářová, E., Csöllei, J., Mokřý, P., Vančo, J., Vanko, M.: Acta Facult. Pharm. Univ. Com., 2008; 55, 122–128.
5. Waisser, K., Dražková, K., Čižmárik, J., Kaustová, J.: Folia Microbiol., 2003; 48, 45–50.
6. Malík, I., Sedlářová, E., Csöllei, J., Andriamainty, F., Čižmárik, J., Kečkéšová, S.: Acta Facult. Pharm. Univ. Com., 2007; 54, 136–145.

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 symposií, 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