Oxadiazines are interesting and promising heterocyclic compounds. A diversity of biological effects is associated with oxadiazines bearing heteroatoms at 1,2,4 or 1,3,4 positions. 6H-1,2,4-oxadiazine-3,5-(2H,4H)-dione, the 6-oxa analogue of uracil, has been shown to significantly inhibit growth in several bacterial strains while not being highly inhibitory to mammalian cells (1). 1,3,4-Oxadiazine derivatives exhibit cardiovascular, antibacterial, plant growth regulating, miticidal and nematocidal, acricidal, insecticidal and anticonvulsive activities (2, 3). In addition, oxadiazines are useful intermediates in the synthesis of tenidap prodrugs or β-lactam antibiotics, in particular in the synthesis of carbapenems and penems (4, 5).

Novel 1,2,5-oxadiazine derivatives – Synthesis and in vitro biological studies

A new synthetic approach to the 1,2,5-oxadiazine ring system is described. 2-Substituted or 2,4-disubstituted 2H-1,2,5-oxadiazine-3,6(4H,5H)-dione derivatives were prepared by cyclisation of hydroxamic acids derived from N-(1-benzotriazolylcarbonyl)-amino acids. The structures of the synthesised compounds were fully characterised by IR, 1H and 13C NMR spectroscopy and elemental analysis. The aim of this study was to evaluate biological activity of the newly synthesised oxadiazine derivatives. Cytotoxic and cyostatic activities were tested on two cell lines (HeLa and GMK) and evaluated by MTT-test. Two human DNA viruses (adenovirus and herpesvirus) and two human RNA viruses (coxsackievirus and echovirus) were used in the antiviral test. Selected biological studies indicated that 2-phenyl-2H-1,2,5-oxadiazine-3,6(4H,5H)-dione and 4-benzyl-2-phenyl-2H-1,2,5-oxadiazine-3,6(4H,5H)-dione statistically significantly inhibited cell growth. A minor antiviral effect was observed upon adenovirus, herpesvirus and enteroviruses.

Keywords: 1,2,5-oxadiazine derivatives, synthesis, cytotoxic effect, cyostatic effect, antiviral activity, cytopathic effect

Oxadiazines are interesting and promising heterocyclic compounds. A diversity of biological effects is associated with oxadiazines bearing heteroatoms at 1,2,4 or 1,3,4 positions. 6H-1,2,4-oxadiazine-3,5-(2H,4H)-dione, the 6-oxa analogue of uracil, has been shown to significantly inhibit growth in several bacterial strains while not being highly inhibitory to mammalian cells (1). 1,3,4-Oxadiazine derivatives exhibit cardiovascular, antibacterial, plant growth regulating, miticidal and nematocidal, acricidal, insecticidal and anticonvulsive activities (2, 3). In addition, oxadiazines are useful intermediates in the synthesis of tenidap prodrugs or β-lactam antibiotics, in particular in the synthesis of carbapenems and penems (4, 5).
On the other hand, 1,2,5-oxadiazines are not very common heterocyclic systems. A comprehensive review of their synthesis and reactivity has been reported by Smalley (6). The promising therapeutic potential of this class of compounds prompted us to synthesise and biologically screen several 1,2,5-oxadiazine derivatives. Our synthetic approach involves hydroxamic acids as the starting compounds, which themselves have been of multifold biological and pharmacological interest (7–9).

Herein we present a new synthetic approach to 1,2,5-oxadiazines derivatives and their biological activity on mammalian cells. Furthermore, their antiviral activity against adenoviruses, herpes viruses and enteroviruses was investigated.

EXPERIMENTAL

Materials and analytical methods

Melting points were determined on a Boëtius Microheating Stage (Franz Küstner Nachf. KG, Germany) and were uncorrected. IR spectra were recorded on a FTIR Perkin Elmer Paragon 500 spectrometer (Perkin Elmer, UK). $^1$H and $^{13}$C NMR spectra were recorded on a Varian Gemini 300 spectrometer (Varian, USA), operating at 300 and 75.5 MHz for the $^1$H and $^{13}$C nuclei, respectively. Samples were measured in DMSO-d$_6$ solutions at 20 °C in 5 mm NMR tubes. Chemical shifts (δ) in ppm were referred to TMS. Coupling constants (J) in Hz, were observed through three bonds.

For TLC, silica gel plates Kieselgel 60 F$_{254}$ (Merck, Germany) and the following solvent mixtures were used: cyclohexane/ethyl acetate/methanol (3:1:0.3), cyclohexane/ethylacetate (1:1), chloroform/methanol (9:1). Spots were visualised by short-wave UV light and iodine vapour. Column chromatography was performed on silica gel (Kemika, Croatia), 0.063–0.200 mm, with dichloromethane/methanol (9:1) or cyclohexane/ethyl acetate (1:1) as eluent.

Amino acids were purchased from Kemika (Croatia). N-Phenylhydroxylamine was prepared by reduction of nitrosobenzene with L-ascorbate (11). All solvents were of analytical grade purity and were dried prior to use.

Syntheses

$\text{N-}(1\text{-benzotriazolylcarbonyl})\text{-amino acids (N-Btc-amino acids, 1a-c).}$ – N-Btc-glycine (1a), N-Btc-L-alanine (1b) and N-Btc-L-phenylalanine (1c) were synthesised following the procedure published previously (12).

$\text{N-}(1\text{-benzotriazolylcarbonyl})\text{-amino acid chlorides (2a-c).}$ – Chlorides 2 were prepared from N-Btc-amino acids 1a-c and thionyl chloride (13).

$\text{Hydroxamic acids 3a-c.}$ – To a suspension (2a) or solution (2b,c) of 3–5 mmol chloride 2 in 55–80 mL cold toluene (0 °C), a solution of 3–5 mmol N-phenylhydroxylamine (PHA) in 45–75 mL toluene and a solution of 5.5 mmol N-methylmorpholine (MM) in 30 mL toluene were added simultaneously and dropwise. The reaction mixture was stirred at room temperature for 3–24 h.

Product 3a partially precipitated from the reaction mixture, together with MM hydrochloride. The precipitate was filtered off and dissolved in a water/ethyl acetate mix-
ture. The organic layer was extracted with water, twice with cold 1% HCl solution and twice with water, then dried over anhydrous sodium sulphate, filtered and evaporated to give 0.525 g (35%) of 3a. An additional amount (0.315 g, 21%) of 3a was isolated from toluene mother liquor by an analogous extraction procedure.

Products 3b and 3c were isolated as follows. The reaction mixture was extracted with water, three times with 1% HCl solution and two times with water. The organic layer was dried over anhydrous sodium sulphate, filtered and evaporated under reduced pressure. The analytically pure 3b was obtained by purification on silica gel column (eluent: dichloromethane/methanol 9:1) and additional titration with ethyl acetate and cyclohexane. Pure product 3c was obtained by recrystallisation with ether/cyclohexane mixture.

1,2,5-Oxadiazine-3,6-dione derivatives (4a,c). – To a solution of 1.6 mmol of hydroxamic acid 3a or 3c in 130 mL acetone, 5 mL of 10% solution of sodium carbonate was added. The reaction mixture was stirred at room temperature for 2.5 h. Precipitated inorganic salt was filtered off and the mother liquor was evaporated. The residue was dissolved in dichloromethane and extracted twice with cold 1% HCl solution and twice with water. The organic layer was dried over anhydrous sodium sulphate, filtered and evaporated under reduced pressure. Products 4a and 4c were purified by column chromatography (eluent: cyclohexane/ethyl acetate 1:1).

Cell lines

Green monkey kidney cells (GMK; ATCC CCL-26, USA) and human cervical carcinoma cells (HeLa; ATCC CCL-2, USA) were used. Monolayer cell cultures were grown in a modified eagle medium (DMEM; Dulbecco minimum essential medium, Difco, USA) supplemented with 10% inactivated foetal calf serum (FS; Gibco BRL, UK), 2 mmol L⁻¹ L-glutamine, 10 mmol L⁻¹ Hepes (pH 7.4) and antibiotics (100 U mL⁻¹ penicillin, 100 mg mL⁻¹ streptomycin), and were incubated at 37 °C and 5% of CO₂.

MTT-test

MTT-test was used to measure the mitochondrial activity in cells (14). When added, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-2H-tetrazolium bromide (MTT) formed formazane crystals in live cells and cells in an early apoptosis cycle (programmed cell death). Formazane crystals dissolved when dimethyl sulphoxide (DMSO) was added. This enabled the spectrophotometric reading at 570 nm. According to the MTT-test results, it was possible to determine if a tested compound acted as a cytotoxic agent, cytostatic agent or if it stimulated cell growth. The tested compound had a cytotoxic effect on cells (CT) if spectrophotometric values after 72 h of incubation were below the initial control values measured after 24 h of incubation. Values at 72 h of incubation below the control values measured after 72 h of incubation indicated the inhibitory effect (IC) of cell growth, i.e., cytostatic effect, whereas values at 72 h of incubation higher than control values measured after 72 h of incubation meant that the tested compound stimulated cell growth (proliferative effect).

Acyclovir (Ac) was used as a positive control compound at the same concentrations as 4a and 4c. Each compound was dissolved in DMSO to produce 0.1 mol L⁻¹ solution
and further diluted in DMEM to concentrations of $10^{-4}$ mol L$^{-1}$ ($c_1$), $10^{-5}$ mol L$^{-1}$ ($c_2$) and $10^{-6}$ mol L$^{-1}$ ($c_3$). The final concentration of DMSO in different dilutions of compounds was less than 0.1% and at that concentrations it did not influence the cell growth.

**Antiviral assay**

Five different viral titres (V-1 to V-5, relative titres) obtained by serial dilution (1:2 for adenovirus and herpesvirus, and 1:10 for enteroviruses) were used to test the inhibition of the cytopathic effect (CPE) (15). HeLa and GMK cells were seeded at $10^5$ cells per mL on 24-well microtitre plates (Becton Dickinson, USA). Viral infection was performed on one-day-old confluent cell monolayers with addition of the tested compounds at different concentrations. The CPE was followed by an optical microscope 24 h after infection for enteroviruses (coxsackievirus and echovirus) and 48 h for adenovirus and herpesvirus. The CPE of enteroviruses was observed on GMK cells, while CPE of adenovirus and herpesvirus was observed on HeLa cells (10).

**Statistics**

The results are shown as the percentage of CPE inhibition compared to CPE without compounds on each plate. The results were statistically analysed on a personal computer by ANOVA test and shown graphically. The criteria for statistical significance was $p < 0.05$. The concentration values that inhibit 50% of cell growth ($IC_{50}$) were calculated using the linear regression model.

**RESULTS AND DISCUSSION**

**Syntheses**

The starting compounds for the synthesis of hydroxamic acids 3 were N-Btc-amino acids (1a-c). They were prepared from the appropriate amino acids and 1-benzotriazole carboxylic acid chloride and converted into the corresponding chlorides 2 by means of
Table I. Reaction conditions and analytical data of hydroxamic acids 3a-c and 1,2,5-oxadiazine-3,6-dione derivatives 4a,c

<table>
<thead>
<tr>
<th>Compound No.</th>
<th>R</th>
<th>Solvent</th>
<th>Time (h)</th>
<th>Yield (%)</th>
<th>M.p. (°C)</th>
<th>Molecular formula</th>
<th>CHN analysis (%)</th>
<th>IR (KBr) ( \nu_{\max} ) (cm(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>3a</td>
<td>H</td>
<td>Toluene</td>
<td>24(^a)</td>
<td>56</td>
<td>147–149</td>
<td>( \text{C}<em>{15}\text{H}</em>{13}\text{N}<em>{5}\text{O}</em>{3} ) (311.29)</td>
<td>57.88 4.21 22.50</td>
<td>3342, 3186, 2876, 1731, 1687, 1659, 1594, 1530, 1496, 1447, 1396, 1301, 1286, 1236, 1090, 1004, 753, 691, 609</td>
</tr>
<tr>
<td>3b</td>
<td>CH(_3)</td>
<td>Toluene</td>
<td>3(^a)</td>
<td>15</td>
<td>132–135</td>
<td>( \text{C}<em>{16}\text{H}</em>{15}\text{N}<em>{5}\text{O}</em>{3} ) (325.32)</td>
<td>59.07 4.65 21.53</td>
<td>3252, 2930, 1734, 1712, 1675, 1593, 1529, 1450, 1380, 1285, 1235, 1136, 1072, 1034, 936, 840, 755, 695</td>
</tr>
<tr>
<td>3c</td>
<td>CH(_2)C(_6)H(_5)</td>
<td>Toluene</td>
<td>22(^a)</td>
<td>31</td>
<td>130–133</td>
<td>( \text{C}<em>{22}\text{H}</em>{19}\text{N}<em>{5}\text{O}</em>{3} ) (401.42)</td>
<td>65.83 4.77 17.45</td>
<td>3241, 3030, 2925, 1740, 1658, 1594, 1522, 1496, 1448, 1388, 1287, 1233, 1087, 1043, 928, 834, 750, 696, 615, 573</td>
</tr>
<tr>
<td>4a</td>
<td>H</td>
<td>Acetone</td>
<td>2.5(^b)</td>
<td>13</td>
<td>115–117</td>
<td>( \text{C}<em>{6}\text{H}</em>{12}\text{N}<em>{2}\text{O}</em>{3} ) (192.17)</td>
<td>56.25 4.19 14.58</td>
<td>3280, 3163, 1747, 1692, 1590, 1495, 1445, 1407, 1365, 1137, 1104, 1006, 763, 691, 539</td>
</tr>
<tr>
<td>4c</td>
<td>CH(_2)C(_6)H(_5)</td>
<td>Acetone</td>
<td>2.5(^b)</td>
<td>28</td>
<td>96–98</td>
<td>( \text{C}<em>{16}\text{H}</em>{14}\text{N}<em>{2}\text{O}</em>{3} ) (282.29)</td>
<td>68.08 5.00 9.92</td>
<td>3265, 3169, 1772, 1688, 1594, 1490, 1365, 1130, 988, 729, 692</td>
</tr>
</tbody>
</table>

\(^a\) 1 h at 0 °C, the remaining period at room temperature.

\(^b\) Room temperature.
thionyl chloride (12, 13). In the next reaction step, acyl chlorides reacted with N-phenylhydroxylamine and gave the following hydroxamic acids: N'-hydroxy-N'-phenyl-N-Btc-glycine amide (3a), N'-hydroxy-N'-phenyl-N-Btc-L-alanine amide (3b), and N'-hydroxy-N'-phenyl-N-Btc-L-phenylalanine amide (3c) (Scheme 1). It is worth pointing out that the analogous reactions of N-Btc-amino acid chlorides with hydroxylamine hydrochloride in our hands failed to give the corresponding hydroxamic acids (toluene or acetonitrile were used as solvents and triethylamine or N-methylmorpholine as hydrogen chloride acceptors).

Under basic conditions, hydroxamic acids readily underwent cyclisation to afford 1,2,5-oxadiazine-3,6-diones (4) (Scheme 1). The substituent on C-4 depends on the starting amino acid, while the substituent on C-2 atom depends on the hydroxylamine used in the previous reaction step. Two oxadiazine derivatives, viz. 2-phenyl-2H-1,2,5-oxadiazine-3,6(4H,5H)-dione (4a) and 4-benzyl-2-phenyl-2H-1,2,5-oxadiazine-3,6(4H,5H)-dione (4c), were isolated and analysed, while the corresponding L-alanine derivative (4b) was only detected by TLC.

Spectral assignment and CHN analysis of the synthesised new compounds 3 and 4 confirmed their structures. IR spectra of 3 showed the absorption maxima of carbonyl group bound to benzotriazole at 1731–1739 and the hydroxamic acid carbonyl group at 1658–1687 cm⁻¹. In the IR spectra of oxadiazine derivatives, two carbonyl absorptions were visible: carbonyl at position 3 at 1747–1772 cm⁻¹ and carbonyl at position 6 at 1688–1692 (amide I) and 1590–1594 cm⁻¹ (amide II).

¹H NMR spectra of products 1a-c confirmed presence of carboxylic acid group at 13.15–12.00, NH group at 9.42–9.34, aromatic protons at 8.15–7.50 and C-2 atom at 4.50–4.04 ppm. ¹³C spectra confirmed presence of two carbonyl groups: carboxylic acid carbonyl at 173.28–172.18 and carbonyl bound to benzotriazole at 149.44–148.87 and, finally, chiral C-2 atom at 54.98–41.84 ppm.

¹H NMR spectra of products 3a-c revealed presence of N-hydroxyl group at 11 ppm, instead of carboxy group. Hydroxamic acid carbonyl was shifted to lower ppm values (167.80–165.25 ppm) than carboxylic acid carbonyl, while carbonyl bound to benzotriazole, aromatic C-atoms and C-2 had practically the same values as in compounds 1a-c.

¹H NMR spectra of products 4a,c confirmed presence of NH group at 8.84–8.72, aromatic protons at 7.56–7.27 and C-2 atom at 4.49–4.10 ppm. ¹³C spectra confirmed presence of two carbonyl groups at 164.85–164.56 and 152.83–152.69, aromatic C-atoms at 137.39–122.41 and C-2 atom at 55.48–44.20 ppm.

Reaction conditions, yields and analytical data of the newly synthesised compounds are presented in Table I and NMR spectral data are given in Table II.

**Biological studies**

The synthesized oxadiazines were used in biological studies. Values obtained by the MTT-test were all below control values 72 h following incubation (Figs. 1 and 2). No growth inhibition of GMK cells was observed for compounds 4a and 4c at concentration c₃, but at concentrations c₁ and c₂ a statistically significant growth inhibition of GMK cells was observed, ranging from 40% to 60% (Fig. 1). Similarly, a statistically significant growth inhibition of 75% of HeLa cells was observed for 4a at concentration c₁ (Fig. 2),
Table II. \(^1\)H NMR and \(^{13}\)C NMR data of Btc-amino acids 1a-c, hydroxamic acids 3a-c and 1,2,5-oxadiazine-3,6-dione derivatives 4a,c

<table>
<thead>
<tr>
<th>Compound No.</th>
<th>R</th>
<th>(^1)H NMR (^{13})C NMR</th>
</tr>
</thead>
<tbody>
<tr>
<td>1a</td>
<td>H</td>
<td>12.00 (s, 1H, OH), 9.42 (t, 1H, 3, (J = 5.7) Hz), 8.14 (dd, 2H, 2', 5', (J = 8.2) Hz), 7.67 (t, 1H, 3', (J = 7.7) Hz), 7.50 (t, 1H, 4', (J = 7.5) Hz), 4.04 (d, 2H, 2, (J = 5.6) Hz)</td>
</tr>
<tr>
<td>1b</td>
<td>5CH(_3)</td>
<td>12.85 (s, 1H, OH), 9.34 (d, 1H, 3, (J = 7.7) Hz), 8.14 (dd, 2H, 2', 5', (J = 9.00) Hz), 7.67 (t, 1H, 3', (J = 7.3) Hz), 7.50 (t, 1H, 4', (J = 7.3) Hz), 4.50-4.43 (m, 1H, 2), 1.50 (d, 3H, 5, (J = 7.3) Hz)</td>
</tr>
<tr>
<td>1c</td>
<td>5CH(_2)</td>
<td>13.15 (s, 1H, OH), 9.34 (d, 1H, 3, (J = 8.3) Hz), 8.15 (d, 1H, 2', (J = 8.3) Hz), 8.06 (d, 1H, 5', (J = 8.3) Hz), 7.65 (t, 1H, 3', (J = 7.7) Hz), 7.49 (t, 1H, 4', (J = 7.7) Hz), 7.32 (d, 2H, 7, 11, (J = 7.3) Hz), 7.23 (t, 2H, 8, 10, (J = 7.5) Hz), 7.14 (t, 1H, 9, (J = 7.2) Hz), 4.72-4.65 (m, 1H, 2), 3.29 (s, 2H, 5)</td>
</tr>
</tbody>
</table>
Table II, continuations

|   | H     | 10.90 (s, 1H, OH), 9.33 (t, 1H, 3, J = 5.8 Hz), 8.22 (t, 2H, 2', 5', J = 9.3 Hz), 7.77-7.69 (m, 2H, 2'', 6''), 7.58 (t, 2H, 3', 4', J = 7.7 Hz), 7.41 (t, 2H, 3'', 5'', J = 8.0 Hz), 7.18 (t, 1H, 4'', J = 7.4), 4.47 (d, 2H, 2, J = 5.7 Hz) | 167.29 (1), 149.47 (4), 145.42 (1'), 141.23 (1''), 131.16 (6'), 130.09 (5'), 128.51 (3'', 5''), 125.60 (4''), 124.73 (4''), 119.82 (3') 119.18 (2'', 6''), 113.40 (2''), 42.51 (2) |
|   | 5CH₃  | 11.00 (s, 1H, OH), 8.80 (s, 1H, 3), 7.92-7.91 (m, 2H, 2', 5', J = 9.3 Hz), 7.56-7.33 (m, 7H, 3', 4', 2''-6''), 4.34 (q, 1H, 2), 1.37 (d, 3H, 5) | 167.80 (1), 153.91 (4), 139.91 (1'), 138.19 (1''), 130.21 (5'), 129.86 (6'), 128.91 (3'', 5''), 126.37 (4''), 126.23 (4''), 123.84 (3), 121.34 (2'', 6'') 115.93 (2''), 50.50 (2), 16.49 (5) |
|   | 8.84 (s, 1H, 3) 8.23-7.20 (m, 14H, arom.), 4.50-4.45 (m, 1H, 2), 2.60-2.50 (m, 2H, 5) | 165.25 (1), 152.47 (4), 148.44 (1'), 140.00 (6), 135.78 (1''), 130.34 (6''), 129.84 (5), 129.16 (3'', 5''), 127.91 (7, 8, 10, 11), 125.62 (9, 4'), 124.38 (3'', 4'', 6''), 115.63 (2''), 55.95 (2), 38.77 (5) |
|   | H     | 8.72 (s, 1H, 3), 7.56-7.34 (m, 5H, arom.), 4.10 (s, 2H, 2) | 164.56 (1), 152.83 (4), 137.39 (1'), 129.14 (3'', 5''), 127.56 (4''), 122.41 (2'', 6''), 44.20 (2) |
|   | 8.84 (s, 1H, 3), 7.51-7.27 (m, 10H, arom.), 4.49 (t, 1H, 2) 3.19-3.05 (m, 2H, 5) | 164.85 (1), 151.69 (4), 137.09 (1'), 135.59 (6), 129.78 (8, 10), 129.13 (3'', 5''), 128.48 (7, 11), 127.95 (4''), 127.10 (9), 123.16 (2'', 6'') 55.48 (2), 37.34 (5) |

*a NMR data for compounds 1a-c are included as well since they have not been reported until now.  
*b DMSO-d₆ solution, δ (ppm)
while cell proliferation was observed at concentrations \( c_2 \) and \( c_3 \). Statistically significant growth inhibition of HeLa cells was observed also for \( 4c \), ranging from 50% to 75% (Fig. 2). The higher growth inhibition of the tested compounds on HeLa cells, compared to GMK cells, was probably due to the programmed cell death phenomena permanently present in HeLa cells (data not shown) (16). Contrary to the tested compounds, acyclovir stimulated both GMK and HeLa cell growth at all concentrations (Figs. 1 and 2).

As the cytostatic effect of \( 4a \) and \( 4c \) was observed, \( IC_{50} \) values (17) were calculated and are presented in Table III. A compound \( 4a \) showed higher cytostatic effect on GMK than on HeLa cells, while \( 4c \) was more cytotoxic on HeLa than on GMK cells.

Results of the antiviral assay (percentage of CPE inhibition) are shown in Table IV. The inhibition of CPE (approximately 5–20%) for \( 4a \) was observed at concentration \( c_1 \) for all tested viruses, while the inhibition of coxsackievirus CPE (approximately 5–15%) was observed only at concentrations \( c_2 \) and \( c_3 \). Compound \( 4c \) inhibited CPE of adenovirus at concentration \( c_1 \) (approximately 20%) as well as CPE of herpesvirus and enteroviruses at

<table>
<thead>
<tr>
<th>Cell line</th>
<th>( IC_{50} ) (mol L(^{-1}))</th>
<th>( 4a )</th>
<th>( 4c )</th>
</tr>
</thead>
<tbody>
<tr>
<td>GMK</td>
<td>1.125 \times 10^{-4}</td>
<td>5.800 \times 10^{-5}</td>
<td></td>
</tr>
<tr>
<td>HeLa</td>
<td>0.897 \times 10^{-5}</td>
<td>0.824 \times 10^{-4}</td>
<td></td>
</tr>
</tbody>
</table>
concentrations $c_1$, $c_2$ and $c_3$ (approximately 5–20%). A satisfactory inhibitory effect ranging from 20 to 40% of acyclovir, as expected (18), was observed on herpesvirus at all tested concentrations and virus titres.

Poor CPE inhibition of 4a and 4c observed for all tested viruses (maximum 20%) and the lack of CPE inhibition at minor virus titres and at lower concentrations of the tested compounds was probably not an antiviral effect but more likely their inhibitory effect upon cell growth (shown previously by the MTT-test). Thus, the lack of CPE inhibition at low virus titres, where it is usually possible to observe the inhibition of CPE for antiviral compounds (e.g. acyclovir), may confirm the previous statement.

Table IV. Cytopathic effect (CPE) on cell culture of different viruses\(^a\) upon 4a, 4c and acyclovir (Ac)\(^b\) compared to control (solvent DMEM)

<table>
<thead>
<tr>
<th>Virus/titre</th>
<th>CPE inhibition (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>4a</td>
</tr>
<tr>
<td></td>
<td>$c_3$</td>
</tr>
<tr>
<td>Adenovirus 7</td>
<td></td>
</tr>
<tr>
<td>V-1</td>
<td>0</td>
</tr>
<tr>
<td>V-2</td>
<td>0</td>
</tr>
<tr>
<td>V-3</td>
<td>10</td>
</tr>
<tr>
<td>V-4</td>
<td>0</td>
</tr>
<tr>
<td>V-5</td>
<td>0</td>
</tr>
<tr>
<td>Herpesvirus 1</td>
<td></td>
</tr>
<tr>
<td>V-1</td>
<td>0</td>
</tr>
<tr>
<td>V-2</td>
<td>0</td>
</tr>
<tr>
<td>V-3</td>
<td>0</td>
</tr>
<tr>
<td>V-4</td>
<td>0</td>
</tr>
<tr>
<td>V-5</td>
<td>0</td>
</tr>
<tr>
<td>Echovirus 7</td>
<td></td>
</tr>
<tr>
<td>V-1</td>
<td>0</td>
</tr>
<tr>
<td>V-2</td>
<td>0</td>
</tr>
<tr>
<td>V-3</td>
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<tr>
<td>V-4</td>
<td>0</td>
</tr>
<tr>
<td>V-5</td>
<td>0</td>
</tr>
<tr>
<td>Coxsackievirus B5</td>
<td></td>
</tr>
<tr>
<td>V-1</td>
<td>0</td>
</tr>
<tr>
<td>V-2</td>
<td>15</td>
</tr>
<tr>
<td>V-3</td>
<td>15</td>
</tr>
<tr>
<td>V-4</td>
<td>5</td>
</tr>
</tbody>
</table>

\(^a\) Viruses titres V-1 to V-5: serial dilution 1:2 for adenovirus 7 and herpesvirus 1 and 1:10 for echovirus 7 and coxsackievirus B5.

\(^b\) $c_1 = 10^{-4}$ mol L$^{-1}$, $c_2 = 10^{-5}$ mol L$^{-1}$, $c_3 = 10^{-6}$ mol L$^{-1}$. 

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CONCLUSIONS

1,3,5-Oxadiazine-3,6-diones can be successfully prepared by cyclisation of the appropriate hydroxamic acids. Since various methods for the preparation of a wide array of N-monosubstituted hydroxylamines are available, this procedure constitutes a new general method for construction of the 1,3,5-oxadiazine ring system. Selected biological studies indicated that oxadiazines 4a and 4c had no cytotoxic effect on HeLa and GMK cells at tested concentrations, but they had a pronounced cytostatic effect, i.e., they inhibited cell growth in the 25–75% range, depending on the compound, cell line and concentration. A minor inhibition of CPE on adenovirus, herpesvirus and enteroviruses, for compounds 4a and 4c, was observed. These preliminary findings indicate that potential antiviral drugs could be found within oxadiazine derivatives. Thus, further investigation should be undertaken.

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Novi derivati 1,2,5-oksadiazina – Sinteza i biološko djelovanje u in vitro uvjetima


Ključne riječi: 1,2,5-oksadiazin derivati, sinteza, citotoksični učinak, citostatski učinak, protuvirusni učinak, citopatski učinak

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