Synthesis of Heterocyclic and Non-heterocyclic Compounds Derived from Novel 2-Furanones and Evaluation of their Anti-viral Activity

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ABSTRACT

Objective: The aim of the present study is to synthesize novel 2-furanones due to their biological activities and also their ability to be converted to several biologically active heterocyclic and non-heterocyclic compounds. Methods: Two novel 2-furanone derivatives were synthesized and used to prepare fourteen heterocyclic and non-heterocyclic compounds. Structures of all the newly synthesized compounds were confirmed by elemental and spectral analysis including Mass, IR and 1H-NMR spectroscopy. Evaluation of antiviral activity of selected examples of the obtained compounds was performed using two gastroenteric viruses: adenovirus and rotavirus. Results: Two pyridazinone derivatives namely; 3-(3,4-Dichlorophenyl)-5-[(2-methoxyphenyl)methyl]-1H-pyridazin-6-one (7a) and 3-(3,4-Dichlorophenyl)-5-[(4-fluorophenyl)methyl]-1H-pyridazin-6-one (7b) showed high activities against rotavirus. Conclusions: Compounds 7a and 7b can be considered as promising anti-rotaviral agents needing further investigations and clinical studies.

Keywords: Furanone; Hydrazide; Pyridazinone; Rotavirus; Adenovirus

INTRODUCTION

Viruses are from the prime causes of diseases worldwide; viral gastroenteritis is a serious viral infection which affects millions of individuals around the world, most of them are children. The viral gastroenteritis constitutes around 21–40% of infectious diarrhea cases in developed countries 1,2. The infection may occur due to different viruses, for example, coxsackievirus, adenovirus, and rotavirus. Rotavirus is the leading cause of severe gastroenteritis in the pediatric population worldwide 3 adenovirus is the second most common virus causing gastroenteritis in young children, 4 and coxsackievirus is considered the least dangerous cause of gastroenteritis among the three viruses 5. In spite of the fact that in the recent years, various anti-viral agents have been produced, there is a need to create new and safe antiviral drugs.

2-Furanones; well-known heterocyclic derivatives; had attracted a great attention during the last decade due to facile ring opening and conversion to other heterocycles; pyrrolones, pyridazinones, pyrazoles and oxadizoles. These heterocycles acquired an obvious medicinal interest as antimicrobial 6-8, anti-inflammatory 8-16, antimycobacterial 18,19 and anti-cancer agents 20-22.

Literature is also enriched with furanone derivatives and other heterocyclic and non-heterocyclic compounds derived from them having antiviral activities 23-25, as shown in Figure 1; 2-furanone derivative A showed remarkable activity comparable to that of zanamivir 23, and furanone derivative B which is reported as potent anti-adenoviral agent. 26 Both hydrazide C and
Figure 1. Reported potent antiviral derivatives.

pyridazinone D were reported to have promising antiviral activities against anti-avian influenza virus H5N1 virus.  Flurinated pyridazinone C were reported as multi-functional antiviral agents comparable to Ribavirin. Oxazolone derivatives D were reported as promising antiviral agents comparable to that of acyclovir.

The previously mentioned antiviral activity stimulates our attention to continue our previous work to synthesize new antiviral agent against gastroenteric viral infection.

MATERIAL AND METHODS

Synthesis of lead compounds

All commercial chemicals used as starting materials and reagents in this study were purchased from Merck (Darmstadt, Germany) and were of reagent grade. All melting points were uncorrected and measured using Electro-thermal IA 9100 apparatus (Shimadzu, Japan); IR spectra were recorded as potassium bromide pellets on a Perkin-Elmer 1650 spectrophotometer (USA), Faculty of Science, Cairo University, Cairo, Egypt. 1H-NMR spectra were determined on a Varian Mercury (300 MHz) spectrometer (Varian UK) and chemical shifts were expressed as ppm against TMS as internal reference (The Main Chemical warfare Laboratories, Almaza, Cairo, Egypt). Mass spectra were recorded on 70 eV (EI Ms-QP 1000 EX, Shimadzu, Japan), Faculty of Science, Cairo University, Cairo, Egypt. Microanalyses were performed using Vario, Elementar apparatus (Shimadzu, Japan), Organic Microanalysis Unit, Faculty of Science, Cairo University, Cairo, Egypt. Column Chromatography was performed on (Merck) Silica gel 60 (particle size 0.06-0.20 mm). All the listed compounds are new except compound 1 was previously reported.

General procedure for the synthesis of compounds 2a,b:

A mixture of compound 1 (0.03 mol) and equimolar amount of aromatic aldehyde was refluxed in acetic anhydride (15 mL) with triethylamine (3- 4 drops) for 4 h. After completion of reaction, the product was filtered, washed with ethanol and recrystallized from ethanol to obtain compounds 2a,b.

5-(3,4-Dichlorophenyl)3-(2-methoxophenyl)methylene]-furan-2-one (2a)

Yield: 52%; m.p.: 218-220 °C; IR (KBr) ν (cm⁻¹): 1753 (C=O); MS (EI) m/z: 350 (M+4, 4.35%), 348 (M+2, 29.8%), 346 (M⁺, 44.87%), 173 (benzoyl fragment, 100%); 1H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.8 (s, 3H, OCH₃), 7.0-8.1 (m, 9H, Ar-H+ CH methylene); Anal. Calcd. for C₁₈H₁₂Cl₂O₃ (347.19): C, 62.27; H, 3.48%. Found: C, 62.57; H, 3.15%.

5-(3,4-Dichlorophenyl)3-[(4-flurophenyl)methylene]-furan-2-one (2b)

Yield: 34%; m.p.: 240-242 °C; IR (KBr) ν (cm⁻¹): 1752 (C=O); MS (EI) m/z: 338 (M+4, 3.9%), 336 (M⁺, 27.66%), 334 (M⁺, 41.69%), 173 (benzoyl, 100%); 1H-NMR (DMSO-d6, 300 MHz) δ (ppm): 7.3-8.1 (m, 9H, Ar-H+ CH methylene); Anal. Calcd. for C₁₈H₁₃Cl₂FO₂ (335.13): C, 60.92; H, 2.71%. Found: C, 60.72; H, 2.64%.
General procedure for the synthesis of compounds 2a,b

A solution of the furanone derivatives 2a,b (0.01 mol) and ammonium acetate (7.7g, 0.1 mol) in acetic acid (10 mL) was refluxed for 3h. The reaction mixture was left to cool at room temperature and the product obtained was filtered off, recrystallized from ethanol to give compounds 3a,b.

5-(3,4-Dichlorophenyl)-3-[2-methoxyphenyl]methylene]-1H-pyrrol-2-one (3a)

Yield: 30%; m.p.: > 300 °C; IR (KBr) v (cm⁻¹): 3314 (NH), 1769 (C=O); MS (EI) m/z: 349 (M+4, 11.9%), 347 (M+2, 67.2%), 345 (M⁺, 100%); H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.9 (s, 3H, OCH3), 6.9-8.2 (m, 9H, Ar-H+ CH methylene), 10.7 (s, 1H, NH, D2O-exchangeable); Anal. Calcd. for C18H13Cl2FNO2 (442.29): C, 65.17; H, 4.10; N, 3.17%. Found: C, 65.59; H, 4.32; N, 3.47%.

General procedure for the synthesis of compounds 5a-c

A solution of the furanone 2a,b (0.01 mol) and phenyl hydrazine 3ml in Na ethoxide (10ml) was refluxed for 3h. The product obtained was filtered, washed with water and recrystallized from ethanol to give compounds 5a,b.

6-(3,4-dichlorophenyl)4-[(2-methoxyphenyl)methylene]-1-phenyl-1,4-dihydropyridazin-3(2H)-one (5a)

Yield: 80%; m.p.: >300°C; IR (KBr) v (cm⁻¹): 3311 (NH), 1727 (C=O); MS (EI) m/z: 436 (M⁺, 0.14%); H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.8 (s, 3H, OCH3), 6.8-8.5 (m, 15H, Ar-H + CH methylene + NH-D2O exchangeable); Anal. Calcd. for C23H16Cl2FNO2 (437.30): C, 65.92; H, 4.15; N, 6.41%. Found: C, 65.42; H, 4.28; N, 6.34%.

General procedure for the synthesis of compounds 4a-c

To a solution of the furanone 2a,b (0.01 mol) in absolute ethanol (20 mL), benzylamine (1.07 mL, 0.01 mol) was added and the reaction mixture was refluxed for 5h. The product was filtered off, washed with ethanol and finally recrystallized from a suitable solvent to give the amides 4a,b.

N-Benzyl-2-[(2-methoxyphenyl)methylene]-4-(3,4-dichlorophenyl)-4-oxo-butanamide (4a)

Yield: 45%; m.p.: 250-252 °C; IR (KBr) v (cm⁻¹): 3275 (NH), 1711, 1742 (2c=O); MS (EI) m/z: 453 (M⁺, 3.1%); H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.7 (s, 2H, CH2-CO), 3.9 (s, 3H, OCH3), 4.2 (dd, 2H, CH2-NH), 7 (s, 1H, NH, D2O-exchangeable), 6.9-7.6 (m, 13H, Ar-H + CH methylene); Anal. Calcd. for C18H13Cl2NO3 (454.32): C, 66.09; H, 4.66; N, 3.08%. Found: C, 66.25; H, 4.37; N, 3.42%.

N-Benzyl-2-[(4-fluorophenyl)methylene]-4-(3,4-dichlorophenyl)-4-oxo-butanamide (4b)

Yield: 30%; m.p.: 174-176 °C; IR (KBr) v (cm⁻¹): 3245 (NH), 1678, 1642 (2c=O); MS (EI) m/z: 441 (M⁺, 3.22%); H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.3 (s, 2H, CH2-CO), 4.2 (dd, 2H, CH2-NH), 7.0-8.0 (m, 14H, Ar-H + CH methylene + NH, D2O-exchangeable); Anal. Calcd. for C18H16Cl2FNO2 (442.29): C, 65.17; H, 4.40; N, 3.17%. Found: C, 65.59; H, 4.32; N, 3.47%.

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Scheme A. Synthesis of compounds 2a,b - 5a,b.

4-(3,4-Dichlorophenyl)-2-[(4-flurophenyl)methylene]-4-oxo-butanehydrazide (6b)
Yield: 70%; m.p.: 180-182 °C; IR (KBr) v (cm\(^{-1}\)): 3279 (NH), 3250 (NH\(_2\)), 1692, 1656 (2 C=O); MS (EI) m/z: 368 (M+2, 1.79%), 366 (M\(^+\), 2.67%), 337 (M+2 – NHNH\(_2\), 39%), 335 (M–NHNH\(_2\), 62%); \(^1\)H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.2 (s, 2H, CH\(_2\)), 4.4 (s, 2H, NH\(_2\)), D\(_2\)O-exchangeable), 6.8 (s, 1H, NHCO, D\(_2\)O-exchangeable), 7.1-7.6 (m, 8H, Ar-H + CH-methylene); Anal. Calcd. for C\(_{17}\)H\(_{13}\)Cl\(_2\)FN\(_2\)O\(_2\) (367.20): C, 55.61; H, 3.57; N, 7.63%. Found: C, 55.82; H, 3.21; N, 7.42%.

3-(3,4-Dichlorophenyl)-5-[(2-methoxyphenyl)methyl]-1H-pyridazin-6-one (7a)
Yield: (1\(^{st}\) method: 69%, 2\(^{nd}\) method: 65%); m.p.: 208-210 °C; IR (KBr) v (cm\(^{-1}\)): 3218 (NH), 1649 (C=O); MS (EI) m/z: 364 (M+4, 7.8%), 362 (M+2, 45.67%), 360 (M\(^+\), 70.36%); \(^1\)H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.71 (s, 2H, CH\(_2\)), 3.76 (s, 3H, OCH\(_3\)), 6.8-7.9 (m, 8H, Ar-H), 13.2 (s, 1H, NH-pyridazinone, D\(_2\)O-exchangeable); Anal. Calcd. for C\(_{18}\)H\(_{14}\)Cl\(_2\)N\(_2\)O\(_2\) (361.22): C, 59.85; H, 3.91; N, 7.76%. Found: C, 59.74; H, 3.52; N, 7.42%.

General procedure for the synthesis of compounds 7a,b
Method 1: A solution of hydrazides 6a,b (0.01 mol) in HCl/AcOH (1:3) was refluxed for 3 h. The solid that separated after concentration and cooling was recrystallized from ethanol to obtain compounds 7a,b.
Method 2: To a solution of the furanones 2a,b (0.01 mol) in absolute ethanol (20 mL), hydrazine hydrate (3.5 mL, 0.11 mol) was added. The reaction mixture was refluxed for 4 h, then cooled and poured onto ice water. The product obtained 7a,b was filtered off, washed with hexane.

3-(3,4-Dichlorophenyl)-5-[(4-flurophenyl)methyl]-1H-pyridazin-6-one (7b)
Yield: (1\(^{st}\) method: 44%, 2\(^{nd}\) method: 40%); m.p.: 212-214 °C; IR (KBr) v (cm\(^{-1}\)): 3187 (NH), 1647 (C=O); MS (EI) m/z: 352 (M+4, 12.2%), 350 (M+2, 65.76%), 348 (M\(^+\), 100%); \(^1\)H-NMR (DMSO-d6, 300 MHz) δ (ppm): 3.8 (s, 2H, CH\(_2\)), 7.0-8.1 (m, 8H, Ar-H), 13.27 (s, 1H, NH-pyridazinone, D\(_2\)O-exchangeable); Anal. Calcd. for C\(_{17}\)H\(_{11}\)Cl\(_2\)FN\(_2\)O\(_2\)(349.18): C, 58.48; H, 3.18; N, 8.02%. Found: C, 58.56; H, 3.25; N, 8.42%.

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Scheme B. Synthesis of compounds 6a,b - 9a,b.

**General procedure for the synthesis of compounds 8a,b**

To a solution of the hydrazides 6a,b (0.01 mol) in dry benzene (20 mL), benzoyl chloride (1.4 mL, 0.01 mol) was added. The reaction mixture was heated under reflux for 2 h. The solvent was evaporated, and the solid obtained was washed thoroughly with ethanol, drained, and recrystallized from the suitable solvent to give compounds 8a,b.

2-benzoyl-5-[(2-methoxyphenyl)methylene]-3-(3,4-dichlorophenyl)-1H-pyridazin-6-one (8a)

Yield: 35%; m.p.: 177-179 °C; IR (KBr) υ (cm⁻¹): 3217 (NH), 1790,1694 (2 C=O); MS (EI) m/z: 465 (M⁺, 5%), 105 (benzoyl, 100%); ¹H-NMR (DMSO-d₆, 300 MHz) δ (ppm): 3.8 (s, 3H, OCH₃), 6.9-8 (m, 14H, Ar-H + CH methylene), 11.2 (s, 1H, NH, D₂O-exchangeable); Anal. Calcd. for C₂₅H₁₈Cl₂N₂O₃ (465.32): C, 64.25; H, 4.31; N, 5.99%. Found: C, 64.51; H, 4.16; N, 5.64%.

2-benzoyl-5-[(4-fluorophenyl)methylene]-3-(3,4-dichlorophenyl)-1H-pyridazin-6-one (8b)

Yield: 36%; m.p.: 266-268 °C; IR (KBr) υ (cm⁻¹): 3186 (NH), 1710,1672 (2 C=O); MS (EI) m/z: 454 (M⁺, 4.25%), 452 (M⁺, 6.25%); ¹H-NMR (DMSO-d₆, 300 MHz) δ (ppm): 7-8.2(m, 14H, Ar-H+ CH methylene), 11.2(s, 1H, NH, D₂O-exchangeable); Anal. Calcd. for C₂₄H₁₆Cl₂F₁N₂O₂ (453.28): C, 63.59; H, 3.34; N, 6.18%. Found: C, 63.24; H, 3.17; N, 6.55%.

**General procedure for the synthesis of compounds 9a,b**

A solution of hydrazides 6a,b (0.01 mol) and carbon disulphide (3ml) in pyridine (10ml) was refluxed for 3h. The reaction mixture was left to cool at room temperature and poured onto ice water; the product obtained was filtered, washed with water and recrystallized from ethanol to give compounds 9a,b.

1-(3,4-dichlorophenyl)-4-(2-methoxyphenyl)-3-(2-thioxo-4,5-dihydro-1,3,4-oxadiazol-5-yl)but-3-en-1-one (9a)

Yield: 45%; m.p.: 136-138 °C; IR (KBr) υ (cm⁻¹): 3203 (NH), 1651 (C=O), 1244 (C=S); MS (EI) m/z: 420 (M⁺, 1.47%); ¹H-NMR (DMSO-d₆, 300 MHz) δ (ppm): 3.7 (s, 3H, OCH₃), 3.8 (s, 2H, CH₂), 6.8-8.7(m, 8H, Ar-H+ CH methylene), 13.2(s, 1H, NH, D₂O-exchangeable); Anal. Calcd. for C₁₉H₁₄Cl₂N₂O₃S (421.23): C, 54.17; H, 3.35; N, 6.65%. Found: C, 54.25; H, 3.24; N, 6.17%.

1-(3,4-dichlorophenyl)-4-(4-fluorophenyl)-3-(2-thioxo-4,5-dihydro-1,3,4-oxadiazol-5-yl)but-3-en-1-one (9b)

Yield: 50%; m.p.: 191-193 °C; IR (KBr) υ (cm⁻¹): 3284 (NH), 1771 (C=O), 1224 (C=S); MS (EI) m/z: 408 (M⁺, 0.15%); ¹H-NMR (DMSO-d₆, 300 MHz) δ (ppm): 3.8 (s, 2H, CH₂), 7.09-8.09 (m, 8H, Ar-H+ CH methylene), 13.3(s, 1H, NH, D₂O-exchangeable); Anal. Calcd. for C₁₈H₁₁Cl₂F₁N₂O₂S (409.19): C, 52.83; H, 2.71; N, 6.85%. Found: C, 52.54; H, 2.75; N, 6.34%.

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Biological evaluation of anti-viral activity

Cytotoxicity assay
All samples (100 mg) were dissolved in 500 µL of ethanol or acetic acid. Cell monolayers Hep2 and MA104 (obtained from The Holding Company for Biological Products & Vaccines VACSERA, Egypt) were trypsinized, washed with culture medium and plated in a 96-well flat bottomed plate with 5 X 10³ cells per well for both cell lines. After 24 h incubation, each diluted (Greiner-Bio-One, Germany) tested material (10 fold dilutions of decontaminated samples which 12 µL of 100x of antibiotic, antinmucotic mixture was added to 500 µL of each sample) was added to the appropriate wells and the plates were incubated for further 48 h at 37°C in a humidified incubator with 5% CO2. The supernatants were removed from the wells and cell viability was evaluated using microscopical examination, trypan blue and the MTT technique [33-35]. The results are obtained from triplicate assays with at least 5 extract concentrations. The percentage of cytotoxicity is calculated as [(A-B)/A] X 100, where A and B are the OD492 of untreated and of treated cells, respectively.

Antiviral test
Non-toxic dose of each tested compound was used in the in vitro antiviral screening method was used to estimate the inhibition of the cytopathic effect (CPE) of the pure compound on MA104 and HEP-2 cell monolayers infected with rotavirus Wa strain with initial titre 1 X 10⁶ PFU/mL (ATCC VR2018) and adenovirus type 7 with initial titre 1 X 10³ PFU/mL (obtained by Dr. Ali Fahmy, VACSERA, EGYPT) using the endpoint titration technique (EPTT). [34] Confluent monolayers of MA104 and HEP-2 cells were grown in 96-well microtiter plates, which were infected with serial tenfold dilutions of rotavirus Wa strain and adenovirus type 7 suspensions, respectively. The viruses were allowed to adsorb for 60 min at 37°C. Then, serial twofold dilutions of the test compounds in maintenance medium, supplemented with 2% serum and antibiotic, were added. The plates were incubated at 37°C, and the viral cytopathic effect was recorded by light microscopy after 2 to 8 days. Virus suspensions are characterized by their virus titres, which are expressed as the smallest amount of virus capable of producing a reaction in the host cells. The antiviral activity is expressed as a reduction factor (RF), being the ratio of the viral titres in the virus control and in the presence of the maximal non-toxic dose of test substance.

MTT assay (antiviral colorimetric assay):
Both MA104 and Hep2 cell monolayers were grown in 96-well microtiter plates. Dilutions of the extracts, prepared as described above for the EPTT assay, were added 1 h before viral infection. Ten infectious doses of virus were added to each well and incubated at 37°C in humidified 5% CO2 atmosphere for 48 h. Controls consisted of untreated infected, treated uninected and untreated uninfected cells. Cell viability was evaluated by the MTT colorimetric technique. [35] Briefly, the supernatants were removed from the wells and 28 µL of an MTT (Sigma) solution (2 mg/mL in PBS) was added to each well. The plates were incubated for 1.5 h at 37°C, and 130 µL of DMSO was added to the wells to dissolve the MTT crystals. The plates were placed on a shaker for 15 min and the optical density was determined at 492 nm (OD492) on a multiwell spectrophotometer. The 50% cytotoxic concentration (CC₅₀) of the test extract is defined as the concentration that reduces the OD492 of treated uninfected cells to 50% of that of untreated uninfected cells. The 50% antiviral effective concentration, i.e., 50% inhibitory concentration of the viral effect (IC₅₀) is expressed as the concentration that reduces the absorbance of infected cells to 50% when compared to infected cells and control cells. The percent protection is calculated as [(A-B)/C] X 100, where A, B and C are the OD492 of untreated infected, untreated infected, and untreated uninfected cells, respectively.

Data analysis
CC₅₀ and IC₅₀ for each compound were obtained from dose-effect-curves. The CC₅₀ and IC₅₀ are the average of four assays with 5 concentrations within the inhibitory range of the compound. The therapeutic index (i.e., selective index) is defined as CC₅₀/IC₅₀.

RESULTS AND DISCUSSION

Chemistry
4-Oxo-butanedioic acid derivative 1 was synthesized as reported before [29] and it is used to prepare 2-furanone derivatives 2a,b by reaction with aromatic aldehydes in acetic anhydride following modified Perkin reaction conditions. [7, 11, 12, 18, 19, 24] as revealed in scheme A.

2-Furanone derivatives 2a,b is useful as starting materials in the synthesis of several heterocyclic and non-heterocyclic derivatives. When they were reacted, separately, with ammonium acetate; pyrrol-2-one derivatives 3a,b were produced. 4-Oxo-butanamides 4a,b were synthesized by refluxing 2-furanones 2a,b with benzyl amine and 1-phenyl-pyridazinone 5a,b were also prepared from 2-furanones by refluxing with phenyl hydrazine according to the reported procedure [19,21].

Pyridazinone derivatives 7a,b can be prepared by stirring of 2-furanone derivatives 2a,b with hydrazine hydrate to obtain hydrazide derivatives 6a,b which can be cyclized by refluxing in HCl/AcOH to obtain the desired pyridazinone derivatives 7a,b. They can be

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prepared directly by refluxing 2-furanone derivatives 2a,b with hydrazine hydrate.

Hydrazide derivatives 6a,b were refluxed with benzoyl chloride to afford 2-benzoyl-pyridazinone derivatives 8a,b.

Finally, oxadiazol-thione derivatives 9a,b were prepared via heating under reflux hydrazide derivatives 6a,b with carbon disulfide in pyridine as revealed in scheme B.

Biological results

Cytotoxicity

Cytotoxicity testing of any chemotherapeutic agent (including antiviral compounds) is a very important step in its biological evaluation, as it should determine that neither acute nor long-term toxicity should occur to the host. Cytotoxicity of the selected compounds was examined on two different cell lines: Hep-2, and MA-104. Table 1 shows the CC50 for all the tested compounds. The CC50 was determined using two different techniques: by counting the number of viable cells and by examining the effect on cell morphology.

Antiviral assay results

The non-toxic doses of the compounds were tested against adenovirus type 7 and rotavirus Wa strain. Considerable antiviral activity was observed with some of the tested compounds against adenovirus type 7 and rotavirus Wa strain with different degrees of activity and specificity. Figure 2 showed the Rotavirus % reduction by the tested compounds, the viral reduction % for the most potent compounds 7b and 7a reached 84% and 79% respectively, while compounds 6a and 6b showed moderate activity with viral reduction 53% and 47% respectively.

Figure 3 showed the Adenovirus % reduction by the tested compounds, compounds 2b, 6b, 7a and 7b showed only lower activities against adenovirus with % viral reduction ranging from 25.5 to 38%.

Table 1. CC50 values of the tested compounds on Hep-2 and MA-104 cell lines

| Compounds | CC50 for Hep-2 cell line (mM) | CC50 for MA-104 cell line (mM) |
|-----------|-------------------------------|-------------------------------|
| 2a        | 166                           | 80.22                         |
| 2b        | 40.76                         | 112                           |
| 6a        | 2.66                          | 138                           |
| 6b        | 3.18                          | 77.3                          |
| 7a        | 48.8                          | 44                            |
| 7b        | 5.6                           | 162                           |

To explain structureactivity relationship (SAR) of the tested compounds from the results; it was revealed that two newly synthesized furanone derivatives are inactive or with low activity against both rotavirus and adenovirus. Upon preparation of the hydrazides, they acquired moderate activity against rotavirus and low activity against adenovirus.

Pyridazinone derivatives, in consistent with what was reported before, acquired high antiviral activity, herein with specificity against rotavirus over adenovirus. It is also noticed the superiority of the fluorinated derivative over the methoxy substituted derivative. Figure 4

CONCLUSION

In conclusion, as presented in this study, novel 2-furanone derivatives were synthesized and used them to prepare novel hydrazides, 2-pyrrolone, 2-pyridazinone and oxadiazole derivatives. 6 of the synthesized compounds were investigated for their anti-viral activity against gastroentericviruses rotavirus and adenovirus; the biological results revealed that pyridazinones 7a and 7b showed promising activity against rotavirus. Thus, these compounds might be promising anti-viral candidates and there is a need for more studies and structural optimization to improve their activity.
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Conflict of interest
Authors would like to declare that there are no relationships or interests that could have direct or potential influence or impart bias on the work.

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