SYNTHESIS AND ANTICANCER ACTIVITY OF IMIDAZOLE DERIVATIVES INCORPORATING CHROMENE MOIETY

Nashwa M. Saleh
Department of Chemistry, Faculty of Science (Girl’s), Al-Azhar University, Nasr City, Cairo, Egypt

ABSTRACT

The oxo-chromene-6-carbaldehyde derivative 1 was used as a starting material for the synthesis of some novel pyran[3,2-b] xanthene-4,7-dione 2 and chromen-oxazol-5-(4H)-one derivatives 4,5. Furthermore, the oxazolone derivative 4 was used as key intermediates to synthesize some novel imidazol derivatives 6-14. The structures of target compounds were confirmed by elemental analyses and spectral data. All the target compounds were subjected to in-vitro antitumor activity against breast and colon human tumor cell lines (MCF7 and HCT). Most of the screened compounds showed interesting cytotoxic activities compared to a reference drug.

Keywords: Chromene, Oxazolone, Imidazole, Antitumor activity.

INTRODUCTION

In recent years, the high therapeutic properties of the imidazole related drugs have been attracting the attention the medicinal chemists to synthesize a large number of novel chemotherapeutic agents. Imidazole drugs have broadened scope in remedying various dispositions in clinical medicines. Medicinal properties of imidazole containing compounds include anticancer (Congio, et al., 2008), antimicrobial, (Aridoss, et al., 2006; Nagarapu, et al., 2008), antibacterial, (Bhandari, et al., 2009) antifungal, (Emami, et al., 2008) and antioxidant (Smith and Reeves, 1987; Bakr, et al., 2011). Encouraged by these observations and in continuation of our previous work to discover new biologically active heterocyclic compounds (Abdel-Aziz, et al., 2010; Abdalla, et al., 2009; Amr, et al., 2009; Abde-Wahab, et al., 2009a; Abde-Wahab, et al., 2009b; Abde-Wahab, et al., 2009c). Lepidiline A and B are the imidazole compounds which exhibit cytotoxicity against various types of human cancer cell line sat micro molar concentrations (Cui, et al., 2003). In addition to imidazole, some other azole and benzazole ring systems such as benzoxazole, benzothiazole, triazole, tetrazole, thiadiazole and thiazoline can be shown as pharmacophore groups which are responsible for the anticancer activity (Kumar, et al., 2002; Hose, et al., 2003; Al-Masoudi, et al., 2006; Perez-Ruixo, et al., 2006; Ozkay, et al., 2010; Padmavathi, et al., 2011). We now tried to apply the Erlenmeyer-Ploechl azalactone synthesis (Erlenmeyer and Liebig, 1893). This latter reaction has often been used with the aim of synthesizing fused pyran-2-ones by combining condensation, with lactonization. For this reason simple one-pot procedures have been developed using 1,3-dicarbonyl compounds, one-carbon synthon (triethylorthoformate, or N,N-dimethylformamide dimethyl-acetal) N-acetyl glycine and large excess of acetic anhydride (Kepe, et al., 1992). Another approach for these reactions made use of a two steps synthesis incorporating the one-carbon synthon either in the CH-acidic compound (Kepe, et al., 1999) or in the azalactone (Nulwad and Shirodkar, 2002). The present study was provoked by the paper of Mulwad and Shirodkar describing the condensation of 3-formyl-4-hyd-roxycoumarin with hippuric acid or N-acetylglucose to form 3-acylamino-2, 5-dioxo-pyrano[3,2-c]benzopyrans (Ivanov, et al., 2005).
RESULT AND DISCUSSION

Chemistry

The synthetic procedures adopted to obtain the target new compounds are showed in Schemes 1-5. Treatment of 7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromene-6-carbaldehyde (Kepe, et al., 1993) 1 with cyclic active methylene as 5,5-dimethylcyclohexane-1,3-dione yielded the condensation product 5-methoxy-2,9,9-trimethyl-8,9,10,10a-tetrahydropyrano[3,2-b]xanthene-4,7-dione 2. The formation of compound 2 is assumed to proceed through the in situ intramolecular cyclization of the non isolable intermediate via nucleophilic addition of the hydroxyl group to the carbonyl group followed by elimination of water, Scheme 1. Structure of compound 2 was confirmed on the basis of elemental analysis and spectral data. IR spectrum of compound 2 revealed the absence of aldehyde and hydroxyl groups, the carbonyl stretching frequency was observed at 1657 cm\(^{-1}\) and the coumarin carbonyl stretching at 1723 cm\(^{-1}\). Also, the \(^1\)HNMR spectrum indicated the presence of a singlet at \(\delta 1.17\) ppm due to \(\text{C}_9-2\text{CH}_3\) protons. The \(\text{C}_2-\text{CH}_3\) protons were observed a singlet at 2.29 ppm. The \(\text{C}_{10}-\text{CH}_2\) and \(\text{C}_8-\text{CH}_2\) protons were observed at 1.5, 2.9 ppm. The \(\text{C}_5-\text{OCH}_3\) protons were observed downfield as singlet at 4.23 ppm. The \(\text{C}_3, 6, 12-\text{H}\) of coumarin protons resonated in the range of 6.13-7.79 ppm. Also, the structure 2 was supported by its mass spectrum which revealed a molecular ion peak at \(m/z 340\ (9.72\%)\) corresponding to the molecular formula \(\text{C}_{20}\text{H}_{20}\text{O}_5\).

![Scheme 1](image)

Scheme 1. Expected scheme for the synthesis of xanthene-4,7-dione derivative from chromene-carbaldehyde.

The present study was provoked by the paper of Mulwad (Ivanov, et al., 2005) 1 which reacted with o-chlorobenzyglycine and acetylglycine 3a, b in acetic acid/acetanhydride in presence of fused sodium acetate under reflux afforded 2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)oxazol-5(4H)-one and 4-(7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-2-methyl-oxazol-5-(4H)-one 4,5, respectively. Scheme 2. The intermediates could not be isolated and underwent spontaneous recyclization via opening of the azalactone ring and formed the fused 2(1H)-pyridone ring. Compounds 4, 5 confirmed by IR spectra which revealed the oxazol carbonyl stretching frequency at 1703, 1710 cm\(^{-1}\), where as the coumarin carbonyl stretching at 1619, 1617 cm\(^{-1}\), while it showed the presence of broad band for OH group stretching at 3423, 3363 cm\(^{-1}\) respectively. Structures 4, 5 were supported by its mass and \(^1\)H NMR spectra. \(^1\)H NMR (DMSO-d_6, 300MHz) of 4 raved a singlet was observed at 2.26 ppm due to \(\text{C}_2-\text{CH}_3\) protons. The \(\text{C}_5-\text{OCH}_3\) protons were observed downfield as singlet at 3.91 ppm.
The C₃,₈–H of coumarin, CH= and CH-aromatic were resonated in the range of δ 6.13-8.19 ppm.

Scheme 2. Expected scheme for the synthesis of oxazole derivatives from chromene-6-carbaldehyde.

OH proton was observed as singlet at 11.34 ppm. Its mass spectrum which revealed a molecular ion peak at m/z 411 (M-1, 5.16 %) corresponding to the molecular formula C₂₁H₁₄ClNO₆. ¹H NMR (DMSO-d₆, 300MHz) of 5 raveled a singlet was observed at 2.23 ppm due to C₂–CH₃ protons. The CH₃ protons of oxazolone were observed a singlet at 3.41 ppm. The C₅-OCH₃ protons were observed downfield as singlet at δ 3.89 ppm. The C₃,₈–H of coumarin and CH= were resonated in the range of δ 6.53-8.21 ppm. The OH proton was observed as singlet at 11.26 ppm. Its mass spectrum which revealed a molecular ion peak at m/z 315 (6.04 %) corresponding to the molecular formula C₁₆H₁₃NO₆. Oxazolone derivative 4 was used as intermediate for the synthesis of the some imidazole derivatives via its reaction with some aromatic and hetero aromatic amines. Thus, Oxazolone derivative 4 reacted with 4-aminoacetophenone, 7-aminothiophenol, sulfaquionoxaline , 2-amino-4-hydroxypyrimidine and 4-aminoantipyrine in glacial acetic acid in presence of fused sodium acetate to yield yield 2-(2-chlorophenyl)-4-(7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1-(4-aryl)-1H-imidazol-5(4H)-one 6-10, respectively, (Scheme 3). The structure of compounds 6-10 were confirmed by their IR, ¹HNMR and Mass spectra. The IR spectra revealed the imidazol carbonyl stretching frequency and the coumarin carbonyl stretching with addition special group for each compounds, its mass spectrum and ¹H NMR spectra respectively in experimental section.
Furthermore, treatment of compound 4 with formamide achieved 2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1H-imidazol-5(4H)-one 11. Compound 4 reacted with hydrazine hydrate afforded 1-amino-2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1H-imidazol-5(4H)-one 12, (Scheme 4). Compound 12 confirmed by IR spectrum which affords the imidazol carbonyl stretching frequency at 1701 cm⁻¹, the coumarin carbonyl stretching at 1664 cm⁻¹ and NH₂ at 3328, 3221. Also, the structure 12 was supported by its mass spectrum which revealed a molecular ion peak at m/z 425 (22.55%) corresponding to the molecular formula C₂₁H₁₆ClN₃O₅.
Scheme 4. Expected scheme for the reaction of oxazole derivatives with formamide and hydrazine hydrate and thiosemicarbazide.

Finally, treatment of oxazoles 4,5 with thiosemicarbazide afford imidazotriazole derivatives 13, 14, respectively. Compounds 13, 14 confirmed by IR spectra which revealed disappear the imidazol carbonyl stretching frequency and appear the coumarin carbonyl stretching frequency at 1632, 1650 and (C=S) at 1568, 1567, respectively and presence of stretching broad band for OH group at 3432, 3426 cm⁻¹. Also, structures 13, 14 were supported by its mass spectrum and ¹H NMR spectra showed in experimental section. The formation of imidazotriazole derivatives 13, 14 is assumed to proceed through the in situ intramolecular cyclization of the non isolable intermediate via nucleophilic addition of the amino group to the carbonyl group followed by elimination of water, (Scheme 5).

BIOLICAL ACTIVITIES

In vitro anticancer screening

Chemotherapy is a major therapeutic approach for the two human cultured cell lines which breast cancer cell and colon cancer cell lines (MCF7, HCT) in comparison to the known anti cancer drugs; doxorubicin and vinblastine as reference drugs, is one of the most effective antitumor agents used to produce regression in acute leukemias, Hodgkin’s disease, and other lymphomas. The cytotoxic effects of the newly synthesized compounds were evaluated in the MCF7 breast cancer cell line with doxorubicin was used as a reference drug in six different concentrations. Also, HCT colon cancer cell line with vinblastine was used as a reference drug in three different concentrations. The primary anticancer assay was performed in accordance with the protocol of the Drug Evaluation Branch (Wasylyk, et al., 2008; Wilson and Bottiglier, 1962). From the results obtained in table1, the relationship between surviving fraction and drug concentration was plotted to obtain the survival curve of the MCF7 breast cancer cell line and another of the HCT colon cancer cell line. The response parameter calculated was the IC₅₀ value which corresponds to the compound concentration causing 50% mortality in net cells as depicted in Table 1. In the new work, seven of the newly synthesized compounds 2, 6, 7, 8, 9, 10, 13, 14 were selected. The most of the screened compounds showed interesting cytotoxic activities compared to a reference drug, according to IC₅₀ (which concentration causing 50% inhibition of cell growth. Chemotherapy is a major therapeutic for the both localized and metastasized cancers. Covering all structural variations in these analogs, being of tetrahydropyrano[3,2-b]xanthene-4,7-dione 2 on colon cell line HCT was more active compound (IC₅₀: 3.1 µg/ml) and was found to be near to a reference drug (Doxorubicin, IC₅₀: 3.7µg/ml). However the activity of compound 2 against MCF7 was (IC₅₀: 4.2 µg/ml) has near to a reference drug (vinblastine IC₅₀: 2.97µg/ml). Imidazole moiety attached to mercaptophenyl derivative 7 (IC₅₀: 6.63 µg/ml) or to imidazotriazole derivatives 13 (IC₅₀: 7.85 µg/ml) and 14 (IC₅₀: 6.9 µg/ml) have moderate activity which have one half that of doxorubicin (IC₅₀: 3.743 µg/ml) against HCT. For MCF7, compound 7 (IC₅₀: 8.5 µg/ml), compound 13 (IC₅₀: 9.2 µg/ml) and compound 14 (IC₅₀: 8.8 µg/ml) which have one third that of vinblastine (IC₅₀: 2.97 µg/ml) according to the presence of mercapto group. The antitumor activity of another compounds containing hetero atom as compound 8 (IC₅₀: 10.5 µg/ml for MCF7, (C₅₀: 11.2 µg/ml for HCT), compound 9 (IC₅₀: 10.9 µg/ml for MCF7, IC₅₀: 12.3 µg/ml for HCT) and compound 10 (IC₅₀: 12 µg/ml for MCF7, IC₅₀: 12.8 µg/ml for HCT) have one forth activity against MCF7 which that of vinblastine (IC₅₀: 2.97µg/ml) and against HCT which have one third that of doxorubicin (IC₅₀: 3.743 µg/ml) respectively. All have significance value according to presence of quinoxaline, pyrimidine and pyrazole moiety. Also compound 6 (IC₅₀: 14 µg/ml for MCF7, IC₅₀: 13.3 µg/ml for HCT) containing acetylphenyl group has less activity
compare to the six compounds but has considering value against to antitumor activity. Furthermore, all testing compounds gave antitumor activity against colon cancer cell HCT more than the activity against breast cancer cell MCF7. Finally, the decreasing order of growth inhibitory activities against breast cancer cell MCF7 and colon cancer cell HCT in comparison to the known anticancer drugs are $2>14>13>7>8>9>10>6$. Table (1).

**CONCLUSIONS**

In this study, seven of the newly synthesized compounds $2, 6, 7, 8, 9, 10, 13, 14$ were selected to evaluate their in vitro growth inhibitory activities against breast cancer cell MCF7 and colon cancer cell HCT in comparison to the known anticancer drugs: doxorubicin and vinblastine as reference drugs. In vitro growth inhibitory activities of imidazole derivatives against MHF7 and HCT cell lines revealed significant potential antitumor activity. Best results were gained by compound $2$ since it showed approximately similar potency against MCF7 and HCT respectively. Chromone derivatives containing Imidazole moiety attached to mercaptophenyl $7$ or to imidazotriazole derivatives $13, 14$ is moderate activity according to the presence of mercapto group. Hetero compounds as $7, 8$ and $9$ have signification value according to presence of quinoxaline, pyrimidine and pyrazole groups. Also, compound $6$ containing acetylphenyl group has considering value to antitumor activity. There is provided chromone derivatives of the general formula which an antitumor pharmaceutical composition comprising as an effective ingredient the novel chromone derivatives.

**EXPERIMENTAL**

**General methods**

Melting points were determined on a Stuart melting point apparatus. IR spectra were recorded on a Shimadzu-440 IR spectrophotometer using the KBr technique (Shimadzu, Japan). NMR spectra were measured on a Bruker Avance 300 (300Hz) in DMSO-$d_6$ as a solvent, using tetramethylsilane (TMS) as an internal standard. The mass spectra were recorded on Shimadzu GCMS-QP-1000EX mass spectrometers at 70 eV. Elemental analyses were carried out by the Micro-analytical Research Centre, Cairo University. Analytical results for C, H and N were within ±0.4% of the calculated values. All reagents were of commercial quality and used without purification, follow up of the reaction and checking the purity of the compounds.

7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromene-6-carbaldehyde $1$ was prepared according to the literature procedures (Kepe, et al., 1993).

5-methoxy-2,9,9-trimethyl-8,9,10,10a-tetrahydropyran-3,2-b]xanthene-4,7-dione $2$.

A mixture of $1$ (0.01 mol) and cyclic active methylene compound as 5, 5-dimethylcyclohexane-1,3-dione (0.01 mol) in ethanol (50 mL) and catalytic amount of triethylamine (0.5 mL) was heated under reflux for 3 h. The reaction mixture was poured onto ice water, collected, dried and solid was crystallized from the proper solvent to give $2$ as yellow crystals. Yield: 80%; m.p. 166-168°C; IR (KBr, cm$^{-1}$): disappear of OH and the presence of two C=O groups at 1657, 1723 in added to 2953 (CH aliph.), 3076 (CH arom.), MS, m/z (%): 340 [M] (9.72), 52.3 (100), $^1$H NMR (DMSO-$d_6$, 300MHz): $\delta$ 1.17 [s, 6H, 2CH$_3$], 1.5 [s, 2H, C$_{10}$H], 1.98 [s, 3H, CH$_3$], 2.9 [s, 2H, C$_8$H], 3.9 [s, 3H, OCH$_3$], 6.13-7.79 [m, 4H, Ar-H]. Anal. Calcd. for C$_{20}$H$_{20}$O$_5$: C, 70.57%; H, 5.92%; O, 23.50%; Found: C, 70.64%; H, 5.99%; O, 23.56%. 
2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene) oxazol-5(4H)-one 4.

A mixture of 1 (0.01 mol), o-chlorobenzoylglycine (0.01 mol) and fused sodium acetate (0.03 mol) in acetic acid (10 mL) and acetic anhydride (10 mL) was heated on a steam bath for 3h. The reaction mixture was poured into a cold mixture of ethanol-water (1:1 (50 mL) and stirred for 15 min. The solid product was filtered off, washed of water, dried and recrystallized from the proper solvent to give 4 as yellow crystals. Yield: 80%; m.p. 190-192°C; IR (KBr, cm⁻¹): 3423 (br, OH), 3010 (CH arom.), 2984 (CH aliph.), 1703, 1619 (2C=O), MS, m/z (%): 411 [M-1] (5.16), 68.5 (100). ¹H NMR (DMSO-d₆, 300MHz): ¹J 1.26 [s, 3H, CH₃], 3.93 [s, 3H, OCH₃], 6.13-8.19 [m, 7H, Ar-H], 10.34 [s, 1H, OH]. Anal. Calcd. for C₂₁H₁₄ClN₂O₆: C, 61.25%; H, 3.43%; N, 3.40%; Found: C, 61.34%; H, 3.46%; N, 3.45%.

4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-2-methoxyazol-5(4H) one 5.

A mixture of 1 (0.01 mol), acetyl glycine (0.01 mol) and fused sodium acetate (0.03 mol) in acetic acid (10 mL) and acetic anhydride (10 mL) was heated on a steam bath for 3h. The reaction mixture was then poured into a cold mixture of ethanol-water (1:1 (50 mL) and stirred for 15min. The solid product was filtered off, washed of water, dried and recrystallized from the proper solvents to give 5 as yellow crystals. Yield: 90%; m.p. 170-172°C; IR (KBr, cm⁻¹): 3363 (br, OH), 3008 (CH arom.), 2994 (CH aliph.), 1710, 1617 (2C=O), MS, m/z (%): 315 [M] (6.04), 166 (100). ¹H NMR (DMSO-d₆, 300MHz): ¹J 1.19 [s, 3H, CH₃], 1.41 [s, 3H, CH₃], 3.93 [s, 3H, OCH₃], 6.53-8.21 [m, 7H, Ar-H, =CH], 10.26 [s, 1H, OH]. Anal. Calcd. for C₁₅H₁₃NO₆: C, 60.95%; H, 4.16%; N, 4.44%; Found: C, 60.87%; H, 4.06%; N, 4.39%.

General procedure 2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)substituted-1H-imidazol-5(4H)-one 6-10.

A mixture of 4 (0.01 mol) and aromatic amines namely: 4-amino-acetophenone i, O-aminothiophenol ii or hetero-aromatic amines as sulfamethoxypyridazine iii, 2-amino-4-hydroxy-pyrimidine iv and 4-aminoantipyrine v (0.01 mol) and fused sodium acetate (0.03 mol) in glacial acetic acid (30 mL) was heated under reflux for 1h, then cooled. The reaction mixture was poured onto cold water. The solid products were filtered off, washed of water, dried and recrystallized from the proper solvents to give 6-10 respectively.

1-(4-acetylphenyl)-2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1H-imidazol-5(4H)-one 6.

Brown crystals; Yield: 60%; m.p.270-272°C; IR (KBr, cm⁻¹): 3362 (br, OH), 3078 (CH arom.), 2920 (CH aliph.), 1730, 1665 (2C=O), MS, m/z (%): 528 [M-1] (5.15), 138 (100). ¹H NMR (DMSO-d₆, 300MHz): ¹J 1.81 [s, 3H, CH₃], 2.24 [s, 3H, CH₃], 3.89 [s, 3H, COCH₃], 6.32-8.21 [m, 11H, Ar-H, =CH], 10.56 [s, 1H, OH]. Anal. Calcd. for C₂₉H₂₁ClN₂O₆: C, 65.85%; H, 4.00%; N, 5.30%; Found: C, 65.80%; H, 3.93%; N, 5.22%.

2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1-(2-mercaptophenyl)-1H-imidazol-5(4H)-one 7.

Black crystals; Yield: 65%; m.p.240-242°C; IR (KBr, cm⁻¹): 3444 (br, OH), 3078 (CH arom.), 2924 (CH aliph.), 1729, 1645 (2C=O), SH (1523), MS, m/z (%): 518 [M] (7.10), 138 (100), ¹H NMR (DMSO-d₆, 300MHz): ¹J 1.62 [s, 3H, CH₃], 3.91 [s, 3H, OCH₃], 5.04 [s, 1H, SH], 6.53-8.91 [m, 11H, Ar-H, =CH], 10.56 [s, 1H, OH]. Anal. Calcd. for C₂₇H₁₉ClN₂O₅S: C, 62.49%; H, 3.69%; N, 5.40%; Found: C, 62.43%; H, 3.62%; N, 5.24%.
4-(2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-5-oxo,4,5-dihydropyrazol-1-yl)-N-(quinoxalin-2-yl)benzenesulfonylamide 8.

Yellow crystals; Yield: 55%; m.p. 170-172°C; IR (KBr, cm⁻¹): 3441 (br, OH), 3360 (NH), 3080 (CH arom.), 2920 (CH aliph.), 1731, 1666 (2C=O), 1310, 1170 (SO₂), MS, m/z (%): 692 [M-2] (38), 139 (100), ¹H NMR (DMSO-d₆, 300MHz): a 1.96 [s, 3H, CH₃], 3.99 [s, 3H, OCH₃], 5.1 [s, 1H, CH-quinoxaline], 6.94-8.91 [m, 11H, Ar-H, =CH], 10.48 [s, 1H, OH], 11.56 [s, 1H, NH=SO₃]. Anal. Calcd. for C₃₅H₂₄ClN₅O₅S: C, 60.56%; H, 3.49%; N, 10.09%; Found: C, 60.48%; H, 3.42%; N, 9.96%.

2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1-(4-hydroxypyrimidin-2-yl)-1H-imidazol-5(4H)-one 9.

Yellow crystals; Yield: 60%; m.p. 200-202°C; IR (KBr, cm⁻¹): 3431 (br, OH), 3009 (CH arom.), 2920 (CH aliph.), 1730, 1649 (2C=O), MS, m/z (%): 505 [M+1] (24), 69 (100), ¹H NMR (DMSO-d₆, 300MHz): a 1.57 [s, 3H, CH₃], 3.95 [s, 3H, OCH₃], 6.26-6.32 [d, 2H, CH-pyrimidine], 6.79-7.91 [m, 7H, Ar-H, =CH], 10.10, 10.56 [2s, 2H, 2OH]. Anal. Calcd. for C₂₅H₁₇ClN₄O₆: C, 59.47%; H, 3.39%; N, 11.10%; Found: C, 59.42%; H, 3.21%; N, 10.96%.

4-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-4-oxo-4H-chromen-6-yl)methylene)-5-oxo,4,5-dihydropyrazol-1-yl)-1,5-dimethyl-2-phenyl-1,2-dihydropyrazol-3-one 10.

Yellow crystals; Yield: 48%; m.p. 210-212°C; IR (KBr, cm⁻¹): 3433 (br, OH), 3023 (CH arom.), 2914 (CH aliph.), 1717, 1643 (2C=O), MS, m/z (%): 598 [M+2] (13.9), 139 (100), ¹H NMR (DMSO-d₆, 300MHz): a 1.14, 1.23 [2s, 6H, 2CH₃], 3.81 [s, 3H, N-CH₃], 3.96 [s, 3H, OCH₃], 6.32-8.96 [m, 12H, Ar-H], 10.56 [s, 1H, OH]. Anal. Calcd. for C₃₂H₂₅ClN₄O₆: C, 64.38%; H, 4.22%; N, 9.38%; Found: C, 64.27%; H, 4.16%; N, 9.29%.

2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1H-imidazol-5(4H)-one 11.

A solution of 4 (0.01 mol.) in formamide (10 ml.) was refluxed for 5h. The solid obtained was crystallized from dioxane to give 11 as black crystals. Yield: 65%; m.p. 280-282°C; IR (KBr, cm⁻¹): 3414 (br OH, NH), 3060 (CH arom.), 2922 (CH aliph.), 1640 (C=O), ¹H NMR (DMSO-d₆, 300MHz): a 1.63 [s, 3H, CH₃], 3.36 [s, 3H, OCH₃], 6.13-7.97 [m, 7H, Ar-H, =CH], 10.02 [s, 1H, OH], 11.19 [s, 1H, NH]. Anal. Calcd. for C₂₁H₁₅ClN₄O₅: C, 61.40%; H, 3.68%; N, 6.82%; Found: C, 61.38%; H, 3.62%; N, 6.76%.

1-amino-2-(2-chlorophenyl)-4-((7-hydroxy-5-methoxy-2-methyl-4-oxo-4H-chromen-6-yl)methylene)-1H-imidazol-5(4H)-one 12.

A mixture of 4 (0.01 mol) and hydrazine hydrate (0.02 mol) in ethanol (20 ml) was refluxed for 3hrs, then allowed to cool. The solid precipitate was collected and recrystallized from the proper solvents to give 12 as yellow crystals. Yield: 60%; m.p. Up to 300°C; IR (KBr, cm⁻¹): 3414, 3328, 3221 (br OH, NH₂), 3060 (CH arom.), 2922 (CH aliph.), 1732, 1640 (C=O), MS, m/z (%): 425 [M] (8.25), 166 (100), ¹H NMR (DMSO-d₆, 300MHz): a 1.9 [s, 3H, CH₃], 3.39 [s, 3H, OCH₃], 5.8 [s, 2H, NH₂], 6.23 [s, 1H, =CH], 7.24-8.32 [m, 8H, Ar-H], 10.98 [s, 1H, OH]. Anal. Calcd. for C₂₁H₁₆ClN₄O₅: C, 59.23%; H, 3.79%; N, 9.87%; Found: C, 59.32%; H, 3.73%; N, 9.46%.

General procedure for Imidazo[5,1-c][1,2,4]triazol-7-ylidene)derivatives 13,14.

To a solution of compound 4 or 5 (0.01mol) in dimethylformamide (20 ml) was added an equimolar amount of thiosemicarbazide (0.01 mol). The reaction mixture was heated under reflux for 8h, left over night, partially concentrated and cooled. The solid products were
filtrated off, washed of water, dried and recrystallized from the proper solvents to give 13, 14 respectively.

**6-((5-(2-chlorophenyl)-3-mercaptop-7H-imidazo[5,1-c][1,2,4]triazol-7-ylidene)methyl)-7-hydroxy-5-methoxy-2-methyl-4H-chromen-4-one 13.**

Brown crystals; Yield: 40%; m.p. 220-222°C; IR (KBr, cm\(^{-1}\)): 3414, 3326, 3286 (br OH, NH), 3074 (CH arom.), 2952 (CH aliph.), 1634 (C=O), 1145 (C=S), MS, m/z (%): 467 [M+1] (28.25), 138 (100) \(^1\)H NMR (DMSO-\(d_6\), 300 MHz): \(\delta\) 1.9 [s, 3H, CH\(_3\)], 3.34 [s, 3H, OCH\(_3\)], 6.03-7.49 [m, 7H, Ar-H, =CH], 10.04 [s, 1H, OH], 11.12 [s, 1H, NH]. Anal. Calcd. for C\(_{22}\)H\(_{15}\)ClN\(_4\)O\(_4\): C, 56.59%; H, 3.24%; N, 12.00%; Found: C, 56.64%; H, 3.46%; N, 12.17%.

**7-hydroxy-5-methoxy-2-methyl-6-((5-methyl-3-thioxo-2,3-dihydroimidazo[5,1-c][1,2,4]triazol-7-ylidene)methyl)-4H-chromen-4-one 14.**

Brown crystals; Yield: 40%; m.p. 198-200°C; IR (KBr, cm\(^{-1}\)): 3415, 3331 (br OH, NH), 3076 (CH arom.), 2972 (CH aliph.), 1654 (C=O), 1567 (C=S), MS, m/z (%): 368 [M-2] (8.25), 166 (100), \(^1\)H NMR (DMSO-\(d_6\), 300 MHz): \(\delta\) 1.27 [s, 3H, CH\(_3\)], 1.48 [s, 3H, CH\(_3\)], 3.71 [s, 3H, OCH\(_3\)], 6.57[d, 1H, =CH], 7.52 [s, 2H, 2CH], 10.45 [s, 1H, OH], 11.06 [s, 1H, NH]. Anal. Calcd. for C\(_{17}\)H\(_{14}\)N\(_4\)O\(_4\)S: C, 55.13%; H, 3.81%; N, 15.13%; Found: C, 55.09%; H, 3.75%; N, 15.03%.

**In-vitro anticancer evaluation**

**Materials and methods**

The human tumor cell lines (MCF7 and HCT) were available at the regional center for mycology & biotechnology at Al-Azhar University, Cairo, Egypt. Mammalian cell lines: MCF7 and HCT cells (human cell line of a well differentiated breast and colon carcinoma isolated from a breast or colon biopsy of a male Caucasian aged 15 years) were obtained from the American Type Culture Collection (ATCC). Chemicals Used: Dimethyl sulfoxide (DMSO), crystal violet and trypan blue dye were purchased from Sigma (St. Louis, Mo., USA). Fetal Bovine serum, DMEM, RPMI-1640, HEPES buffer solution, L-glutamine, gentamycin and 0.25% Trypsin-EDTA were purchased from Lonza. Crystal violet stain (1%): It composed of 0.5% (w/v) crystal violet and 50% methanol then made up to volume with d\(H_2\)O and filtered through a Whatmann No.1 filter paper.

**Procedure**

The cells were propagated in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% heat-inactivated fetal bovine serum, 1% L-glutamine, HEPES buffer and 50 µg/ml genta-mycin. All cells were maintained at 37°C in a humidified atmosphere with 5% CO\(_2\) and were subcultured two times a week. Cell toxicity was monitored by determining the effect of the test samples on cell morphology and cell viability.

**Cytotoxicity evaluation using viability assay** (Mosmann, 1983 and Vijayan, 2004): For cytotoxicity assay, the cells were seeded in 96-well plate at a cell concentration of 1\(\times\)10\(^4\) cells per well in 100µl of growth medium. Fresh medium containing different concentrations of the test sample was added after 24 h of seeding. Serial two-fold dilutions of the tested chemical compound were added to confluent cell monolayers dispensed into 96-well, flat-bottomed microtiter plates (Falcon, NJ, USA) using a multichannel pipette. The microtiter plates were incubated at 37°C in a humidified incubator with 5% CO\(_2\) for a period of 48 h. Three wells were used for each concentration of the test sample. Control cells were incubated without test sample and with or without DMSO. The little percentage of DMSO present in the wells (maximal 0.1%) has no effect on the cell line. After incubation of the cells for 24 h at 37°C, various concentrations of sample (50, 25, 12.5, 6.25, 3.125 & 1.56 µg)
were added, and the incubation was continued for 48 h and viable cells yield was determined by a colorimetric method. In brief, after the end of the incubation period, media were aspirated and the crystal violet solution (1%) was added to each well for at least 30 minutes. The stain was removed and the plates were rinsed using tap water until all excess stain is removed. Glacial acetic acid (30%) was then added to all wells and mixed thoroughly, and then the absorbance of the plates were measured on Microplate reader (TECAN, Inc.), using a test wavelength of 490 nm. All results were corrected for background absorbance detected in wells without added stain. Treated samples were compared with the cell control in the absence of the tested compounds. All experiments were carried out in triplicate. The relation between surviving fraction and drug concentration was plotted to get the survival curve of each tumor cell line. The concentration required for 50% inhibition of cell viability (IC$_{50}$) was calculated and compared with the reference drug vinblastine and the results are given in table 1.

**Table 1.** In-vitro anticancer evaluation against human tumor breast and colon cell lines (MCF7 and HCT).

<table>
<thead>
<tr>
<th>Cpd. No.</th>
<th>IC$_{50}$ $^{a,b}$ (µg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MF7</td>
</tr>
<tr>
<td>2</td>
<td>3.1</td>
</tr>
<tr>
<td>6</td>
<td>14</td>
</tr>
<tr>
<td>7</td>
<td>8.5</td>
</tr>
<tr>
<td>8</td>
<td>10.5</td>
</tr>
<tr>
<td>9</td>
<td>10.9</td>
</tr>
<tr>
<td>10</td>
<td>12</td>
</tr>
<tr>
<td>13</td>
<td>9.2</td>
</tr>
<tr>
<td>14</td>
<td>8.8</td>
</tr>
<tr>
<td>Doxorubicin</td>
<td>......</td>
</tr>
<tr>
<td>Vinblastine</td>
<td>2.97</td>
</tr>
</tbody>
</table>

$^{a}$IC$_{50}$, compound concentration required to inhibit tumor cell proliferation by 50%.
$^{b}$Values are means of three experiments.

**REFERENCE**


تحضير ودراسة خواص الأورام المسرطنة لمشتقات الايميدازول المحتويه علي جزيء الكومارين

نشوى مصطفي صالح نادى

الكيمياء - كلية العلوم (بنات) جامعة الأزهر - القاهرة

تحضير و توصيف مركبات جديدة غير متجانسة الحلقة كمشتقات اييميدازول [1-2-3-4-5-6-7-8] انترازولينيدين (2) عن طريق تكافف 7- هيدروكسي- 2- مييثيل- 4- أووكووكومارين-5- كربالديهد (1) مع مركب المحتوى على مجموعة الميثيلين النشطة و مشتق الأوكسارول-4-إيثيل-1- أون (5) من تفاعلي المشتق (1). مع إثريكوربيزنول جلايسيين أو ميثيل جلايسيين في وجود أسيتات الصوديوم في التوالي. واستخدم المشتق (6-7-8) كمركب أولي في تطبيقات تطبيقات جديدة مختلفة من إعداد مركبات نجدة لجزئي الايميدازول. وتم تحضير مشتقات مختلفة من الإيميدازول (8-10) بتفاعل المشتق (4) مع بعض مركبات الأيمين الهيدرات والترويديس كاريبيدز لينتج مشتقات متمايز ايميدازول و هيدروزاورين و ايميدازول (11) على التوالي. فقد تم في هذا البحث دراسة الترسبي البيناني لكذ تلك المركبات الجديدة باستخدام طرق التحليل الكيفي والكيمياء والكيمياء النوى المانتط. وكذلك أظهرت الدراسة فاعلية المركبات الجديدة كمضادات للأورام السرطانية التي حققت نسبة موت للخلايا السرطانية تمثل إلى الكمية القياسية المستخدمة عند تركزات 50 ميكروجرام / مللي. واظهرت فاعلية المركبات الجديدة تناجع نوع من الخلايا السرطانية الإنسانية وهي خلايا السرطانية الثديية مستخدمة مادة فيلاستين كمستحضر قياسي والخلايا السرطانية للقولون مستخدمة مادة الدوكسيروبسن كمستحضرات قياسية لقياس فاعلية هذه المركبات الجديدة. مما يوضح إمكانية الاستفادة من هذه المركبات في المجال التطبيقي كمضادات للأورام السرطانية.