

Synthesis, Characterize and Antimicrobial Study of New Chalcones and Pyrazole Derivatives from Progesterone

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Abstract: *The aim of this work synthesized a new series of chalcones and pyrazole derivatives from the progesterone, the new steroidal synthesized by the claisen-Schmidt condensation reaction, were react progesterone with different aromatic substituted aldehyde in presence sodium hydroxide afforded chalcones derivatives (1-6). Cyclized the chalcone derivatives by reaction with hydrazine hydrate to afforded pyrazole derivatives (7-12). All synthesized compounds were characterized by measurement melting point, FT-IR spectral, Elemental Analysis and some of them by ¹H-NMR spectral and Antimicrobials activities.*

Keyword: Progesterone, Chalcone, Pyrazole, Antimicrobial, Biological activity.

1. Introduction

Progesterone is one of the most important hormones of the steroidal pregnan [1], and play a major part in female pregnancy[2]. Varying responses to transcription activity monitors it[3] also progesterone brain is derived from the steroidogenic endocrine gland or from local synthesis by neural cells [4]. Progesterone is steroid hormone, a steroid is any of the group of natural or synthetic, fat-soluble, organic compound belonging to the class of the lipids and characterized by molecular are four fused ring to tailing 17 carbon atoms: three six carbon ring and one five carbone ring fused to gather and have three dimensional configuration [5]. Chalcone is important group of the natural product for the different synthesis class, chalcone are easy synthesis by the reacting of two aromatic ring [6]. Chalcone have several biological activities such as anti-inflammatory [7], antimicrobial agent [8], antioxidant [9], antiplatelet [10], and anticancer[11, 12]. These activities are larger attributed due to β unsaturated ketone moiety, chalcone important their ability to act as an intermediate for synthesis activity heterocyclic compound such as pyrazole derivatives in this work of research. pyrazol ring is a prominent structural motif found in numerous pharmaceutically active compound, pyrazol important biological active such anti-inflammatory, postmenopausal and osteoporosis [13]

2. Experimental

The melting points were determined in open capillary tubes on a Gallen Kamp melting point apparatus and were uncorrected. The FT-IR Spectra of prepared derivatives were taken on Shimadzu-2N, FTIR-8400S, Elemental Analysis %, ¹H-NMR Spectra of some prepared derivatives were recorded on a Varian-Mercury 300MHZ Spectrometer, d₆-DMSO use as a solvent in ¹H-NMR Spectra.

General procedure for preparation of chalcones (1-6)

A solution of progesterone (0.01 mol) in 30ml absolute ethanol, (0.02 mol) substituted aromatic aldehyde in presence 40% NaOH, the mixture stirring for 4hrs. at room temperature and left over night, after that acidification with dilute hydrochloric acid and filtration afforded solid product and recrystallization from ethanol. The properties of compounds (1-6) show in the table 1.

General procedure for preparation pyrazole derivatives (7-12)

A mixture of steroidal chalcones (1-6) (0.01 mol) with (0.02 mol)hydrazine hydrate in 30 ml of ethanol was refluxed for 5 hrs., after that cooling the formed solid product was collected by filtration washed with ethanol, dried and crystallization from ethanol afforded pyrazole derivatives (7-12), and recrystallization from ethanol. The properties of compounds (7-12) show in the table 1.

Table 1: Physical and spectral data of synthesized compounds:

NO	Molecular Formula	Yield %	Mp. °C	Element analysis calculate / Found %	FT-IR cm^{-1}
1	C ₃₅ H ₃₈ O ₄	70	220-222	C 80.43/80.40 H 7.33/7.30 O 12.24/12.20	1712-1672 C=O, 1560 C=C, 3401 OH, 2970 C-H aroma., 2897 C-H aliph.
2	C ₃₇ H ₄₂ O ₄	65	202- 204	C 80.69/80.65 H 6.93/6.90 O 11.62 /11.60	1730-11710 C=O, 1570 C=C, 3070 C-H aroma., 2890 C-H aliph.
3	C ₃₅ H ₃₆ Br ₂ O ₂	65	177- 179	C 64.83/64.80 H 5.61/5.59 Br 24.64/24.6, O 4.93/4.90	1701-1689 C=O, 1597 C=C, 3061 C-H aroma., 2941 C-H aliph., 755 C-Br.
4	C ₃₅ H ₃₆ N ₂ O ₆	75	200d	C 72.39/72.35 H 6.25/6.20 N 4.8/4.5 O 16.53/16.5	1747-1676 C=O, 1589 C=C, 3053 C-H aroma., 2856 C-H aliph., 1566 NO ₂ asymm. 1356 NO ₂ symm.
5	C ₃₅ H ₃₈ O ₄	68	198-200	C 80.43/80.40 H 7.33/7.30 O 12.24/12.20	1720-1705 C=O, 1580 C=C, 3410 OH, 3040 C-H aroma., 2889 C-H aliph.
6	C ₃₅ H ₃₆ N ₂ O ₆	70	147- 149	C 72.39/72.35 H 6.25/6.20 N 4.8/4.5, O 16.53/16.5	1710-1665 C=O, 1580 C=C, 3086 C-H aroma., 2939 C-H aliph., 1521 NO ₂ asymm. 1357 NO ₂ symm.
7	C ₃₆ H ₄₀ N ₄ O ₂	60	210- 212	C 77.11/77.1 H 7.19/7.14 N 9.99/9.94 O 5.71 /5.65	3242 NH, 3401 OH, 3073 C-H aroma., 2882 C-H aliph., 1620 C=N, 1587 C=C.
8	C ₃₈ H ₄₄ N ₄ O ₂	60	197-199	C 77.52/77.46 H 7.53/7.50 N 9.52/9.50, O 5, 45 /5.40	3342 NH, 3053 C-H aroma., 2852 C-H aliph., 1618 C=N, 1597 C=C.
9	C ₃₆ H ₃₈ Br ₂ N ₄	65	212-214	C 62.98/62.93 H 5.58/5.50 N 8.16/8.12, Br 23.28 /23.24	3362 NH, 3030 C-H aroma., 2953 C-H aliph., 1639 C=N, 1527 C=C, C-Br 798.
10	C ₃₆ H ₃₈ N ₆ O ₄	70	125-127	C 69.88/69.84 H 6.19/6.15 N 13.58/13.54 O 10.34 /10.30	3296 NH, 3055 C-H aroma., 2993 C-H aliph., 1602 C=N, 1552 C=C, 1535 NO ₂ asymm. 1342 NO ₂ symm.
11	C ₃₆ H ₄₀ N ₄ O ₂	70	235-237	C 77.11/77.1 H 7.19/7.14 N 9.99/9.94 O 5.71 /5.65	3254 NH, 3410 OH, 3053 C-H aroma., 2982 C-H aliph., 1616 C=N, 1580 C=C.
12	C ₃₆ H ₃₈ N ₆ O ₄	65	116-118	C 69.88/69.84 H 6.19/6.15 N 13.58/13.54 O 10.34 /10.30	3304 NH, 3076 C-H aroma., 2939 C-H aliph., 1616 C=N, 1593 C=C, 1514 NO ₂ asymm. 1329 NO ₂ symm.

Table 2: Chemical Schiff's ¹H-NMR Spectra of compounds (7-10):

NO	¹ H-NMR (DMSO-d ₆) δ ppm
7	11.2(s, 1H, NHpyrazole), 9.2(s, 1H, OH), 6.4(s, 1H, CH=ethylen), 1.2(t, 2H, CH ₂ -CH ₂ cyclopentane), 2.6(d, 2H, CH-CH ₂ methylen), 2.1(s, 1H, CH cyclo hexane,), 1.1(s, 3H, CH ₃ methyl), 8.0-8.5(m, CH protons aromatic)
8	10.3(s, 1H, NHpyrazole), 3.9(s, 3H, OCH ₃), 6.7(s, 1H, CH=ethylen), 1.4(t, 2H, CH ₂ -CH ₂ cyclopentane), 2.6(d, 2H, CH-CH ₂ methylen), 2.1(s, 1H, CH cyclo hexane), 1.46(s, 3H, CH ₃ methyl), 7.0-7.8(m, CH protons aromatic)
9	11.1(s, 1H, NHpyrazole), 6.5(s, 1H, CH= ethylen), 1.7(t, 2H, CH ₂ -CH ₂ cyclopentane), 2.5(d, 2H, CH-CH ₂ methylen), 2.1(s, 1H, CH cyclo hexane), 1.04(s, 3H, CH ₃ methyl), 7.2-7.8(m, CH protons aromatic)
10	10.3(s, 1H, NHpyrazole), 6.9(s, 1H, CH=C), 1.2(t, 2H, CH ₂ -CH ₂ cyclopentane), 2.6(d, 2H, CH-CH ₂ methylen), 2.2(s, 1H, CH cyclo hexane), 1.04(s, 3H, CH ₃ methyl), 7.9-8.1(m, CH protons aromatic)

4. Biological Activity

The antimicrobial screening data show that the compounds exhibit antimicrobial properties and it is important to note that new derivatives exhibit more inhibitory effect than original molecule clear that zone of inhibition against the gram positive, gram negative and antifungal act as more powerful. The newly synthesized steroidal chalcones and pyrazole derivatives were tested for Staphylococcus aureus and Streptococcus pneumonia as gram positive and Pseudomonas aeruginosa and E.Coli as gram negative, the antifungal select the albicans and A. niger, the solvent use DMSO as a negative control. The gar disc diffusion method was followed in this screening and the Ampicillin was used as standard drug.

Some of the tested show no significant effect against the used antimicrobial, but the compound have electron-withdrawing group show good activity against and some of derivatives show moderate activity against. The result shows in the table 3.

Table 3: Antimicrobial activity of Synthesized compounds

Comp.(100ng/ml)	Zone of Inhibition (mm)					
	Gram positive		Gram negative		Anti Fungal	
	Streptococcus pneumonia	Staphylococcus aureus	Pseudomonas aeruginosa	E.Coli	C. albicans	A. niger
1	19	20	19	20	22	17
2	---	14	24	17	20	---
3	19	---	15	15	21	---
4	20	24	20	20	18	21
5	30	24	18	---	20	---
6	20	19	---	20	19	17
7	18	22	20	15	---	18
8	---	14	15	18	12	---
9	--	---	--	24	---	20
10	20	24	18	15	20	19
11	17	19	15	19	15	18
12	19	20	20	20	19	20
Ampicillin (100ng/ml)	20	22	20	23	21	20
Solvent control DMSO	---	---	---	---	---	---

5. Conclusions

In this my work described in this review indicated the synthesis new chalcone and pyrazole derivatives and the resulted in the produced with high Antibacterial and antifungal activity. Thus research also can act as important tool for medical chemists to develop new steroidal compounds possessing heterocyclic moiety that could be better agents in term efficiency and safety.

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