

Synthesis of New Oxazepine and Thiazolidine Compounds derived from Pyrimidine-2(1H)-one

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Abstract

In this paper the compounds (11-20) (methyl pyrimidine - 2(1H)-one and others phenyl pyrimidine -2(1H)- one) that had been already prepared from α , β unsaturated carbonyl compounds that are called chalcones, these compounds are usually prepared from the reaction of different aldehydes (4-methoxy benzaldehyde, 2-nitro benzaldehyde, 3-nitrobenzaldehyde, 4-N,N—dimethyle amino benzaldehyde, benzaldehyde, 4-nitro benzaldehyde) with different ketones (acetone, acetophenone, 2-nitro acetophenone, 3-nitro acetophenone) After preparing and purifying them a suitable mesurment for physical, chemical and spectroscopic properties had been made to get chalcones that reacted with urea under known chemical conditions to get the pyrimidinone compounds that we need. Pyrimidine compounds had been reacted with two aromatic amines (2,4-dinitro aniline and 4- amino acetophenone) using glacial acetic acid as catalyst in absolute ethanol giving a new compounds of schiff's bases (21-40). New thiazolidine 4-one (41-50) had been synthesized from the reacting of Schiff's bases (21-30) with thioglycolic acid in absolute ethanol. 1,3-oxazepine derivatives. (51-60) had been prepared from reaction between Schiff's bases (31-40) and malic anhydrid in absolut ethanol. The structures of the Synthesized compounds had been estimated by IR, ¹H- NMR and some physical data.

Keywords: pyrimidine, Schiff's bases, Thiazolidine, 1,3- oxazepine

تحضير مركبات اوكسازيبين وثيازولدين جديدة مشتقة من مركبات بيريميدين-2(1H)- اون

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الخلاصة

في هذا البحث تم تحضير المركبات (11-20) (مثيل بيريميدين-2(1H)- اون ومركبات اخرى من فنيل بيريميدين-2(1H) - اون والتي تم تحضيرها مسبقا من مركبات كاربونيل α و β غير المشبعة والتي يطلق عليها اسم الجالكونات (1-10) والتي تحضر من مفاعلة الديهايدات مختلفة (4- ميثوكسي بنزالديهايد، 2- نايتروبنزالديهايد، 3- نايتروبنزالديهايد، 4- نايتروبنزالديهايد، N,N-ثنائي مثل امينو بنزالديهايد، بنزالديهايد) مع كيتونات مختلفة (اسيتون، اسيتوفينون، 2-

نايترواسيتوفينون , 3- نايترواسيتوفينون) وبعد تحضير هذه المركبات وتثبيتها واجراء قياسات لخواصها الفيزيائية والكيميائية والطيفية التي تمكنا من اجراءها للحصول على الجالكونات والتي تم مفاعلها مع اليوريا وحسب الظروف الكيميائية المعروفة للحصول على مركبات البيريميدينون المطلوبة. مركبات البيريميدين تم مفاعلها مع الامينات الاروماتية (2،4- ثنائي نايتروانيلين و 4- امينو اسيتوفينون) باستخدام حامض الخليك الثلجي كحفاز بوجود الايثانول المطلق ليعطي مركبات جديدة من قواعد شيف (40-21). مركبات جديدة من ثايازولدين-4- أون (50-41) تم تحضيرها من مفاعلة قواعد شيف (30-21) مع حامض الثايوكلايكوليك في الايثانول المطلق. حضرت مشتقات 1،3- اوكسازيبين (60-51) من مفاعلة قواعد شيف (40-31) مع ماليك انهيدريد في الايثانول المطلق. تم تشخيص المركبات المحضرة باستخدام طيف الاشعة تحت الحمراء وطيف الرنين النووي المغناطيسي وبعض الخواص الفيزيائية.

الكلمات المفتاحية: بيريميدين، قواعد شيف، ثايازولدين، اوكسازيبين.

INTRODUCTION

Pyrimidines were used as substrates in the preparation of many schiff's bases which undergo ring closure (cyclo addition and replacement reaction) [1] to prepare thia- zolidine derivatives which have various pharmacological activities such as antibacterial [2], anti-fungal [3], anti-Cancer [4], anti con-vulsant [5] and herbicidal actions [6]. 1,3-oxazipene derivatives also Synthesized from the reaction of Schiff's bases with maleic anhydrid these compounds have been found to exhibit biological activites including anti-bacterial, antifungal [7,8], anticancer [9].

EXPERIMENTAL

All reagents and compounds are from Fluka and BDH. Melting points are measured using: Electro thermal IA 9100 melting points apparatus type (not corrected). FT-IR Spectra were recorded on BRUKER ALPHLA FTIR Germany. ¹H-NMR spectra were recorded by NMREADY 60 PRO. Using DMSO-d6 as a solvent in college of education and science Ibn al-haitham in Baghdad/Iraq.

Synthesis of chalcones (1-10) [10,11]

0.1 mole of benzaldehyde or one of its compensations dissolved in 10 ml of EtOH was added to 0.1 mole of acetone or acetophenone or one of its compensations that had been also dissolved in 10 ml of EtOH ,the mixture was stirred in aqueous alkali (40% NaOH) in room temprature for 5 hours. Table (1) involves physical properties as reported in the literature[12]

Synthesis of Pyrimidine-2(1H)-one (1-10) [13]:

(0.01) mole of chalcones (1-10) had been dissolved in EtOH abs. This was added to (0.01) mole of urea which dissolved in 20 ml of EtONa then reflux the mixture for 12 hours. The solvent had been concentrated and poured into ice cold water with constant stirring then it was neutralize with acid Solid produce filtered and washed with water. Table (2) involves the physical properties.

Synthesis of Schiff's Bases (21-40) [14]:

(0.01) mole of pyrimidines compounds (11-20) was added to (0.01) mole of aromatic amine (2,4-dinitro aniline , 4- amino acetophenon in abs. EtOH (25) ml with some drops of glacial acetic acid is had been stirred for 4 hours . Table (3,4) involves the physical properties.

Synthesis of Thiazolidine Derivatives (41-50) [15-17]:

(0.001) mole of Schiff's bases (21-30) in abs. EtOH (15 ml) was added to (0.001) mole of thioglycolic acid, this mixture was refluxed for 5 hours, and treated with potassium bicarbonate to produce a compound, then product filtered. Table (5) includes the physical properties.

Synthesis of 1,3- Oxazepine Derivatives (51-60)[18]:

Mixture of Schiff's bases (31-40) (0.0004) mole was added to (0.0006 mole) of maleic anhydride in (20) ml of abs. Ethanol was refluxed for 6 hours. The product was filtered and washed with water. Table (6) involves the physical properties

Table (1)The Physical Properties of Compound (1-10)

Comp. No.	R	R ₁	Molecular formula	M. Wt	m.p ^o c	Colour	Yield %
1	4-OCH ₃	CH ₃	C ₁₁ H ₁₂ O ₂	176	66-68	Yellow	89
2	4-NO ₂	C ₆ H ₅	C ₁₅ H ₁₁ NO ₃	253	151-153	Light Brown	81
3	3-NO ₂	C ₆ H ₅	C ₁₅ H ₁₁ NO ₃	253	129-131	Yellow	64
4	4-N(CH ₃) ₂	C ₆ H ₅	C ₁₇ H ₁₇ NO	251	63-65	Orange	65
5	H	3-NO ₂ C ₆ H ₄	C ₁₅ H ₁₁ NO ₃	253	118-120	Brown	71
6	2-NO ₂	C ₆ H ₅	C ₁₅ H ₁₁ NO ₃	253	131-134	Yellow	75
7	H	C ₆ H ₅	C ₁₅ H ₁₂ O	208	123-126	Yellow	88
8	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₁₅ H ₁₀ N ₂ O ₅	294	177-179	Brown	81
9	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₁₅ H ₁₀ N ₂ O ₅	294	181-184	Dark Brown	73
10	H	2-NO ₂ C ₆ H ₄	C ₁₅ H ₁₁ NO ₃	253	84-86	Brown	63

Table (2)The Physical Properties of Compound (11-20)

Comp. No.	R	R ₁	Molecular formula	M. Wt	m.p. °c	Colour	Yield %
11	4-OCH ₃	CH ₃	C ₁₂ H ₁₂ N ₂ O ₂	216	253-255	Creamy	72
12	4-NO ₂	C ₆ H ₅	C ₁₆ H ₁₁ N ₃ O ₃	293	336-339	Dark Brown	63
13	3-NO ₂	C ₆ H ₅	C ₁₆ H ₁₁ N ₃ O ₃	293	297-299	Pale Brown	77
14	4-N(CH ₃) ₂	C ₆ H ₅	C ₁₈ H ₁₇ N ₃ O	291	151-154	Pale Brown	60
15	H	3-NO ₂ C ₆ H ₄	C ₁₆ H ₁₁ N ₃ O ₃	293	263-265	Dark Brown	86
16	2-NO ₂	C ₆ H ₅	C ₁₆ H ₁₁ N ₃ O ₃	293	285-288	Brown	81
17	H	C ₆ H ₅	C ₁₆ H ₁₂ N ₂ O	248	222-225	Yellow	85
18	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₁₆ H ₁₀ N ₄ O ₅	338	285-289	Dark Brown	76
19	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₁₆ H ₁₀ N ₄ O ₅	338	315-318	Dark Brown	70
20	H	2-NO ₂ C ₆ H ₄	C ₁₆ H ₁₁ N ₃ O ₃	293	253-255	Brown	80

Table (3)The Physical Properties of Compound (21-30)

Comp. No.	R	R ₁	Molecular formula	M. Wt	m.p °c	Colour	Yield %
21	4-OCH ₃	CH ₃	C ₂₄ H ₁₇ N ₅ O ₅	329	101-104	Light green	90
22	4-NO ₂	C ₆ H ₅	C ₂₀ H ₁₅ NO ₂	410	68-70	Brown	81
23	3-NO ₂	C ₆ H ₅	C ₂₄ H ₁₄ N ₄ O ₃	410	68-71	Dark Brown	76
24	4-N(CH ₃) ₂	C ₆ H ₅	C ₂₄ H ₁₈ N ₄ O ₃	407	62-65	Brown	73
25	H	3-NO ₂ C ₆ H ₄	C ₂₆ H ₂₃ N ₄ O	410	82-85	Dark Brown	63
26	2-NO ₂	C ₆ H ₅	C ₂₄ H ₁₈ N ₄ O ₃	410	277-280	Brown	77
27	H	C ₆ H ₅	C ₂₄ H ₁₈ N ₄ O	365	162-165	Yellow	88
28	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₂₄ H ₁₈ N ₄ O ₃	455	88-90	Brown	78
29	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₂₄ H ₁₈ N ₄ O ₃	455	95-96	Light orange	59
30	H	2-NO ₂ C ₆ H ₄	C ₂₄ H ₁₉ N ₃ O	410	75-78	Dark Yellow	91

Table (4) The Physical Properties of Compound (31-40)

Comp. No.	R	R ₁	Molecular formula	M. Wt	m.p ^o c	Colour	Yield %
31	4-OCH ₃	CH ₃	C ₁₈ H ₁₂ N ₅ O ₅	378	118-120	Light Yellow	85
32	4-NO ₂	C ₆ H ₅	C ₂₂ H ₁₄ N ₆ O ₆	458	272-275	Light Brown	67
33	3-NO ₂	C ₆ H ₅	C ₂₂ H ₁₄ N ₆ O ₆	458	171-174	Pale Brown	65
34	4-N(CH ₃) ₂	C ₆ H ₅	C ₂₂ H ₁₂ N ₆ O ₆	456	148-151	Light green	89
35	H	3-NO ₂ C ₆ H ₄	C ₂₂ H ₁₄ N ₆ O ₆	458	168-170	Light green	63
36	2-NO ₂	C ₆ H ₅	C ₂₂ H ₁₄ N ₆ O ₆	458	167-170	Dark Yellow	72
37	H	C ₆ H ₅	C ₂₂ H ₁₅ N ₅ O ₄	413	169-172	Yellow	82
38	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₂₂ H ₁₃ N ₇ O ₈	503	206-209	Dark Brown	75
39	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₂₂ H ₁₃ N ₇ O ₈	503	217-220	Dark Brown	81
40	H	2-NO ₂ C ₆ H ₄	C ₂₂ H ₁₄ N ₆ O ₆	458	159-161	Brown	66

Table (5) The Physical Properties of Compound (41-50)

Comp. No.	R	R ₁	Molecular formula	M. Wt	mp ^o c	Colour	Yield %
41	4-OCH ₃	CH ₃	C ₂₂ H ₁₇ N ₃ O ₃ S	403	180-183	Creamy	79
42	4-NO ₂	C ₆ H ₅	C ₂₆ H ₂₀ N ₄ O ₄ S	484	284-286	Light Brown	81
43	3-NO ₂	C ₆ H ₅	C ₂₆ H ₂₀ N ₄ O ₄ S	484	275-278	Brown	78
44	4-N(CH ₃) ₂	C ₆ H ₅	C ₂₈ H ₂₅ N ₄ O ₂ S	481	273-275	Yellow	85
45	H	3-NO ₂ C ₆ H ₄	C ₂₆ H ₂₀ N ₄ O ₄ S	484	103-105	Light green	81
46	2-NO ₂	C ₆ H ₅	C ₂₆ H ₂₀ N ₄ O ₄ S	484	245-248	Dark Brown	80
47	H	C ₆ H ₅	C ₂₆ H ₂₁ N ₃ O ₂ S	439	76-78	Yellow	88
48	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₂₆ H ₂₀ N ₄ O ₄ S	484	88-90	Brown	66
49	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₂₆ H ₂₀ N ₄ O ₄ S	529	76-78	Yellow	91
50	H	2-NO ₂ C ₆ H ₄	C ₂₆ H ₂₀ N ₄ O ₄ S	484	95-98	Dark Brown	73

Table (6) The Physical Properties of Compound (51-60)

Comp. No.	R	R ₁	Molecular formula	M. Wt	mp °c	Colour	Yield %
51	4-OCH ₃	CH ₃	C ₂₀ H ₁₇ N ₅ O ₈	483	95-98	Light Yellow	65
52	4-NO ₂	C ₆ H ₅	C ₂₄ H ₁₆ N ₆ O ₉	532	111-114	Brown	55
53	3-NO ₂	C ₆ H ₅	C ₂₄ H ₁₆ N ₆ O ₉	532	333-335	Pale Brown	71
54	4-N(CH ₃) ₂	C ₆ H ₅	C ₂₆ H ₂₂ N ₆ O ₇	530	85-87	Red	83
55	H	3-NO ₂ C ₆ H ₄	C ₂₄ H ₁₆ N ₆ O ₉	532	148-151	Light green	67
56	2-NO ₂	C ₆ H ₅	C ₂₄ H ₁₆ N ₆ O ₉	532	327-330	Dark Brown	74
57	H	C ₆ H ₅	C ₂₄ H ₁₇ N ₅ O ₇	487	226-229	Yellow	81
58	4-NO ₂	3-NO ₂ C ₆ H ₄	C ₂₄ H ₁₅ N ₅ O ₁₁	577	267-269	Brown	85
59	4-NO ₂	2-NO ₂ C ₆ H ₄	C ₂₄ H ₁₅ N ₅ O ₁₁	577	287-290	Dark Brown	87
60	H	2-NO ₂ C ₆ H ₄	C ₂₄ H ₁₆ N ₆ O ₉	532	181-184	Light Brown	77

RESULTS AND DISCUSSIONS

New schiff's bases were synthesized by reactions of pyrimidines compounds with different aromatic amine in abs. EtOH with some drops of glacial acetic acid. The FTIR spectra of Schiff's bases compounds (21-40) showed the absence of peak of carbonyl but a new peak was appeared at (1588-1656)cm⁻¹ which is attributed to a new (C=N) group as in [19]. Table (7,8) showed some spectral data. Figure (1), (3) showed IR spectrum for compounds (29, 35).

Thiazolidine compounds were prepared from the reaction of Schiff's bases (21-30) with thioglycolic acid in abs. EtOH. FTIR spectrum showed new peak appeared at (669-776) which is attributed to a new (C-S) group and another peak appeared at (1555-1643) cm⁻¹ is attributed to (C=N) group as in [20], Table (9) referred to some spectral data. Figure (5), showed IR spectrum for compound (49). 1,3- Oxazepine compound prepared from the reaction of Schiff's bases (31-40) with maleic anhydride in abs. EtOH. FT-IR spectrum showed a new peak at (1684-1711) cm⁻¹ that is attributed to (O-C) Lactane and another peak appeared at (1620_1654) cm⁻¹ attributed to (N - C=O) Lactane. Table (9) showed some spectral data [18]. Figure (7) showed IR spectrum for compound (54).

The ¹H-NMR spectrum showed these bands:

Compound (31); ¹H-NMR (DMSO, 60MHz): δ = 7.18 - 8.77(m, 7H, Ar-H), 7.02 (s,C=C-H,1H), 5.39 (b, N-H,1H),1.59(s,O-CH₃), 1.1 (s , -CH₃, 3H). [19].

Figure (2) showed NMR spectrum of compound (31).

Compound (29): ¹H-NMR (DMSO,60MHz): δ= 6.43-7.68 (dd, Ar-H), 5.94 (m,5H,Ar-H), 2.4 (b,1H, N-H).

Figure (2) showed NMR spectrum of compound (29).

Compound (49):¹H-NMR (DMSO,60MHz): δ= 6.43-7.68 (dd,8H,Ar-H), 5.95 (m,5H,Ar-H,C=C-H), 2.83 (b,¹H,N-H), 2.4-2.67(b,5H,CH₃,CH₂),[20].

Figure (6) showed NMR spectrum of compound (49).

Compound (54):¹HNMR (DMSO,60MHz): δ= 6.99-8.77(m,12H, Ar-H), 6.14-6.30 (d,3H,C=C-H), 3.01(s,1H, N-H), 1.16(s,3H,CH₃),

Figure (8) showed NMR spectrum of compound (54).

Table (7) Some Spectral Data of Compounds (21-30)

Comp. No	R	R ₁	N-H	C-H Ar	C-H Alph.	C=O	C=N	Others
21	4-OCH ₃	CH ₃	3387	3097	2851	1700	1638	1551 C=C 1170 C-O-C Asy 1021 C-O-C Sym
22	4-NO ₂	C ₆ H ₅	3320	3115	2905	1684	1642	1542 NO ₂ Asy 1351 NO ₂ Sym 1580 C=C
23	3-NO ₂	C ₆ H ₅	3352	3052	2928	1710	1655	1485 NO ₂ Asy 1351 NO ₂ Asy 1588 C=C
24	4-N(CH ₃) ₂	C ₆ H ₅	3216	3033	2917	1675	1588	1520 C=C
25	H	3-NO ₂ C ₆ H ₄	3218	3081	2908	1685	1639	1553 NO ₂ Asy 1356 NO ₂ Sym 1578 C=C
26	2-NO ₂	C ₆ H ₅	3219	3058	2968	1730	1655	1588 C=C 1526 NO ₂ Asy 1345 NO ₂ Sym
27	H	C ₆ H ₅	3353	3105	2999	1725	1656	1599 C=C
28	4-NO ₂	3-NO ₂ C ₆ H ₄	3343	3110	2942	1686	1637	1578 C=C 1524 NO ₂ Asy 1404 NO ₂ Sym
29	4-NO ₂	2-NO ₂ C ₆ H ₄	3214	3018	2854	1718	1640	1551 C=C 1510 NO ₂ Asy 1354 NO ₂ Sym
30	H	2-NO ₂ C ₆ H ₄	3363	3031	2922	1686	1642	1588 C=C 1524 NO ₂ Asy 1406 NO ₂ Sym

Table (8) Some Spectral Data of Compounds (31-40)

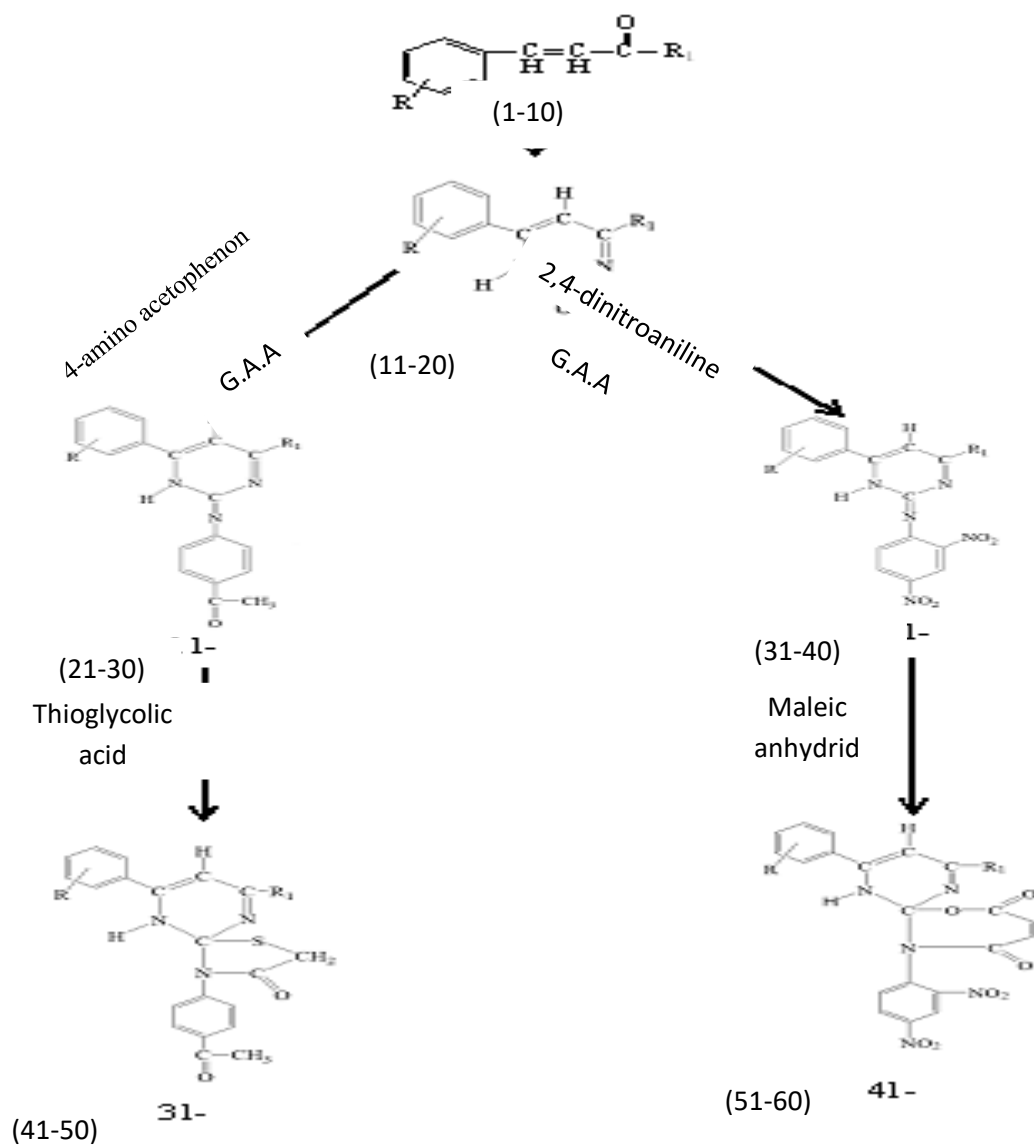
Comp. No.	R	R ₁	N-H	C-H Ar	C-H alph.	C=N	Others
31	4-OCH ₃	CH ₃	3329	3158	2985	1630	1403 COC Asy 1251 C-O-C Sym 1550 NO ₂ Asy 1333 NO ₂ Sym
32	4-NO ₂	C ₆ H ₅	3353	3090	2850	1640	1526 NO ₂ Asy 1326 NO ₂ Sym
33	3-NO ₂	C ₆ H ₅	3325	3099	2845	1620	1504 NO ₂ Asy 1326 NO ₂ Sym
34	4-N(CH ₃) ₂	C ₆ H ₅	3326	3096	2902	1592	1504 NO ₂ Asy 1379 NO ₂ Sym
35	H	3-NO ₂ C ₆ H ₄	3324	3099	2935	1618	1490 NO ₂ Asy 1325 NO ₂ Sym
36	2-NO ₂	C ₆ H ₅	3326	3125	2850	1605	1556 NO ₂ Asy 1328 NO ₂ Sym
37	H	C ₆ H ₅	3175	3072	2833	1623	1507 NO ₂ Asy 1330 NO ₂ Sym
38	4-NO ₂	3-NO ₂ C ₆ H ₄	3331	3166	2878	1627	1555 NO ₂ Asy 1328 NO ₂ Sym
39	4-NO ₂	2-NO ₂ C ₆ H ₄	3326	3024	2940	1633	1552 NO ₂ Asy 1326 NO ₂ Sym
40	H	2-NO ₂ C ₆ H ₄	3326	3043	2953	1621	1504 NO ₂ Asy 1324 NO ₂ Sym

Table (9) Some Spectral Data of Compounds (41- 50)

Comp. No	R	R ₁	N-H	C-H Ar	C-H Alph.	C=O	C=N	C=C	C=S	Others
41	4-OCH ₃	CH ₃	3200	3030	2850	1684	1578	1460	673	1109 cocAsy 1027coc Sym
42	4-NO ₂	C ₆ H ₅	3268	3015	2919	1640	1574	1490	676	1498 NO ₂ Asy 1340 NO ₂ Sym
43	3-NO ₂	C ₆ H ₅	3383	3075	2980	1655	1595	1479	665	1488 NO ₂ Asy 1354 NO ₂ Sym
44	4-N(CH ₃) ₂	C ₆ H ₅	3216	3091	2981	1684	1573	1488	673	-----
45	H	3-NO ₂ C ₆ H ₄	3215	3129	2910	1702	1555	1468	686	1504 NO ₂ Asy 1378 NO ₂ Sym
46	2-NO ₂	C ₆ H ₅	3442	3109	2920	1685	1576	1458	776	1374 NO ₂ Sym 1203 NO ₂ Sym
47	H	C ₆ H ₅	3374	3051	2950	1700	1610	1578	669	-----
48	4-NO ₂	3-NO ₂ C ₆ H ₄	3264	3119	2870	1675	1633	1569	672	1389 NO ₂ Sym 1212 NO ₂ Sym
49	4-NO ₂	2-NO ₂ C ₆ H ₄	3319	3080	2830	1695	1641	1578	669	1425 NO ₂ Sym 1357 NO ₂ Sym
50	H	2-NO ₂ C ₆ H ₄	3374	3120	2920	1685	1643	1569	672	1389 NO ₂ Sym 1222 NO ₂ Sym

Table (10) Some Spectral Data of Compounds (41-50)

Comp. No.	R	R ₁	N-H	C-H Ar	C-H Alph.	$\overset{\text{O}}{\parallel}$ -C- Lacton	$\overset{\text{O}}{\parallel}$ -C- Lactam	C=N	Others
51	4-OCH ₃	CH ₃	3328	3055	2930	1703	1621	1579	1501 NO ₂ Asy 1327 NO ₂ Sym 1418 C=C Ar 1170 C-O-C Asy 1064 C-O-C Sym
52	4-NO ₂	C ₆ H ₅	3285	3045	2855	1695	1630	1559	1504 NO ₂ Asy 1328 NO ₂ Sym 1416 C=C Ar
53	3-NO ₂	C ₆ H ₅	3351	3061	2830	1695	1644	1605	1509 NO ₂ Asy 1324 NO ₂ Sym 1416 C=C Ar
54	4-N(CH ₃) ₂	C ₆ H ₅	3326	3021	2918	1711	1620	1576	1506 NO ₂ Asy 1325 NO ₂ Sym 1427 C=C Ar
55	H	3-NO ₂ C ₆ H ₄	3327	3103	2930	1712	1620	1578	1501 NO ₂ Asy 1325 NO ₂ Sym 1422 C=C Ar
56	2-NO ₂	C ₆ H ₅	3251	3056	2969	1708	1654	1587	1525 NO ₂ Asy 1345 NO ₂ Sym 1457 C=C Ar
57	H	C ₆ H ₅	3316	3174	2908	1700	1653	1570	1507 NO ₂ Asy 1301 NO ₂ Sym 1457 C=C AR
58	4-NO ₂	3-NO ₂ C ₆ H ₄	3354	3080	2968	1684	1635	1552	1524 NO ₂ Asy 1388 NO ₂ Sym 1461 C=C Ar
59	4-NO ₂	2-NO ₂ C ₆ H ₄	3293	3085	2918	1690	1640	1553	1510 NO ₂ Asy 1377 NO ₂ Sym 1459 C=C Ar
60	H	2-NO ₂ C ₆ H ₄	3281	3078	2855	1688	1651	1563	1501 NO ₂ Asy 1328 NO ₂ Sym 1458 C=C Ar



R= 4-OCH₂, 4-NO₂, 4-N(CH₃)₂, 2-NO₂, H

R₁= CH₃, 3-NO₂C₆H₅, 2-NO₂C₆H₅

Scheme (1) illustrates the prepared compounds (1-60)

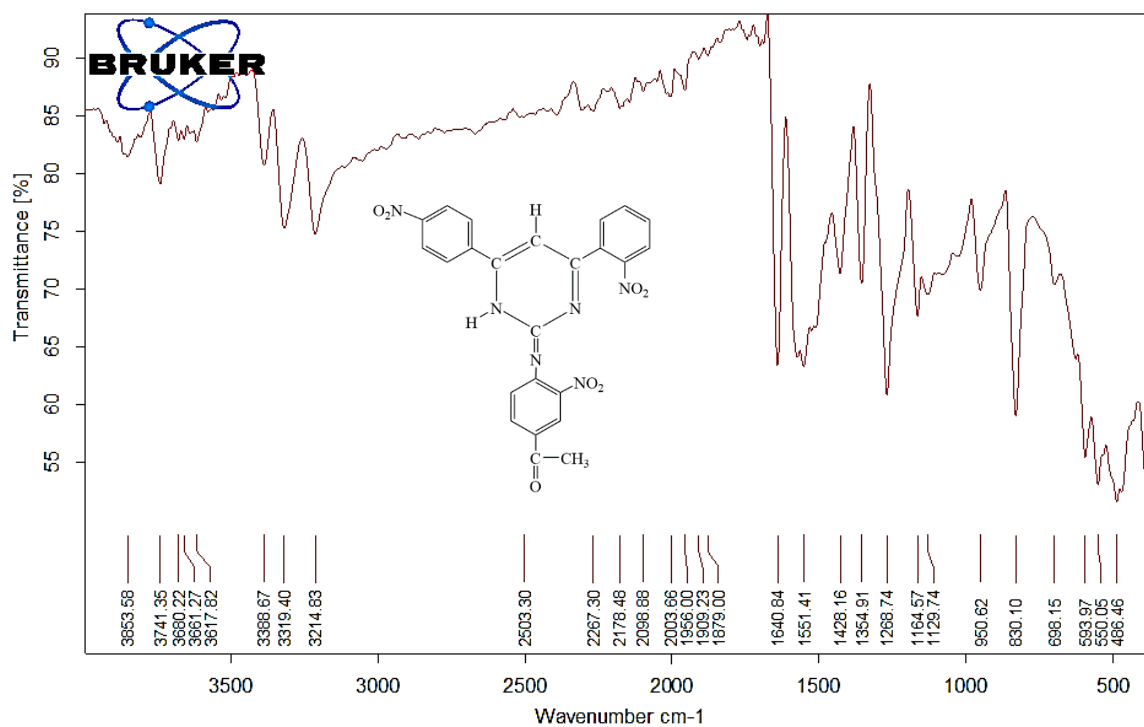


Fig (1): IR spectrum of the compound (29)

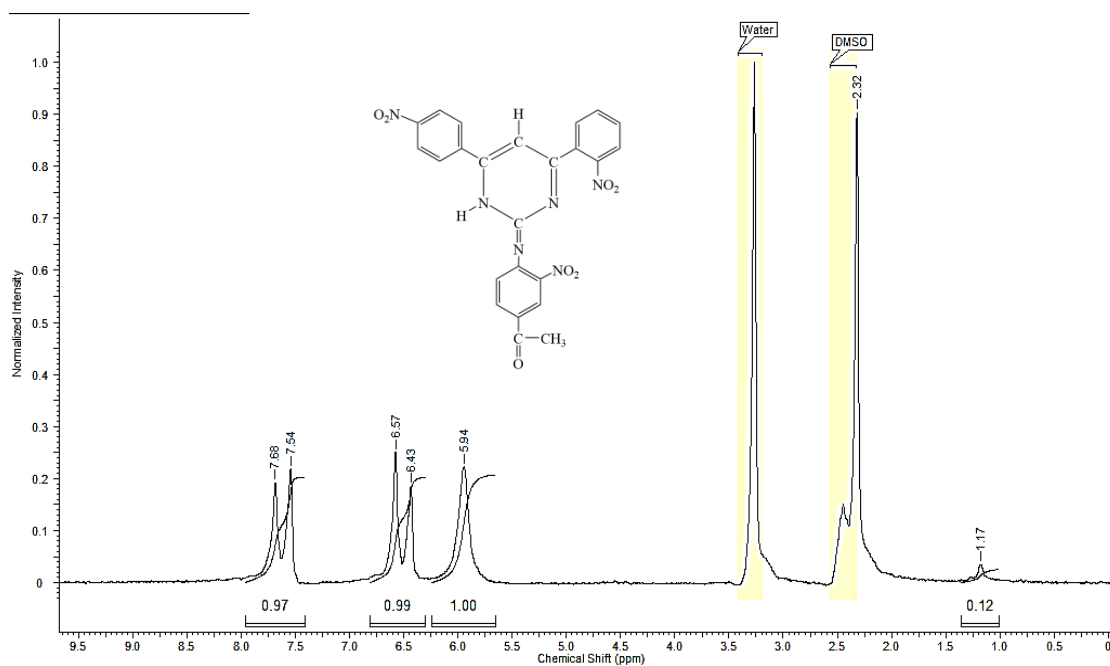


Fig (2): NMR spectrum of the compound (29)

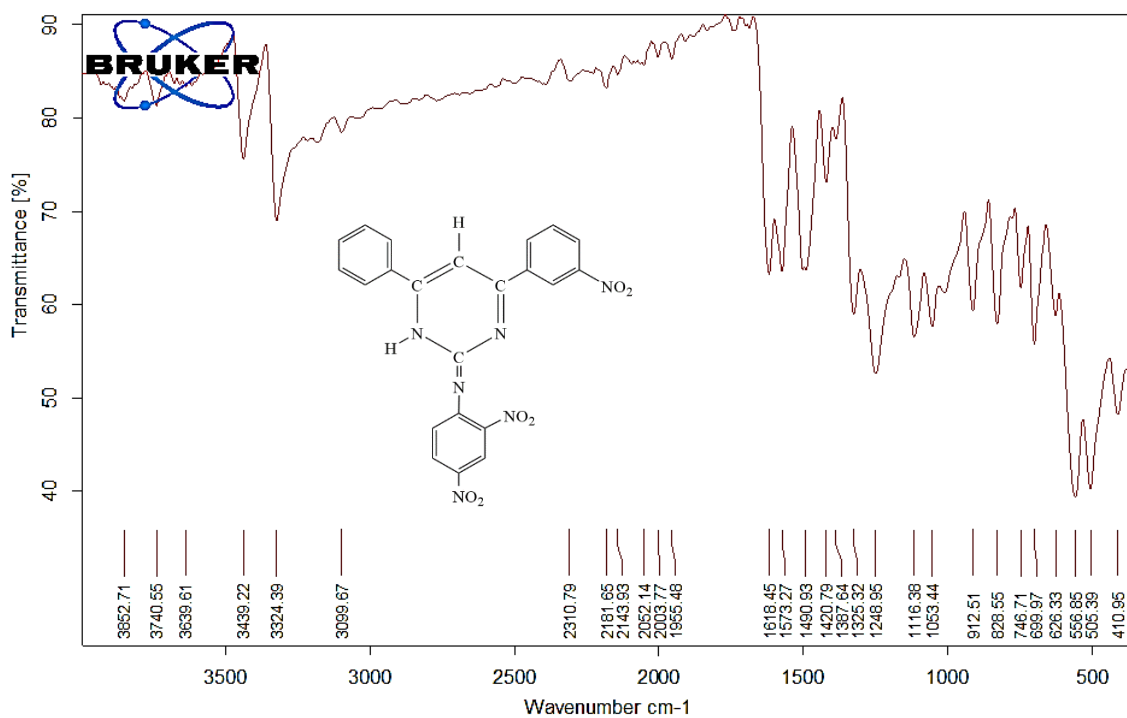


Fig (4): IR spectrum of the compound (35)

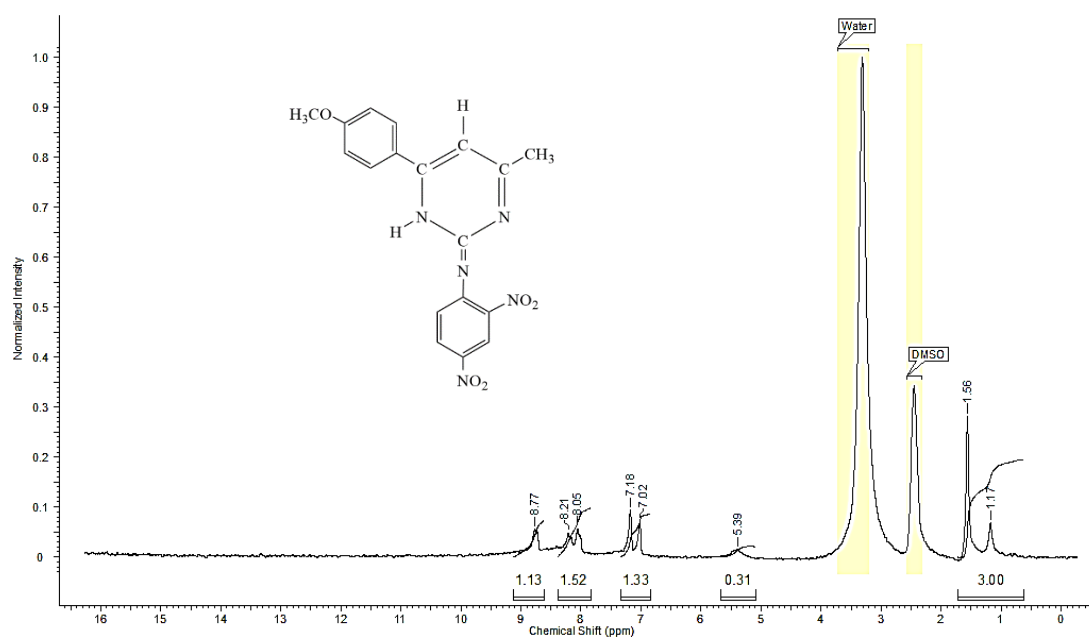


Fig (4): NMR spectrum of the compound (31)

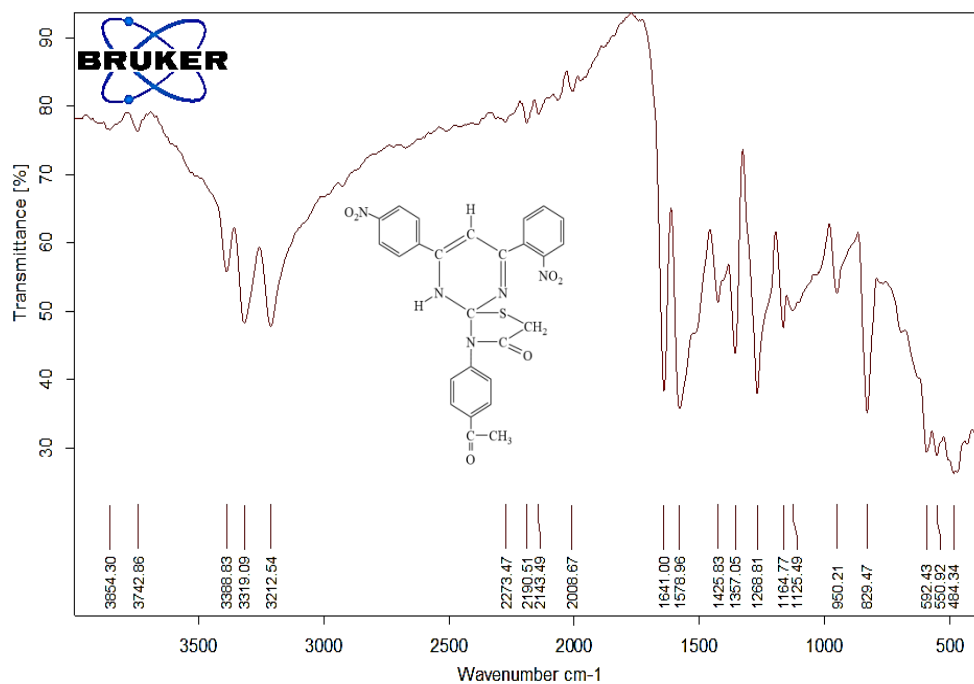


Fig (5): IR spectrum of the compound (49)

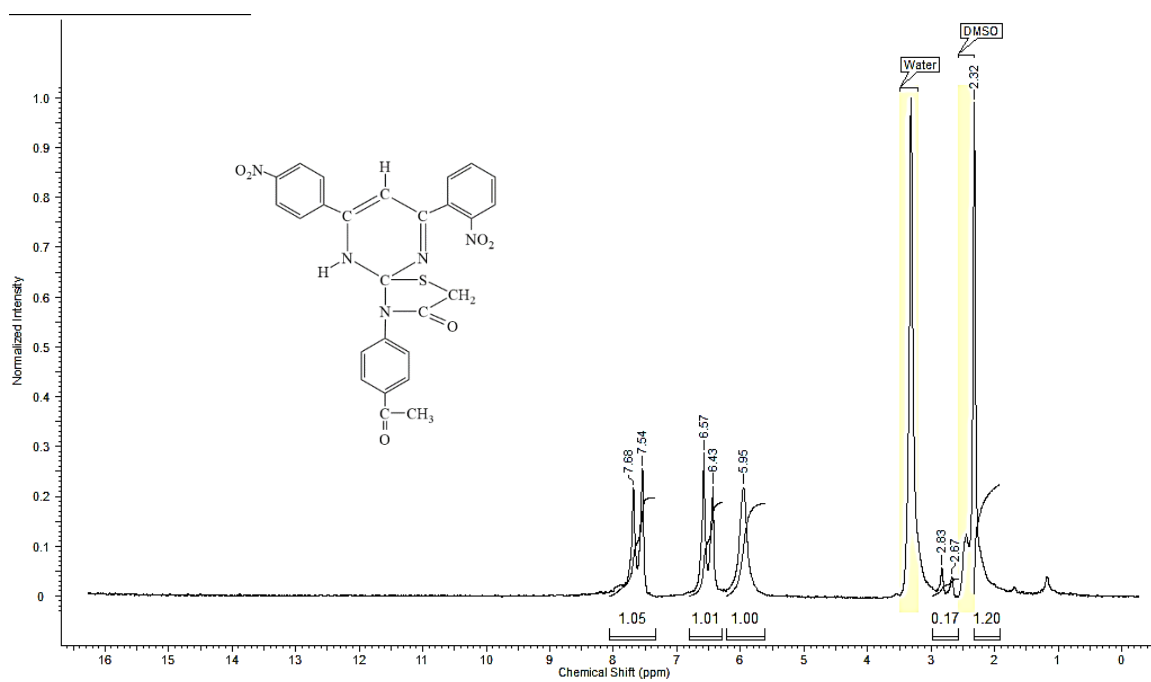


Fig (6): NMR spectrum of the compound (49)

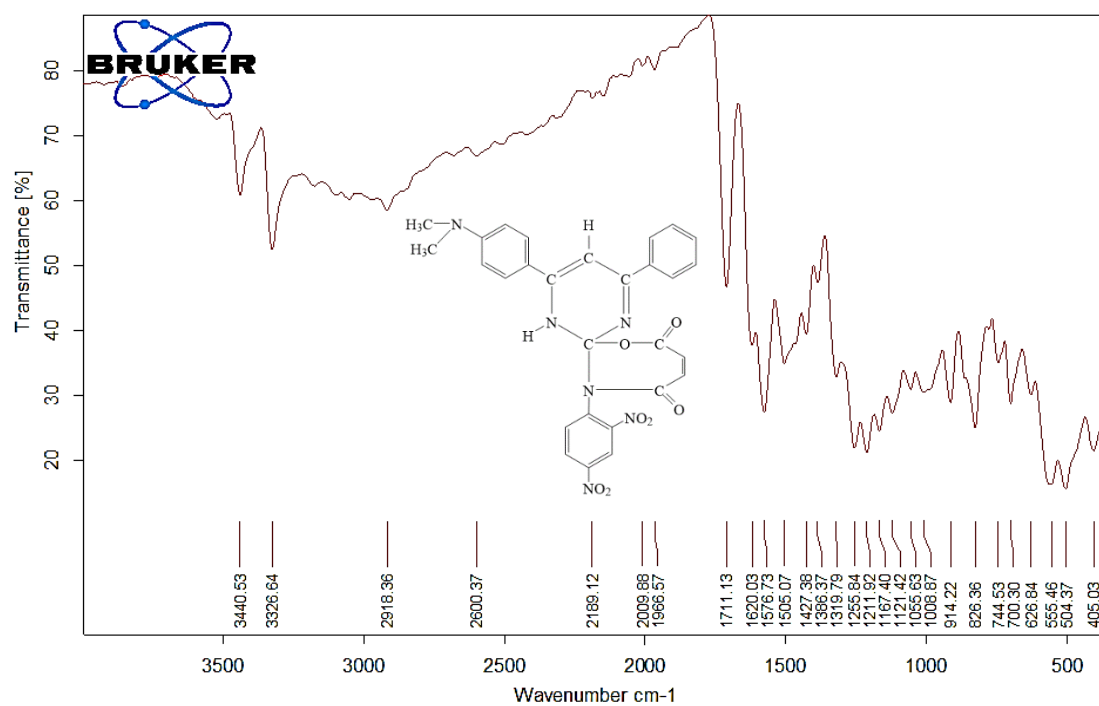


Fig (7): IR spectrum of the compound (54)

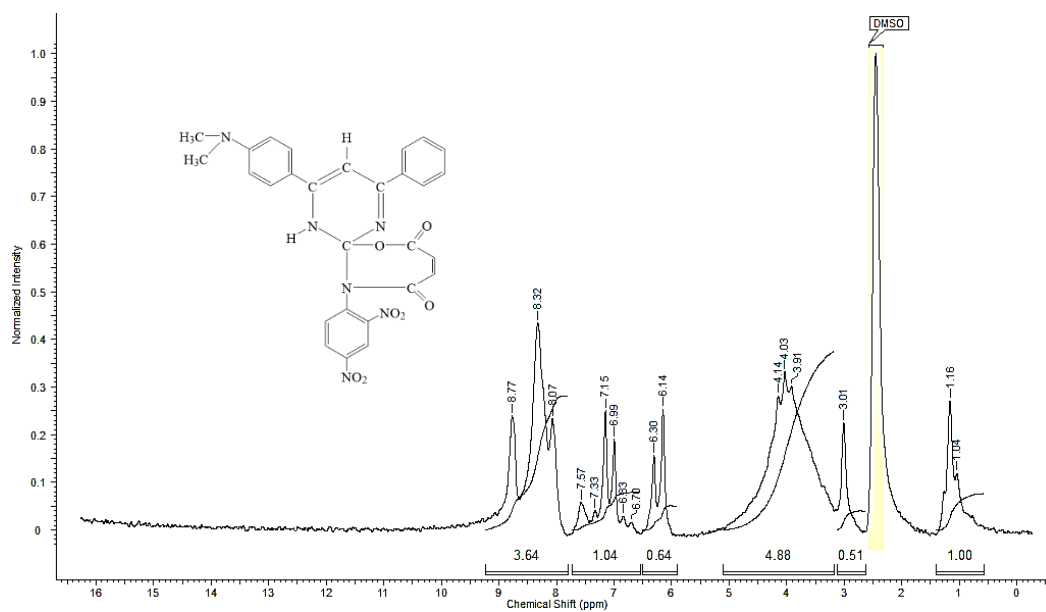


Fig (8): NMR spectrum of the compound (54)

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