


## SYNTHESIS AND SPECTRAL IDENTIFICATION OF SOME TRANSITION METAL COMPLEXES WITH NEW LIGAND OF N-(5-SULFANYL-1,3,4-THIADIAZOL-2-YL)BENZENE SULFONAMIDE

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| <p><b>*For Correspondence:</b><br/> <sup>1</sup>Technical institute of shatra, <sup>2</sup> Thi-qar University, College of pharmacy, <sup>3</sup>University of Basrah, Polymer research centre. <sup>123</sup>Iraq.</p> | <p><b>ABSTRACT</b><br/>           1, 3, 4-thiadiazole–benzenesulfonamide compounds are important due to these are have versatile pharmacological activity such as gonorrhea, scarlet fever, blood poisoning ,tonsillitis ,sinus infection ,urinary tract infection, antimicrobial , Antibacterial, Anti- thyroid, Anticancer ,Anti-tumor, Diuretic, anti-inflammatory, antiviral, antifungal, anti-malarial, anti-HIV, Anticancer . In our present study Some Transition Metal Complexes which are Co(III), Fe(III), Cr(III), Cu(II), Ni(II) with New Ligand of N-(5-Sulfanyl-1,3,4-Thiadiazol-2-yl)Benzene Sulfonamide have been synthesized and identify by using the spectral methods of <sup>1</sup>HNMR , Mass ,IR, as well as Molar conductance and C,H,N . It may be concluded for all the synthesized complexes ligand acts as a bidentate and coordinated through thiadiazole nitrogen and sulfonamide oxygen atoms. It's supported by the appearance of a band corresponding to the metal–nitrogen and the metal–Oxygen stretching vibration at 600–608 cm<sup>-1</sup> and 485-488 cm<sup>-1</sup> in the complexes. The physical and chemical data suggested the octahedral geometry for all complexes except for Cu and Ni complexes which where tetrahedral consequently.</p> <p><b>KEY WORDS:</b> Identification, complexes, Sulphonamide, Synthesis, thiadiazole.</p> |
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### INTRODUCTION

The history of discovery and development of sulphonamide is very interesting. In 1908 Ghlemo has been successfully prepared (Sulpha drug). 1935 Sulphonamide was established by G. Dhomagk as antimicrobial drugs compounds. He has been suggested that the biological activity of sulphonamide is due to its metabolic product 4-aminobenzene sulphonamide. These were containing sulphonamido group (-SO<sub>2</sub>NH<sub>2</sub>-) which was responsible for the higher activity of Sulphonamide drugs [1]. The antimicrobial activity of Sulphonamide extends to treatment of infections caused by gram (+) and gram (-) microorganisms cocci and bacilli, fungi, mycobacteria, some large viruses and protozoa. These were the first effective chemo therapeutic agents of antibacterial drugs has been used for treatment of bacterial infection of human. Furthermore

used in the treatment of many disease like gonorrhoea, scarlet fever, blood poisoning, tonsillitis, sinus infection, urinary tract infection [2]. There are several Authors reports on the synthesis of Sulpha drug.

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**Scheme (1)**

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sulfa drug as pharmacological agents with a wide variety of biological activity such as antimicrobial, Antibacterial, Anti-thyroid, Anticancer, Anti-tumor, Diuretic, [3] anti-inflammatory, [4] antiviral, antifungal, antimalarial, anti-Hiv, Anticancer [5].

The most common method used for synthesis of sulfa compound is by the reaction between alkyl or aryl sulfonyl halide, with amines. In present basic medium [6].

However, in present work we synthesized *N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl)benzenesulfonamide Ligand with some transition metal complexes of Cu(II), Ni(II), Co(III), Fe(III), Cr(III), Which are expected to be highly pharmacological activity of these Complexes compared to it ligand [7]

## 2. Materials

1) hydrazinecarbothioamide 2) methanedithione CS<sub>2</sub>  
3) ethanol 4) anhydrous sodium carbonate Na<sub>2</sub>CO<sub>3</sub> 5) hydrochloric acid HCl 6) pyridine 7) benzenesulphonyl chloride.

## 3. Measurements:-

the FTIR spectra in the range (4000-200) cm<sup>-1</sup> were recorded as CsI discs using a Shimadzu FTIR spectrophotometer . Mass spectra were recorded in the range (0-900) m/z on a 5973 network mass selective detector . Elemental analysis C , H , N were carried out on a Thermo finigan flash analyzer , molar conductance measurements were made in anhydrous DMSO at 25oC using Inolabcond 720 professional benchtop meter ,.The 1H nuclear magnetic resonance spectra were recorded on a Mercury-300BB NMR 300 spectrometer, DOSO-d6 used as solvent. Melting points were determined in Melting points apparatus U.k.

## 4. Experimental work:-

### 4.1 Preparation of Ligand

#### 1) Synthesis of 5-amino -1, 3, 4-thiadiazole-2-thiol :-

hydrazinecarbothioamide (.0.045 mole) was suspended in absolute ethanol (50ml) in (R.B.F) round bottom flask (500ml), with Na<sub>2</sub>CO<sub>3</sub> anhydrous sodium carbonate (0.025mole) and CS<sub>2</sub> (0.131mole) were then added respectively with continues stirring. The reactant mixture was refluxed for 6 hours; the reaction mixture was then allowed to cool at room temperature and filtered. The filtrate was evaporated under vacuum then added cold distilled water (90 ml), finally acidification with concentrated HCl drop by drop, white –yellowish precipitate was formed, and the white-yellowish precipitate was collected by filtration, and washed with distilled water, re-crystallized by using hot distilled water. The percentage of yield (82 %), melting point m.p. 234 – 236 C , [8] .

#### 2)Synthesis of *N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl) benzenesulfonamide :-

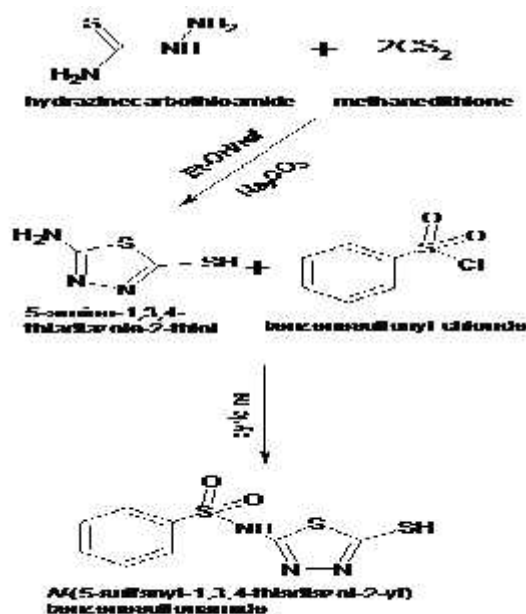
*N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl)Benzenesulfonamide synthesized by Refluxing the mixture of 2 gm of the Synthesis of 5-amino -1, 3, 4-thiadiazole-2- thiol with 6 gm of benzenesulphoyl chloride and 10 ml of pyridine for 45 min. pour the reaction mixture into 20 ml of cold distilled water and stirrer until the wanted product crystallizes .then filter of the solid and recrystallize it from ethanol. percentage of yield ( 62 % ), melting point m.p. 228 – 230 C [9] .

## 4.2 Preparation of complexes:-

By the reaction between the The hydrated metal chloride salts of Ni(II) , Cu(II),Cr(III), Fe(III), Co(III), (0.02 mol) was added to solution of the *N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl)Benzenesulfonamide (0.02mol) in hot absolute ethanol (75 ml) and the mixture was refluxed on a water bath for 3 hours and the solvent was evaporated by concentrated the mixture to half of the original volume and then cooled . The isolated complexes were filtered and washed several times with ethanol and finally dried in air. The physical appearance, percentage of yield, melting point was listed in tablet 1 [10].

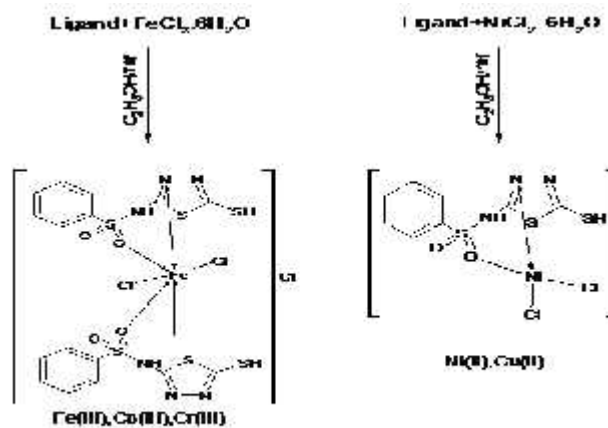
## 5. Present work:-

### 5.1 Preparation of Ligand:-



Scheme (2)

### 5.2 Preparation of complexes :-

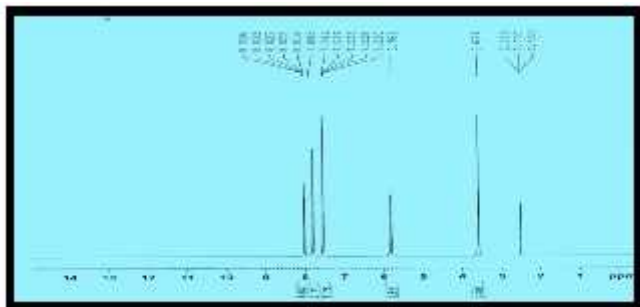


Scheme (3)

## RESULTS & DISCUSSION

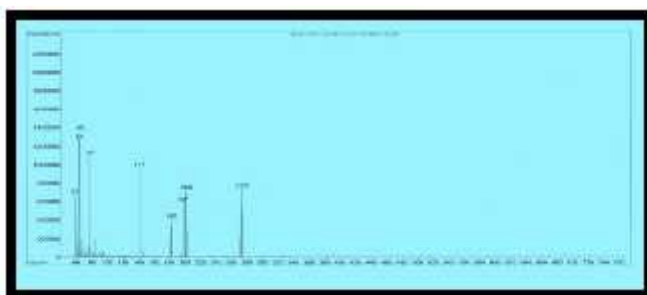
<sup>1</sup>H-NMR Spectra data of the free ligand reported in figure (1). The physical properties, Molecular weight, Molecular formula, and molar conductance data and elemental analysis of the ligand and its complexes tabulated in table (1) and table (2) and (3)

**Figure (1) NMR spectra of the ligand**



The Mass Spectra as shown in a figure (2,3,4) tabulated in table (4). While the Infra-Red Spectroscopy as shown in a figure (5,6) tabulated in table (5).

**Figure (2) mass spectra of Ligand**



**Table 1. physical properties and Molecular weight Molecular formula, data of the ligand and its complexes.**

| No | Formula   | M.Wt | Colour          | M.P °C | % Yield |
|----|---|------|-----------------|--------|---------|
| 1  | [C <sub>8</sub> H <sub>7</sub> N <sub>3</sub> O <sub>2</sub> S <sub>3</sub> ] | 273  | Yellowish white | 228    | 62%     |
| 2  | [Ni(L)Cl <sub>2</sub> ]   | 403  | Deep Green      | 240    | 65%     |
| 3  | [Cu(L)Cl <sub>2</sub> ]   | 408  | Blue            | 238    | 68%     |
| 4  | [Cr(L) <sub>2</sub> Cl <sub>2</sub> ]Cl                                       | 705  | Green           | 251    | 70%     |
| 5  | [Fe(L) <sub>2</sub> Cl <sub>2</sub> ]Cl                                       | 709  | Pale brown      | 256    | 72%     |
| 6  | [Co(L) <sub>2</sub> Cl <sub>2</sub> ]Cl                                       | 712  | Deep brown      | 254    | 67%     |

the Analytical and spectra data (mass spectra, <sup>1</sup>H NMR, IR, elemental analyses C, H, N and Molar conductance of all synthesized Ligand and its complexes were appeared a good agreement between the The calculated values and experimental values and appeared full agreement with the proposed structure .

### 6.1 <sup>1</sup>H-NMR

The <sup>1</sup>H-NMR of the synthesized Ligand the following peaks appeared: at 3.621 (Singlet, 1H, SH), and 5.7 (Singlet, 1H, NH) and 7.520–8.036 (multiplet, 5H, Ar-H).

The NMR Spectra give strong support for the composition of the ligand ,so all the protons are at their expected region The number of protons calculated from integration curves and the recorded chemical shifts in figure (1)[11]

### 6.2 Mass spectra:-

The mass spectra of the ligand shows a molecular ion peak fragment [M/Z] at 273 ,200,215,197,182,141,90,77,65, and 59 due to fragments ion [C<sub>8</sub>H<sub>7</sub>N<sub>3</sub>O<sub>2</sub>S<sub>3</sub>]<sup>+</sup>, [C<sub>7</sub>H<sub>6</sub>NO<sub>2</sub>S<sub>2</sub>]<sup>+</sup>, [C<sub>7</sub>H<sub>6</sub>N<sub>3</sub>O<sub>2</sub>S]<sup>+</sup>, [C<sub>7</sub>H<sub>6</sub>N<sub>2</sub>O<sub>2</sub>S]<sup>+</sup>, [C<sub>6</sub>H<sub>5</sub>O<sub>2</sub>S]<sup>+</sup>, [C<sub>6</sub>H<sub>5</sub>]<sup>+</sup>, [C<sub>5</sub>H<sub>5</sub>]<sup>+</sup>, [CHNS]<sup>+</sup>. Consequently as showed in Figure(2).

The mass spectra of the complex [Cr(L)<sub>2</sub>Cl<sub>2</sub>]Cl shows a molecular ion peak at m/z at 705,669,634,598 which is correspond to the fragment ion peak of [Cr(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup> Cl , [Cr(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup>, [Cr(L)<sub>2</sub>Cl]<sup>+</sup>, [Cr(L)<sub>2</sub>]<sup>+</sup>. Consequently. As showed in Figure (3).

The mass spectra of the complex [Co(L)<sub>2</sub>Cl<sub>2</sub>]Cl shows a molecular ion peak at m/z at 712,677,641,606 which is correspond to the fragment ion peak of [Co(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup> Cl , [Co(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup>, [Co(L)<sub>2</sub>Cl]<sup>+</sup>, [Co(L)<sub>2</sub>]<sup>+</sup>. Consequently.

The mass spectra of the complex [Fe(L)<sub>2</sub>Cl<sub>2</sub>]Cl shows a molecular ion peak at m/z 709,674,639,603 which is correspond to the fragment ion peak of [Fe(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup> Cl , [Fe(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup>, [Fe(L)<sub>2</sub>Cl]<sup>+</sup>, [Fe(L)<sub>2</sub>]<sup>+</sup>. Consequently. The mass spectra of the complex [Ni(L)Cl<sub>2</sub>]<sup>+</sup> shows a molecular ion peak at m/z 403 , 367,332 correspond to the fragment ion peak of [Ni(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup>, [Ni(L)<sub>2</sub>Cl]<sup>+</sup>, [Ni(L)<sub>2</sub>]<sup>+</sup>. Consequently. As showed in Figure (4).

The mass spectra of the complex [Cu(L)Cl<sub>2</sub>]<sup>+</sup> shows a molecular ion peak at m/z 403 , 367,332 correspond to the fragment ion peak of [Cu(L)<sub>2</sub>Cl<sub>2</sub>]<sup>+</sup>, [Cu(L)<sub>2</sub>Cl]<sup>+</sup>, [Cu(L)<sub>2</sub>]<sup>+</sup>. Consequently.[12]

**Table 2. molar conductance data of all complexes measurements were made in anhydrous DMSO at 25°C ,Concentration 10<sup>-4</sup> at 298K .**

| NO | Formula                                 | Λ <sub>m</sub> (S.cm <sup>2</sup> .mol <sup>-1</sup> ) | Electrolyte Type |
|----|---|--|------------------|
| 1  | [Ni(L)Cl <sub>2</sub> ]                 | 11.9   | Non Electrolyte  |
| 2  | [Cu(L)Cl <sub>2</sub> ]                 | 13.5   | Non Electrolyte  |
| 3  | [Cr(L) <sub>2</sub> Cl <sub>2</sub> ]Cl | 22.1   | 1:1              |
| 4  | [Fe(L) <sub>2</sub> Cl <sub>2</sub> ]Cl | 27.8   | 1:1              |
| 5  | [Co(L) <sub>2</sub> Cl <sub>2</sub> ]Cl | 24.2   | 1:1              |

**Table 3 . Elemental analysis of Ligand : C H N**

| Theoretical Data  |       |        |
|-------------------|-------|--------|
| C                 | H     | N      |
| 35.15%            | 2.58% | 15.37% |
| Experimental Data |       |        |
| C                 | H     | N      |
| 35.28%            | 2.64% | 15.25% |

| Table 4 . The mass spectrum of ligand and its Complexes |  |               |
|---|--|---------------|
| No  | Ion  | Molecular Ion |
| 1   | [C <sub>8</sub> H <sub>7</sub> N <sub>3</sub> O <sub>2</sub> S <sub>3</sub> ] <sup>+</sup> | 273           |
| 2   | [C <sub>7</sub> H <sub>6</sub> NO <sub>2</sub> S <sub>2</sub> ] <sup>+</sup>               | 200           |
| 3   | [C <sub>7</sub> H <sub>6</sub> N <sub>3</sub> O <sub>2</sub> S] <sup>+</sup>               | 197           |
| 4   | [C <sub>7</sub> H <sub>6</sub> N <sub>2</sub> O <sub>2</sub> S] <sup>+</sup>               | 182           |
| 5   | [C <sub>6</sub> H <sub>5</sub> O <sub>2</sub> S] <sup>+</sup>                              | 141           |
| 6   | [C <sub>6</sub> H <sub>5</sub> ] <sup>+</sup>  | 77            |
| 7   | [C <sub>5</sub> H <sub>5</sub> ] <sup>+</sup>  | 65            |
| 8   | [CHNS] <sup>+</sup>  | 59            |
| 9   | [Cr(L)2Cl2] <sup>+</sup> Cl  | 705           |
| 10  | [Cr(L)2Cl2] <sup>+</sup>   | 669           |
| 11  | [Cr(L)2Cl] <sup>+</sup>  | 634           |
| 12  | [Cr(L)2] <sup>+</sup>  | 598           |
| 13  | [Fe(L)2Cl2] <sup>+</sup> Cl  | 709           |
| 14  | [Fe(L)2Cl2] <sup>+</sup>   | 674           |
| 15  | [Fe(L)2Cl] <sup>+</sup>  | 639           |
| 16  | [Fe(L)2] <sup>+</sup>  | 603           |
| 17  | [Co(L)2Cl2] <sup>+</sup> Cl  | 712           |
| 18  | [Co(L)2Cl2] <sup>+</sup>   | 677           |
| 19  | [Co(L)2Cl] <sup>+</sup>  | 641           |
| 20  | [Co(L)2] <sup>+</sup>  | 606           |
| 21  | [Cu(L)2Cl2] <sup>+</sup>   | 408           |
| 22  | [Cu(L)2Cl] <sup>+</sup>  | 372           |
| 23  | [Cu(L)2] <sup>+</sup>  | 337           |
| 24  | [Ni(L)2Cl2] <sup>+</sup>   | 403           |
| 25  | [Ni(L)2Cl] <sup>+</sup>  | 367           |
| 26  | [Ni(L)2] <sup>+</sup>  | 332           |

### 7.1 Infra-Red Spectroscopy:-

The FTIR spectrum for synthesized Ligand *N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl)benzene sulfonamide shows A characteristic stretching absorption bands at 3300 cm<sup>-1</sup>, 3027cm<sup>-1</sup>,2760 cm<sup>-1</sup>,1635 cm<sup>-1</sup> and 1480 cm<sup>-1</sup>, which were assigned to  $\nu$  N-H,  $\nu$ C-H Aro,  $\nu$ S-H,  $\nu$ C=N of ring and  $\nu$ S=O consequently. The C=N stretching vibrations are important to predict the bonding mode of the ligand ,these bands shift lower wavelength in the spectra of complexes compare with ligand, observed changes are the evidences of complexion had happened [13]. The IR data of the Ligand and complexes are shown in Table (5) and figure (5,6). The Characteristics groups exhibited by the ligands

Table5. Infra-Red Spectroscopy absorption bands of ligand and its complexes.

| No | Compound      | $\nu$ N-H | $\nu$ C-H | $\nu$ S-H | $\nu$ C=N  |
|----|---------------|-----------|-----------|-----------|------------|
| 1  | Ligand        | 3380      | 3055      | 2811      | 1605       |
| 2  | [Fe(L)2Cl2]Cl | 3390      | 3070      | 2825      | 1612       |
| 3  | [Co(L)2Cl2]Cl | 3390      | 3070      | 2825      | 1612       |
| 4  | [Cr(L)2Cl2]Cl | 3390      | 3070      | 2825      | 1612       |
| 5  | [Ni(L)Cl2]    | 3390      | 3070      | 2825      | 1612       |
| 6  | [Cu(L)Cl2]    | 3390      | 3070      | 2825      | 1612       |
|    |               | $\nu$ S=O | $\nu$ M-N | $\nu$ M-O | $\nu$ M-Cl |
| 1  | Ligand        | 1445      | -----     | -----     | -----      |
| 2  | [Fe(L)2Cl2]Cl | 1460      | 600       | 485       | 305        |
| 3  | [Co(L)2Cl2]Cl | 1460      | 603       | 488       | 307        |
| 4  | [Cr(L)2Cl2]Cl | 1460      | 604       | 486       | 306        |
| 5  | [Ni(L)Cl2]    | 1460      | 608       | 487       | 309        |
| 6  | [Cu(L)Cl2]    | 1460      | 602       | 485       | 302        |

and complexes.[13]

Figure (3) mass spectra of Cr(L)2Cl2]Cl

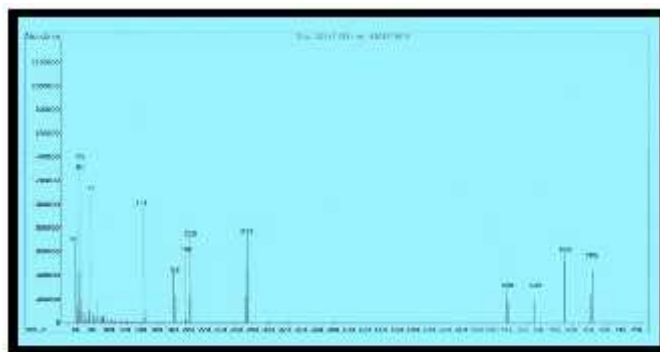


Figure (4) Mass spectra of the  $[\text{Ni}(\text{L})_2\text{Cl}_2]$

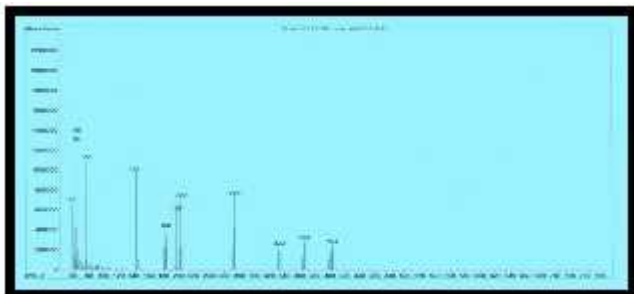


Figure (5) IR spectrum of the ligand

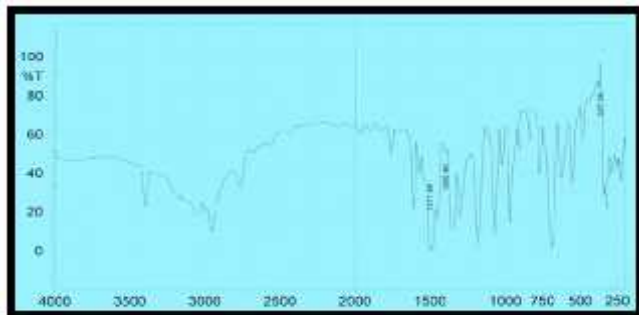
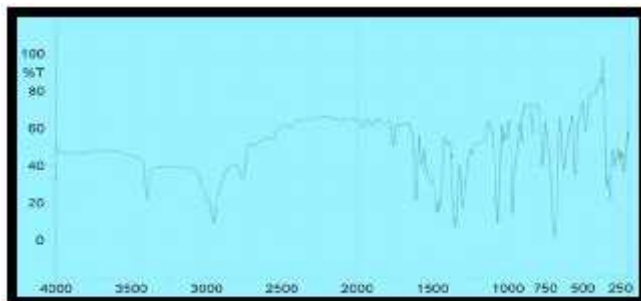


Figure (6) IR spectrum of the  $[\text{Fe}(\text{L})_2\text{Cl}_2]\text{Cl}$



## CONCLUSION

In the present work, a series of Co(III), Fe(III), Cr(III), Ni(II), Cu(II) complexes with new ligand (L), have been prepared and characterized on the basis of IR, <sup>1</sup>HNMR, Mass spectroscopic as well as by elemental analyses C, H, N and Molar conductance. According to all and the physicochemical measurements as the prepared complexes, we can suggested the chemical configuration for the complexes. The ligand Synthesis of *N*-(5-sulfanyl-1,3,4-thiadiazol-2-yl) benzenesulfonamide was successfully synthesized. The ligand was treated to different transition metal salt to form the corresponding complexes as shown in the scheme (3). It may be concluded that the ligand coordinate through sulfonamide oxygen and thiadiazole nitrogen atoms. This view is further supported by the appearance of a band corresponding to the metal–nitrogen, metal–oxygen stretching vibration at 542–563 and 500–600  $\text{cm}^{-1}$  respectively in the complexes. (Cr (III), Fe (III) and Co (III) leading to the formation octahedral geometry complexes

while the Cu (II) and Ni (II) atoms leading to the formation tetrahedral geometry complexes.

## REFERENCES

1. Alka L.Gupta, Text book of medicinal chemistry, 2nd edition 2008, pp152.
2. Dr. Jayashree Ghosh. Text book of pharmaceutical chemistry, 3rd edition 2010, pp135.
3. Gomathi Vellaiswamy and Selvameena ramaswamy. Synthesis, spectral characterization and antimicrobial screening of novel Schiff bases from sulfa drugs. *Int J Pharm Pharm Sci*, Vol 6, Issue 1, 487-491.
4. Shashikant R Pattan<sup>1</sup>\*, R L Hullolikar, Nachiket. S. Dighe<sup>1</sup>, B.N.Ingalagi<sup>2</sup>, M.B. Hole<sup>1</sup>; SYNTHESIS AND EVALUATION OF SOME NEW PHENYL THIAZOLE DERIVATIVES FOR THEIR ANTI-INFLAMMATORY ACTIVITIES;. *Pharm. Sci. & Res.* Vol.1 (4), 2009, 96-102.
5. Sham m sondhi, Amrender dhar dwivedi, Jaiveer singh and pp Gupta: synthesis and anti-inflammatory activity evaluation of some sulfanamide and amidine derivatives of 4-aryl-3-(2 or 4-picolyl)-2-imino-4-thiazolines; *Indian journal of chemistry*; Vol.49B, August 2010; pp.1076-1082
6. J.Dharuman, Chemistry of synthetic drug, 2nd edition, pp251.
7. Sebastián Bellú, Estela Hure, Marcela Trapé and Marcela Rizzotto\*; THE INTERACTION BETWEEN MERCURY(II) AND SULFATHIAZOLE; *Quim. Nova*, Vol. 26, No. 2, 188-192, 2003.
8. Zaid Hassan Abood Rahman Tama Haiwal; Synthesis of Some New Schiff Bases, Tetrazole and 1,3-Oxazepine Derivatives Containing Azo Group and 1,3,4 -Thiadiazole Moiety; *Journal of Babylon University/Pure and Applied Sciences/ No.(1)/ Vol.(21): 2013*
9. G B Talagadadevi, B R Vancha, S Anusuri, M. Rao. Jampani\*; Synthesis of *N*-[4-((5-methyl-1, 3, 4-thiadiazol-2-yl)sulfamoyl)phenyl]sulfamoyl]phenylamine: An Impurity in the Antibacterial Drug Sulfamethizole; *International Journal of Pharmaceutical Sciences and Drug Research* 2012; 4(2): 134-136.
10. Enemose, Edith, A., Akporhonor, E.E. and Osakwe, S.A. Cu (II) and Ni(II) Complexes of Sulfamethazine Mixed with Pyrimethamine: Synthesis, Characterization and Antimicrobial Study; *Chemistry and Materials Research*; Vol.6 No.6, 2014 .
11. Mostafa M. Ghorab<sup>1</sup>, Fatma A. Ragab<sup>2</sup>, Helmy I. Heiba<sup>3</sup>\*, Hebaallah M. Agha<sup>3</sup>; Synthesis of Some Novel Sulfonamides Containing Biologically Active Alkanoic Acid, Acetamide, Thiazole, and Pyrrole Moieties of Expected Antitumor and Radiosensitizing Activities; *J. Basic. Appl. Chem.*, 1(2)8-14, 2011.
12. Safaa N Abdou<sup>1</sup>, Abeer A Faheim<sup>1, 2</sup> and Abdel-Nasser MA Alaghaz<sup>3, 4</sup>\*; Synthesis, Spectral Characterization, Cyclic Voltammetry, Molecular Modeling and Catalytic Activity of Sulfa-Drug Divalent Metal Complexes; *Current Synthetic and Systems Biology*; Volume 2 • Issue 2 • 1000112.
13. V. A. JAGTAP<sup>1</sup>\*, E. JAYCHANDRAN<sup>2</sup>, G.M SREENIVASA<sup>2</sup> AND B.S.SATHE<sup>1</sup>; SYNTHESIS OF SOME FLUORO SUBSTITUTED SULPHONAMIDE BENZOTHIAZOLE COMPRISING THIAZOLE FOR ANTI-MICROBIAL SCREENING; *International Journal of Pharma and Bio Sciences*; ISSN 0975-6299 Vol.1/Issue-4/Oct-Dec.2010.