



Microwave-Assisted Synthesis of Schiff Bases and Evaluation of Their Antibacterial Activities

Manoj Kumar Batra

Department of Chemistry, S.P.C Government College, Ajmer-305001, Rajasthan, India

Abstract: a new Schiff bases was synthesized using microwave-assisted synthesis and conventional condensation with aromatic aldehydes. Microwave-assisted organic synthesis (MAOS) is the study of chemical reactions under the effect of microwave radiation. Microwaves radiation have high energy electric fields and will generally heat any substance containing mobile electric charges, such as polar molecules in a solvent or conducting ions in a solid. In recent year the synthesis of Schiff bases under influence of microwave irradiation was found much easier and faster than conventional heating. The synthesis in Microwave irradiation in solvent free or lower solvent conditions are good method for reduce the pollution, lowering the cost and increase the product together with simplicity in processing and handling.

Keywords: microwave, Schiff base, antibacterial properties.

1. Introduction

Microwave-assisted synthesis is green chemical method, the application of microwave-assisted is useful technology in organic synthesis because it is simple, sensitive, reducing the hazard, often possible to reduce reaction times to a few minutes under solvent free or lower solvent and increase the yields and easier work up as compared to conventional methods. In Schiff bases the carbon-nitrogen double bond (azomethine group) plays important role in synthetic reaction in organic chemistry and which imports in elucidating the mechanism of rasemination and transamination reactions in biological system. Schiff base ligands are considered “privileged ligands” because they are easily prepared by the condensation between aldehydes and imines. Stereogenic centers or other elements of chirality (planes, axes) can be introduced in the synthetic design. Schiff base ligands are able to coordinate many

different metals [1–5], and to stabilize them in various oxidation states. The Schiff base complexes have been used in catalytic reactions [6] and as models for biological systems [7,8]. Many copper complexes of Schiff bases were prepared [9–14]. It has been reported that the structure of the substituent bonded to the imino nitrogen affects the coordination geometry of the complex [15]. During the past two decades, considerable attention has been paid to the chemistry of the metal complexes of Schiff bases containing nitrogen and other donors [16–21]. This may be attributed to their stability, biological activity [22] and potential applications in many fields such as oxidation catalysis [23], electrochemistry [24]. The complexes make these compounds effective and stereospecific catalysts for oxidation, reduction and hydrolysis and they show biological activity, and other transformations of organic and inorganic chemistry [25]. It is well known that some drugs have higher activity when administered as metal complexes than as free ligands [25]. In the present paper the conventional and economical microwave assisted synthesis of Schiff bases of Schiff bases.

2. Experimental:

All the chemical and solvents used were of A R grade and were used without further purification.

Preparation of Schiff base legend (L):

The Schiff base ligand (L) was prepared by the following general method. This was done by the condensation of 20 ml of vanillin (0.03 g, 10 mmol) with 2-aminophenol (0.022 g, 10 mmol) in ethanol (1:1 molar ratio). The mixture was stirred for 4 h.

Antibacterial Studies: Evaluation of antimicrobial activity of all compounds *in vitro* was carried out by paper disc method against bacteria including *E. coli*, *S. aureus*, *M. luteus*, and *B. licheniformis*. Ofloxacin was additionally tested as positive control.

3. Result and Discussion

Antibacterial Studies: Evaluation of antimicrobial activity of all compounds *in vitro* was carried out by paper disc method against bacteria including *E. coli*, *S. aureus*, *M. luteus*, and *B. licheniformis*. Ofloxacin was additionally tested as positive control. The Disc Diffusion method [26], [27] was used to determine the antimicrobial activities of the Schiff bases using standard procedure of 6 mm disc were prepared from Whatman's filter paper no. 1. Silver nanoparticles coated Schiff base solutions of varying concentrations ranging from 100, 500, 1000 ppm was prepared. Nutrient agar was prepared, sterilized and used as the growth medium for the culture of microorganisms; 20 ml of the sterilized medium was poured into each sterilized Petri dish, covered and allowed to solidify. 16 hour old broth cultures of the specified microorganisms were used for testing antibacterial activity [28]. The sample, control

and standard treated discs were air dried at room temperature, to remove any residual solvent which might interfere with the determination, sterilized and inoculated. These plates were initially placed at low temperature for 1 hour so as to allow the maximum diffusion of compounds from the test disc into the agar plate and later incubated at 37°C for 24 h in case bacteria [29], after which the zone of inhibition could be easily observed. The data represent the values of three replicates and are evaluated as mean \pm SEM values were determined and are shown in table 1. The significance level of all compounds were ($P < .001$), ($*P < .01$).

The antibacterial activity was evaluated by tube dilution method which depends on the inhibition of growth of a microbial culture in a uniform solution of antibiotic in a fluid medium that is favourable to its rapid growth in the absence of the antibiotic [30]. In this method minimum inhibitory concentration MIC of the test compounds was determined.

Their MIC values are then table 2.

Table-1: Antimicrobial activity of AgNps anchored with Schiff base:

Conc.(ppm)	<i>E. coli</i> (-)	<i>S. aureus</i> (+)	<i>M. luteus</i> (+)	<i>E. lichenformis</i> (+)
100	21(\pm .205)	20(\pm .450)	21 (\pm .209)	20 (\pm .205)
500	32(\pm .568)	32(\pm .016)	29(\pm .805)	28(\pm .548)
1000	38(\pm .650)	39(\pm .360)	36(\pm .036)	38(\pm .025)

Table 2: MIC value (in mg/ml) of silver nanoparticle anchored with Schiff base

<i>E. coli</i> (-)	<i>S. aureus</i> (+)	<i>M. luteus</i> (+)	<i>E. lichenformis</i> (+)
0.26	0.35	0.31	0.26

4. Conclusion

The Antibacterial activity of the synthesized Schiff base compounds was studied using disc diffusion method and the concentration was fixed using Minimum inhibitory concentration (MIC) method. The antibacterial study revealed that all compounds showed little to excellent activity as compared to standard drug Ofloxacin.

References

- [1] S.M.E. Khalil, Chem. Papers 54 (2000) 12.
- [2] A.H. Osman, Transition Met. Chem. 31 (2006) 35.
- [3] S.A. Sallam, Transition Met. Chem. 31 (2006) 46.
- [4] M. Cindrić, N. Strukan, V. Vrdoljak, T. Kajfež, B. Kamenar, Croatica Chim. Acta 76 (2003) 157.
- [5] C. Sousa, C. Freire, B. de Castro, Molecules 8 (2003) 894.
- [6] D.E. Hamilton, R.S. Drago, A. Zombeck, J. Am. Chem. Soc. 109 (1987) 374.
- [7] D. Chen, A.E. Martel, Inorg. Chem. 26 (1987) 1026.
- [8] J. Costamagna, J. Vargas, R. Latorre, A. Alvarado, G. Mena, Coord. Chem. Rev. 119 (1992) 67.
- [9] M.J. Samide, D.G. Peters, J. Electroanal. Chem. 443 (1998) 95.
- [10] M. Kato, Y. Muto, Coord. Chem. Rev. 92 (1988) 45.
- [11] J. Losada, I. Del Peso, L. Beyer, Inorg. Chem. Acta 301 (2001) 107.
- [12] M.L.P. Santos, I.A. Bagatin, E.M. Pereina, A.M.C. Ferreira, J. Chem. Soc. Dalton Trans. (2001) 838.
- [13] T.N. Rahmonui, S.D. Sid, N. Chenah, O.B. Baitich, Synt. React. Inorg. Met. Org. Chem. 29 (1999) 79.
- [14] M.A. Martinez, R. Aquiler-Saloma, N.M. Ruvalcaba, R.C. Rosado, A.N. Vazquez, V.G. Vidales, A.Z. Dehesa, R.A. Toskano, S.H. Ortega, G.J.M. Fernandez, J. Chem. Soc. Dalton Trans. (2001) 2346.
- [15] L.T. Yildirm, K.C. Emregul, R. Kurtaran, O. Atakol, Cryst. Res. Technol. 37 (2002) 1344.
- [16] S.S. Djebbar, B.O. Benali, J.P. Deloume, Polyhedron 16 (1997) 2175.
- [17] P. Bhattacharyya, J. Parr, A.T. Ross, J. Chem. Soc. Dalton (1998) 3149.
- [18] L. He, S.H. Gou, Q.F. Shi, J. Chem. Crystallogr. 29 (1999) 207.
- [19] J.C. Wu, N. Tang, W.S. Liu, M.Y. Tan, A.S.C. Chan, Chin. Chem. Lett. 12(2001) 757.
- [20] D. Chatterjee, A. Mitra, J. Coord. Chem. 57 (2004) 175.
- [21] G. Minu, H. Bhowon, A. Li Kam Wah, M.O. Dosieah Ridana, D. Ramalingum Lacour, Synth. React. Inorg. Metal-Org. Chem. 34(2004) 1.
- [22] C.M. Liu, R.G. Xiong, X.Z. You, Y.J. Liu, K.K. Cheung, Polyhedron 15(1996) 4565.
- [23] S.S. Djebbar, B.O. Benali, J.P. Deloume, Transition Met. Chem. 23 (1998) 443.

- [24] Y.J. Hamada, IEEE Trans. Electron Devices 44 (1997) 1208.
- [25] R. Ramesh, M. Sivagamasundari, Synth. React. Inorg. Met.-Org. Chem. 33(2003) 899.
- [26] D. Singh, S. Sharma, R. Rani, S. Mishra, and R. Sharma, "Kaempferol-7-O-glucoside and their antimicrobial screening isolate from *Cassia renigera* wall," Int J. Pharm Clin Res, vol. 3, pp. 30-34, Apr. 2011.
- [27] M. L. Delignette-Muller and J. P. Flandrois, "An accurate diffusion method for determining bacterial sensitivity to antibiotics," J. Antimicrob Chemother, vol. 34, no. 1, pp. 73-81, Jul. 1994.
- [28] C. H. Sridevi, K. Balaji, A. Naidu, and R. Sudhakaran, "Antimicrobial evaluation and synthesis of some phenyl pyrazolo benzothiazolo quinoxaline derivatives," J. Chem, vol. 6, no. 3, pp. 866-870, Jul. 2009.
- [29] C. H. Sridevi, M. Kannan, G. Abhinayani, and N. Sravya, "Designing and biological evaluation of new benzimidazole compounds," Chem Sci Trans, vol. 2, no. 3 pp. 922-926, Nov. 2013.
- [30] B. Suman, S. Neha, K. Anu, and K. Sunil, "Design, synthesis, characterization and computational studies on benzamide substituted Mannich bases as novel, potential antibacterial agents," The Scientific World J., pp. 1-9, Jan 2014.