Synthesis, Characterization and Screening of Some Schiff Bases as Potential Antimicrobial Agents

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Authors’ contributions

The chemical part was carried by author SB, whereas biological part is was done by SC. Both authors read and approved the final manuscript.

ABSTRACT

Schiff bases derived from 4-amino antipyrene were prepared, and IR and NMR spectral analysis characterized their structure. The Schiff bases produced were (1) 4-((4-chlorobenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS-1], (2) 4-((4-hydroxybenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS-2], (3)1,5-dimethyl-2phenyl-4-((3-phenylallylidene)amino) -1H-pyrazole-3(2H)-one. [SBS-3], (4)4-((4-hydroxy-3-methoxybenzylidene)amino)-1,5- dimethyl-1H-pyrazole-3(2H)-one. [SBS-4], (5) 4-(benzylideneamino)-1,5 -dimethyl-1H-pyrazole-3(2H)-one. [SBS-5], (6) 4-((furan-2-ylmethelene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS-6] and (7) 4-((4-methoxybenzylidene)amino)-1,5- dimethyl-1H-pyrazole-3(2H)-one. [SBS-7]. The antibacterial activity was studied against S. subfava NCIM 2178, B. megaterium ATCC 9885, P. pseudoalcaligenes ATCC 17440, P. vulgaris NCTC 8313, C. freundii ATCC 10787 and E. aerogenes ATCC 13048. The antibacterial activity was done using Agar Ditch method. The antibacterial activity was evaluated in two polar solvents DMSO and DMF. A differential effect of the compounds extracted in a particular solvent (DMSO/DMF) inhibited different bacteria to a different level. It supports the earlier conclusion that antibacterial activity is dependent on molecular
1. INTRODUCTION

The molecular manipulation of promising lead compounds is still a major line of approach to new drugs. Molecular manipulations involves the efforts to combine the separate groups of similar activity into one compound, thus making structural changes into the compound leading to changes in the biological activity.

Schiff bases are considered as very important class of organic compounds which have wide applications such as dyes, liquid crystals, corrosion inhibitors, complexing agents etc [1-6].

Further, Schiff bases have been reported to demonstrate a wide range of pharmacological activities which include antibacterial, antifungal, anti-HIV, antitumor, anti-inflammatory, antipyretic, etc [7-16].

A variety of Schiff bases have been synthesized from various compounds, the main aim being in search of new lead molecules. Savich et al. [17] synthesized Schiff bases from 2-hydroxy-1-naphthaldehyde with arylamides. Fioravanti et al. [18] observed antibacterial and antifungal activity in few Schiff bases derived from N-heteroaryl benzylamines. Shetye et al. [19] have reported the synthesis of Schiff bases derived from 3-phenyl salicylaldehyde and studied their biological properties. Yang et al. [20] prepared Schiff bases by using salicylaldehyde and aryl amines. Saghatforoush et al. [21] have prepared tetradentate Schiff bases derived from amino thio ether pyridine and salicylaldehyde derivatives and studied their metal complexes. Schiff bases from 4-amino-N-carbamimidoyl benzene sulfonamide have been synthesized by Singh et al. [22] and their antimicrobial activities have also been studied. Amanullah et al. [23] synthesized some novel Schiff bases and studied their cytotoxic, antibacterial activity and physico-chemical properties. Deshpande et al. [24] have synthesized some heterocyclic Schiff bases.

Considering the wide range of biological activities of Schiff bases, in the present work, some new Schiff bases have been synthesized from 4-amino antipyrene with different side chains and to evaluate their potency as anti bacterial agents.

2. EXPERIMENTAL

The following Schiff bases have been synthesized:

1. 4-((4-chlorobenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS1]
2. 4-((4-hydroxybenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS2]
3. 1,5-dimethyl-2-phenyl-4-((3-phenyllallylidene)amino)-1H-pyrazole-3(2H)-one. [SBS3]
4. 4-((4-hydroxy-3-methoxybenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS4]
5. 4-(benzylideneamino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS5]
6. 4-((furan-2-ylmethene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS6]
7. 4-((4-methoxybenzylidene)amino)-1,5-dimethyl-1H-pyrazole-3(2H)-one. [SBS7]

2.1 Experimental Procedure

Equimolar mixture of 4-amino antipyrine and different aldehydes were dissolved in methanol and the mixture was refluxed for 10-15 hours at 65-70°C. The mixture was then poured on crushed ice, filtered and dried.

2.2 Synthesis Scheme

Keywords: Schiff bases; 4-amino antipyrene; antibacterial activity; DMSO; DMF; drug designing.
2.3 Antibacterial Activity

The in vitro antibacterial activity of the synthesized compounds was evaluated against two Gram positive bacteria (S. subfava NCIM 2178 and B. megaterium ATCC 9885) and four Gram negative bacteria (P. pseudoalcaligenes ATCC 17440, P. vulgaris NCTC 8313, C. freundii ATCC 10787 and E. aerogenes ATCC 13048). The antibacterial activity was done by the Agar ditch method using Mueller Hinton Agar No.2 as the nutrient medium [27]. The plates were incubated for 24 h at 37°C. 0.2 ml of the activated strain was inoculated in Mueller Hinton Agar. Mueller Hinton Agar was then poured in the Petri dishes and allowed to solidify. After solidification of the media, 0.85 cm ditch was made in the plates using a sterile cork borer and these were completely filled with the test solution. The plates were incubated for 24 h at 37°C. The mean value obtained for the three wells was used to calculate the zone of growth inhibition of each sample. The controls were maintained for each bacterial strain and each solvent, where pure solvent was inoculated into the well. The inhibition zone formed by these compounds against the particular test bacterial strain determined the antibacterial activities of the synthetic compounds.

3. RESULTS AND CONCLUSION

Seven compounds were synthesized and IR and NMR spectral data analysis confirmed their molecular structure. Their data are given below:

SBS1: \(^1H\ NMR (\delta ppm)\): 2.50 (3H, C-CH\(_3\)), 3.18-3.19 (3H, N-CH\(_3\)), 7.26-7.50 (9H, Ar-H), 8.19-8.22 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3051 (Ar, C=H str.), 2951 (C-H str. (asym)), 2866 (C-H str. (sym)), 1710 (C=O str.), 1627 (C=N str.), 1519 (aromatic C=C str.), 1474 (alkane C-H def. (asym.)), 738 (C-Cl str.).
Mass(m/e): 326, 290, 234, 216, 188, 121, 107.

SBS2: \(^1H\ NMR (\delta ppm)\): 2.42 (3H, C-CH\(_3\)), 3.16 (3H, N-CH\(_3\)), 4.58 (1H, OH), 6.67-7.52 (9H, Ar-H), 8.84 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3630 (OH str), 3030 (Ar, C-H str.), 2954 (C-H str. (asym)), 2871 (C-H str. (sym)), 1714 (C=O str.), 1638 (C=N str.), 1555 (aromatic C=C str.), 1479 (alkane C-H def. (asym.)), 847 (mono sub. aryl ring).
Mass(m/e): 308, 277, 217, 199, 188, 121, 107.

SBS3: \(^1H\ NMR (\delta ppm)\): 2.45 (3H, C-CH\(_3\)), 3.14 (3H, N-CH\(_3\)), 5.58-5.62 (2H, CH=CH), 7.45-8.2 (10H, Ar-H), 8.32 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3058 (Ar, C-H str.), 3021 (CH=CH str.), 2941 (C-H str. (sym)), 2869 (C-H str. (sym)), 1717 (C=O str.), 1619 (C=N str.), 1519 (aromatic C=C str.), 1477 (alkane C-H def. (asym.)), 849 (mono sub. aryl ring).
Mass (m/e): 318, 289, 225, 203, 188, 167, 121, 107.

SBS4: \(^1H\ NMR (\delta ppm)\): 2.46 (3H, C-CH\(_3\)), 3.10 (3H, N-CH\(_3\)), 3.74 (3H, OCH\(_3\)), 5.4 (1H, OH), 8.91-7.31 (8H, Ar-H), 9.6 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3629 (O-H str), 3034 (Ar, C-H str.), 2944 (C-H str.), 1699 (C=O str.), 1540 (Ar, C=C str.), 1620 (C=N str.), 1497 (Ar, C=C str.), 1479 (C-H ben), 1161 (C-O-C str.), 1053 (C-O str.), 845 (mono sub. aryl ring).
Mass (m/e): 338, 307, 322, 202, 188, 139, 121, 107.

SBS5: \(^1H\ NMR (\delta ppm)\): 2.49 (3H, C-CH\(_3\)), 3.15 (3H, N-CH\(_3\)), 7.39-7.66 (10H, Ar-H), 9.76 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3071 (Ar, C-H str.), 2943 (C-H str.), 2867 (C-H str.), 1610 (Ar, C=C str.), 1642 (C=N str.), 1519 (Ar, C=C str.), 1512 (Ar, C=C str.), 1490 (C-H ben), 1427 (C-H ben), 1377 (C-H ben), 1265 (C-C str.), 844 (mono sub. aryl ring).
Mass (m/e): 292, 276, 202, 188, 171, 121, 107.

SBS6: \(^1H\ NMR (\delta ppm)\): 2.47 (3H, C-CH\(_3\)), 3.45 (3H, N-CH\(_3\)), 6.46-6.56 (8H, Ar-H), 7.45 (1H, N=CH).
\textit{IR (KBr, cm}\(^{-1}\)): 3028 (Ar, C-H str.), 2993 (C-H str.), 1725 (C=O str.), 1545 (Ar, C=C str.), 1629 (C=N str.), 1507 (Ar, C=C str.), 1480 (C-H ben), 1070 (C-O str), 841 (mono sub. aryl ring).
Mass (m/e): 282, 265, 202, 188, 121.

SBS7: \(^1H\ NMR (\delta ppm)\): 2.46 (3H, C-CH\(_3\)), 3.10 (3H, N-CH\(_3\)), 3.65 (3H, OCH\(_3\)), 6.44-7.69 (9H, Ar-H), 9.66 (1H, N=CH).
The antibacterial activity against the Gram negative bacteria *C. freundii* and *P. vulgaris* in both polar solvents DMSO and DMF are shown in Fig. 2. Here an entirely different trend was observed. Only three compounds i.e. SBS1, SBS2 and SBS3 in DMF showed antibacterial activity against *C. freundii* and maximum activity was shown by SBS3. All other compounds in both the solvents did not show any activity against *C. freundii*. All the seven compounds showed activity against *P. vulgaris* extracted in DMSO while none of the compounds exhibited antibacterial activity in DMF. Maximum activity was shown by SBS1, closely followed by SBS2 and SBS3 while minimum activity was shown by SBS6 and SBS7. This again firmly proves and confirms our earlier conclusion that there is a direct and strong relation between structures of the compound, solvent in which it is extracted and the bacterial strain against which it is being studied.

The antibacterial activity against *P. pseudoalkaligenes* and *E. aerogenes* are shown in Fig. 3. All the seven compounds synthesized from 4-amino antipyrene extracted in DMSO could not inhibit the growth of either of these two Gram negative bacteria.

The compound extracted in DMF inhibited both these bacteria but a differential effect was envisaged. All the compounds extracted in DMF could inhibit *P. pseudoalkaligenes* but each compound had a different level of inhibition. Maximum inhibitory activity was shown by SBS4 and SBS5 followed by SBS3, SBS6 and SBS7 respectively. Minimum activity was by SBS1 and SBS2. In *E. aerogenes*, only SBS3 showed some inhibition while all the other compounds showed no activity.

### Table 1. Analytical and physical data of the synthesized compounds

| Compound code | Molecular formula | Molecular weight (gm/Mol) | M.P °C | % Yield | Rf value |
|---------------|-------------------|--------------------------|--------|---------|----------|
| SBS-1         | C_{18}H_{16}N_{3}OCl | 325.5                    | 193    | 63      | 0.62     |
| SBS-2         | C_{18}H_{17}N_{3}O_{2} | 307                     | 197    | 59      | 0.64     |
| SBS-3         | C_{20}H_{19}N_{3}O | 317                     | 98     | 52      | 0.61     |
| SBS-4         | C_{19}H_{19}N_{3}O_{3} | 337                    | 194    | 67      | 0.30     |
| SBS-5         | C_{18}H_{17}N_{3}O | 291                     | 163    | 56      | 0.60     |
| SBS-6         | C_{16}H_{15}N_{3}O_{2} | 281                    | 216    | 58      | 0.38     |
| SBS-7         | C_{16}H_{19}N_{3}O_{2} | 321                    | 159    | 59      | 0.48     |

Acetone: Benzene (1:9)
Fig. 1. Antibacterial activity of synthesized compounds against gram positive bacteria
Fig. 2. Antibacterial activity of synthesized compounds against gram negative bacteria
In the present work, among the seven synthesized compounds and the two solvents used, the best activity was shown by SBS3 in DMF solvent. Hence, it can be concluded that cinnamaldehyde as side chain with 4-amino antipyrine as central ligand and DMF as solvent are the best in inhibiting the studied bacterial strains. This once again proves the earlier conclusion that antibacterial activity is dependent on molecular structure of the compound, solvent used and the bacterial strain under consideration [25,26]. Such screening of various organic compounds and identifying active agents is the need of the hour; because successful prediction of lead molecule and drug like properties at the onset of drug discovery will pay off later in drug development.
COMPETING INTERESTS

Authors have declared that no competing interests exist.

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