SYNTHESIS AND ANTIMICROBIAL ACTIVITIES OF SOME 4-(6-METHYL-2-NAPHTHYL)-5,6-DIHYDRO-6-(SUBSTITUTED PHENYL)-4H-1,3-OXAZINE-2-AMINES

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ABSTRACT

Some 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-4H-1,3-oxazine-2-amines have been synthesised by hydroxyapatite catalyzed solvent-free cyclization of 6-methyl-2-naphthyl chalcones and urea under microwave irradiation. The yields of the oxazine were more than 85%. The synthesised oxazine amines have been characterized by their physical constants, analytical and spectroscopic data. The antimicrobial activities of all prepared 4-(6-methyl-2-naphthyl)-5, 6-dihydro-6-(substituted phenyl)-4H-1,3-oxazine-2-amines have been evaluated using Bauer-Kirby disc diffusion method with a variety of bacterial and fungal strains.

Keywords: Oxazine-2-amines, Enones, Hydroxyapatite, Green synthesis, Antimicrobial activities.

1. INTRODUCTION

The unsaturated 1,3-oxazine and their derivatives are important bio-active six membered heterocyclic compounds which possess one oxygen and one nitrogen atom [1,2]. Oxazines exist many isomeric structures such as 1, 2 or 1, 3 or 1, 4 oxazines [3] depend upon the relative position of these two atoms and the double bond. These oxazines were medicinally important due to the presence of oxygen, nitrogen heteroatoms along with double bonds in their structural moieties [4]. The important medicinal activities of these oxazines are anti-bacterial, anti-fungal [4-6], anti-plasmodial [7], anti-cancer [8], anti-depressants [9], cytotoxicity [10], anti-osteoplastic [11], anti-tumour [12], anti-oxidant [13], anti-tuberculosis [14], anti-neoplastic [15], antagonists [16], anti-

inflammatory [17], anti-infectants [18], IKK kinase beta [19] and PTP-1B inhibition [20]. These oxazine derivatives were applied for improving the super resolution microscope [21], synthesis of eosinophils [22], identification and separation of neutrophils [23]. Many oxazine derivatives were used as dyes [24]. Nowadays scientists, organic chemists are interested for solvent-free synthesis [4, 25-31] of organic substrates. Numerous solvent assisted and solvent-free synthetic methods were available for synthesis of oxazine derivatives [4, 25-27, 32,]. Hetero Diels-Alder reaction [4], ring closure [33], Betti base induced condensation [27], Mannich type condensation-cyclization [5] and cyclization of chalcones [6] were used for synthesis of oxazine derivatives [25] have synthesised some benzoxazine/ oxazine fused isoquinolines and naphthyridines by solvent-

free method [25]. They have synthesised some 1,3-oxazine derivatives by solvent-assisted method from chalcones and urea [6]. More than 75% yield of dihydro-2H-benzo- and naphtho-1, 3-oxazine derivatives were prepared by Mathew et al.[5] using eco-friendly method. Efficient synthesis of some 1,3-oxazine-4-thiones were synthesised by N-methyl imidazole promoted solvent-free conditions. Sapkal et al., have studied the role of ammonium acetate for solvent-free synthesis of 1, 3-disubstituted-2,3-dihydro-1H-naphthyl oxazines [27]. Within the above view, there is no information available in the literature for the solvent-free synthesis of oxazine amine-2-derivatives including 4-(6-methyl-2-naphthyl)-6-(substituted phenyl) 4H-1,3-oxazine-2-amines by cyclization of enones. Therefore the authors have taken effort to synthesized some 4-(6-methyl-2-naphthyl)-6-(substituted phenyl) 4H-1,3-oxazine-2-amines by solvent-free cyclization of enones and urea under microwave irradiation and studied the antimicrobial activities.

2. EXPERIMENTAL

2.1. GENERAL

All chemicals were used in this study were purchased from Sigma-Aldrich and Merck Chemical companies. Mettler FP51 melting point apparatus was used for determining the melting point of all synthesized oxazine-2-amines in open glass capillaries and are uncorrected. The AVATAR-300 Fourier transform spectrophotometer was used for recording infrared spectra (KBr, 4000-400 cm⁻¹) of all oxazines in KBr disc. The Bruker AV400 series type NMR spectrometer was utilized for recording NMR spectra of all oxazines, operating at 400MHz for ¹H and 100 MHz for ¹³C spectra in CDCl₃ solvent

using TMS as internal standard. Mass spectra of all synthesised oxazines were recorded on SHIMADZU mass spectrometer using chemical ionization technique.

2.2. PREPARATION OF HYDROXYAPATITE AND 6-METHYL-2-NAPHTHYL CHALCONES

Hydroxyapatite catalyst and 6-methyl-2-naphthyl chalcones were prepared according to literature procedure [34].

2.3. SYNTHESIS OF 4-(6-METHYL-2-NAPHTHYL)-5,6-DIHYDRO-6-(SUBSTITUTED PHENYL)-4H-1,3-OXAZINE-2-AMINES

An appropriate equi-molar quantities of 6-methyl-2-naphthyl chalcones (2 mmol), urea (2 mmol) and 0.4 g of hydroxyapatite were taken in a 50 mL beaker, closed with the lid. This mixture was subjected to microwave irradiation for 2-4 minutes at 650W (Samsung Grill, GW73BD Microwave oven, 230V A/c, 50Hz, 2450Hz, 100-750W (IEC-705), (Scheme 1). After completion of the reaction, dichloromethane (20 mL) was added, followed by simple filtration. The solution was concentrated and purified by recrystallization. The synthesized oxazines were characterized by their physical constants, IR, ¹H and ¹³C NMR and Mass spectral data.

Scheme. 1. Synthesis of 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-4H-1,3-oxazine-2-amines.

3. RESULTS AND DISCUSSION

The authors attempts made to synthesize oxazine-2-amine derivatives by cyclization of chalcones possess electron with-drawing as well as electron donating group as substituents, urea and in the presence of a catalyst hydroxyapatite using microwave irradiation. Hence the authors have synthesized some substituted 1,3-oxazine derivatives by the cyclization of 2 mmol of 6-methyl-2-naphthyl chalcones, 2 mmol of urea under microwave irradiation with 0.4 g of hydroxyapatite catalyst at 550W for 4-6 minutes (Scheme 1). During the course of this reaction hydroxyapatite catalyses cyclization between 6-methyl-2-naphthyl chalcones and urea followed by rearrangement gave the 1,3-oxazine amines. The yield of the oxazine in this reaction is more than 80%. The 6-methyl-2-naphthyl chalcones containing electron donating substituent (OCH₃) gave higher yields than electron-withdrawing (halogens, NO₂) substituents. Further we have investigated this cyclization reaction with equimolar quantities of the styryl 6-methyl-2-naphthyl ketone (entry 1) and urea under the same condition as above. In this reaction the obtained yield was 90%. The effect of catalyst on this reaction was studied by varying the catalyst quantity from 0.1 g to 1 g. As the catalyst quantity is increased from 0.1 g to 0.4 g, the percentage of yield of product is increased from 84 to 90%. Further increase in the catalyst amount beyond 0.4 g, there is no significant increase in the percentage of the product. The effect of catalyst loading is shown in Figure 1. The optimum quantity of catalyst loading was found to be 0.4 g. The physical constants, analytical and mass spectral data are summarized in Table 1. The effect of solvents on the yield was also studied with methanol, ethanol, dichloromethane and tetrahydrofuran from each component of the

catalyst (entry 1). Similarly the effect of microwave irradiation was studied on each component of the catalyst. The effect of solvents on the yield of oxazine derivatives was presented in Table 2. From the table highest yield of oxazine obtained from the cyclization of 6-methyl-2-naphthyl chalcones and urea with the catalyst hydroxyapatite in microwave irradiation.

Fig.1. Effect of catalyst loading.

The infrared and nmr spectroscopic data of selective oxazine-2-amine derivatives are summarized as follows:

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-phenyl-4H-1,3-oxazine-2-amine (1): FTIR (KBr): 3538 (NH), 1587 (C=N), 1212 (C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.321 (s, 1H, NH₂), 2.655 (dd, 1H, H₄), 2.345 (dd, 1H, H₅), 2.217 (dd, 1H, H_{5'}), 4.256 (dd, 1H, H₆), 3.575 (s, 3H, CH₃), 6.521-7.378 (m, 11H, Ar-H) ppm.; ¹³C NMR (CDCl₃-d₆, TMS) δ : 165.34 (C₂), 52.76 (C₄), 47.34 (C₅), 65.68 (C₆), 25.98 (CH₃), 125.34-142.87 (Ar-C) ppm

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(3-aminophenyl)-4H-1,3-oxazine-2-amine (2): FTIR (KBr): 3560 (NH), 1615(C=N), 1224(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆,

TMS) δ : 2.229(s, 1H, NH₂), 2.532 (dd, 1H, H₄), 2.443(dd, 1H, H₅), 2.211(dd, 1H, H_{5'}), 3.353 (dd, 1H, H₆), 3.243 (s, 3H, CH₃), 6.234-7.265(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.32(C₂), 51.36 (C₄), 47.26(C₅), 66.54 (C₆), 24.65(CH₃), 126.54-139.67 (Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-aminophenyl)-4H-1,3-oxazine-2-amine (3): FTIR(KBr): 3555(NH), 1620(C=N), 1212(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.223 (s, 1H, NH₂), 2.556 (dd, 1H, H₄), 2.454(dd, 1H, H₅), 2.265 (dd, 1H, H_{5'}), 4.371(dd, 1H, H₆), 3.234(s, 3H, CH₃), 6.234-7.276 (m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.43(C₂), 51.76 (C₄), 47.35(C₅), 66.76 (C₆), 24.65(CH₃), 126.34-139.89 (Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(3-bromophenyl)-4H-1,3-oxazine-2-amine (4): FTIR(KBr): 3540(NH), 1612(C=N), 1224(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.218(s, 1H, NH₂), 2.324(dd, 1H, H₄), 2.265dd, 1H, H₅), 2.267 (dd, 1H, H_{5'}), 4.360(dd, 1H, H₆), 3.334 (s, 3H, CH₃), 6.234-7.267(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.46 (C₂), 51.56 (C₄), 47.65(C₅), 66.76 (C₆), 25.33(CH₃), 124.87-139.90 (Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(3-chlorophenyl)-4H-1,3-oxazine-2-amine (5): FTIR(KBr): 3550(NH), 1615(C=N), 1230(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.256(s, 1H, NH₂), 2.332 (dd, 1H, H₄), 2.416(dd, 1H, H₅), 2.265 (dd, 1H, H_{5'}), 4.376(dd, 1H, H₆), 3.142(s, 3H, CH₃), 6.256-7.276 (m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.67(C₂), 51.98 (C₄), 47.02(C₅), 66.47 (C₆), 24.12(CH₃), 126.65-139.87 (Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-chlorophenyl)-4H-1,3-oxazine-2-amine (6): FTIR(KBr): 3550(NH), 1605(C=N), 1223(C-O-C)cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.243 (s, 1H, NH₂), 2.343 (dd, 1H, H₄), 2.422(dd, 1H, H₅), 2.232(dd, 1H, H_{5'}), 4.368(dd, 1H, H₆), 2.932(s, 3H, CH₃), 6.243-7.276 (m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.56(C₂), 51.87 (C₄), 47.87(C₅), 66.72 (C₆), 24.11(CH₃), 126.65-139.87(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-dimethylaminophenyl)-4H-1,3-oxazine-2-amine (7): FTIR(KBr): 3530(NH), 1592(C=N), 1206(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.211(s, 1H, NH₂), 2.330 (dd, 1H, H₄), 2.420(dd, 1H, H₅), 2.232(dd, 1H, H_{5'}), 4.328(dd, 1H, H₆), 3.203(s, 6H, N(CH₃)₂), 3.154(s, 3H, CH₃), 26.265-7.218 (m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.80 (C₂), 51.45 (C₄), 47.78(C₅), 66.98 (C₆), 44.25(N(CH₃)₂), 24.55(CH₃), 126.19-139.76(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-hydroxyphenyl)-4H-1,3-oxazine-2-amine (8): FTIR(KBr): 3540(NH), 1615(C=N), 1211(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.219 (s, 1H, NH₂), 2.340 (dd, 1H, H₄), 2.454(dd, 1H, H₅), 2.287(dd, 1H, H_{5'}), 4.376(dd, 1H, H₆), 3.198(s, 3H, CH₃), 6.234-7.254(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.64 (C₂), 51.54 (C₄), 47.65(C₅), 63.76 (C₆), 44.76 (N(CH₃)₂), 24.67(s, 3H, CH₃), 126.64-139.76(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-methoxyphenyl)-4H-1,3-oxazine-2-amine (9): FTIR(KBr): 3540(NH), 1624(C=N), 1218(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆,

TMS) δ : 2.217 (s, 1H, NH₂), 2.334(dd, 1H, H₄), 2.454(dd, 1H, H₅), 2.265(dd, 1H, H_{5'}), 4.327(dd, 1H, H₆), 4.233(s, 3H, OCH₃), 3.326(s, 3H, CH₃), 6.233-7.253(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.54 (C₂), 51.26 (C₄), 47.37(C₅), 66.76 (C₆), 64.76(s, 3H, OCH₃), 24.98(s, 3H, CH₃), 126.32-139.66(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-methylphenyl)-4H-1,3-oxazine-2-amine (10): FTIR(KBr): 3538(NH), 1616(C=N), 1217(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.219 (s, 1H, NH₂), 2.322 (dd, 1H, H₄), 2.436(dd, 1H, H₅), 2.226(dd, 1H, H_{5'}), 4.347(dd, 1H, H₆), 2.833 (s, 3H, CH₃), 2.513 (s, 3H, CH₃), 6.288-7.272(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.78 (C₂), 51.89 (C₄), 47.45(C₅), 66.76 (C₆), 25.87(CH₃), 25.43(CH₃), 114.65-145.87(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(2-nitrophenyl)-4H-1,3-oxazine-2-amine (11): FTIR(KBr): 3555(NH), 1622(C=N), 1214(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.265 (s, 1H, NH₂), 2.333 (dd, 1H, H₄), 2.465(dd, 1H, H₅), 2.265(dd, 1H, H_{5'}), 4.365(dd, 1H, H₆), 3.164 (s, 3H, CH₃), 6.243-7.276(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 64.54(C₂), 51.65 (C₄), 47.87(C₅), 66.34 (C₆), 26.87(s, 3H, CH₃), 114.65-145.76(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(3-nitrophenyl)-4H-1,3-oxazine-2-amine (12): FTIR(KBr): 3558(NH), 1622(C=N), 1218(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.218 (s, 1H, NH₂), 2.339(dd, 1H, H₄), 2.465(dd, 1H, H₅), 2.242(dd, 1H, H_{5'}), 4.354(dd, 1H, H₆), 3.056 (s, 3H, CH₃), 6.544-7.265(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.68(C₂),

51.42 (C₄), 47.56(C₅), 66.87 (C₆), 24.78(s, 3H, CH₃), 124.88-145.98(Ar-C)ppm.

4-(6-Methyl-2-naphthyl)-5,6-dihydro-6-(4-nitrophenyl)-4H-1,3-oxazine-2-amine (13): FTIR(KBr): 3555(NH), 1633(C=N), 1213(C-O-C) cm⁻¹. ¹H NMR (CDCl₃-d₆, TMS) δ : 2.220 (s, 1H, NH₂), 2.336(dd, 1H, H₄), 2.464(dd, 1H, H₅), 2.265(dd, 1H, H_{5'}), 4.376(dd, 1H, H₆), 3.123 (s, 3H, CH₃), 6.536-7.276(m, 10H, Ar-H) ppm. ¹³C NMR (CDCl₃-d₆, TMS) δ : 164.76(C₂), 51.78 (C₄), 47.87(C₅), 66.37 (C₆), 4.78(s, 3H, CH₃), 124.25-145.998(Ar-C)ppm.

3.1. ANTIMICROBIAL ACTIVITIES

3.1.1 ANTIBACTERIAL ACTIVITIES

Antibacterial activities of all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substitutedphenyl)-oxazine-2-amines were evaluated using Kirby-Bauer [35] disc diffusion technique. In each Petri plate about 0.5 mL of the test bacterial sample was spread uniformly over the solidified Mueller Hinton agar using sterile glass spreader. Then the discs with 5 mm diameter made up of Whatmann No.1 filter paper, impregnated with the solution of the compound were placed on the medium using sterile forceps. The plates were incubated for 24 h at 37°C by keeping the plates upside down to prevent the collection of water droplets over the medium [36]. After 24 h, the plates were visually examined and the diameter values of the zone of inhibition were measured. Triplicate results were recorded by repeating the same procedure. The anti-bacterial activities of all prepared 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substitutedphenyl)-oxazine-2-amines have been evaluated against two gram positive pathogenic strains *Staphylococcus aureus*, *Enterococcus faecalis* and four gram negative

strains *Escherichia coli*, *Klebsiella* species, *Pseudomonas* and *Proteus vulgaris*. The disc diffusion technique was followed using the Kirby-Bauer [35] method, at a concentration of 250 µg/mL. Ampicillin and Streptomycin were used as the standard drugs. The measured anti-bacterial activities of all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-substituted phenyl-oxazine-2-amines were presented in Table 3. The oxazine-2-amines 4-6 showed maximum zone of inhibition against *Escherichia coli*, with greater than 20 mm compared to the oxazines 3, 8-10 are active in 8-12 mm of zone of inhibition. The oxazines 1, 2, 7, 11-13 are inactive. The 1,3-oxazine-2-amine 6 only effective against *S. aureus* within 20-24 mm of zone of inhibition. Compounds 7, 11-13 were moderately active within 13-19 mm of zone of inhibition. The oxazines 1, 5, 8-10 were active within 8-12 mm of zone of inhibition. The oxazine derivatives 4-6 were more active against *Pseudomonas* showing greater than 20 mm zone of inhibition and the other derivatives 7, 8, and 13 were showed the zone of inhibitions between 12-19 mm. The oxazines 2-3, and 11 were showed active with in 8-12 mm of zone of inhibition. The oxazine-2-amines 6 was effective within 20-24 mm of zone of inhibition against *Klebsiella* species. The oxazine derivative 6 was active against *P. vulgaris* greater than 20mm of zone of inhibition. The other compounds are less effective. The oxazines 14 and 16 showed activities against *E. faecalis* when they are screened with 20-24 mm zone of inhibition.

3.1.2 ANTI-FUNGAL SENSITIVITY ASSAY

Antifungal sensitivity assay all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines (entries 1-13) were performed using Kirby-Bauer disc diffusion

technique. PDA medium was prepared and sterilized as above. It was poured (ear bearing heating condition) in the Petri-plate which was already filled with 1 mL of the fungal species. The plate was rotated clockwise and counter clock-wise for uniform spreading of the species. The discs were impregnated with the test solution. The test solution was prepared by dissolving 15 mg of the oxazine in 1 mL of DMSO solvent. The medium was allowed to solidify and kept for 24 h. Then the plates were visually examined and the diameter values of zone of inhibition were measured. Triplicate results were recorded by repeating the same procedure. The study of antifungal activities of all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines has been done with *Candida albicans* as the fungal strain using the disc diffusion technique and the other two strains *Penicillium* species and *Aspergillus niger*, the dilution method was adopted. The drug dilution was kept as 50 g/mL. Griseofulvin has been taken as the standard drug. The observed antifungal activities of all all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines were presented in Table 4. The study of antifungal activities of all 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines against *C. albicans*, showed that the three compounds 11-13 were effective with 20 mm as the zone of inhibition in 250 µg/disc while oxazine-2-amines 5, 6 and 10 were active with 13-19 mm zone of inhibition and the compound 1,4, 8 and 9 were shows least active with 8-12 mm zone of inhibitions. The oxazine derivatives 5, 6, 11-13 were more visible against *Penicillium* species, in the development of the fungal colony and 2-3 colonies are recorded for the compound 10. The inhibition of oxazine-2-amines against *A. niger* was more active in five compounds 5, 6,

11-13. The compounds 2-4 and 7 were active with 13-19 mm zone of inhibitions. Presence of amino, dimethylamino, methoxy, methyl, dimethyl, chloro, bromo and nitro

substituents are responsible for antimicrobial activities of 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines.

TABLE 1: ANALYTICAL, PHYSICAL CONSTANTS, YIELD AND MASS FRAGMENT OF 4-ARYL-5,6-DIHYDRO-6(SUBSTITUTED PHENYL)-4H-1,3-OXAZINE-2-AMINES

Entry	X	M.F.	M.W.	Yield (%)	m.p.(°C)	Mass (m/z)
1	H	C ₂₁ H ₂₀ N ₂ O	317	90	117-118	317[M ⁺], 311, 255, 175, 160, 157, 91, 77, 43, 42, 31, 30, 16
2	3-NH ₂	C ₂₁ H ₂₁ N ₃ O	332	89	107-108	332[M ⁺], 316, 315, 274, 255, 190, 160, 157, 134, 98, 92, 91, 77, 58, 32, 31, 16, 15
3	4-NH ₂	C ₂₁ H ₂₁ N ₃ O	332	89	95-96	347[M ⁺], 316, 274, 255, 190, 160, 157, 98, 92, 91, 77, 31, 16, 15
4	3-Br	C ₂₁ H ₁₉ BrN ₂ O	395	87	117-118	395[M ⁺], 397[M ²⁺], 316, 301, 255, 175, 157, 91, 79, 77, 45, 42, 31, 16,
5	3-Cl	C ₂₁ H ₁₉ ClN ₂ O	351	86	131-143	351[M ⁺], 353[M ²⁺], 316, 350, 335, 308, 272, 255, 209, 194, 157, 11, 94, 91, 84, 77, 58, 43, 42, 35, 31, 16
6	4-Cl	C ₂₁ H ₁₉ ClN ₂ O	366	86	123-124	351[M ⁺], 353[M ²⁺], 335, 308, 272, 255, 111, 94, 91, 84, 77, 42, 35, 31, 16,
7	4-N(CH ₃) ₂	C ₂₃ H ₂₅ N ₃ O	360	88	115-116	360[M ⁺], 359, 345, 344, 331, 255, 218, 157, 120, 102, 99, 91, 77, 44, 43, 31, 30, 16, 15, 14,
8	4-OH	C ₂₁ H ₂₀ N ₂ O ₂	333	87	100-101	333[M ⁺], 332, 331, 318, 316, 255, 197, 175, 157, 127, 91, 77, 43, 42, 31, 17, 16
9	4-OCH ₃	C ₂₂ H ₂₂ N ₂ O ₂	347	92	125-126	347 [M ⁺], 322, 304, 255, 205, 198, 164, 157, 109, 107, 84, 77, 58, 43, 42, 31, 15, 16
10	4-CH ₃	C ₂₂ H ₂₂ N ₂ O	331	91	107-108	331[M ⁺], 330, 315, 288, 255, 240, 175, 157, 127, 91, 84, 77, 43, 42, 31, 16, 15,
11	2-NO ₂	C ₂₁ H ₁₉ N ₃ O ₃	362	85	117-118	362[M ⁺], 361, 346, 331, 319, 275, 255, 198, 185, 179, 164, 157, 136, 127, 99, 91, 77, 58, 46, 43, 31, 16
12	3-NO ₂	C ₂₁ H ₁₉ N ₃ O ₃	362	86	127-128	362[M ⁺], 361, 346, 331, 255, 179, 164, 157, 136, 127, 99, 91, 77, 46, 43, 31, 16
13	4-NO ₂	C ₂₁ H ₁₉ N ₃ O ₃	362	85	109-110	362[M ⁺], 361, 346, 331, 319, 275, 255, 185, 179, 164, , 136, 127, 91, 77, 58, 43, 31, 16

TABLE 2: THE EFFECT OF SOLVENTS IN CONVENTIONAL HEATING AND WITHOUT SOLVENT IN MICROWAVE IRRADIATION ON YIELD OF OXAZINE AMINE (ENTRY 1).

Solvents				Microwave irradiation
MeOH	EtOH	DCM	THF	
61	60	64	65	90

MeOH=Methanol; EtOH=Ethanol;

DCM=Dihloro-methane;

THF=Tetrahydrofuron

Disc size: 6.35 mm; Duration: 24-45 h;

Standard: Ampicillin (30-33 mm) and

Streptomycin(20-25 mm); Control:

Methanol; ---: No activities; $\pm$: Active(8-12

mm); +: Moderately active(13-19 mm); ++:

Active(20-24 mm).

TABLE 3: THE OXAZINE-2-AMINES 4-6 SHOWED MAXIMUM ZONE OF INHIBITION AGAINST ESCHERICHIA COLI, WITH GREATER THAN 20 MM COMPARED TO THE OXAZINES 3, 8-10 ARE ACTIVE IN 8-12 MM OF ZONE OF INHIBITION. THE OXAZINES 1, 2, 7, 11-13 ARE INACTIVE.

Entry	X	<i>E. coli</i>	<i>S. aureus</i>	<i>Pseudomonas</i>	<i>Klebsiella</i>	<i>P. vulgaris</i>	<i>E. faecalis</i>
1	H	---	$\pm$	---	$\pm$	---	---
2	3-NH ₂	---	++	+	---	$\pm$	$\pm$
3	4-NH ₂	$\pm$	---	+	++	---	
4	3-Br	++	+	++	++	+	++
5	3-Cl	++	$\pm$	++	+	+	+
6	4-Cl	++	++	++	+	++	++
7	4-N(CH ₃) ₂	---	---	$\pm$	$\pm$	---	$\pm$
8	4-OH	$\pm$	---	$\pm$	$\pm$	$\pm$	$\pm$
9	4-OCH ₃	$\pm$	$\pm$	---	---	$\pm$	---
10	4-CH ₃	$\pm$	$\pm$	$\pm$	$\pm$	$\pm$	$\pm$
11	2-NO ₂	---	+	+	---	$\pm$	---
12	3-NO ₂	---	+	---	$\pm$	---	$\pm$
13	4-NO ₂	---	+	$\pm$	---	+	

TABLE 4: ANTIFUNGAL ACTIVITIES OF 4-(6-METHYL-2-NAPHTHYL)-5,6-DIHYDRO-6(SUBSTITUTED PHENYL)-4H-1,3-OXAZINE-2-AMINES

Entry	X	Disc diffusion technique (250 µg/mL)	Drug dilution method (50 µg/mL)	
		<i>Candida albicans</i>	<i>Penicillium</i>	<i>Aspergillus niger</i>
1	H	---	---	---
2	3-NH ₂	$\pm$	$\pm$	$\pm$
3	4-NH ₂	$\pm$	$\pm$	$\pm$
4	3-Br	---	$\pm$	+
5	3-Cl	+	++	+
6	4-Cl	$\pm$	++	+
7	4-N(CH ₃) ₂	$\pm$	$\pm$	$\pm$
8	4-OH	---	---	---
9	4-OCH ₃	---	---	+
10	4-CH ₃	+	+	---
11	2-NO ₂	++	++	++
12	3-NO ₂	++	++	++
13	4-NO ₂	++	++	++

Standard: Griseofulvin and Gentamycin; Duration : 72 h; Control: Methanol; Medium: Potato dextrose agar; ++: No fungal colony; +: One fungal colony; $\pm$: Two-three fungal colonies; ---: Heavy fungal colony.

4. CONCLUSIONS

Some unsaturated 1,3-oxazine-2-amine derivatives including 4-(6-methyl-2-naphthyl)-5,6-dihydro-6-(substituted phenyl)-oxazine-2-amines have been synthesised by solvent-free cyclization of 6-methyl-2-naphthyl chalcones and urea in presence of hydroxyapatite catalyst under microwave irradiation. The yields of these oxazines were more than 85%. This synthetic methodology offers non-hazardous, solvent-free cyclization, easy-workup procedure, shorter reaction time and better yields. These oxazine derivatives were characterized by their physical constants, spectral data and these data supports the compounds. The antimicrobial activities of these oxazine-2-amines were evaluated using Bauer-Kirby method. The amino,

dimethylamino, methoxy, methyl, dimethyl, chloro, bromo and nitro substituents responded for the antibacterial and antifungal activities.

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التخليق والأنشطة المضادة للميكروبات لبعض

{ 4- (6-ميثيل-2-نفثيل) -5،6-ثنائي هيدرو-6- (فينيل مستبدله) -4 يد- 1،3-اوكسازين-2- أمين }

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الملخص:

بعض { 4- (6-ميثيل-2-نفثيل) -5،6-ثنائي هيدرو-6- (فينيل مستبدله) -4 يد- 1،3-اوكسازين-2- أمين } تم تخليقها باستخدام المذيبات المحفزة الهيدروكسابتية والخالية من حلقات 6-ميثيل-2-نفثيل جالكون واليوربا تحت أشعة الميكروويف، وكانت نواتج الاوكسازين أكثر من 85%. تم تحديد خواص أوكسازين الأمين المخلق والتي تميزت بثبات خواصها الفيزيائية وكذلك بنيانها الطيفية. تم تقييم الأنشطة المضادة للميكروبات لكل التحضيرات { 4- (6-ميثيل-2-نفثيل) -5،6-ثنائي هيدرو-6- (فينيل مستبدله) -4 يد- 1،3-اوكسازين-2- أمين } باستخدام طريقة الإنتشار الباور-كيريبي القرصي مع مجموعة متنوعة من السلالات البكتيرية والفطرية.