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Review on Antimicrobial Activity of Imidazole



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ABSTRACT

Imidazole is two nitrogen-containing heterocycles observed in many natural as well as synthetic compounds. Imidazole has a unique structure that makes it quite abundant in natural compounds like purine, histamine, histamine and nucleic acid. Due to its polar nature Imidazole is quite useful in optimizing the lead structures. Imidazole shows several biological properties including antimicrobial properties. Here we are reporting a review on the antimicrobial property of some Imidazole derivatives.



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INTRODUCTION

Imidazole is a five-member heterocycle having polar nature with water and polar solvent solubility. Imidazole exist in two tautomeric forms as hydrogen on the nitrogen atom can exist on any one of nitrogen atom as shown in figure no 1.

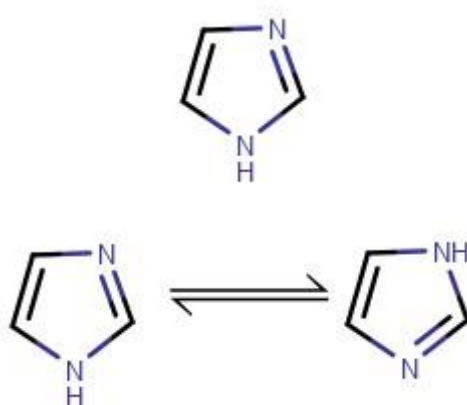
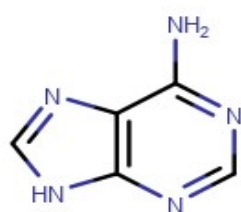
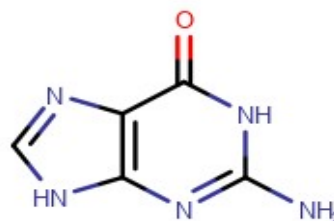


Figure no 1: Imidazole Nucleus

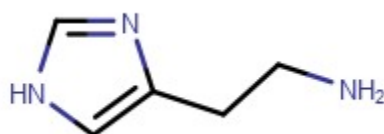
Biological applications of the Imidazole are well established and Imidazole is observed in many biological compounds like nucleic acid and histamine as shown in figure no 2.



Adenine



Guanine



Histamine

Figure no 2: Imidazole nucleus containing natural compounds

Imidazole has been an important component of many medicinal compounds which are shown in figure no 3.

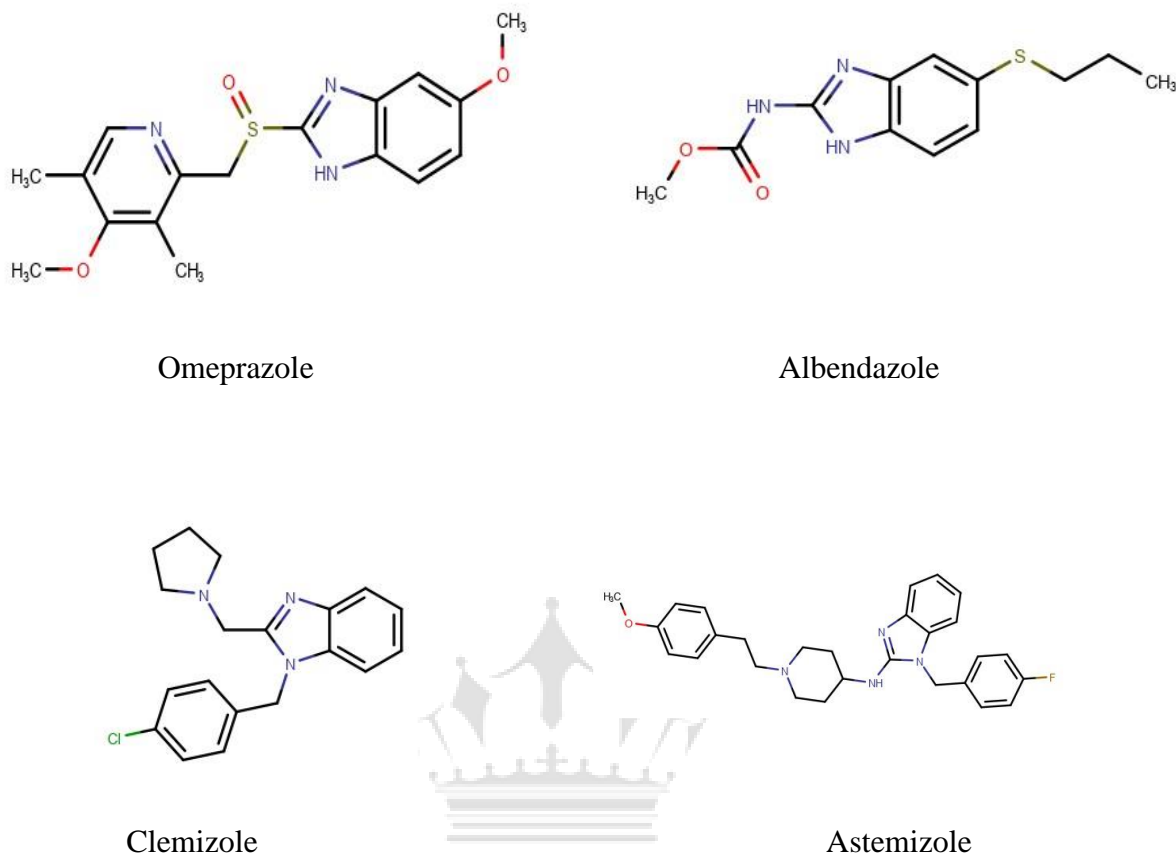
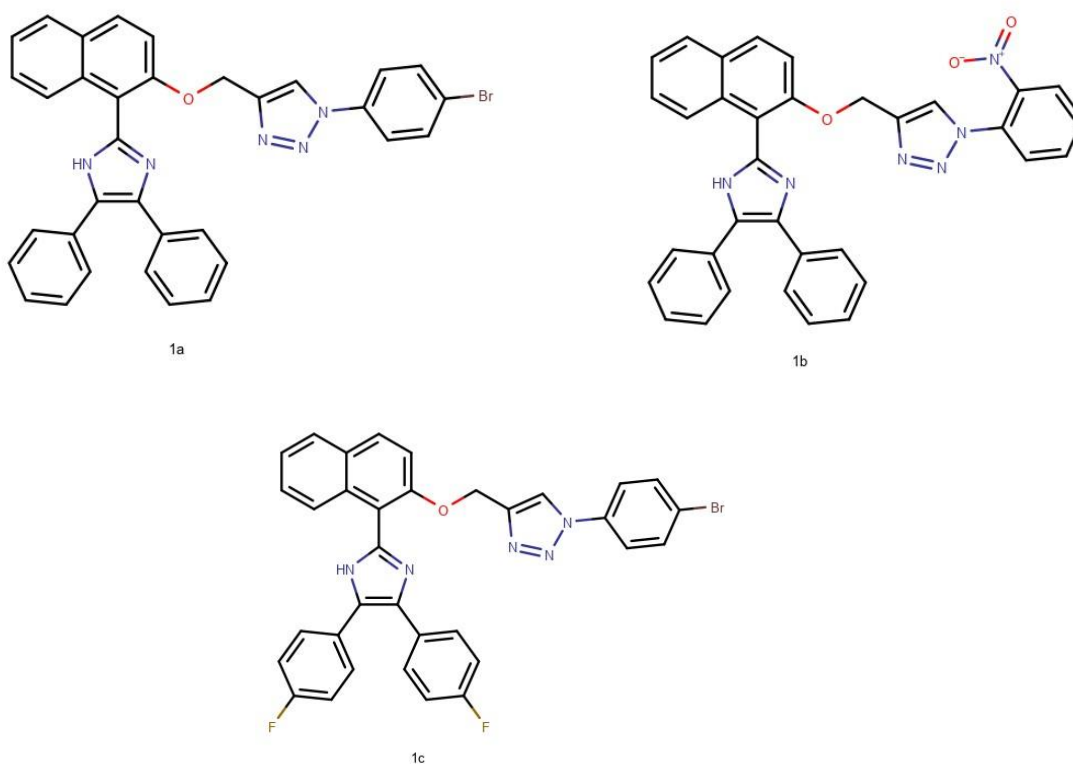


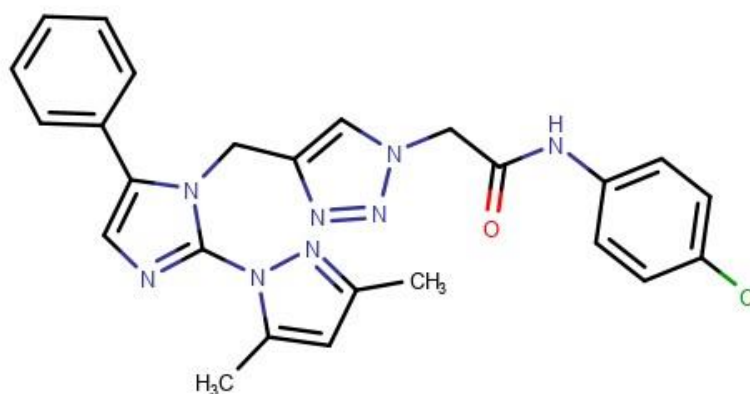
Figure no 3: Benzimidazole nucleus containing drugs

Number of compounds with potent antimicrobial activity with Imidazole as important pharmacophore is reported. Here we have summarized some antimicrobial applications of the imidazole.

Verma et. al. (2019) reported the antimicrobial activity of 2,4,5-Trisubstituted-1H-Imidazole-Triazole . 1-(4-Bromophenyl)-4-(((1-(4,5-diphenyl-1H-imidazol-2-yl) naphthalen-2-yl)oxy)methyl)-1H-1,2,3-triazole (1a), 4-(((1-(4,5-Diphenyl-1H-imidazol-2-yl)naphthalen-2-yl)oxy)methyl)-1-(2-nitrophenyl)-1H-1,2,3-triazole(1b), 4-(((1-(4,5-Bis(4-fluorophenyl)-1H-imidazol-2-yl)naphthalen-2-yl)oxy)methyl)-1-(4-bromophenyl)-1H-1,2,3-triazole (1c) are the potent compounds observed in the series.



Verma et. al. (2020) reported the antimicrobial activity of pyrazole-imidazole-triazole hybrids with their molecular docking. N-(4-Chlorophenyl)-2-(4-((2-(3,5-dimethyl-1H-pyrazol-1-yl)-5-phenyl-1H-imidazole-1-yl)methyl)-1H-1,2,3-triazol-1-yl)acetamide (2) is the potent compounds observed in the series.



2

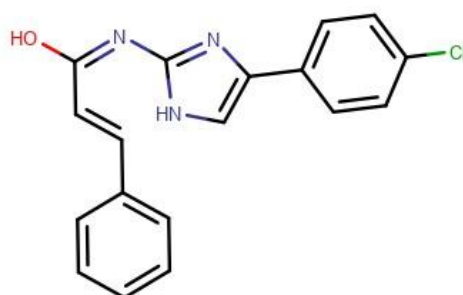
Sta, czek et. al. (2020) reported antimicrobial activity of hydrazones (3) which are synthesized using 3-oxide-1H-imidazole-4-carbohydrazides. 1-Benzyl-5-methyl-N0-[(Z)-(5-nitro-2-furyl)methylidene]-1H-imidazole-4-carbohydrazone 3-oxide, 1,5-Dimethyl-N0-[(Z)-(5-nitro-

2-furyl)methylidene]-1H-imidazole-4-carbohydrazone 3-oxide, 1-Cyclohexyl-5-methyl-N0-[(Z)-(5-nitro-2-furyl)methylidene]-1H-imidazole-4-carbohydrazone 3-oxide was found to be potent compounds from the developed series.



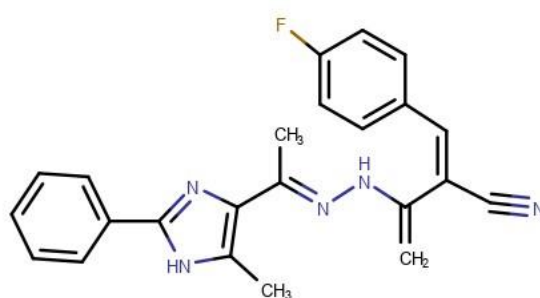
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Amido-linked pyrrolyl and pyrazolyl-oxazoles, thiazoles, and imidazoles have been developed by Padmavathi et. al. (2011). Results indicated chlorosubstituted imidazolyl cinnamate (4) was found to be a lead compound.



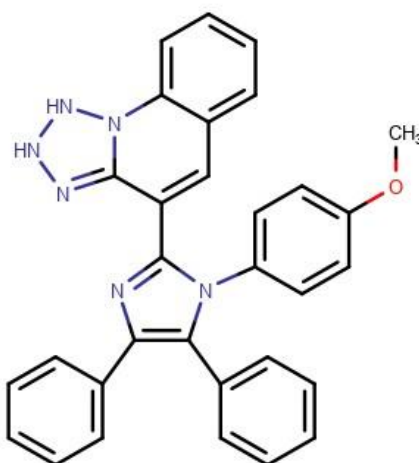
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Abdel-Wahab et. al.(2020) reported the development of imidazole-based heterocycles as antimicrobial, antioxidant, anti-hemolytic, and cytotoxic. 2-Cyano-3-(4-fluorophenyl)-N0-[1-(5-methyl-2-phenyl-1Himidazol-4-yl)ethylidene]acrylohydrazone (5) is the promising antimicrobial agent.



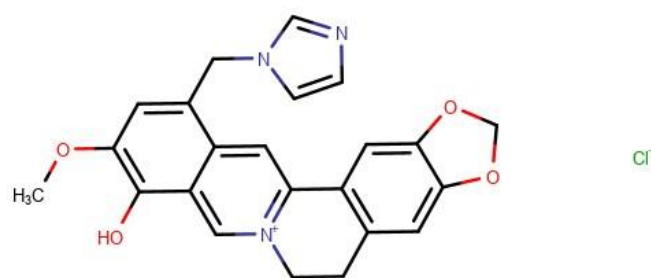
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Kathrotiya et. al. (2012) reported development of the tetrazolo[1,5-a]-quinoline based imidazoles as antimicrobial agents using molecular iodine. 1-(4-methoxyphenyl)-4,5-diphenyl-2-{{1H,2H-[1,2,3,4]tetrazolo[1,5-a]quinolin-4-yl}}-1H-imidazole (6) was found to be more promising antimicrobial agent.



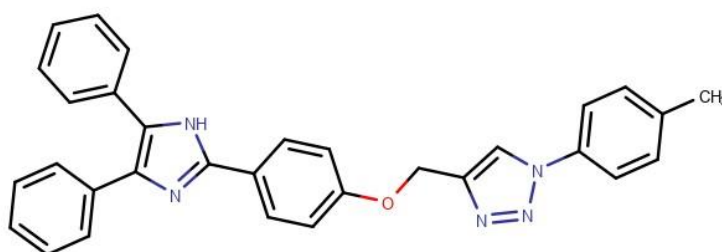
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Cheng-He Zhou et. al. (2016) reported the development of berberine imidazoles as antimicrobial agents. 12-((1H-imidazole-1-yl)methyl)-9-hydroxy-10-methoxy-5,6-dihydro-[1,3]dioxolo[4,5-g]isoquinolino[3,2-a] isoquinolin-7-ium chloride(7) was found to be promising agent.



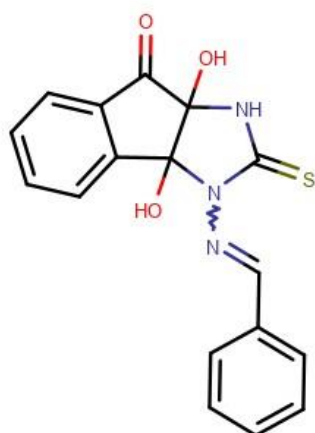
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Verma et. al.(2019) developed triazole containing triaryl-1H-imidazole (8) as antimicrobial agents and reported their docking studies.



8

Saini et. al. (2017) developed 3-(arylideneamino)-3a,8adihydroxy-1,3,3a,8a-tetrahydroindeno[1,2-d]imidazole-2,8-diones and their 2-thioxo derivatives as antimicrobial agents.



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SUMMARY:

Imidazole is one most promising heterocyclic agents utilized as a synthetic intermediate or scaffold. Various Imidazole derivatives have good antimicrobial activity, so the Imidazole nucleus will be attractive heterocycles in antimicrobial drug design.

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