

“Designing Synthesis and Biological Evaluation Of pyridazinone Derivative”

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I. INTRODUCTION

Chemistry is the branch of science in which study of the molecules and their transformation; it deals with their composition, properties of substance element and molecule. Chemistry is the science of matter and it occurs in three forms: solid, liquid and gas.

Pharmaceutical chemistry is a science that gives the information of drug molecules, their preparation, in vivo and vitro studies, the properties of drugs. In this, the study of the method of quality control and the condition of their usage. Or this is also called the chemistry of drugs.

Pharmaceutical chemistry is important for the quality of drug molecules and it should be safe for consumer and products. Pharmaceutical chemistry is the study of chemical properties of compounds. In this, the study of drug designing, synthesizing and developing of drug molecules in pharmaceutical chemistry. This is also called Medicinal chemistry. In which the study, identification, synthesis and development of new chemical drug molecules which have therapeutic activity. It also studies existing drugs and their biological effects and their quantitative structure-activity relationship (QSAR).

Pharmaceutical Chemistry is the science of drugs, in this, the development of drug entities. It also covers drug discovery, delivery and there are parts of biomedical analysis. In this, the study of pharmacokinetics, and pharmacodynamics properties of drug molecules. Pharmaceutical chemistry work is based on the laboratory manual.

The chemistry of heterocyclic compounds is a continuously exploring field of organic chemistry. The synthesis of heterocyclic compounds has attracted the attention of scientists because of the biological importance of them.

Pharmaceutical chemistry involves remedies and cures and for disease conditions, analytical techniques, quality assurance, pharmacology, metabolism, absorption and chemistry of the drug. Pharmaceutical chemistry leads to careers in biotechnology, organic chemistry and drug development, pharmaceutical companies, quality assurance, research facilities, and more.

Chemistry is the science of matter and its structure change properties is an important part of pharmaceutical science and it includes acids and bases. And other organic compounds, chemicals like soap, detergents, dyes, polymers and metals etc. In simple words, the scope of chemistry is everywhere and every time.

Medicinal chemistry is a discipline that covers the design, development, and synthesis of pharmaceutical drug molecules. This is a process in which scientists from chemistry or synthetic organic chemistry, pharmacology, medicinal chemistry and other biological sciences study the drug molecule. It is the evaluation of the properties of drugs.

The use of plants, animals and minerals as medicine from many years ago. New drug discovery methods are useful nowadays for the synthesis of new molecules. New molecules with potential pharmaceutical effects are called hit compounds. These are natural compounds synthesized by computational chemistry and from a screening method.

The hit compound is improved pharmacokinetic, pharmacodynamic, pharmacological effects of compounds from biotechnology. Then identify lead compound. The lead compound is further optimized to be a drug entity that is safe to use in human in clinical trials study.

Heterocyclics have the largest of classical divisions of organic chemistry and are of many importance in biological and industrial fields.

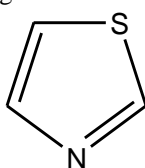
The majority of pharmaceuticals and biologically active agrochemicals are heterocyclic while countless additives and modifiers used in industrial applications ranging from cosmetics, reprography, information storage and plastics are also heterocyclic in nature. One striking structural feature inherent to heterocyclics, which is exploited to great advantage by the drug industry, lies in their ability to prepare substituents around a core scaffold in defined three-dimensional representations.

For more than a century, heterocyclic have constituted one of the largest areas of research in organic chemistry. Heterocyclic have contributed to the development of society from a biological and industrial point of view as well as to the understanding of life processes to improve the quality of life. Among the approximately 20 million chemical compounds identified by the end of the second millennium, more than two-thirds are fully or partially aromatic and approximately half are heterocyclic. (wonder nucleus) which posses almost all types of biological activities

Heterocyclic is a branch of organic chemistry which deals with the cyclic compounds in which at least one hetero atom is a member of its rings. Nitrogen, oxygen and sulphur are the common hetero atoms present in the heterocyclic compounds. Heterocyclic compounds also known as ring compounds and cyclic compounds. Heterocyclic compounds is said as cyclic compounds or ring compounds because its structure having one rings.

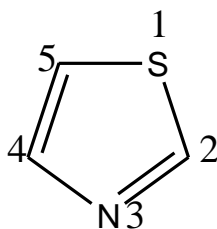
Now a day's heterocyclic compounds having a great demands in the market of pharmaceutical company because of its great importance against the variety of common disease. Therefore day by day interested has been increases for synthesis of heterocyclic compounds. Heterocyclic compounds is biological active compounds. Heterocyclic compounds having antifungal, anti-inflammatory antibacterial, antioxidant, anticonvulsant, anti allergic activity.

Thiazoles is a example of heterocyclic compound. Thiazoles is a five-member heterocyclic compounds in which sulfur and nitrogen atom is the member of the rings atom.



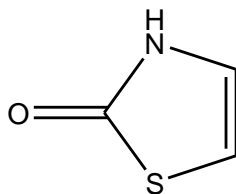
Thiazole

Nomenclature—Thiazole is also known as 1,3-thiazole because it contains sulfur atom in 1 position and nitrogen atom at 3 position in its structure



Thiazole

Derivatives --Thiazolone is an oxygenated derivative of thiazole.



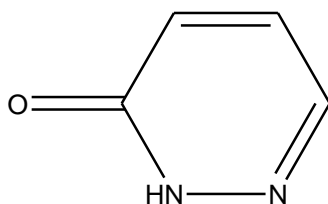
Thiazolone

Importance—thiazolone and its derivatives having a greater importance to curing many chronic diseases especially bacterial, fungal. Bacterial and fungal resistivity is always a challenging problem because of the multi drug resistivity against pathogen is increasing day by day. To overcome these problems is only by formation of new compounds with active nucleus to deal with resistivity against pathogens.

Pyridazinone:

Various pyridazin-3(2H) - ones have attracted considerable attention as they are endowed with a variety of pharmacological activities. These derivatives represent one of the most active classes of compounds possessing a wide spectrum of biological activity ranging from cardiovascular properties, anti-inflammatory, anti-diabetic, antidepressant, analgesic, anti-AIDS, anticancer, antimicrobial and anticonvulsant activities. As a result a number of pyridazinone derivatives have reached clinical trial level as cardio tonic and antihypertensive drug. In present study, a review of different biological activities such as cardiovascular, anti-inflammatory, analgesic, antinociceptive, antiasthmatic, antidiabetic, antidepressant, anticonvulsant, anti-HIV-1, antiproliferative, antimicrobial and insecticidal activities have been dealt in detail. The pyridazinones have attracted a

great deal of attention because of the wide spectrum of their pharmaceutical and agrochemical activities.¹² Pyridazinone are six-member heterocyclic compounds, 2 nitrogen atoms are present at adjacent positions. Pyridazin-3-one, a saturated or unsaturated form of pyridazine with carbonyl group on third carbon, has been considered as a magic moiety (wonder nucleus) which posses almost all types of biological activities.¹³



Pyridazinone

Pyridazine has been assumed to be a planar molecule for which two Kekule structures are given as mentioned. Pyridazine is a resonance hybrid in which the greater contribution is made by the structure containing = NN= configuration. The 1, 2 diazine systems that contain a carbonyl group in the ring are called pyridazinone. This nitrogen heterocycle is included in chemicals with a wide range of biological activities and can also be used to link other pharmacophoric groups. Pyridazine is colorless liquid and boiling point 208 °C, density 1.1015, melting point -8 °C.⁴

The synthesis of novel pyridazinone derivatives and investigation of their chemical and biological activities have gained more importance in recent years. Pyridazinone derivatives have established a variety of pharmacological activities most of them are related to cardiovascular effects. In this field a number of compounds such as zardaverine/imidazole, bemoradan, indolidan, pimobendan are few examples of pyridazinones that are active as cardiostimulant agents/ platelet.¹⁻⁹ Literature survey revealed that substituted pyridazinones have reported to possess

Pharmacological activities such as antidepressant, antihypertensive, antithrombotic, antifungal, antibacterial, antimicrobial, analgesic, anti-inflammatory, anti-feedant, anti-platelet, anticancer, diuretics, anti-HIV, vasodilator and other anticipated biological and pharmacological properties.

Aminopyrine is a classical drug with strong analgesic and anti-inflammatory activities but reported to have side effects. A series of pyridazinone derivatives are structurally related to aminopyrine with a pyrazolone ring. Among these 4-alkoxy-2-methyl-5-morpholino-3(2H)-pyridazinones have been found to have strong analgesic and anti-inflammatory activities. In the present scenario, a large number of medications acting through different mechanisms for the treatment of hypertension are available. Even the one has to admit that blood pressure of majority of hypertensive patients is inadequately controlled, partly because the treatment is not conducted intensively enough, but partly also because the medication(s) are not taken as prescribed. Lowering blood pressure in hypertensive patients requires therefore not only a broad choice of effective and well-tolerated medications, but also skills to motivate them to comply lifelong with the treatment. Hypertension is the most common cardiovascular disease. The definition of hypertension, therefore using any specific cut off point is arbitrary. The studies on the hydralazine group drugs led to the synthesis of many pyridazinone derivatives with a wide activity spectrum on cardiovascular system¹⁻³. Pyridazinone derivatives, a class of compounds containing the N-N bond, exhibit a wide range of pharmacological activities such as antidepressant⁴, antihypertensive⁵⁻⁸, antithrombotic⁹, anticonvulsant¹⁰, cardiostimulant¹¹, antibacterial¹², diuretics¹³, anti-HIV¹⁴ and anticancer¹⁵. Some pyridazinone derivatives like indolidan¹⁶, bemoradan¹⁷, pimobendan¹⁸, levosimendan¹⁹ (antihypertensive), already approved in the clinical market. The current work describes the synthesis of some new substituted pyridazine derivatives with encouraging antihypertensive activity by non-invasive method using Tail Cuff method.

The main objective present research work to synthesis, characterization and biological evaluation of pyridazine derivatives. All the synthesized compounds were obtained in good yield by optimizing various synthetic procedures. The structures of the compounds were established by elemental analysis, IR, ¹H-NMR and Mass spectral data analysis. To study the different synthesized derivative by using different analytical parameters. And also to find out the good pharmacologically active synthesized compounds.

Chemistry of Pyridazinone and Pyridazine:

Pyridazine is one of the three possible isomeric diazines. The diazines are a group of compounds formally derived from benzene by the replacement of two of the ring carbon atom by nitrogen. It was obtained as early as 1886 by Fischer and was first synthesized by Tauber in 1895. The nitrogen atoms pyridazine displays properties different from the isomeric diazine.

Pyridazine is a planer six membered ring and it is the represented as a resonance hybrid of two structure (1a) and (1b) with a greater contribution from the canonical structure (1a) (Figure: 8).

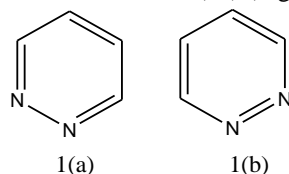


Figure : 8

This is supported by the result of microwave spectroscopy, electron diffraction data & X-ray crystallographic analysis, which all indicate that the N-N bond has single bond character. Bond length and bond angle have been also calculated by microwave spectroscopy, electron diffraction.

Six possible reduced pyridazines, 1, 2(4), 1, 4(5) and 4, 5(6) dihydropyridazines are known (Figure: 9).

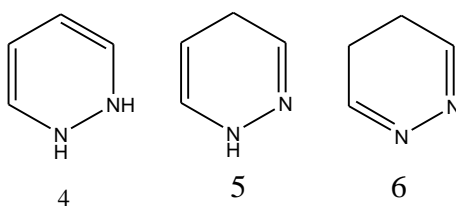


Figure : 9

Appropriately substituted pyridazines exhibits tautomerism. This 3 & 4- hydroxyl pyridazines (7) and (8) exist predominantly in the oxo form.²⁻⁴

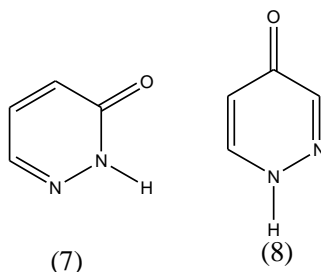


Figure : 10

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