

**BIOLOGICAL IMPORTANCE OF TRIAZOLE DERIVATIVES**

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Article Received on 06/05/2023

Article Revised on 27/05/2023

Article Accepted on 17/06/2023

**ABSTRACT**

Heterocyclic compounds especially triazole derivatives play a very important role in pharmacological fields such as antimicrobial, anticonvulsant, analgesic insecticidal, anticancer, activities. Many researchers and scientists have synthesized triazole derivatives and discussed various biological activities. On the basis of above observation, researches has foccus on synthetic plan of triazole derivatives and evaluation their biological activity.

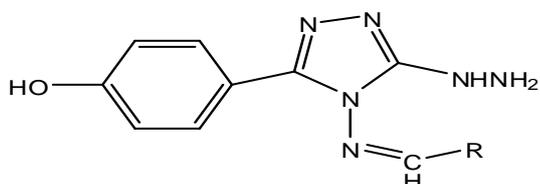
**KEYWORDS:** Triazoles, biological activity, antimicrobial, anti-inflammatory, anticancer activity.

**INTRODUCTION**

Chemical formula of triazole is  $C_2H_3N_3$ . It is five membered rings which contain two carbon and three nitrogen atoms. A special class of heterocyclic compounds of triazole possesses wide range of biological properties. From the earlier study, it is revealed that triazole nucleus possesses different pharmacological properties such as anticonvulsant<sup>[1-2]</sup>, antimicrobial<sup>[3-10]</sup>, anticancer<sup>[11-13]</sup>, antituberculosis<sup>[14]</sup>, antibacterial<sup>[15-19]</sup>, antifungal<sup>[20-23]</sup>, anti-inflammatory<sup>[24-27]</sup>, antioxidant<sup>[28]</sup> activity etc. Some common drugs such as fluconazole, terconazole, nefazodone, rizatripton, etoperidone, posaconazole, anastrozole etc. are the example of standard drugs which contain triazole nucleus. This review article focused the work on triazoles that explained biological activity reported by many scientists in field of chemistry.

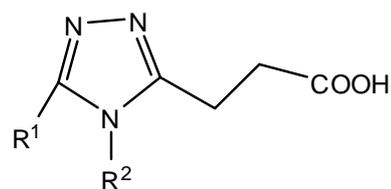
**BIOLOGICAL ASPECTS OF TRIAZOLES**

Synthesis, characterization and biological activity of some novel triazole derivatives have been synthesized by Arora et al.<sup>[29]</sup> and screened for antimicrobial activity. Antibacterial activity evaluated against *E. coli*, *B. pseudomonas aeruginosa*, *C. bacillus subtili*, *D. staphylococcus aureus* and compared with ciprofloxacin. These compounds also showed antifungal activity against *A. niger*.



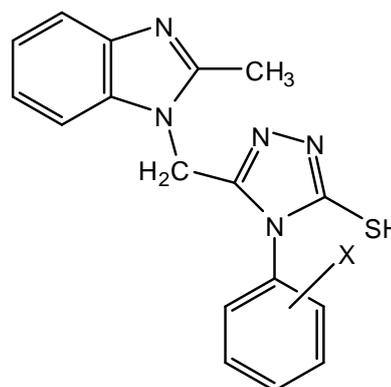
R = Benzaldehyde, salicylaldehyde, anisaldehyde, p-hydroxybenzaldehyde, m-nitrobenzaldehyde, p-dimethylaminobenzaldehyde, formaldehyde.

Paprocka et al.<sup>[30]</sup> have synthesized new 1,2,4-triazole derivatives containing propionic acid moiety and evaluated biological activity of these compounds such as antimicrobial, anthelmintic and anti-inflammatory activity.



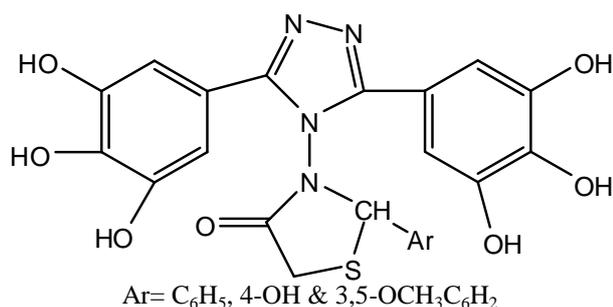
R<sup>1</sup> = 2-pyridyl, 4-pyridyl, Ph  
R<sup>2</sup> = 2-pyridyl, Ph, 4-PhCH<sub>3</sub>, 4PhNO<sub>2</sub>

Ansari et al.<sup>[31]</sup> have prepared synthesis and biological activity of some triazole bearing benzimidazole derivatives and screened for antimicrobial activity against different bacteria and fungi then compared by standard drugs ampicilline and amphotericin B.

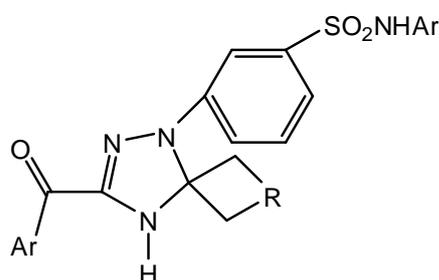


X = 2-CH<sub>3</sub>, 2-C<sub>2</sub>H<sub>5</sub>, 2-C<sub>3</sub>H<sub>7</sub>, 2-OH, 4-OH, 4-OCH<sub>3</sub>, 2-Cl, 4-Cl, 4-NH<sub>2</sub>

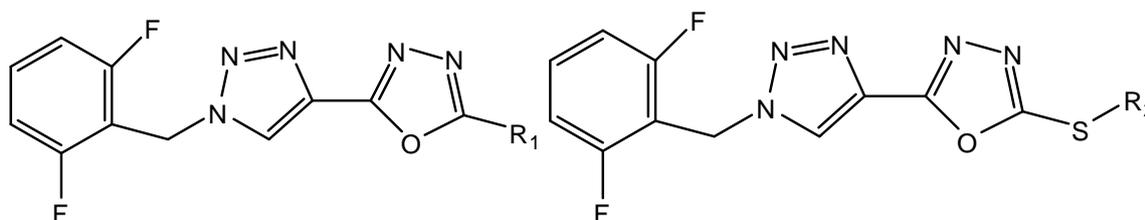
F.A. Nashaan and M.S. Al-Rawi<sup>[32]</sup> have synthesized Design, synthesis and biological activity of new thiazolidine-4-one derived from symmetrical 4-amino 1,2,4-triazole.



Dalloul et. al.<sup>[33]</sup> have synthesized spiro1,2,4-triazole derivatives and showed antimicrobial activity against various bacteria *E. coli*, *S. aureus*, *K. Protius* then compared with tetracycline and various fungi *C. albicans*, *A. niger* then compared with fluconazole.



Ar/R = (a) 2-thiazolyl / (CH<sub>2</sub>)<sub>2</sub>, (b) 2-pyrimidinyl / (CH<sub>2</sub>)<sub>2</sub>, (c) thiazolyl / (CH<sub>2</sub>)<sub>3</sub>, (d) 2-pyrimidinyl / (CH<sub>2</sub>)<sub>3</sub>, (e) 2-thiazolyl / (CH<sub>2</sub>)<sub>4</sub>, (f) 2-pyrimidinyl / (CH<sub>2</sub>)<sub>5</sub>, (g) 2-thiazolyl / CH<sub>2</sub>CHMeCH<sub>2</sub>, (h) 2-pyrimidinyl / CH<sub>2</sub>CHMeCH<sub>2</sub>, (i) 2-thiazolyl / CH<sub>2</sub>CH(t-bu)CH<sub>2</sub>, (j) 2-pyrimidinyl / CH<sub>2</sub>CH(t-bu)CH<sub>2</sub>  
Ar = CH<sub>3</sub>, C<sub>2</sub>H<sub>5</sub>, 2-furyl, 2-thianyl, 2-Naphthyl



R<sub>1</sub> = Phenyl, 2-tolyl, 3-chlorophenyl, 4-chlorophenyl, 4-fluorophenyl, 2 & 4-dichlorophenyl, 3 & 4-dimethoxyphenyl  
R<sub>2</sub> = Ethyl, isopropyl, n-butyl, isopentyl, benzyl

## CONCLUSION

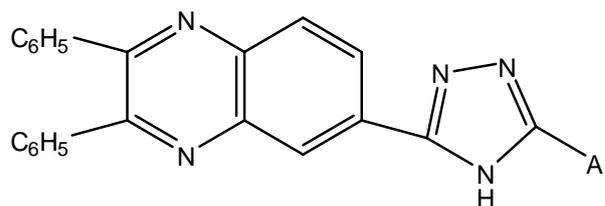
Triazole nucleus is a part of heterocyclic compounds which plays an important role in preparation of drugs. In this review, we focused on biological importance of triazole derivatives such as anti-inflammatory, anticonvulsant, antibacterial, antimicrobial, anticancer activities etc. The potential power triazole derivatives in

Synthesis and biological evaluation of 3-amino-1,2,4-triazole derivatives as potential anticancer compounds have been synthesized by Grytsai et al.<sup>[34]</sup>



R = H, 2-CH<sub>3</sub>, 2-OH, 2-OC<sub>2</sub>H<sub>5</sub>, 2-NO<sub>2</sub>, 2-Cl, 2-Br, 3-NO<sub>2</sub>, 3-Br, 4-CH<sub>3</sub>, 4-Cl, 2,5-diCl, 3,4-diCl, 3,4-di(CH<sub>3</sub>), 2-OH-5-CH<sub>3</sub>, 2-OH-6-CH<sub>3</sub>, 2-OH-4-NO<sub>2</sub>, 2-OH-4-Cl, 2-OMe-5-NO<sub>2</sub>, 2-Cl-5-NO<sub>2</sub>, 2,4-diCl

Synthesis and biological evaluation of some novel quinoxaliny triazole derivatives have been prepared by Mazumder et al.<sup>[35]</sup> They showed antifungal activity against *C. albicans* and *A. niger* then compared with fluconazole.



Ar = C<sub>6</sub>H<sub>5</sub>, 3-NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>, 3-OCH<sub>3</sub>C<sub>6</sub>H<sub>4</sub>, 3-OCH<sub>3</sub>&4-OHC<sub>6</sub>H<sub>3</sub>, 4-FC<sub>6</sub>H<sub>4</sub>, 4-ClC<sub>6</sub>H<sub>4</sub>, 2-ClC<sub>6</sub>H<sub>4</sub>, 4-OCH<sub>3</sub>C<sub>6</sub>H<sub>4</sub>, 2,4-Cl<sub>2</sub>C<sub>6</sub>H<sub>3</sub>, 3,4,5(OCH<sub>3</sub>)C<sub>6</sub>H<sub>2</sub>

Jadhav et al.<sup>[36]</sup> have prepared 1,4-disubstituted 1,2,3-triazole derivatives and evaluated biological activity as antimicrobial agents. Antibacterial activity screened against different bacteria such as *E. coli*, *P. aeruginosa*, *B. subtilis*, *S. pyogenes*, *K. pneumonia*, *S. aureus*, *K. terrigena* and compared by chloramphenicol. Antifungal activity evaluated against various fungi such as *C. albicans*, *T. viride*, *A. flavus*, *A. brasiliense* and compared by nystatin.

field of pharmacological property, we can plan for the synthesis and evaluation of new drugs of triazole derivatives by possessing of possible substituent.

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