

RNI No. RAJHIN/2007/26996
ISSN 0976-7452 Raaso
Refreed Journal

रासो

अन्तर्विषयी त्रैमासिक शोध पत्रिका

वर्ष-18, अंक-2, मार्च अप्रैल मई 2025





RNI No. RAJHIN/2007/26996
ISSN 0976-7452Raaso
Refreed Journal

रासो

इतिहास, संस्कृति, साहित्य एवं दर्शन की त्रैमासिक शोध पत्रिका

वर्ष-18

अंक - 2

मार्च-अप्रैल-मई-2025

सम्पादक

डॉ. दिनेश चन्द्र शर्मा

निदेशक

कॉम्पिटिशन प्लस संस्थान, ग्रुप ऑफ इन्स्टीट्यूशन्स, अजमेर (राजस्थान)

संरक्षक

* प्रो. जे. पी. शर्मा - पूर्व कुलपति, मोहनलाल सुखाड़िया विश्वविद्यालय, उदयपुर (राजस्थान)

सलाहकार मण्डल

- * प्रो. विभा उपाध्याय - प्रोफेसर, इतिहास एवं भारतीय संस्कृति विभाग, राजस्थान विश्वविद्यालय, जयपुर (राज.)
- * नर्मदा प्रसाद उपाध्याय - प्रसिद्ध साहित्यकार तथा संस्कृति चिंतक, इन्दौर (मध्य प्रदेश)
- * प्रो. टी.के. माथुर - पूर्व अध्यक्ष, इतिहास विभाग महर्षि दयानन्द सरस्वती विश्वविद्यालय, अजमेर (राज.)
- * डॉ. रमेश अग्रवाल - पूर्व स्थानीय सम्पादक, दैनिक भास्कर, अजमेर (राजस्थान)
- * डॉ. गौरव बिस्सा - सहआचार्य, विभागाध्यक्ष, प्रबंध अध्ययन विभाग, राजकीय अभियांत्रिकी महाविद्यालय, बीकानेर (राज.)
- * डॉ. बीना शर्मा - पूर्व विभागाध्यक्ष, हिन्दी विभाग, सावित्री कन्या महाविद्यालय, अजमेर
- * प्रो. शोभा आर. मिश्रा - आचार्य एवं विभागाध्यक्ष, दर्शनशास्त्र विभाग, माधव महाविद्यालय, उज्जैन
- * डॉ. वेद प्रकाश विद्यार्थी - पूर्व सहायक निदेशक, केन्द्रीय हिन्दी निदेशालय, नई दिल्ली।

☆ उप सम्पादक

डॉ. रीता पारीक

प्राचार्य

हुकुमचन्द नोबल इंस्टीट्यूट ऑफ साइंस एण्ड टेक्नोलॉजी, अजमेर

☆ प्रबन्ध सम्पादक

डॉ. मृत्युंजय शर्मा

सह-आचार्य

हुकुमचन्द नोबल इंस्टीट्यूट ऑफ साइंस एण्ड टेक्नोलॉजी, अजमेर

☆ सहायक सम्पादक

डॉ. निष्काम शर्मा

☆ शब्द संयोजन

मुकेश कुमार माहेश्वरी

☆ सम्पादकीय/सचिव

व्यवस्थापकीय कार्यालय

सृजन संस्थान,

आचमन, वेद विहार,

लोहाखान, अजमेर-06

दूरभाष : 0145-2426610

e-mail : rasoajmjournal@gmail.com

☆ प्रकाशक :

पुष्पा शर्मा

सृजन संस्थान,

द्वारा आचमन पब्लिकेशन्स,

अजमेर

☆ मुद्रक :

मैसर्स आर्य प्रिन्टर्स,

केसरगंज, अजमेर

☆ कम्प्यूटराइज्ड सेटिंग :

श्रीनिवास प्रिन्टोग्राफिक्स

132, पंचवटी कॉलोनी,

अजमेर दूरभाष : 0145-2442701

सहयोग दर

* वार्षिक सदस्यता - 500 रु. (चार अंक)

संस्थाओं हेतु - 600 रु. (चार अंक)

कृपया बैंक/ड्राफ्ट/मनीऑर्डर श्रीमती पुष्पा शर्मा, प्रकाशक, आचमन पब्लिकेशन्स,
अजमेर- 305006 के नाम भेजें।

प्रकाशक-श्रीमती पुष्पा शर्मा, 51/1509, वेद विहार, लोहाखान, अजमेर द्वारा मैसर्स सृजन संस्थान, अजमेर
की ओर से प्रकाशित एवं मुद्रक-श्रीमती उषा गर्ग द्वारा मैसर्स आर्य प्रिन्टर्स, केसरगंज, अजमेर से मुद्रित।

CHEMICAL DEVELOPMENT OF ANTIBIOTICS AND THEIR IMPACT

Ashwani Sharma

Asstt. Professor (Science & Technology Department)

Hukumchand Noble Institute of Science & Technology, Ajmer

Introduction

The advent of antibiotics stands as one of the most significant milestones in medical science. Before their discovery, infections like pneumonia, tuberculosis, and sepsis were often fatal. Antibiotics, derived from natural sources or synthesized chemically, are compounds that inhibit or destroy the growth of microorganisms-particularly bacteria.

The term "antibiotic" was first coined by Selman Waksman in 1942, referring to substances produced by microorganisms that inhibit the growth of others. However, with advancements in chemical synthesis, many antibiotics today are fully or partially synthetic, offering enhanced potency, stability, and broader activity.

The chemical development of antibiotics not only revolutionized medicine but also profoundly impacted agriculture, industry, and global health outcomes.

"Antibiotic" is from antibiosis, meaning against life. Substances derived from a microorganism or produced synthetically (Sulfonamides & Quinolones) to kill or suppress the growth of other microorganisms.

Classification of Antibiotics

Antibiotics are classified by several ways:

- 1 On the basis of mechanism of action
- 2 On the basis of spectrum of activity
- 3 On the basis of mode of action.

Early History of Antibiotic Discovery

The foundation of antibiotic chemistry began with Alexander Fleming's discovery of penicillin in 1928. Fleming observed that the *Penicillium notatum* mold secreted a substance that inhibited *Staphylococcus* bacteria. However, it was not until the early 1940s that Howard Florey and Ernst Chain successfully purified and mass-produced penicillin, marking the dawn of the antibiotic era.

शोध-पत्र में उल्लिखित सन्दर्भ एवं विचारों के लिए लेखक स्वयं उत्तरदायी है, सम्पादक नहीं।

Following this discovery, the 1940s-1960s became known as the "Golden Age of Antibiotics," characterized by the discovery of many new antibiotic classes such as:

Streptomycin (the first effective treatment for tuberculosis)

* Tetracyclines

* Chloramphenicol

* Erythromycin

* Cephalosporins

These discoveries were made possible through systematic screening of soil microorganisms and the application of organic and analytical chemistry to isolate and modify active compounds.

Chemical Structure and Classification of Antibiotics

Antibiotics are structurally diverse compounds, classified based on their chemical backbone and mechanism of action. Key classes include:

(1) β -Lactam Antibiotics- Contain a β -lactam ring structure essential for activity.

Examples: Penicillins, Cephalosporins, Carbapenems.

Mechanism: Inhibit bacterial cell wall synthesis by binding to penicillin-binding proteins (PBPs).

(2) Aminoglycosides Contain amino sugars linked by glycosidic bonds.

Example: Streptomycin, Gentamicin.

Mechanism: Bind to the 30S ribosomal subunit, interfering with protein synthesis.

(3) Tetracyclines- Four-ring (tetracyclic) structure.

Mechanism: Block tRNA binding to the 30S ribosome, halting protein synthesis.

(4) Macrolides- Characterized by large macrocyclic lactone rings.

Example: Erythromycin, Azithromycin.

Mechanism: Inhibit protein synthesis by binding to the 50S ribosomal subunit.

(5) Quinolones and Fluoroquinolones- Synthetic compounds with a quinolone nucleus.

Mechanism: Inhibit DNA gyrase and topoisomerase IV, blocking bacterial DNA replication.

(6) Glycopeptides- Large, complex molecules (e.g., Vancomycin).

Mechanism: Inhibit cell wall synthesis by binding to D-Ala-D-Ala termini of peptidoglycan precursors.

Each chemical structure provides unique pharmacological properties, allowing scientists to modify functional groups for better absorption, resistance prevention,

Classes of Antibiotics -main classes of antibiotics are:

- * Beta-Lactams
- * Macrolides & Ketolides
- * Penicillins
- * Aminoglycosides
- * Cephalosporins
- * Fluoroquinolones
- * Carbapenems
- * Tetracyclines, Amphenicols
- * Monobactams reduced toxicity.
- * Development of Antibiotic Resistance

A major challenge that emerged from antibiotic usage is bacterial resistance. Resistance arises through genetic mutations or acquisition of resistance genes via plasmids. Key mechanisms include:

- * Enzymatic degradation (e.g., b-lactamases breaking the b-lactam ring)
- * Altered target sites (mutation in ribosomal proteins)
- * Efflux pumps expelling the drug from bacterial cells
- * Reduced permeability of bacterial membranes
- * The overuse and misuse of antibiotics in medicine and agriculture have accelerated resistance evolution.
- * Multidrug-resistant pathogens.

Antibiotic Resistances and Cross Resistances

Antibiotic resistance is the phenomenon that susceptibility of pathogenic microorganisms to antibiotic becomes lower or even loses after the microorganisms contact with antibiotic many times. When the bacteria show resistance to one antibiotic, they are also resistant to some other antibiotics. This phenomenon is called cross antibiotic resistance.

Environmental Impact of Antibiotics

The widespread use of antibiotics has led to environmental contamination through pharmaceutical waste, hospital effluents, and agricultural runoff.

Antibiotic residues in soil and water promote the emergence of resistant bacteria in the environment. Therefore, green chemistry approaches are being developed to synthesize biodegradable antibiotics and improve waste treatment systems.

Recent Advances and Future Directions

Modern antibiotic research combines chemical synthesis, genomics, and artificial intelligence.

Promising trends include:

Novel antibiotic scaffolds discovered through computational modeling and genome mining.

Antimicrobial peptides (AMPs) as alternatives to conventional antibiotics.

CRISPR-based antibacterial agents for targeted bacterial gene disruption.

Phage therapy in combination with chemical antibiotics.

Synthetic biology for designing microbes that produce new antibiotic molecules.

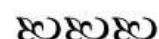
Additionally, the concept of antibiotic stewardship encourages responsible use of these vital drugs to sustain their effectiveness.

Conclusion

The chemical development of antibiotics stands as a triumph of scientific innovation and human ingenuity. From the natural compounds discovered in soil microbes to sophisticated synthetic derivatives designed in laboratories, antibiotics have saved countless lives and reshaped modern medicine. However, the misuse and overuse of these powerful agents have triggered a global crisis of antibiotic resistance.

The future of antibiotic development lies in multidisciplinary approaches -combining chemistry, biology, data science, and environmental sustainability- to create next-generation antimicrobials. Through continued research and responsible application, antibiotics will remain a cornerstone of healthcare and an enduring testament to the power of in improving life on Earth.

References



1. Fleming, A. (1929). *On the Antibacterial Action of Cultures of a Penicillium*. *British Journal of Experimental Pathology*, 10(3), 226-236.
2. Waksman, S. A. (1942). *History of the Word Antibiotic*. *Journal of Bacteriology*, 52(3), 141-148.
3. Katzung, B. G., et al. (2021). *Basic and Clinical Pharmacology* (15th ed.).

McGraw-Hill Education.

4. Walsh, C. (2003). *Antibiotics: Actions, Origins, Resistance*. ASM Press.
5. Wright, G. D. (2012). *Antibiotics: A New Hope*. *Communications in Chemistry*, 3(6), 15-22.
6. Demain, A. L., & Elander, R. P. (1999). *The Beta-Lactam Antibiotics: Past, Present, and Future*. *Antonie van Leeuwenhoek*, 75(1), 5-19.
7. Ventola, C. L. (2015). *The Antibiotic Resistance Crisis: Causes and*
8. *Threats*. *Pharmacy and Therapeutics*, 40(4), 277-283.
9. Chopra, I., & Roberts, M. (2001). *Tetracycline Antibiotics: Mode of Action and Resistance Mechanisms*. *Microbiology and Molecular Biology Reviews*, 65(2), 232-260.
10. Silver, L. L. (2011). *Challenges of Antibacterial Discovery*. *Clinical Microbiology Reviews*, 24(1), 71-109.
11. World Health Organization (WHO). (2022). *Global Antimicrobial Resistance and Use Surveillance Report*.