



INDIAN JOURNAL OF PRACTICAL PEDIATRICS



- **IJPP is a quarterly subscription journal of the Indian Academy of Pediatrics committed to presenting practical pediatric issues and management updates in a simple and clear manner**
- **Indexed in EM Care, CABI Publishing, Scopus**

Vol.24 No.1 (JAN.- MAR.) 2022

Dr.S.Thangavelu
Editor-in-Chief

Dr.T.L.Ratnakumari
Executive Editor

CONTENTS

TOPIC OF INTEREST - "ANTIMICROBIALS - I"

Overview of antibiotics	5
Dhanya Dharmapalan	
Pharmacodynamics and pharmacokinetics of antimicrobial therapy - Clinical application	11
Bhupendra Kumar Gupta, Sudhir Mishra	
Side effects of antimicrobials and preventive strategies	17
Bindusha S, Santhosh Kumar A	
Antimicrobials for perinatal and neonatal infections	29
Suman Rao PN, Swapnik Kandepi	
Choice of Anti-staphylococcal therapy	37
Upendra S Kinjawadekar, Rajendra Vaidya	
Antituberculous therapy - Current practice	44
Gowrishankar NC	
Antimalarial drug therapy	52
Ritabrata Kundu, Aniruddha Ghosh	
Rational antimicrobial therapy in office practice	58
Palaniraman R	
Prophylactic antimicrobials	61
Padmasani Venkat Ramanan, Shalini Sharma	
Newer antibiotics	70
Dhanalakshmi K, Lakshan Raj S	
GENERAL ARTICLE	
Career guidance for pediatricians	80
Julius Xavier Scott, Nisha Kalaiarasan	

Journal Office and address for communications: Dr. S.Thangavelu, Editor-in-Chief, Indian Journal of Practical Pediatrics, 1A, Block II, Krsna Apartments, 50, Thamizh Salai (Halls Road), Egmore, Chennai - 600 008. Tamil Nadu, India. Tel.No. : 044-28190032 E.mail : ijpp_iap@rediffmail.com

DRUG PROFILE**Therapy of acne vulgaris 86**

Jeerson C. Unni

RADIOLOGY**Introduction to chest X-ray 91**

Kasivisalakshi KP, Raveendran J, Tamilarasi V, Babu M

CASE REPORT**Malignant infantile osteopetrosis 96**

Preethi N, Latha Kanchi Parthasarathy, Ramya Uppuluri, Devimeenakshi K

Sporadic hemiplegic migraine in a child 98

Arulmozhi T, Suba Rajinikanth

CASE VIGNETTE**An unusual case of foreign body aspiration in an infant 100**

Anil Kumar P, Vishnu Priya R

Orchitis as an unusual manifestation in multisystem inflammatory syndrome in children 102

Kalpana S, Shobha SR

LEARNING TOGETHER – OSCE 103**PICTURE QUIZ 109****ANNEXURE 110****ADVERTISEMENTS 113,116,117,118****NEWS AND NOTES 16,69,101,102****CLIPPINGS 10,36,43,51,69,79,85,101,108****FOR YOUR KIND ATTENTION**

- * The views expressed by the authors do not necessarily reflect those of the sponsor or publisher. Although every care has been taken to ensure technical accuracy, no responsibility is accepted for errors or omissions.
- * The claims of the manufacturers and efficacy of the products advertised in the journal are the responsibility of the advertiser. The journal does not own any responsibility for the guarantee of the products advertised.
- * Part or whole of the material published in this issue may be reproduced with the note "Acknowledgement" to "Indian Journal of Practical Pediatrics" without prior permission.
- * The write up should be in accordance with the recommendations of Central IAP particularly with issues involving National Programmes like Immunization, Public Health Programs and Nutrition.
- * NOTE: Many trade names of the vaccines are included in the text for the sake of clarity.

EDITORIAL BOARD

Published by Dr. S.Thangavelu, Editor-in-Chief, IJPP, on behalf of Indian Academy of Pediatrics, from 1A, Block II, Krsna Apartments, 50, Thamizh Salai (Halls Road), Egmore, Chennai - 600 008. Tamil Nadu, India and Printed by Mr. D.Ramanathan, at Alamu Printing Works, 9, Iyyah Street, Royapettah, Chennai - 600 014.

ANTIMICROBIALS - I

OVERVIEW OF ANTIBIOTICS

***Dhanya Dharmapalan**

Abstract: Antibiotics are used for treating bacterial infections. They are classified based on origin (synthetic or natural), mechanism of action, type of action (bacteriostatic or bactericidal) and spectrum (narrow or broad). Majority of antibiotics used in clinical practice such as cephalosporins, carbapenems, fluoroquinolones have broad spectrum of activity. Broad spectrum antibiotics have disadvantages of alteration of host microbiome and selection of resistance. To overcome the problem of antibiotic resistance, antibiotic surveillance and antibiotic stewardship measures were recognized as a policy by the World Health Organization, as a component of which a new classification of antibiotics called as Access, Watch, Reserve (AWaRe) was introduced. This classification helps in guiding the selection of antibiotic and prevention of their abuse and overuse.

Keywords: Antibiotics, Spectrum, Access Watch

Reserve Classification.

Points to Remember

- **First generation cephalosporins have narrow spectrum and possess strong activity against Gram positive bacteria.**
- **Beta lactam / beta-lactamase inhibitors cross the blood brain barrier sub-optimally and hence should not be used in treating meningitis.**
- **Moxifloxacin and Levofloxacin should be reserved for use in multidrug resistant TB.**
- **Recent evidence suggest a similar efficacy for bacteriostatic and bactericidal antibiotics in treating infections.**
- **Narrow spectrum antibiotics should always be chosen over broad spectrum antibiotics whenever causative infection is identified.**
- **AWaRe classification of antibiotics by the WHO comprises of three categories - Access, Watch and Reserve, mainly based on the antibiotic resistance threshold.**
- **Antibiotic use as first and second line agents for common infections, needs to be improved from the 'Access' category of AWaRe Classification.**

References

1. WHO. The 2019 WHO AWaRe classification of antibiotics for evaluation and monitoring of use. Geneva: World Health Organization; 2019. (WHO/EMP/IAU/2019.11). Available from <https://www.who.int/publications/i/item/WHOEMPIAU2019.11>. Accessed on 20 February 2022.
2. Sarkar P, Yarlagadda V, Ghosh C, Haldar J. A review on cell wall synthesis inhibitors with an emphasis on glycopeptide antibiotics. *Medchemcomm* 2017; 8(3):516-533. Published 2017 Jan 26. doi:10.1039/c6md00585c
3. Bui T, Preuss CV. Cephalosporins. [Updated 2021 Aug 31]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2022 Jan-. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK551517/> Accessed on 20 February 2022.
4. Papp-Wallace KM, Endimiani A, Taracila MA, Bonomo RA. Carbapenems: past, present, and future.

* Consultant in Pediatric Infectious Diseases,
Apollo Hospitals,
Navi Mumbai.
email: drdhanyaroshan@gmail.com

- Antimicrob Agents Chemother 2011; 55(11):4943-4960. doi:10.1128/AAC.00296-11.
5. Zavascki AP, Goldani LZ, Li J, Nation RL. Polymyxin B for the treatment of multidrug-resistant pathogens: a critical review. *J Antimicrob Chemother* 2007; 60(6):1206-1215. doi: 10.1093/jac/dkm357. Epub 2007 Sep 17. PMID: 17878146.
 6. Krause KM, Serio AW, Kane TR, Connolly LE. Aminoglycosides: An Overview. *Cold Spring Harb Perspect Med* 2016; 6(6):a027029. Published 2016 Jun 1. doi:10.1101/cshperspect.a027029.
 7. Shutter MC, Akhondi H. Tetracycline. [Updated 2022 Jan 19]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2022 Jan-. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK549905>. Accessed on 20 February 2022.
 8. Hooper DC. Mechanisms of action and resistance of older and newer fluoroquinolones. *Clin Infect Dis* 2000; 31 Suppl 2:S24-28. doi: 10.1086/314056. PMID: 10984324.
 9. Patel K, Goldman JL. Safety Concerns Surrounding Quinolone Use in Children. *J Clin Pharmacol* 2016; 56(9):1060-1075. doi:10.1002/jcph.715
 10. Löfmark S, Edlund C, Nord CE. Metronidazole is still the drug of choice for treatment of anaerobic infections. *Clin Infect Dis* 2010; 50 Suppl 1:S16-23. doi: 10.1086/647939. PMID: 20067388.
 11. Loeffler AM. Uses of rifampin for infections other than tuberculosis. *Pediatr Infect Dis J* 1999; 18(7):631-632. doi: 10.1097/00006454-199907000-00012. PMID: 10440440.
 12. Wald-Dickler N, Holtom P, Spellberg B. Busting the Myth of “Static vs Cidal”: A Systemic Literature Review. *Clin Infect Dis* 2018; 66(9):1470-1474. doi: 10.1093/cid/cix1127. PMID: 29293890; PMCID: PMC5905615.
 13. Melander RJ, Zurawski DV, Melander C. Narrow-Spectrum Antibacterial Agents. *Med Chem Comm* 2018; 9(1):12-21. doi:10.1039/C7MD00528H.

ANTIMICROBIALS - I

PHARMACODYNAMICS AND PHARMACOKINETICS OF ANTIMICROBIAL THERAPY - CLINICAL APPLICATION

***Bhupendra Kumar Gupta**
****Sudhir Mishra**

Abstract: Emergence of antimicrobial resistance and poor clinical outcome consequently, are mostly because of inappropriate drug choice and suboptimal dosing. Strategies for better clinical outcomes include selection of an appropriate antibiotic and optimization of antimicrobial dosing regimen. Hence, one must primarily understand the antimicrobial pharmacodynamics and pharmacokinetics of a particular drug to decide on the dosing regimen. Pharmacodynamics denotes the mechanism of action of the drug such as a drug's molecular, biochemical and physiologic effects. Pharmacokinetics deals with absorption, distribution, metabolism and excretion of the drug simplified and abbreviated as ADME.

Keywords: Pharmacodynamics, Pharmacokinetics,

Minimal inhibitory concentration, Antimicrobial resistance.

Points to Remember

- *Antimicrobial therapy for the treatment of infection not only require appropriate choice of antimicrobial agent but also appropriate dosing regimen, route and duration. To decide this, site of infection, type of bacteria and age of patients are other important variables.*
- *Dose optimization of antimicrobial agent requires understanding of pharmacokinetic and pharmacodynamic properties.*
- *Antimicrobial efficacy of concentration dependent antimicrobial agents (e.g. aminoglycoside, quinolones) depends on the peak concentration (C max) to MIC ratio.*
- *Antimicrobial efficacy of time dependent antimicrobial agents (e.g. β -lactam, vancomycin) depends on the percentage of time that free plasma concentration of antimicrobial agent is maintained above MIC.*
- *Dose adjustment is required in certain situations, renal impairment being the most important of them.*

References

1. Manjunath P Pai, Mackenzie L Cottrell, Angela DM Kashuba, et al. Pharmacokinetics and Pharmacodynamics of Anti-infective Agents. Bennett JE, Dolin R, Blaser MJ, Eds. Mandell, Douglas and Bennett's Principles and Practice of Infectious Diseases. 8th edn. Saunders; Philadelphia. 2015; pp252-262.
2. Barbour A, Scaglione F, Derendorf H. Class-dependent relevance of tissue distribution in the interpretation of anti-infective pharmacokinetic/pharmacodynamics indices. Int J Antimicrob Agents 2010; 35:431-438.
3. Zanger UM, Schwab M. Cytochrome P450 enzymes in drug metabolism: regulation of gene expression, enzyme activities, and impact of genetic variation. Pharmacol Ther 2013; 138:103-141.
4. Bebitođlu BT, Ođuz E, Nuhođlu Ç, Dalkýlýç AEK, Çirtlik P, Temel F, et al. Evaluation of potential drug-drug

* Assistant Professor and Neonatologist

** Chief Consultant and HOD,
Department of Pediatrics,
Tata Main Hospital and Manipal Tata Medical College,
Jamshedpur.
email: drsudhir@tatasteel.com

- interactions in a pediatric population. *Turk Pediatr Ars* 2020; 55:30-38.
5. Levison ME. Pharmacodynamics of antimicrobial agents. Bactericidal and postantibiotic effects. *Infect Dis Clin North Am* 1995; 9:483-495.
 6. Mouton JW, Dudley MN, Cars O, Derendorf H, Drusano GL. Standardization of pharmacokinetic/pharmacodynamic (PK/PD) terminology for anti-infective drugs: an update. *J Antimicrob Chemother* 2005; 55:601-607.
 7. Lacy MK, Nicolau DP, Nightingale CH, Quintiliani R. The pharmacodynamics of aminoglycosides. *Clin Infect Dis* 1998; 27:23-27.
 8. Lode H, Borner K, Koeppe P. Pharmacodynamics of fluoroquinolones. *Clin Infect Dis* 1998; 27:33-39.
 9. Kasiakou SK, Lawrence KR, Choulis N, Falagas ME. Continuous versus intermittent intravenous administration of antibacterials with time-dependent action: a systematic review of pharmacokinetic and pharmacodynamic parameters. *Drugs* 2005; 65:2499-2511.
 10. Zhanel GG, Hoban DJ, Harding GKM. The Postantibiotic Effect: A Review of in Vitro and in Vivo Data. *DICP* 1991; 25:153-163. doi:10.1177/106002809102500210.
 11. Bush K. Synergistic Antibiotic Combinations. In: Fisher JF, Mobashery S, Miller MJ. (eds) *Antibacterials. Topics in Medicinal Chemistry*, 2017 vol 25. Springer, Cham. https://doi.org/10.1007/7355_2017_23.
 12. Blot SI, Pea F, Lipman J. The effect of pathophysiology on pharmacokinetics in the critically ill patient-concepts appraised by the example of antimicrobial agents. *Adv Drug Deliv Rev* 2014; 77:3-11.
 13. Felton T, Troke PF, Hope WW. Tissue penetration of antifungal agents. *Clin Microbiol Rev* 2014; 27:68-88.
 14. Gilbert B, Robbins P, Livornese LL Jr. Use of antibacterial agents in renal failure. *Infect Dis Clin North Am* 2009; 23:899-924.
 15. Aloy B, Launay-Vacher V, Bleibtreu A, Bortolotti P, Faure E, Filali A, et al. Antibiotics and chronic kidney disease: Dose adjustment update for infectious disease clinical practice. *Med Mal Infect* 2020; 50:323-331.
 16. Rahul Chanchlani, Aditi Sinha. Drug dosing in renal failure. RN Srivastava and Arvind Bagga, editor. *Pediatric nephrology*, Ed 5. India: Jaypee Brothers. 2016; pp535-540.
 17. Bûdingen FV, Gonzalez D, Tucker AN, Derendorf H. Relevance of Liver Failure for Anti-Infective Agents: From Pharmacokinetic Alterations to Dosage Adjustments. *Ther Adv Infect Dis* 2014; 2:17-42. doi: 10.1177/2049936113519089.
 18. Halilovic J, Heintz BH. Antibiotic dosing in cirrhosis. *Am J Health Syst Pharm* 2014; 71:1621-1634.

ANTIMICROBIALS - I

SIDE EFFECTS OF ANTIMICROBIALS AND PREVENTIVE STRATEGIES

***Bindusha S**
****Santhosh Kumar A**

Abstract: Antibiotics though life-saving drugs, may have side effects varying in severity from trivial to life-threatening. The side effects include local irritant effects, systemic toxicity, drug hypersensitivity reactions, drug interactions and intestinal dysbiosis. Intestinal dysbiosis has widespread and long lasting effects in a person including the risk of development of diseases with immunological basis, including asthma, allergic diseases and diabetes mellitus. Antibiotics are considered as societal drugs, their side effects are not restricted only to the treated individual as their use is an important modifiable factor that can result in the development of drug resistant bacteria and therapeutic failure.

Keywords: Side effects, Drug resistance, Antimicrobial stewardship programme, Drug hypersensitivity

reactions, Drug interactions, Intestinal dysbiosis.

Points to Remember

- *The side effects of antibiotics are not restricted to the person who consumed the drug, the effect also extends to the society by the development of drug resistance.*
- *Systemic side effect can affect every organ system leading to serious consequences.*
- *Though hypersensitivity reactions are rare, sometimes they may be life threatening.*
- *Inappropriate use of antibiotics is the major cause for development of drug resistance which will lead to deleterious consequences in the individual and the society at large.*
- *Antibiotics are lifesaving drugs and side effects are unavoidable. But there are strategies to prevent or limit the side effects.*
- *Rational antibiotic prescription practices and antibiotic stewardship programs are the only measures to curtail the rapidly increasing menace of drug resistance.*

References

1. Mohsen S, Dickinson JA, Somayaji R. Update on the adverse effects of antimicrobial therapies in community practice. *Can Fam Physician* 2020; 66(9):651-659.
2. Cunha BA. Antibiotic side effects. *Med Clin North Am* 2001; 85(1):149-185.
3. Carey PJ. Drug-Induced Myelosuppression. *Drug Saf* 2003; 26(10):691-706.
4. Alam S, Mushtaq M. Antibiotic Associated Diarrhea in Children. *Indian Pediatr* 2009; 46(6): 491-496.
5. Yan T, Goldman RD. Probiotics for antibiotic-associated diarrhea in children. *Can Fam Physician* 2020; 66(1): 37-39.
6. Sammons JS, Toltzis P, Zaoutis TE. Clostridium difficile Infection in Children. *JAMA Pediatr.* 2013;167(6):567-573. doi:10.1001/jamapediatrics.2013.441
7. Shim JO. Clostridium difficile in Children: To Treat or Not

* Associate Professor,
Department of Pediatrics,
Govt. Medical College,
Thiruvananthapuram.

** Professor,
Department of Pediatrics,
Azeezia Medical College, Kollam.
email: drasanthoshkumar@gmail.com

- to Treat? *Pediatr Gastroenterol Hepatol Nutr* 2014; 17(2):80-84.
8. Allen UD. Clostridium difficile in paediatric populations. *Pediatr Child Health* 2014; 19(1):43-48.
 9. Singhal T, Shah N, Yewale V, Prabhu S. IAP Specialty Series on Rational Antimicrobial Practice In Pediatrics. Third edn. Jaypee Brothers Medical Publishers; 2019.
 10. Rezaei NJ, Bazzazi AM, Alavi SAN. Neurotoxicity of the antibiotics: A comprehensive study. *Neurol India* 2018;66(6):1732.
 11. Rosenberg CR, Fang X, Allison KR. Potentiating aminoglycoside antibiotics to reduce their toxic side effects. *PLOS ONE* 2020;15(9): e0237948.
 12. Black FO, Pesznecker S, Stallings V. Permanent gentamicin vestibulotoxicity. *Otol Neurotol*. 2004 Jul;25(4):559-69. doi: 10.1097/00129492-200407000-00025. PMID: 15241236.
 13. Ellis SJ, Cleverley JR, Müller NL. Drug-Induced Lung Disease. *Am J Roentgenol* 2000; 175(4):1019-1024.
 14. Blumenthal KG, Peter JG, Trubiano JA, Phillips EJ. Antibiotic allergy. *Lancet* 2019; 393(10167):183-198.
 15. Regateiro FS, Marques ML, Gomes ER. Drug-Induced Anaphylaxis: An Update on Epidemiology and Risk Factors. *Int Arch Allergy Immunol* 2020; 181(7): 481-487.
 16. Choudhary S, McLeod M, Torchia D, Romanelli P. Drug Reaction with Eosinophilia and Systemic Symptoms (DRESS) Syndrome. *J Clin Aesthetic Dermatol* 2013; 6(6):31-37.
 17. Sharifzadeh S, Mohammadpour AH, Tavanaee A, Elyasi S. Antibacterial antibiotic-induced drug reaction with eosinophilia and systemic symptoms (DRESS) syndrome: a literature review. *Eur J Clin Pharmacol* 2021; 77(3):275-289.
 18. Neuman H, Forsythe P, Uzan A, Avni O, Koren O. Antibiotics in early life: dysbiosis and the damage done. *FEMS Microbiol Rev* [Internet]. 2018 Jun 25 [cited 2021 Dec 20]; Available from: <https://academic.oup.com/femsre/advance-article/doi/10.1093/femsre/fuy018/5045017>.
 19. Langdon A, Crook N, Dantas G. The effects of antibiotics on the microbiome throughout development and alternative approaches for therapeutic modulation. *Genome Med* 2016;8:39.
 20. Slykerman RF, Thompson J, Waldie KE, Murphy R, Wall C, Mitchell EA. Antibiotics in the first year of life and subsequent neurocognitive outcomes. *Acta Paediatr* 2017;106(1):87-94.
 21. Dancer S. How antibiotics can make us sick: The less obvious adverse effects of antimicrobial chemotherapy. *Lancet Infect Dis* 2004; 4:611-619.

ANTIMICROBIALS - I
ANTIMICROBIALS FOR PERINATAL AND NEONATAL INFECTIONS

***Suman Rao PN**
****Swapnik Kandepi**

Abstract: *Antimicrobials are the most commonly used therapeutics in the neonatal intensive care units, as neonatal sepsis is the third leading cause of neonatal mortality. The signs of neonatal sepsis are non-specific and accurate diagnostic tests are not available immediately. With dynamic neonatal pharmacokinetics, attention to the dose and frequency of the antimicrobials are important. With increasing antibiotic resistance, the search for an ideal empiric antibiotic is still on. This review will include the antimicrobials most commonly used in the management of perinatal and neonatal infections.*

Keywords: *Neonatal sepsis, Management, Infection,*

Perinatal infection.

Points to Remember

- *Antimicrobials are the most frequently used therapeutic agents in neonates.*
- *Empiric antibiotic therapy for neonatal sepsis should be based on local antibiograms.*
- *Empiric therapy should be of sufficiently narrow spectrum and from the “Access category” of “WHO AWaRe antibiotics”.*
- *Antifungal and antiviral therapy are warranted in a select group of neonates.*

References

1. Schulman J, Benitz WE, Profit J, Lee HC, Dueñas G, Bennett MV, et al. Newborn antibiotic exposures and association with proven bloodstream infection. *Pediatrics* 2019; 144(5):e20191105.
2. Hauge C, StålsbyLundborg C, Mandaliya J, Marrone G, Sharma M. Up to 89% of neonates received antibiotics in cross-sectional Indian study including those with no infections and unclear diagnoses. *ActaPaediatr* 2017; 106(10):1674-1683.
3. Dong Y, Speer CP. Late-onset neonatal sepsis: recent developments. *Arch Dis Child - Fetal Neonatal Ed* 2015; 100(3):F257-263.
4. Agarwal R, Chaurasia S, JeevaSankar M, Yadav CP, Arya S, Kapil A, et al. Characterisation and antimicrobial resistance of sepsis pathogens in neonates born in tertiary care centres in Delhi, India: a cohort study. *Lancet Glob Heal* 2016; 4(10):e752-760.
5. Schrag SJ, Farley MM, Petit S, Reingold A, Weston EJ, Pondo T, et al. Epidemiology of invasive early-onset neonatal sepsis, 2005 to 2014. *Pediatrics* 2016; 138(6):e20162013.
6. Shane AL, Sánchez PJ, Stoll BJ. Neonatal sepsis. *Lancet* 2017; 390:1770-1780.
7. Simonsen KA, Anderson-Berry AL, Delair SF, Dele Davies H. Early-onset neonatal sepsis. *ClinMicrobiol Rev* 2014; 27(1):21-47.
8. Russell AB, Sharland M, Heath PT. Improving antibiotic prescribing in neonatal units: time to act. *Arch Dis Child - Fetal Neonatal Ed* 2012; 97(2):F141-146.

* Professor

** Senior Resident,
 Department of Neonatology,
 St. John's Medical College Hospital, Bengaluru.
 email: raosumanv@gmail.com

9. WHO. Pocket Book of Hospital Care for Children. Guidelines for the Management of Common Childhood Illnesses. Geneva:World Health Organization 2013; 125-143. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK154447/>
10. Metsvaht T, Ilmoja ML, Parm Ü, Maipuu L, Merila M, Lutsar I. Comparison of ampicillin plus gentamicin vs. penicillin plus gentamicin in empiric treatment of neonates at risk of early onset sepsis. *Acta Paediatrica, International Journal of Paediatrics*. 2010; 99(5):665-672.
11. Cantey JB, Wozniak PS, Sánchez PJ. Prospective surveillance of antibiotic use in the neonatal intensive care unit: results from the SCOUT study. *Pediatr Infect Dis J* 2015; 34(3):267-272.
12. Tziella C, Borghesi A, Serra G, Stronati M, Corsello G. Antimicrobial therapy in neonatal intensive care unit. *Ital J Pediatr* 2015; 41(1):27.
13. Bizzarro MJ, Shabanova V, Baltimore RS, Dembry LM, Ehrenkranz RA, Gallagher PG. Neonatal sepsis 2004-2013: the rise and fall of coagulase-negative staphylococci. *J Pediatr* 2015; 166(5):1193-1199.
14. Stockmann C, Spigarelli MG, Campbell SC, Constance JE, Courter JD, Thorell EA, et al. Considerations in the pharmacologic treatment and prevention of neonatal sepsis. *Paediatr Drugs* 2014; 16(1):67-81.
15. Korang SK, Safi S, Nava C, Gordon A, Gupta M, Greisen G, et al. Antibiotic regimens for early-onset neonatal sepsis. *Cochrane database Syst Rev* 2021; 5(5):CD013837.
16. Korang SK, Safi S, Nava C, Greisen G, Gupta M, Lausten-Thomsen U, et al. Antibiotic regimens for late-onset neonatal sepsis. *Cochrane database Syst Rev* 2021; 5:CD013836.
17. Tewari VV, Jain N. Monotherapy with Amikacin or Piperacillin-Tazobactam Empirically in Neonates at Risk for Early-Onset Sepsis: A Randomized Controlled Trial. *J Trop Pediatr* 2014; 60(4):297-302.
18. Keij FM, Kornelisse RF, Hartwig NG, Reiss IKM, Allegaert K, Tramper-Stranders GA. Oral antibiotics for neonatal infections: a systematic review and meta-analysis. *J Antimicrob Chemother* 2019; 74(11):3150-3161.
19. Duby J, Lassi ZS, Bhutta ZA. Communitybased antibiotic delivery for possible serious bacterial infections in neonates in low and middleincome countries. *Cochrane Database Syst Rev* 2019; 4(4):CD007646.
20. Fuchs A, Bielicki J, Mathur S, Sharland M, Van Den Anker JN. Reviewing the WHO guidelines for antibiotic use for sepsis in neonates and children. *Pediatr Int Child Health* 2018; 38(1):S3-S15.
21. Esaiassen E, Fjalstad JW, Juvet LK, Van Den Anker JN, Klingenberg C. Antibiotic exposure in neonates and early adverse outcomes: a systematic review and meta-analysis 2017; 72(7):1858-1870.
22. Hsia Y, Lee BR, Versporten A, Yang Y, Bielicki J, Jackson C, et al. Use of the WHO Access, Watch and Reserve classification to define patterns of hospital antibiotic use (AWaRe): an analysis of paediatric survey data from 56 countries. *Lancet Glob Heal* 2019; 7(7):e861-871.
23. World Health Organization. AWaRe classification. Available from: <https://www.who.int/publications/i/item/2021-aware-classification>.
24. Darlow CA, da Costa RMA, Ellis S, Franceschi F, Sharland M, Piddock L, et al. Potential Antibiotics for the Treatment of Neonatal Sepsis Caused by Multidrug-Resistant Bacteria. *Pediatr Drugs* 2021; 23(5):465-484.
25. Fanaroff AA, Fanaroff JM. Advances in Neonatal Infections. *Am J Perinatol* 2020; 37(1):S5-9.
26. Parikh TB, Nanavati RN, Patankar CV, Rao PNS, Bisure K, Udani RH, et al. Fluconazole Prophylaxis against Fungal Colonization and Invasive Fungal Infection in Very Low Birth Weight Infants. *Indian Pediatr* 2007; 44(11):830-837.
27. Ferreras-Antolín L, Sharland M, Warris A. Management of Invasive Fungal Disease in Neonates and Children. *Pediatr Infect Dis J* 2019; 38(6):S2.
28. Kimberlin DW, Jester PM, Sánchez PJ, Ahmed A, Arav-Boger R, Michaels MG, et al. Valganciclovir for Symptomatic Congenital Cytomegalovirus Disease. *N Engl J Med* 2015; 372(10):933-943.

ANTIMICROBIALS - I

CHOICE OF ANTI-STAPHYLOCOCCAL THERAPY

***Upendra S Kinjawadekar**
****Rajendra Vaidya**

Abstract: *Staphylococcus aureus* is considered to be the most virulent amongst all staphylococci. It is known to elude antimicrobial therapy by adopting various strategies posing therapeutic dilemmas for clinicians, particularly in intensive care settings. The coagulase negative staphylococci like *S. Hemolyticus*, *S. Saprophyticus*, *S. Epidermidis* are relatively less pathogenic unless indwelling devices are present. When complicated or invasive *S. aureus* infection is suspected, blood culture and culture of sample from potential source are essential before starting empirical treatment due to the increased prevalence of methicillin resistant staphylococcus aureus. Disruption of skin, immune compromised conditions, malnutrition, burns, scabies or post varicella lesions are more prone for *S. aureus* infections.

Keywords: *Staphylococcus aureus*, Methicillin sensitive staphylococcus aureus, Methicillin resistant staphylococcus aureus, Coagulase negative staphylococcus.

Points to Remember

- *Staphylococcus aureus* coagulase positive is considered the most virulent amongst all gram-positive genus staphylococci.
- Blood culture and sample from potential focus of infection are a must in suspected moderate/severe staphylococcal infections.
- Surface cultures should be avoided as it reflects contamination rather than true infection.
- MSSA is a *S.aureus* isolate with an oxacillin MIC ≤ 2 mcg/mL whereas MRSA is *S.aureus* isolate with an oxacillin MIC ≥ 4 mcg/mL.
- Antibiotic choice will differ while treating MSSA or MRSA infections and understanding the local prevalence of MRSA as well as inducible clindamycin resistance is essential.

References

1. Catherine Liu, Arnold Bayer, Sara E Cosgrove, Robert S Daum, Scott K Fridkin, Rachel J Gorwitz, Sheldon L Kaplan, Adolf W Karchmer, Donald P Levine, Barbara E Murray, Michael J Rybak, David A Talan, Henry F Chambers, Infectious Diseases Society of America. Clinical practice guidelines by the infectious diseases society of america for the treatment of methicillin-resistant *Staphylococcus aureus* infections in adults and children. Clin Infect Dis 2011; 52(3):e18-55.
2. Robert S Daums; *Staphylococcus aureus*; Principles and Practice of Pediatric Infectious Diseases, 5th edn, ELSEVIER, 2018.
3. Lalitha AV, Rebello G, Chettri S, Reddy M. Demographic and Clinical Profile of Invasive Staphylococcal Infections in Children Admitted to Pediatric Intensive Care Unit: A Retrospective Study. Indian J Crit Care Med 2020; 24(9):890–891.
4. Youngster I, Shenoy ES, Hooper DC, Nelson SB. Comparative evaluation of the tolerability of cefazolin and nafcillin for treatment of methicillin-susceptible *Staphylococcus aureus* infections in the outpatient setting. Clin Infect Dis 2014; 59(3):369-375.
5. Wadstrom T, Mollby R. Studies on extracellular proteins from *Staphylococcus aureus*. VII. Studies on β -haemolysin. Biochim Biophys Acta 1971; 242:308-320.

* Consulting Pediatrician,
Kamlesh Mother and Child Hospital and
Apollo Hospitals, Navi Mumbai.

** Preventive and Primary Care Pediatrician
Vaidya Children Hospital,
Aurangabad.
email: upen228@gmail.com

6. Lina G, Piémont Y, Godail-Gamot F, Bes M, Peter MO, Gauduchon V, et al. Involvement of Panton-Valentine leukocidin-producing *Staphylococcus aureus* in primary skin infections and pneumonia. *Clin Infect Dis* 1999; 29:1128-1132.
7. Gillet Y, Issartel B, Vanhems P, Fournet JC, Lina G, Bes M, et al. Association between *Staphylococcus aureus* strains carrying gene for Panton-Valentine leukocidin and highly lethal necrotizing pneumonia in young immunocompetent patients. *Lancet* 2002; 359:753-759.
8. American Academy of Pediatrics. *Staphylococcus aureus*. In: *Red Book: 2018 Report of the Committee on Infectious Diseases*, 31st ed, Kimberlin DW, Brady MT, Jackson MA, Long SS (Eds), American Academy of Pediatrics, Itasca, IL 2018; p733-746.
9. Fergie J, Purcell K. The treatment of community-acquired methicillin-resistant *Staphylococcus aureus* infection. *Pediatr Infect Dis J* 2008; 27(1):67-68.
10. Liu C, Bayer A, Cosgrove SE, Daum RS, Fridkin SK, Gorwitz RJ, et al. Clinical practice guidelines by the Infectious Diseases Society of America for the treatment of methicillin-resistant *Staphylococcus aureus* infections in adults and children: executive summary. *Clin Infect Dis* 2011; 52:285-292.
11. American Academy of Pediatrics. Tables of antibacterial drug dosages. In: *Red Book: 2021-2024 Report of the Committee on Infectious Diseases*, 32nd ed, Kimberlin DW, Barnett ED, Lynfield R, Sawyer MH (Eds), American Academy of Pediatrics, 2021:678-876.
12. Deville JG, Adler S, Azimi PH, Jantausch BA, Morfin MR, Beltran S, et al. Linezolid versus vancomycin in the treatment of known or suspected resistant gram-positive infections in neonates. *Pediatr Infect Dis J* 2003; 22:S158-S163.
13. Yogev R, Patterson LE, Kaplan SL, Adler S, Morfin MR, Martin A, et al. Linezolid for the treatment of complicated skin and skin structure infections in children. *Pediatr Infect Dis J* 2003; 22:S172-S177.
14. Critchley IA, Blosser-Middleton RS, Jones ME, Thornsberry C, Sahm DF, Karlowsky JA. et al. Baseline study to determine in vitro activities of daptomycin against gram-positive pathogens isolated in the United States in 2000-2001. *Antimicrob Agents Chemother* 2003; 47:1689-1693.
15. Allington DR, Rivey MP. Quinupristin/dalfopristin: a therapeutic review. *Clin ther* 2001; 23(1):24-44.
16. Gordon JJ, Harter DH, Phair JP. Meningitis due to *Staphylococcus aureus* *Am J Med* 1985; 78:965-970.
17. Aguilar J, Urdy-Cornejo V, Donabedian S, Perri M, Tibbetts R, Zervos M. *Staphylococcus aureus* meningitis- case series and literature review. *Medicine* 2010; 89:117-125.
18. Sunavala AJ, Udani V. Antimicrobial Therapy in CNS infections. In: *IAP Specialty series on Rational antimicrobial practice in pediatrics*, 3rd edn; pages 279-291
19. Zumo LA, Singh NN. *Staphylococcal Meningitis*. Medscape, updated. Jun 19, 2018. Accessed on 25th February, 2022.

ANTIMICROBIALS - I
**ANTITUBERCULOUS THERAPY-
CURRENT PRACTICE**

***Gowrishankar NC**

Abstract: Tuberculosis affects all age groups. Treatment for tuberculosis has been standardized having a 2 months intensive phase with four first line drugs followed by a 4 months continuation phase with three first line drugs given daily. Adjunctive steroids are useful in central nervous system disease and pericardial involvement. Adverse reactions to drugs, though uncommon, need to be looked for and managed appropriately. Standard regimens for treatment of resistant tuberculosis in India follow World Health Organization guideline. When there is an infectious pulmonary tuberculosis patient in the family, tuberculosis preventive treatment has to be given to all family members irrespective of age after ruling out active tuberculosis. All children after completion of treatment need to be followed up for two years.

Keywords: Children, Treatment, Tuberculosis, Resistant, Preventive therapy.

Points to Remember

- *Only one treatment schedule is recommended (2HRZE/4HRE) for all newly diagnosed TB whether microbiologically confirmed or clinically diagnosed.*
- *Continuation phase can be extended in neurological, skeletal and disseminated TB.*
- *Children must always be followed up for 2 years after completion of treatment for TB.*
- *Drug resistant TB in children is to be treated with shorter or longer regimen based on extent of disease and severity.*
- *All household contacts of adult index pulmonary TB patients have to be given TB preventive treatment after ruling out active disease, regimen being different for drug sensitive and drug resistant strains.*

References

1. Balachandran A. Anti-tuberculous therapy. In: Essentials of Pediatric Pulmonology. Eds Subramanyam L, Shivbalan So, Gowrishankar NC, Vijayasekeran D, Balachandran A. 3rd edn, Pediatric Pulmonology Foundation of India, Chennai -31. 2008; pp222-225.
2. dos Santos Peixoto A, Araújo RM, Schindler HC, Pimentel LM. Treatment of sensitive tuberculosis-mechanisms of action and resistance. Biomed J Sci Tech Res 2019; 18(4):13715-13718.
3. Central TB Division, MoHFW, Government of India. Training modules (1-4) for programme managers and medical officers; New Delhi, India: July 2020. Available from: www.tbcindia.gov.in. Accessed on 3rd Jan, 2022.
4. Singh V, Parakh A. What is new in the management of childhood tuberculosis in 2020? Indian Pediatr 2020; 57(12):1172-1176.
5. Schutz C, Davis AG, Sossen B, Lai RP, Ntsekhe M, Harley YX, et al. Corticosteroids as an adjunct to tuberculosis therapy. Expert Rev Respir Med 2018;

* Pediatric Pulmonologist and Head-Pediatrics,
Mehta Multispeciality Hospitals India Pvt. Ltd,
Chennai.
email:cugowri@yahoo.com

12(10):881-891.

6. Gafar F, Arifin H, Jurnalys YD, Yani FF, Fitria N, Alffenaar JC, et al. Antituberculous drug-induced liver injury in children: incidence and risk factors during the two-month intensive phase of therapy. *Pediatr Infect Dis J* 2019; 38(1):50-53.
7. Chanchlani R, Anand S, Valecha J, Goyal S, Tiwari A, Gupta V. Paradoxical upgradation reaction in disseminated tuberculosis in takayasu arteritis. *Pediatr Oncall J* 2014; 11:111-112.
8. Barr DA, Coussens AK, Irvine S, Ritchie ND, Hebert K, Choo-Kang B, et al. Paradoxical upgrading reaction in extra-pulmonary tuberculosis: association with vitamin D therapy. *Int J Tuberc Lung Dis* 2017;21(6): 677-683.
9. Central TB Division. MoHFW, Government of India. Guidelines for programmatic management of drug resistant tuberculosis in India 2021; New Delhi, India. March 2021. Available from www.tbcindia.gov.in. Accessed on 13th Jan, 2022.
10. Schaaf H. Diagnosis and management of multidrug-resistant tuberculosis in children: a practical approach. *Indian J Pediatr* 2019; 86:717-724.
11. Gowrishankar NC, Subramanyam L. Drug resistant tuberculosis. In: Partha's Current trends in diagnosis and management in pediatric and adolescent practice. First edn. Eds Parthasarathy A, Alok Gupta eds. Jaypee brothers Medical Publishers, New Delhi. 2021; pp159-162.
12. Shah I, Poojari V, Meshram H. Multi-Drug resistant and extensively-drug resistant tuberculosis. *Indian J Pediatr* 2020; 87(10):833-839.
13. Marais BJ. Newer drugs for tuberculosis prevention and treatment in children. *Indian J Pediatr* 2019; 86(8): 725-731.
14. Central TB Division, MoHFW, Government of India. Guidelines for programmatic management of Tuberculosis preventive treatment in India. New Delhi, India. July 2021. Available from www.tbcindia.gov.in. Accessed on 15th Jan, 2022.

ANTIMICROBIALS - I

ANTIMALARIAL DRUG THERAPY

***Ritabrata Kundu**
****Aniruddha Ghosh**

Abstract: *Malaria is a major worldwide problem and a public health problem of developing countries like India. It is caused by intracellular Plasmodium protozoa transmitted to humans by the bite of female Anopheles mosquitos. Malaria is caused by four species of the genus P.vivax, P.falciparum, P.ovale, P.malariae and the fifth species P.knowlesi primarily an animal pathogen reported to cause malaria in South-East Asia especially, Borneo. The diagnosis is confirmed by identification of the organism in stained peripheral blood smear and treated by antimalarial drugs as per the situation.*

Keywords: *Plasmodium infection, Treatment, Prophylaxis, Antimalarials.*

Points to Remember

- *Treatment regimens have been decided keeping the drug resistance pattern in that particular geographic area.*
- *For uncomplicated vivax malaria chloroquine plus primaquine is effective.*
- *Antimalarial drug combinations are now the recommended modality of treatment for P.falciparum infection.*
- *The following ACTs are presently use, Artesunate (AS) + Sulfadoxine-Pyrimethamine (SP), Artesunate + Mefloquine (MQ), Artemether + Lumefantrine.*
- *Artemether + Lumefantrine is the only form available as oral preparation and is well tolerated and effective against multidrug-resistant falciparum malaria.*

References

1. Country profile: India. World Malaria Report 2018. World Health Organization. <https://www.who.int/malaria/publications/country-profiles/en/>
2. National Strategic Plan for Malaria Elimination (2017-22). Directorate of National Vector Borne Disease Control Program. Directorate General of Health Services. New Delhi: Ministry of Health and Family Welfare, Govt. of India; 2017. <http://www.indiaenvironmentportal.org.in/content/445149/national-strategic-plan-for-malaria-elimination-2017-22/>
3. Kundu R. Malaria. In: Gupta P, Menon PSN, Ramji S, Lodha R eds. PG Textbook of Pediatrics, 2nd edn. New Delhi: Jaypee; 2018; pp1232-1237.
4. Meshnick SR. Artemisinin: Mechanisms of action, resistance and toxicity. Int J Parasitol 2002; 32(13):1655-1660.
5. World Health Organisation (WHO). Guidelines for the treatment of Malaria. 3rd edn. Geneva: World Health Organisation; 2015; pp23-25.
6. Marx A, Pewsner D, Egger M, Nüesch R, Bucher HC, Genton B, et al. Meta-analysis: accuracy of rapid tests for malaria in travellers returning from endemic areas. Ann Intern Med 2005; 142(10):836-846.
7. Moody A. Rapid diagnostic tests for malaria parasites. Clin Microbiol Rev 2002; 15(1):66-78.

* Professor

** Assistant Professor,
Department of Pediatrics,
Institute of Child Health,
Kolkata.
email: rkundu22@gmail.com

8. Mutabingwa TK, Anthony D, Heller A, Hallett R, Ahmed J, Drakeley C, et al. Amodiaquine alone, amodiaquine+ sulfadoxine-pyrimethamine, amodiaquine+ artesunate and artemether-lumefantrine for outpatient treatment of malaria in Tanzanian children: a four-arm randomised effectiveness trial. *Lancet* 2005; 365(9469): 1474-1480.
9. Rediscovering wormwood: qinghaosu for malaria. *Lancet* 1992; 339(8794):649-6451.
10. Adjuik M, Babiker A, Garner P, Olliaro P, Taylor W, White N, International Artemisinin Study Group Artesunate combinations for treatment of malaria: meta-analysis. *Lancet* 2004; 363(9402):9-17.
11. Bassat Q. The use of artemether-lumefantrine for the treatment of uncomplicated *Plasmodium vivax* malaria. *PLoS Negl Trop Dis* 2011; 5(12):e1325.
12. Newton PN, Angus BJ, Chierakul W, Dondorp A, Ruangveerayuth R, Silamut K, et al. Randomised comparison of artesunate and quinine in the treatment of severe falciparum malaria. *Clin Infect Dis* 2003; 37:7-16.

ANTIMICROBIALS - I**RATIONAL ANTIMICROBIAL THERAPY
IN OFFICE PRACTICE*****Palaniraman R**

Abstract: *The most common drug prescribed by a pediatrician in office practice other than paracetamol, is an antibiotic. Ninety percent of infections seen in office practice are viral in origin. Therefore, hardly 10% of children need antibiotics. Simple, safe, first line narrow spectrum antibiotics are more than enough to treat common community acquired bacterial infections. Irrational antibiotic use may contribute to increased incidence of community acquired resistant infections, increase in complications due to partial treatment and increase in cost of treatment.*

Keywords: *Antimicrobials, Children, Outpatients, Office practice, Narrow spectrum antibiotics.*

Points to Remember

- *Majority of infections seen in office practice are viral. Only 10% or less are of bacterial etiology like acute otitis media, dysentery, skin and soft tissue infection etc.*
- *In short, amoxicillin is the drug of choice for respiratory infections, cefixime for gastrointestinal and genitourinary infections, cephalosporins (1st generation) for skin and soft tissue infections and doxycycline for scrub typhus.*
- *Antibiotic resistance in community acquired UTI and skin infections are on the rise.*
- *Rational antibiotic therapy should be based on the available scientific evidence.*

Bibliography

1. Palani Raman R. Choice of antibiotics in office practice. Indian J Pract Pediatr 2010; 12:382-385.
2. Amdekar YK. General Principles of Antimicrobial therapy. In: Nitin Shah, Tanu Singhal. Eds, Rational Antimicrobial practice in Pediatrics, 1st edn, IAP's Specialty series, Jaypee Brothers Medical Publishers (p) Ltd., New Delhi 2006; pp121-124.
3. Alter SJ, Vidwan NK, Sobande PO, Omoloja A, Bennett JS. Common childhood bacterial infections. Curr Probl Pediatr Adolesc Health Care 2011; 41(10): 256-283.
4. Bula-Rudas FJ, Olcott JL. Human and Animal Bites. Pediatr Rev 2018; 39(10):490-500.
5. Edwards-Jones V. Antimicrobial stewardship in wound care. Br J Nurs 2020; 29(15):S10-S16.
6. Gerber JS, Jackson MA, Tamma PD, Zaoutis TE. Committee on infectious diseases, Pediatric Infectious Diseases Society. Antibiotic Stewardship in Pediatrics. Pediatrics 2021; 147(1):e2020040295.

* Senior Child Health Consultant,
CR Women and Child Health Centre, Tindivanam.
email: amvarsh@gmail.com

ANTIMICROBIALS - I

PROPHYLACTIC ANTIMICROBIALS

***Padmasani Venkat Ramanan**
****Shalini Sharma**

Abstract: *Prophylactic use of antimicrobials is the use of an antimicrobial agent to prevent infection. The focus of this review is on everyday situations in which a general pediatrician may have to use prophylactic antimicrobials such as preventing recurrent urinary tract infections, recurrent otitis media, surgical site infections, infective endocarditis, rheumatic fever, malaria in travellers and for contacts of highly contagious diseases. Special situations like immunodeficiency disorders, post-transplant state, pediatric hemato-oncology and intensive care settings are distinct entities and have not been addressed in this article in detail.*

Keywords: *Infection, Children, Chemoprophylaxis, Antimicrobial, Prophylaxis.*

Points to Remember

- *Antimicrobials should be used for prophylaxis only when there is a clear indication and the benefits outweigh the risk.*
- *Active infection should be ruled out before initiating prophylaxis.*
- *The importance of non-pharmacological measures, vaccination and infection control should always be borne in mind as they are superior, safer and more effective in prevention of infections.*
- *Attention must be paid for side effects while using certain drugs for prophylaxis such as azithromycin.*
- *The guidelines should be periodically reviewed in the background of emerging evidence and regional antimicrobial resistance patterns.*
- *Monitoring for adverse effects of the drugs used long term is essential.*

References

1. IAP Standard Treatment Guidelines 2022. <https://iapindia.org/standard-treatment-guidelines>. Accessed on December, 2021.
2. Ansari MS, Shekar PA, Singh C, Joshi SS. The Urological Society of India Guidelines for the management of paediatric urinary tract infection (Executive Summary). *Indian J Urol* 2021; 37(1):10-12. doi: 10.4103/iju.IJU_568_20. PMID: 33850350; PMCID: PMC8033226.
3. NICE guideline [NG112] October 2018. Urinary tract infection (recurrent): antimicrobial prescribing.
4. Granath A. Recurrent Acute Otitis Media: What Are the Options for Treatment and Prevention? *Curr Otorhinolaryngol Rep* 2017; 5(2):93-100. doi: 10.1007/s40136-017-0151-7. Epub 2017 May 9. PMID: 28616364; PMCID: PMC5446546.
5. Berríos-Torres SI, Umscheid CA, Bratzler DW. Healthcare Infection Control Practices Advisory Committee. Centre for Disease Control and Prevention Guideline for the Prevention of Surgical Site Infection, 2017. *JAMA Surg* 2017; 152(8):784-791. doi: 10.1001/jamasurg.2017.0904. Erratum in: *JAMA Surg* 2017; 152(8):803. PMID: 28467526.

* Professor,
Department of Pediatrics,
Sri Ramachandra Institute of Higher Education
and Research, Chennai.

** Pediatric ID specialist,
Manipal Hospital,
Bengaluru.

email: padmasani2001@yahoo.com

6. Centre for Clinical Practice, National Institute for Health and Clinical Excellence. Prophylaxis against infective endocarditis Antimicrobial prophylaxis against infective endocarditis in adults and children undergoing interventional procedures - NICE guideline 64 (2008), London, WC1V 6NA. www.nice.org.uk.
7. Nishimura RA, Otto CM, Bonow RO, Carabello BA, Erwin JP III, Fleisher LA, Jneid H, Mack MJ, McLeod CJ, O Gara PT, Rigolin VH, Sundt TM III, Thompson A. 2017 AHA/ACC Focused Update of the 2014 AHA/ACC Guideline for the Management of Patients With Valvular Heart Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation* 2017; 135(25):e1159-e1195. doi: 10.1161/CIR.0000000000000503. Epub 2017 Mar 15. PMID: 28298458.
8. Gerber MA, Baltimore RS, Eaton CB, Gewitz M, Rowley AH, Shulman ST, Taubert KA. Prevention of rheumatic fever and diagnosis and treatment of acute Streptococcal pharyngitis: a scientific statement from the American Heart Association Rheumatic Fever, Endocarditis, and Kawasaki Disease Committee of the Council on Cardiovascular Disease in the Young, the Interdisciplinary Council on Functional Genomics and Translational Biology, and the Interdisciplinary Council on Quality of Care and Outcomes Research: endorsed by the American Academy of Pediatrics. *Circulation* 2009; 119(11):1541-1551. doi: 10.1161/CIRCULATIONAHA.109.191959. Epub 2009 Feb 26. PMID: 19246689.
9. Chiodini PL, Patel D, Goodyer L, Ranson H. Guidelines for malaria prevention in travellers from the United Kingdom, 2021. London: Public Health England; Advisory Committee on Malaria Prevention (ACMP) March 2021.
10. Singh V, Parakh A. What Is New in the Management of Childhood Tuberculosis in 2020? *Indian Pediatr* 2020; 57(12):1172-1176. PMID: 33318324.
11. Uyeki TM, Bernstein HH, Bradley JS, Englund JA, File TM, Fry AM, Gravenstein S, Hayden FG, Harper SA, Hirshon JM, Ison MG, Johnston BL, Knight SL, McGeer A, Riley LE, Wolfe CR, Alexander PE, Pavia AT. Clinical Practice Guidelines by the Infectious Diseases Society of America: 2018 Update on Diagnosis, Treatment, Chemoprophylaxis, and Institutional Outbreak Management of Seasonal Influenza. *Clin Infect Dis* 2019; 68(6):895-902. doi: 10.1093/cid/ciy874. PMID: 30834445; PMCID: PMC6769232.
12. United States Centers for Disease Control and Prevention. https://www.cdc.gov/meningococcal/about_prevention.html. Page last reviewed: May 31, 2019.
13. Briere EC, Rubin L, Moro PL, Cohn A, Clark T, Messonnier N, Division of Bacterial Diseases, National Center for Immunization and Respiratory Diseases. Prevention and control of haemophilus influenzae type b disease: recommendations of the advisory committee on immunization practices (ACIP). *CDC. MMWR Recomm Rep* 2014; 63(RR-01):1.
14. American Academy of Pediatrics. Red Book: 2018 Report of the Committee on Infectious Diseases, 31st edn, Kimberlin DW, Brady MT, Jackson MA, Long SS (Eds), American Academy of Pediatrics, Itasca, IL 2018.
15. Global Task Force on Cholera Control. Prevention and control of cholera outbreaks: WHO policy and recommendations.
16. Rankine-Mullings AE, Owusu-Ofori S. Prophylactic antibiotics for preventing pneumococcal infection in children with sickle cell disease. *Cochrane Database Syst Rev* 2017; 10(10):CD003427. doi: 10.1002/14651858.CD003427.pub4. Update in: *Cochrane Database Syst Rev* 2021; 3:CD003427. PMID: 28994899; PMCID: PMC6485662.
17. McCavit TL, Gilbert M, Buchanan GR. Prophylactic penicillin after 5 years of age in patients with sickle cell disease: a survey of sickle cell disease experts. *Pediatr Blood Cancer* 2013; 60(6):935-939. doi: 10.1002/psc.24395. Epub 2012 Nov 28. PMID: 23193095.
18. Kuruvilla M, Teresa de la Morena M. Antibiotic Prophylaxis in Primary Immune Deficiency Disorders. *J Allergy Clin Immunol Pract* 2013; 1(6):573-582. doi: 10.1016/j.jaip.2013.09.013.
19. Health Protection Surveillance Centre. Guidelines for the Emergency Management of Injuries and Post-exposure Prophylaxis (PEP). EMI Toolkit. Dublin, Ireland HIV PEP updated June 2018. Available at: www.emitoolkit.ie.
20. National AIDS Control Organisation. Updated Guidelines for Prevention of Parent to Child Transmission (PPTCT) of HIV using Multi Drug Anti-retroviral Regimen in India December, 2013.

ANTIMICROBIALS - I

NEWER ANTIBIOTICS

***Dhanalakshmi K**
****Lakshan Raj S**

Abstract: *Antibiotics have revolutionised medical practice and their over zealous use has resulted in increased incidences of emerging resistant organisms. As newer molecules were identified especially for Gram negative organisms, many clinical trials were conducted. However, only few trials included children between 0 and 18 of age. Newer beta lactamase inhibitors were also recognized and they are first generation beta lactamase inhibitors with beta lactum rings, the second generation beta lactamase inhibitors with diaxabicyclooctane molecules and third generation beta lactamase are boronic acid compounds. Liberal and indiscriminate use of antibiotics have resulted in emergence of antibiotic resistance in the bacteriae with newer mechanisms, which in turn led on to higher medical costs and the increased mortality. Hence, there is a need for newer antibiotics and this article deals with antibiotics found in the last two decades and their uses.*

Keywords: *Antibiotics, Beta lactamase inhibitors, Synthetic aminoglycoside, Tetracyclines, Siderophore, Plazomycin, Ervacycline, Cefiderocol, Glycopeptides, Oxazolidinones.*

Points to Remember

- *Microbiological identification of infections should be the norm and newer drugs should not be used empirically.*
- *Excellent new drugs are available for ESBL producers.*
- *In India, carbapenamase producers are predominantly NDM and OXA-48 and to target this mechanism of resistance, more effective drugs are needed.*
- *Majority of the trials were conducted only in adults hence more data on the pharmacokinetics / pharmacodynamics of drugs are needed in children including newborns.*
- *Newer antibiotics should be reserved only for infections where there are limited therapeutic options.*

References

1. Veeraraghavan B, Pragasam AK, Bakthavatchalam YD, Anandan S, Ramasubramanian V, Swaminathan S, et al. Newer β -Lactam/ β -Lactamase inhibitor for multidrug-resistant gram-negative infections: Challenges, implications and surveillance strategy for India. *Indian J Med Microbiol* 2018; 36:334-343.
2. Tamma PD, Aitken SL, Bonomo RA, Mathers AJ, van Duin D, Clancy CJ. Infectious diseases society of America antimicrobial resistant treatment guidance: gram-negative bacterial infections. *Clin Infect Dis* 2020; 72(7):e169-e183.doi:10.1093/cid/ciaa1478.
3. Shirley M. Ceftazidime-Avibactam: A Review in the Treatment of Serious Gram-Negative Bacterial Infections. *Drugs* 2018; 78:675-692.
4. Karaiskos I, Galani I, Souli M, Giamarellou H. Novel β -lactam- β -lactamase inhibitor combinations: expectations for the treatment of carbapenam-resistant Gram-negative pathogens. *Expert Opin Drug Metab Toxicol* 2019; 15:133-149.
5. Kazmierczak KM, de Jonge BLM, Stone GG, Sahn DF. In vitro activity of ceftazidime/avibactam against isolates of Enterobacteriaceae collected in European countries: INFORM global surveillance 2012-15. *J Antimicrob Chemother* 2018; 73:2782-2788.

* Consultant, Pediatric Infectious diseases,

** Pediatric Registrar,
Kanchi Kamakoti Childs Trust Hospital,
Chennai.

email:drkdhana77@gmail.com

6. Yasmin M, Fouts DE, Jacobs MR, Haydar H, Marshall SH, White R, et al. Monitoring Ceftazidime-Avibactam and Aztreonam Concentrations in the Treatment of a Bloodstream Infection Caused by a Multidrug-Resistant *Enterobacter* sp. Carrying Both *Klebsiella pneumoniae* Carbapenemase-4 and New Delhi Metallo- β -Lactamase-1. *Clin Infect Dis* 2020; 71:1095-1098.
7. Testa R, Cantón R, Giani T, Morosini M-I, Nichols WW, Seifert H, et al. In vitro activity of ceftazidime, ceftaroline and aztreonam alone and in combination with avibactam against European Gram-negative and Gram-positive clinical isolates. *Int J Antimicrob Agents* 2015; 45: 641-646.
8. Yahav D, Giske CG, Grâmatniece A, Abodakpi H, Tam VH, Leibovici L. New β -Lactam- β -Lactamase Inhibitor Combinations. *Clin Microbiol Rev* 2020; 34(1):e00115-20. <https://doi.org/10.1128/cmr.00115-20>
9. Horcajada JP, Montero M, Oliver A, Sorlí L, Luque S, Gómez-Zorrilla S, et al. Epidemiology and Treatment of Multidrug-Resistant and Extensively Drug-Resistant *Pseudomonas aeruginosa* Infections. *Clin Microbiol Rev* 2019;32(4):e00031-19.<https://doi.org/10.1128/CMR.00031-19>
10. Lizza BD, Betthausen KD, Ritchie DJ, Micek ST, Kollef MH. New Perspectives on Antimicrobial Agents: Ceftolozane-Tazobactam. *Antimicrob Agents Chemother* 2021; 65:e0231820.
11. Solomkin J, Hershberger E, Miller B, Popejoy M, Friedland I, Steenbergen J, et al. Ceftolozane/Tazobactam Plus Metronidazole for Complicated Intra-abdominal Infections in an Era of Multidrug Resistance: Results From a Randomized, Double-Blind, Phase 3 Trial (ASPECT-cIAI). *Clin Infect Dis* 2015; 60:1462-1471.
12. Novelli A, Del Giacomo P, Rossolini GM, Tumbarello M. Meropenem/vaborbactam: a next generation β -lactam β -lactamase inhibitor combination. *Expert Rev Anti-Infect Ther*2020; 18:643-655.
13. Smith JR, Rybak JM, Claeys KC. Imipenem-Cilastatin-Relebactam: A Novel β -Lactam- β -Lactamase Inhibitor Combination for the Treatment of Multidrug-Resistant Gram-Negative Infections. *Pharmacotherapy* 2020; 40:343-356.
14. Mansour H, Ouweini AEL, Chahine EB, Karaoui LR. Imipenem/cilastatin/relebactam: A new carbapenem β -lactamase inhibitor combination. *Am J Health Syst Pharm* 2021; 78:674-683.
15. Heo YA. Imipenem/Cilastatin/Relebactam: A Review in Gram-Negative Bacterial Infections. *Drugs* 2021; 81:377-388.
16. Eljaaly K, Alharbi A, Alshehri S, Ortwine JK, Pogue JM. Plazomicin: A Novel Aminoglycoside for the Treatment of Resistant Gram-Negative Bacterial Infections. *Drugs* 2019; 79:243-269.
17. Livermore DM, Mushtaq S, Warner M, Zhang J-C, Maharjan S, Doumith M, et al. Activity of aminoglycosides, including ACHN-490, against carbapenem-resistant *Enterobacteriaceae* isolates. *J Antimicrob Chemother* 2011; 66:48-53.
18. Serio AW, Keepers T, Krause KM. Plazomicin is active against Metallo- β -Lactamase-Producing *Enterobacteriaceae*. *Open Forum Infect Dis* 2019, Volume 6, Issue 4, April 2019, ofz123, <https://doi.org/10.1093/ofid/ofz123>.
19. Shafer KM, Zmarlicka MT, Chahine EB, Piccicacco N, Cho JC. Plazomicin: A Next-Generation Aminoglycoside. *Pharmacotherapy* 2019; 39:77-93.
20. Zhanel GG, Cheung D, Adam H, Zelenitsky S, Golden A, Schweizer F, et al. Review of Eravacycline, a Novel Fluorocycline Antibacterial Agent. *Drugs* 2016; 76:567-588.
21. Scott LJ. Eravacycline: A Review in Complicated Intra-Abdominal Infections. *Drugs* 2019; 79:315-324.
22. Zhao C, Wang X, Zhang Y, Wang R, Wang Q, Li H, et al. In vitro activities of Eravacycline against 336 isolates collected from 2012 to 2016 from 11 teaching hospitals in China. *BMC Infect Dis* 2019 Jun 10; 19(1):508. doi: 10.1186/s12879-019-4093-1.
23. Jacobs MR, Abdelhamed AM, Good CE, Rhoads DD, Hujer KM, Hujer AM, et al. ARGONAUT-I: Activity of Cefiderocol (S-649266), a Siderophore Cephalosporin, against Gram-Negative Bacteria, Including Carbapenem-Resistant Nonfermenters and *Enterobacteriaceae* with Defined Extended-Spectrum β -Lactamases and Carbapenemases. *Antimicrob Agents Chemother* 2019. <https://doi.org/10.1128/AAC.01801-18>.
24. Kresken M, Korte-Berwanger M, Gatermann SG, Pfeifer Y, Pfennigwerth N, Seifert H, et al. In vitro activity of cefiderocol against aerobic Gram-negative bacterial pathogens from Germany. *Int J Antimicrob Agents* 2020; 56:106128.
25. Bassetti M, Echols R, Matsunaga Y, Ariyasu M, Doi Y, Ferrer R, et al. Efficacy and safety of cefiderocol or best available therapy for the treatment of serious infections caused by carbapenem-resistant Gram-negative bacteria (CREDIBLE-CR): a randomised, open-label, multicentre, pathogen-focused, descriptive, phase 3 trial. *Lancet Infect Dis* 2021; 21:226-240.
26. Trecarichi EM, Quirino A, Scaglione V, Longhini F, Garofalo E, Bruni A, et al. Successful treatment with cefiderocol for compassionate use in a critically ill patient with XDR *Acinetobacter baumannii* and KPC-producing *Klebsiella pneumoniae*: a case report. *J Antimicrob Chemother* 2019; 74:3399-3401.
27. Karaiskos I, Souli M, Galani I, Giamarellou H. Colistin: still a lifesaver for the 21st century? *Expert Opin Drug Metab Toxicol* 2017; 13:59-71.
28. Tsuji BT, Pogue JM, Zavascki AP, Paul M, Daikos GL, Forrest A, et al. International consensus guidelines for the

- optimal use of the polymyxins: Endorsed by the American College of Clinical Pharmacy (ACCP), European Society of Clinical Microbiology and Infectious Diseases (ESCMID), Infectious Diseases Society of America (IDSA), International Society for Anti-infective Pharmacology (ISAP), Society of Critical Care Medicine (SCCM) and Society of Infectious Diseases Pharmacists (SIDP). *Pharmacotherapy* 2019; 39:10-39.
29. Pontikis K, Karaiskos I, Bastani S, Dimopoulos G, Kalogirou M, Katsiari M, et al. Outcomes of critically ill intensive care unit patients treated with fosfomycin for infections due to pandrug-resistant and extensively drug-resistant carbapenemase-producing Gram-negative bacteria. *Int J Antimicrob Agents* 2014; 43:52-59.
 30. Albur MS, Noel A, Bowker K, MacGowan A. The combination of colistin and fosfomycin is synergistic against NDM-1-producing Enterobacteriaceae in in vitro pharmacokinetic/pharmacodynamic model experiments. *Int J Antimicrob Agents* 2015; 46:560-567.
 31. Matzi V, Lindenmann J, Porubsky C, Kugler SA, Maier A, Dittrich P, et al. Extracellular concentrations of fosfomycin in lung tissue of septic patients. *J Antimicrob Chemother* 2010; 65:995-998.
 32. Pfausler B, Spiss H, Dittrich P, Zeitlinger M, Schmutzhard E, Joukhardar C. Concentrations of fosfomycin in the cerebrospinal fluid of neurointensive care patients with ventriculostomy-associated ventriculitis. *J Antimicrob Chemother* 2004; 53:848-852.
 33. Vardakas KZ, Legakis NJ, Triarides N, Falagas ME. Susceptibility of contemporary isolates to fosfomycin: a systematic review of the literature. *Int J Antimicrob Agents* 2016; 47:269-285.
 34. Dinh A, Salomon J, Bru JP, Bernard L. Fosfomycin: efficacy against infections caused by multidrug-resistant bacteria. *Scand J Infect Dis* 2012; 44:182-189.
 35. Garnacho-Montero J, Dimopoulos G, Poulakou G, Akova M, Cisneros JM, De Waele J, et al. Task force on management and prevention of *Acinetobacter baumannii* infections in the ICU. *Intensive Care Med* 2015; 41:2057-2075.
 36. Giamarellou H, Poulakou G. Pharmacokinetic and pharmacodynamic evaluation of tigecycline. *Expert Opin Drug Metab Toxicol* 2011; 7:1459-1470.
 37. Burkhardt O, Rauch K, Kaever V, Hadem J, Kielstein JT, Welte T. Tigecycline possibly underdosed for the treatment of pneumonia: a pharmacokinetic viewpoint. *Int J Antimicrob Agents* 2009; 34:101-102.
 38. Bassetti M, Poulakou G, Giamarellou H. Is there a future for tigecycline? *Intensive Care Medicine* 2014; 40:1039-1045.
 39. White BP, Barber KE, Stover KR. Ceftaroline for the treatment of methicillin-resistant *Staphylococcus aureus* bacteremia. *Am J Health Syst Pharm* 2017; 74:201-208.
 40. Kosowska-Shick K, McGhee PL, Appelbaum PC. Affinity of ceftaroline and other beta-lactams for penicillin-binding proteins from *Staphylococcus aureus* and *Streptococcus pneumoniae*. *Antimicrob Agents Chemother* 2010; 54:1670-1677.
 41. Scheeren TWL. Ceftobiprole medocaril in the treatment of hospital-acquired pneumonia. *Future Microbiol* 2015; 10:1913-1928.
 42. Gales AC, Sader HS, Jones RN. Antimicrobial activity of dalbavancin tested against Gram-positive clinical isolates from Latin American medical centres. *Clin Microbiol Infect* 2005; 11:95-100.
 43. Klinker KP, Borgert SJ. Beyond Vancomycin: The Tail of the Lipoglycopeptides. *Clin Ther* 2015; 37:2619-2636.
 44. Boucher HW, Wilcox M, Talbot GH, Puttagunta S, Das AF, Dunne MW. Once-weekly dalbavancin versus daily conventional therapy for skin infection. *N Engl J Med* 2014; 370:2169-2179.
 45. Burdette SD, Trotman RX. Tedizolid: The First Once-Daily Oxazolidinone Class Antibiotic. *Clin Infect Dis* 2014; 61:1315-1321.
 46. Ferrández O, Urbina O, Grau S. Critical role of tedizolid in the treatment of acute bacterial skin and skin structure infections. *Drug Des Devel Ther* 2017; 11:65-82.
 47. Jorgensen SCJ, Mercurio NJ, Davis SL, Rybak MJ. Delafloxacin: Place in Therapy and Review of Microbiologic, Clinical and Pharmacologic Properties. *Infect Dis Ther* 2018; 7:197-217.
 48. Draper MP, Weir S, Macone A, Donatelli J, Trieber CA, Tanaka SK, et al. Mechanism of action of the novel aminomethylcycline antibiotic omadacycline. *Antimicrob Agents Chemother* 2014; 58:1279-1283.
 49. Villano S, Steenbergen J, Loh E. Omadacycline: development of a novel aminomethylcycline antibiotic for treating drug-resistant bacterial infections. *Future Microbiol* 2016; 11:1421-1434.

GENERAL ARTICLE

CAREER GUIDANCE FOR PEDIATRICIANS

***Julius Xavier Scott**
****Nisha Kalaiarasan**

Abstract: *Pediatrics is one of the most sought after branches amongst medical undergraduates. That said, the struggles and difficulties of establishing oneself as a successful pediatrician, after postgraduation, is not child's play. Pediatricians, at various points in life, are left at crossroads, as they even try to shift focus away from Pediatrics, as their career, owing to various compelling circumstances. This article focuses on ways and means to better oneself as a pediatrician and also the spread of other career options available to them.*

Keywords: *Pediatrics practice, Subspecialities, Foreign assignments, Work-life balance.*

Points to Remember

- *Pediatrics, as such is a blessed work irrespective of the place or position.*
- *Immediately after the MD or DNB pediatric i.e a second tier course pursuing a superspeciality course always compromise the life mile stones, which are more precious than the career.*
- *Accepting any compulsory postings, though appear to be stressful, it has its own advantages.*
- *Taking up a foreign assignment as job or postgraduate career depends on the country and one should weigh the pros and cons and also decide whether it is a short term or long term career.*
- *Community practice is not a bad option provide one choose a right place where local support is available.*
- *Whatever the career one choose, basic sacred qualities of a pediatrician can be always performed in all the situations.*
- *Many of the pediatricians balance life mile stones and career options, giving priority to the first one.*

References

1. Latha MS, Chitralakshmi K, Ravindran M, Angeline PR, Kannan L, Scott JX. Knowledge, attitude, and awareness of childhood cancer among undergraduate medical students in South India. *South Asian J Cancer* 2015;4(2):75-77. doi: 10.4103/2278-330X.155680. PMID: 25992346; PMCID: PMC4418087.
2. Anand R, Sankaran PS. Factors influencing the career preferences of medical students and interns: a cross-sectional, questionnaire-based survey from India. *J EducEval Health Prof* 2019; 16:12. doi: 10.3352/jeehp.2019.16.12. Epub 2019 May 15. PMID: 31117329; PMCID: PMC6609296.
3. Hu L, Wu H, Zhou W, Shen J, Qiu W, Zhang R, et al. Positive impact of COVID-19 on career choice in pediatric medical students: a longitudinal study. *TranslPediatr* 2020;9(3):243-252. doi: 10.21037/tp-20-100. PMID: 32775243; PMCID: PMC7347769.

* Head, Division of Pediatric Hematology and Oncology, Sri Ramachandra Institute of Higher Education and Research, Chennai.

email: juliusxscott@sriramachandra.edu.in

** Post Doctoral fellowship in Neonatology, Diploma in Allergy and Asthma, Consultant in Pediatrics and Neonatology, Sri Narayani Hospital and Research Centre, Vellore.

4. Aroskar S. Resident to President - The Chronicles of a Lady Pediatrician. *Indian Pediatr* 2019;56(10):883-884. PMID: 31724550.
5. Sells JM, Sells CJ. Pediatrician and parent: a challenge for female physicians. *Pediatrics* 1989;84(2):355-361. PMID: 2748267.
6. Emmett BM, Dovey SM, Wheeler BJ. After-hours on-call: the effect on paediatricians' spouses and families. *J Pediatr Child Health* 2013;49(3):246-250. doi: 10.1111/jpc.12108. Epub 2013 Feb 17. PMID: 23414341.
7. Accessed from <https://www.nmc.org.in/information-desk/college-and-course-search/> last accessed 14th Feb, 2022.
8. Accessed from <https://mcc.nic.in/mccss/SeatMatrix.aspx> last accessed 25th Feb, 2022.
9. Accessed from <https://www.phoindia.org/cms/iap-pho-fellowship/> last accessed 11th Feb, 2022.
10. Accessed from <https://www.piccindia.com/idpccm-accredited-centres.php> last accessed 8th Feb, 2022.
11. Accessed from <https://nbe.edu.in/IB/Information>. last accessed 10th Feb, 2022.
12. Sneha LM, Ravindran M, Kumar R, Venkatraman P, Scott J, Kannan L. Indian Pediatric Postgraduate's Perspective on Future Career Intentions. *Indian J Pediatr* 2017;84(3):183-187. doi: 10.1007/s12098-016-2268-y. Epub 2016 Dec 7. PMID: 27924467.
13. Dennis D, van Heerden P, Khanna R, Knott C, Zhang S, Calhoun A. The Different Challenges in Being an Adult Versus a Pediatric Intensivist. *Crit Care Explor* 2022;4(3):e0654. doi: 10.1097/CCE.0000000000000654. PMID: 35261983; PMCID: PMC8893297.
14. Kumar S, Ashraf AP, Lteif A, Lynch J, Aye T. Pediatric Endocrinology: Perspectives of Pediatric Endocrinologists Regarding Career Choice and Recruitment of Trainees. *Endocr Pract* 2021;27(7):743-748. doi: 10.1016/j.jepprac.2020.12.003. Epub 2020 Dec 15. PMID: 34132198.
15. Latha MS, Thirugnanasambandam RP, Balakrishnan N, Meghanathan HS, Moorthy A, Venkatraman P, et al. The need of pediatric palliative care education among pediatric postgraduates in South India. *Indian J Pediatr* 2014;81(5):455-459. doi: 10.1007/s12098-013-1295-1. Epub 2014 Jan 11. PMID: 24408397.
16. Accessed from <https://www.residencyprogramslist.com/pediatrics> last accessed 24th Feb, 2022.
17. Accessed from <https://www.rcpch.ac.uk/resources/opportunities-uk-international-doctors/> last accessed 4th Feb, 2022.
18. Accessed from <https://www.medicalboard.gov.au> last accessed 11th Mar, 2022.
19. <https://www.headmedical.com> last accessed 4th Feb, 2022.
20. Accessed from <https://www.msf.org/> last accessed 4th Mar, 2022.

DRUG PROFILE

THERAPY OF ACNE VULGARIS

***Jeesson C.Unni**

Abstract: *Acne vulgaris is common in adolescents though children may also be affected. Topical and systemic drugs are prescribed depending on the severity and stage of the disease in a given child. Pediatricians need to be aware of the various modalities of treatment and must be actively involved in the long drawn therapy. This article reviews the various options and recommendations.*

Keywords: *Acne vulgaris, Benzoylperoxide, Azelaic acid, Retinoids, Topical antibiotics.*

Points to Remember

- *Therapy of acne targets the four factors responsible for lesion formation: increased sebum production, hyperkeratinization, colonization by Propionibacterium acnes and the resultant inflammatory reaction.*
- *Treatment goals include scar prevention, reduction of psychological morbidity and resolution of lesions.*
- *Mild to moderate acne is treated with topical preparations, such as benzoyl peroxide, azelaic acid, retinoids and topical antibiotics.*
- *Benzoyl peroxide is an over-the-counter bactericidal agent that does not lead to bacterial resistance.*
- *Topical retinoids are effective in treating inflammatory and noninflammatory lesions by preventing comedones, reducing existing comedones and targeting inflammation.*
- *Topical and oral antibiotics are effective, more so when combined with topical benzoyl peroxide and/or retinoids which reduces the risk of bacterial resistance.*
- *Moderate to severe inflammatory acne is treated with oral antibacterials, oral isotretinoin or Cycloprindiol (cyproterone acetate with ethinylestradiol) an antiandrogenic drug.*
- *Oral isotretinoin is approved for the treatment of severe recalcitrant acne - requires specialist intervention.*

References

1. Fabbrocini G, Annunziata MC, D'Arco V, De Vita V, Lodi G, Mauriello MC, et al. Acne scars: pathogenesis, classification and treatment. *Dermatol Res Pract* 2010; 2010:893080. doi: 10.1155/2010/893080. Epub 2010 Oct 14.
2. Samycia M, Lam JM. Infantile acne. *CMAJ* 2016; 188(17-18): E540. doi: 10.1503/cmaj.160139.
3. Layton AM. Optimal management of acne to prevent scarring and psychological sequelae. *Am J Clin Dermatol* 2001; 2(3):135-141.

* Editor-in-Chief,
IAP Drug Formulary,
Senior Associate Consultant in Pediatrics,
Aster Medcity, Kochi.
email: jeesson1955@gmail.com

4. Leung AK, Barankin B, Lam JM, Leong KF, Hon KL. Dermatology: how to manage acne vulgaris. *Drugs Context*. 2021; 10:2021-8-6. doi: 10.7573/dic.2021-8-6. ECollection 2021.
5. Bitzer J, Römer T, Lopes da Silva Filho A. The use of cyproterone acetate/ethinylestradiol in hyperandrogenic skin symptoms - a review. *Eur J Contracept Reprod Health Care* 2017; 22(3):172-182.
6. R Costa CS, Bagatin E, Martimbianco ALC, da Silva EMK, Lúcio MM, Magin P, et al. Oral isotretinoin for acne. *Cochrane Database of Systematic Reviews* 2018; 11: Art. No.: CD009435. DOI: 10.1002/14651858. CD009435. pub2.
7. Yang Z, Zhang Y, Lazic Mosler E, Hu J, Li H, Zhang Y, et al. Topical benzoyl peroxide for acne. *Cochrane Database Syst Rev* 2020; 3(3):CD011154. doi: 10.1002/14651858.CD011154.pub2.
8. IAP Drug Formulary. 5th Ed. Eds: Jeelson C Unni, MKC Nair, PSN Menon. Pixel Studio, Cochin. 2019.
9. Singh SK, Chaubey S, Bansal A, Kaur G, Malik DS. Cosmeceutical Aptitudes of Azelaic Acid. *Curr Drug Res Rev* 2021; 13(3):222-229.
10. Hauk L. Acne Vulgaris: Treatment Guidelines from the AAD. *Am Fam Physician*. 2017; 95(11):740-741.
11. Tomiæ I, Mioèiæ S, Pepiæ I, Šimiæ D, Filipoviæ-Grèiæ. Efficacy and Safety of Azelaic Acid Nanocrystal-Loaded In Situ Hydrogel in the Treatment of Acne Vulgaris. *J Pharmaceutics* 2021; 13(4):567. doi: 10.3390/pharmaceutics13040567.
12. Joint Formulary Committee. *British National Formulary for children*. London: BMJ Group and Pharmaceutical Press, 2019-2020; pp777-781.
13. Thielitz A, Abdel-Naser MB, Fluhr JW, Zouboulis CC, Gollnick H. Topical retinoids in acne-an evidence-based overview. *J Dtsch Dermatol Ges* 2008; 6(12): 1023-1031.
14. Eady EA, Cove JH, Holland KT, Cunliffe WJ. Erythromycin resistant propionibacteria in antibiotic treated acne patients: association with therapeutic failure. *Br J Dermatol* 1989; 121(1):51-57.
15. Patel DJ, Bhatia N. Oral Antibiotics for Acne. *Am J Clin Dermatol* 2021; 22(2):193-204.

CASE REPORT

MALIGNANT INFANTILE OSTEOPETROSIS

***Preethi N**

****Latha Kanchi Parthasarathy**

*****Ramya Uppuluri**

******Devimeenakshi K**

Abstract: *Malignant infantile osteopetrosis is a rare congenital disorder characterized by increased bone density due to defective resorption of bone by osteoclasts. Infantile malignant osteopetrosis usually presents in infancy with bone marrow failure leading to hematological abnormalities like pancytopenia and hepatosplenomegaly due to extramedullary hematopoiesis. Though it commonly presents in early infancy, it should be suspected even in neonates presenting with characteristic features.*

Keywords: *Osteopetrosis, Bone marrow, Osteoclast.*

References

1. Wilson CJ, Vellodi A. Autosomal recessive osteopetrosis: diagnosis, management, and outcome. Arch Dis Child 2000; 83(5):449-452.
2. Stark Z, Savarirayan R. Osteopetrosis. Orphanet J Rare Dis 2009; 4(1):1-2.
3. Helfrich MH, Aronson DC, Everts V, Mieremet RHP, Gerritsen EJA, Eckhardt PG, et al, Morphologic features of bone in human osteopetrosis. Bone 1991;12(6):411-419.
4. Diniz G, Olukman O, Calkavur S, Buyukinan M, Altay C. A histologically diagnosed case with infantile osteopetrosis complicated by hypopituitarism. Case reports in pathology. 2015; Article ID 786836, 5 pages, 2015. <https://doi.org/10.1155/2015/786836>.
5. Srinivasan M, Abinun M, Cant AJ, Tan K, Oakhill A, Steward CG. Malignant infantile osteopetrosis presenting with neonatal hypocalcaemia. Arch Dis Child-Fetal and Neonatal Ed 2000; 83(1):F21-23.
6. Mazzolari E, Forino C, Razza A, Porta F, Villa A, Notarangelo LD. A singlecenter experience in 20 patients with infantile malignant osteopetrosis. Am J Hematol 2009; 84(8):473-479.
7. Key Jr LL, Ries WL, Rodriguiz RM, Hatcher HC. Recombinant human interferon gamma therapy for osteopetrosis. J Pediatr 1992;121(1):119-124.
8. Askmyr MK, Fasth A, Richter J. Towards better understanding and new therapeutics of osteopetrosis. Brit J Hematol 2008;140(6):597-609.
9. Key L, Carnes D, Cole S, Holtrop M, Bar-Shavit Z, Shapiro F, et al. Treatment of congenital osteopetrosis with high-dose calcitriol. N Engl J Med 1984;310(7):409-415.

* Resident

**** Professor,
Kilpauk Medical College, Chennai.

** Consultant Neonatologist

*** Hemato-oncologist,
Apollo Children's hospital, Chennai.
email: npreethi2007@gmail.com

CASE REPORT**SPORADIC HEMIPLEGIC MIGRAINE IN A CHILD**

***Arulmozhi T**
****Suba Rajinikanth**

Abstract: *Hemiplegic migraine is a rare form of migraine with aura presenting with reversible weakness. In an emergency room setting with a child having headache and focal deficits it is a diagnostic challenge with an immediate need to rule out underlying life-threatening conditions. Identifying this easily treatable condition will avoid undue anxiety and unnecessary investigations.*

Keywords: *Hemiplegic migraine, Childhood headaches, Focal weakness.*

References

1. Headache Classification Committee of the International Headache Society (IHS). The International Classification of Headache Disorders, 3rd edn. Cephalalgia 2018; 38(1):1-211.
2. Malik AH, Shah PA, Yaseen Y. Prevalence of primary headache disorders in school-going children in Kashmir Valley (North-west India). Ann Indian Acad Neurol. 2012; 15(Suppl 1):S100-103.
3. Mishra D, Sharma A, Juneja M. Recurrent Headache in Pediatric Outpatients at a Public Hospital in Delhi, Indian Pediatrics 2013; 50:775-778.
4. Massano D, Jullian S, Kanagarajah L, Gautier M, Vizeneuve A, Elmaleh M, et al. Headache with focal neurologic signs in children at the emergency department. J Pediatr 2014; 165(2):376-382.
5. Perkins E, Suminaite D, Jackson M. "Cerebellar ataxias: β -III spectrin's interactions suggest common pathogenic pathways." J physiol 2016; 594(16): 4661-4676.
6. Thomsen LL, Eriksen M, Roemer S, Andersen I, Olesen J, Russell M. A population based study of familial hemiplegic migraine suggests revised diagnostic criteria. Brain 2002; 125:1379-1391.
7. Thomsen LL, Ostergaard E, Olesen J, Russell MB. Evidence for a separate type of migraine with aura: sporadic hemiplegic migraine. Neurology 2003; 60(4): 595-601.

* Senior Consultant Pediatric Neurologist

** Senior Consultant Pediatrician and Neonatologist,
MGM Health Care,
Chennai.
email: arul.neurocentre@gmail.com