

Introduction:

Antimicrobial resistance is one of the major global threats is already a well-established fact humanity and even WHO (World Health Organization) express that the mortality of infection due to Methicillin Resistance *Staphylococcus Aureus* (MRSA) may be as high as 64% more than infection due to infection by Methicillin Sensitive *Staphylococcus Aureus* (MSSA)1. Even the infection atypical bacteria are also on the rise so the concern of resistance is also rising2,3.

A benzoquinolizine subclass of fluoroquinolone Levonadifloxacin (intravenous) and alalevonadifloxacin (oral prodrug) were licensed for clinical use in India in 2019. This broad-spectrum antibiotic with active moiety, levonadifloxacin, has high potency against methicillin-resistant *Staphylococcus aureus*, multi-drug resistant pneumococci, and anaerobes which is good news to the medical fraternity of the world4.

Aims and Objectives:

In this study, we tried to analyze in vitro efficacy of Levonadifloxacin in respiratory, urinary, and bloodstream infections from both community and nosocomial sources.

Materials and Methods:

We collected 250 consecutive gram-positive bacterial isolates from both community and hospital-acquired infections, including *Staphylococcus Aureus*, and *Enterococcus* strains, sourced from respiratory tract, urine, and blood samples at the Microbiology department. These isolates underwent antimicrobial susceptibility testing using the Kirby-Bauer disc diffusion method on Mueller-Hinton agar (Hi-Media), with interpretation based on the latest CLSI guidelines.

Results:

All the isolates were sensitive to Levonadifloxacin which were *Staphylococcus* including MRSA, MSSA, and Coagulase negative *Staphylococcus* Species (CONS). All *Enterococcus* isolates were sensitive to Levonadifloxacin except Vancomycin Resistant *Enterococcus* (VRE) which were all resistant to Levonadifloxacin (total no 16).

Conclusion:

In our in vitro analysis we found that Levonadifloxacin is a good drug to be used in MSSA, MRSA, and CONS where needed to be treated. VRE seems to be not a good target. However further studies are needed.

References:

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