

STUDY OF PHENOTYPIC AND GENOTYPIC CHARACTERISATION OF KLEBSIELLA SPECIES ISOLATED FROM WOUND INFECTION WITH SPECIAL REFERENCE TO CARBAPENEM RESISTANCE

INTRODUCTION

Wound infection is one of the most important cause of health care related problem because the resulting wound is being colonised by pathogenic microorganism(1) Infection is major cause of morbidity and mortality in hospitalized patients(2) Gram positive bacteria which was the predominant organism in wound infection is being replaced by gram negative bacteria in recent years(3) And *Klebsiella species* is showing rising trend in wound infection among gram negative bacteria (4). Its prevalence in burns is 34.40%(2),in surgical site infection 18.08% and in diabetic foot ulcer it is 8%(3)

AIM & OBJECTIVES:

AIM: To find the prevalence of *Klebsiella species* from wound infection and to find its molecular characterisation with special reference to carbapenem resistance

OBJECTIVE

- To isolate, characterize and speciate *Klebsiella* from wound infection.
- To identify the antibiotic sensitivity pattern of the *klebsiella* species isolated.
- To detect the Exetended spectrum beta lactamase(ESBL), Amp C and carbapenemase production among the identified *Klebsiella* species.
- To do molecular characterisation of the resistant *Klebsiella* isolates.

MATERIALS AND METHODS

The present study was conducted in the Department of Microbiology, Kilpauk medical college and hospital, Chennai from JUNE 2017 TO MAY 2018. 800 wound samples was processed by standard microbiological techniques to speciate *Klebsiella* species. Antimicrobial susceptibility pattern was identified by Kirby bauer disk diffusion test. ESBL, AMPC and Carbapenemase lactamases was identified by both screening method and confirmatory test. Molecular characterization of carbapenemase resistant *Klebsiella* species was done by PCR.

RESULTS:

Out of 800 wound samples, 113 *Klebsiella* species was isolated. 103 was *Klebsiella pneumoniae* and 10 was *Klebsiella oxytoca*. out of 103 *Klebsiella pneumoniae* isolated from wound infection, 52 (46.01%) was ESBL producer, 23(20.35%) AMPC and 9(7.96%) was carbapenemase producers. Molecular study showed all 9 strains to carry bla_{NDM}. Out of these 9 MBL, 4 strains showed co-existence of ESBL and they carried all 3 gene (bla_{SHV}, bla_{TEM}, bla_{CTX-M}) and another 4 strain carried AMPC lactamases all carrying the AMPC gene.

CONCLUSION

This study shows increased prevalence of *Klebsiella pneumoniae* in wound infection carrying multi drug resistant strain leading to multi drug resistant strain. This could be prevented by performing screening and confirmatory test routinely in microbiological laboratories, and also by following simple measures like hand washing and strict adherence to hospital antibiotic policy the occurrence and spread

of resistant strains can be prevented in wound infection . This will enable in better wound healing, there by preventing mortality and morbidity.

KEYWORDS: Klebsiella pneumoniae, ESBL, AMPC., Carbapenemase, MBL, blaNDM, blaSHV, blaTEM and blaCTX-M