

Complicated Urinary tract infections (cUTI) are a substantial cause of global morbidity, mortality, and healthcare expenditure



For serious and difficult-to-treat infections

KEFUZID-AV[®]
(Ceftazidime 2g + Avibactam 0.5g)



The phase 3 RECAPTURE program compared the efficacy and safety of Ceftazidime-Avibactam (**KEFUZID-AV**) and doripenem in patients with complicated urinary tract infection (cUTI), including acute pyelonephritis.

Methods: Hospitalized adults with suspected or microbiologically confirmed cUTI/acute pyelonephritis were randomized 1:1 to Ceftazidime-Avibactam (**KEFUZID-AV**) 2000 mg/ 500 mg every 8 hours or doripenem 500 mg every 8 hours.

In Results: Of 1033 randomized patients, 393 and 417 treated with ceftazidime-avibactam (**KEFUZID-AV**) and doripenem, respectively, were eligible for the primary efficacy analyses; 19.6% had ceftazidime-nonsusceptible baseline pathogens.

Noninferiority of Ceftazidime-Avibactam (**KEFUZID-AV**) vs. doripenem was demonstrated for the US Food and Drug Administration co-primary endpoints of (1) patient-reported symptomatic resolution at day 5: 276 of 393 (70.2%) vs 276 of 417 (66.2%) patients and (2) combined symptomatic resolution/microbiological eradication at test of cure (TOC): 280 of 393 (71.2%) vs 269 of 417 (64.5%) patients. Microbiological eradication at TOC occurred in 304 of 393 (77.4%) ceftazidime-avibactam (**KEFUZID-AV**) vs 296 of 417 (71.0%) doripenem patients. Both treatments showed similar efficacy against ceftazidime-nonsusceptible pathogens. Ceftazidime-avibactam (**KEFUZID-AV**) had a safety profile consistent with that of Ceftazidime alone.

Conclusions: Ceftazidime-Avibactam (**KEFUZID-AV**) was highly effective for the empiric treatment of cUTI (including acute pyelonephritis), and may offer an alternative to carbapenems in this setting

1. Clinical Infectious Diseases: 2016-63(6): 754-762

For Use of a Registered Medical Practitioner only
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The carbapenem sparing Treatment

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TIGEPOS-50
First glycylicycline for cSSI, cIAIs, HAP
(Tigecline 50 mg Inj, as Iyophilized cake)



For infections caused by ESBL-producing pathogens

NOVOMEROTM
EDTA

(Meropenem 1000 mg/Disodium EDTA 37.5 mg)

OBJECTIVE: To understand the outcomes of patient with various agents in the treatment of ESBL-producing bacteremia and to evaluate the efficacy of meropenem and ethylenediaminetetraacetic acid (EDTA) combination against ESBLs

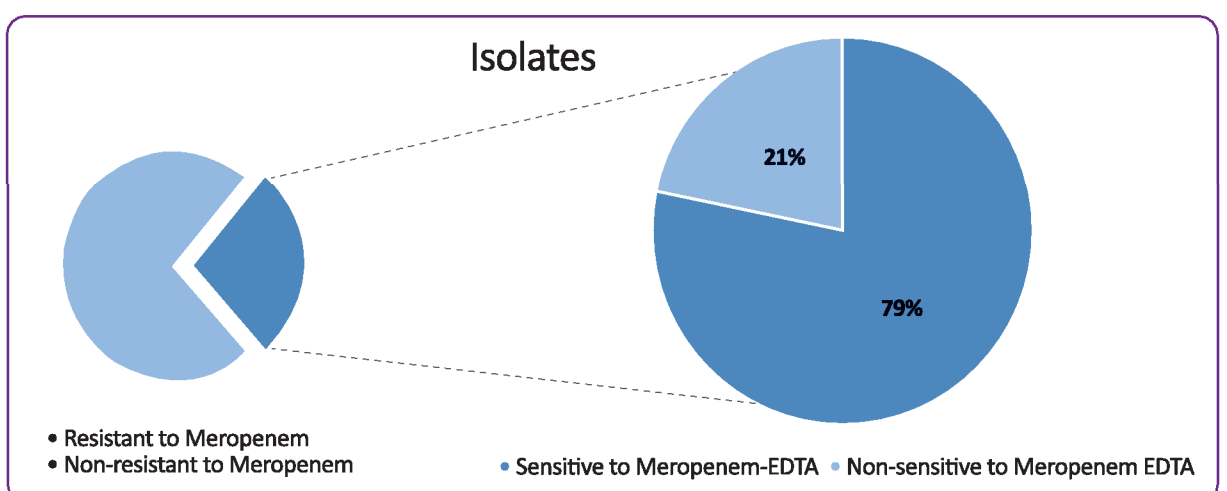


Figure 6: Division of pathogens sensitive to Meropenem and Meropenem-EDTA combination

RESULT: The MIC value of Meropenem-EDTA (0.25) was less than 50% of that of Meropenem (2.45) in sensitive isolates.

CONCLUSION: Meropenem in unification with EDTA can exhibit more potent antimicrobial activity against ESBL producing pathogens than just Meropenem or EDTA alone.

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For Sustained Synergistic Bactericidal Effects

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