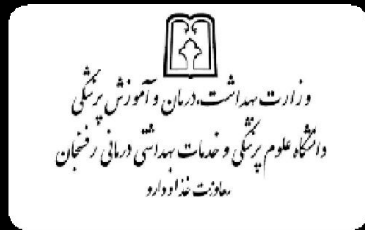



IN THE NAME OF GOD




GUIDELINE OF CARBAPENEMS

Carbapenems are useful for treatment of *moderate to severe* infections, including:

- hospital-acquired pneumonia
- intra-abdominal infections
- complicated urinary tract infections like nosocomial infections and
- bacteremia
- bone and soft tissue infections
- obstetrics and gynecological infections.



Carbapenems other than ertapenem are drugs of choice for treatment of infections caused by multidrug resistant strains of *A.baumannii* that remain susceptible to carbapenems; however, resistance to carbapenems is rapidly increasing.



Meropenem is the only carbapenem approved by FDA for treatment of bacterial meningitis. Imipenem should be avoided because of its propensity to cause seizures.

Mandell, Douglas, and Bennett's
**Principles and Practice of
 INFECTIOUS DISEASES**

TABLE 75-4 Empirical Antibiotic Options for Patients with Severe Sepsis or Septic Shock

	SUSPECTED SOURCE				
	Lung	Abdomen	Skin/Soft Tissue	Urinary Tract	Source Uncertain
Major Community-Acquired Pathogens	<i>Streptococcus pneumoniae</i> <i>Haemophilus influenzae</i> <i>Legionella</i> <i>Chlamydia pneumoniae</i>	<i>Escherichia coli</i> <i>Bacteroides fragilis</i>	<i>Streptococcus pyogenes</i> <i>Staphylococcus aureus</i> Polymicrobial	<i>E. coli</i> <i>Klebsiella</i> species <i>Enterobacter</i> species <i>Proteus</i> spp. Enterococci	
Empirical Antibiotic Therapy	Moxifloxacin or levofloxacin or azithromycin plus cefotaxime or ceftazidime or cefepime or piperacillin-tazobactam	Imipenem or meropenem or doripenem or piperacillin-tazobactam ± aminoglycoside If biliary source: piperacillin-tazobactam, ampicillin-sulbactam, or ceftriaxone with metronidazole	Vancomycin or daptomycin plus either imipenem or meropenem or piperacillin-tazobactam; ± clindamycin (see text)	Ciprofloxacin or levofloxacin (if gram-positive cocci, use ampicillin or vancomycin ± gentamicin)	Vancomycin plus either doripenem or ertapenem or imipenem or meropenem
Major Commensal or Nosocomial Microorganisms	Aerobic gram-negative bacilli	Aerobic gram-negative rods Anaerobes <i>Candida</i> spp.	<i>Staphylococcus aureus</i> (? MRSA) Aerobic gram-negative rods	Aerobic gram-negative rods Enterococci	Consider MDRO if in area of high prevalence Consider echinocandin if neutropenic or indwelling intravascular catheter
Empirical Antibiotic Therapy	Imipenem or meropenem or doripenem or cefepime (if <i>Acinetobacter baumannii</i> or carbapenem-resistant <i>Klebsiella</i> in ICU, add colistin)	Imipenem or meropenem ± aminoglycoside (consider echinocandin)	Vancomycin or daptomycin plus imipenem-cilastatin or meropenem or cefepime, ± clindamycin	Vancomycin plus imipenem or meropenem or cefepime	Cefepime plus vancomycin ± caspofungin

Dosages for intravenous administration (normal renal function):

*Imipenem-cilastatin, 0.5-1.0 g q6-8h

*Meropenem, 1-2 g q8h

*Doripenem, 0.5 g q8h

Piperacillin-tazobactam, 3.375 g q4h or 4.5 g q6h

Vancomycin, load 25-30 mg/kg, then 15-20 q8-12h

Cefepime, 1-2 g q8h

Levofloxacin, 750 mg q24h

*Carbapenems are less susceptible to extended-spectrum β-lactamases; base choice on local resistance pattern.

ICU, intensive care unit; MDRO, multidrug-resistant organisms; MRSA, methicillin-resistant *Staphylococcus aureus*.

For MDRO, resistance usually includes carbapenems.

Ciprofloxacin, 400 mg q8-12h

Moxifloxacin, 400 mg qd

Ceftriaxone, 2.0 g q24h

Caspofungin, 70 mg, followed by 50 mg q24h

Colistin: loading dose = 5 mg/kg body weight. For maintenance dosing,

see University of California, Los Angeles Dosing Protocol:

www.infectiousdiseases-ucla-affiliated.org/Intranet/FILES/ColistinDosing.pdf

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TABLE 70-4 Empirical Antibiotic Options for Patients with Severe Sepsis or Septic Shock

	Suspected Source				
	<i>Lung</i>	<i>Abdomen</i>	<i>Skin/Soft Tissue</i>	<i>Urinary Tract</i>	<i>Meninges</i>
Major community-acquired pathogens	<i>Streptococcus pneumoniae</i> <i>Haemophilus influenzae</i> <i>Legionella</i> <i>Chlamydomphila pneumoniae</i>	<i>Escherichia coli</i> <i>Bacteroides fragilis</i>	<i>Streptococcus pyogenes</i> <i>Staphylococcus aureus</i> Polymicrobial	<i>E. coli</i> <i>Klebsiella</i> species <i>Enterobacter</i> species <i>Proteus</i> species Enterococci	<i>S. pneumoniae</i> <i>Neisseria meningitidis</i> <i>Listeria monocytogenes</i> <i>H. influenzae</i>
Empirical antibiotic therapy	Moxifloxacin <i>plus either</i> cefotaxime <i>or</i> ceftriaxone	Imipenem-cilastatin* <i>or</i> meropenem <i>or</i> piperacillin-tazobactam ± aminoglycoside	Vancomycin <i>plus either</i> imipenem <i>or</i> meropenem <i>or</i> piperacillin-tazobactam	Ciprofloxacin <i>or</i> levofloxacin (If gram-positive cocci, use ampicillin <i>plus</i> gentamicin)	Vancomycin <i>plus</i> ampicillin <i>plus either</i> ceftriaxone <i>or</i> cefepime
Major commensal or nosocomial microorganisms	Aerobic gram-negative bacilli	Aerobic gram-negative rods Anaerobes <i>Candida</i> species	<i>Staphylococcus aureus</i> (? MRSA) Aerobic gram-negative rods	Aerobic gram-negative rods Enterococci	Aerobic gram-negative rods Staphylococci
Empirical antibiotic therapy*	Imipenem-cilastatin* <i>or</i> meropenem <i>or</i> cefepime	Imipenem-cilastatin* <i>or</i> meropenem ± aminoglycoside (Consider amphotericin B)	Vancomycin <i>plus</i> imipenem-cilastatin* <i>or</i> meropenem <i>or</i> cefepime	Vancomycin <i>plus</i> imipenem-cilastatin* <i>or</i> meropenem <i>or</i> cefepime	Cefepime <i>plus</i> vancomycin

Dosages for intravenous administration (normal renal function):

Imipenem-cilastatin, 0.5 g q6h

Meropenem, 1.0 g q8h

Piperacillin-tazobactam, 3.375 g q6h

Vancomycin, 15 mg/kg q12h (if meningitis, 25 mg/kg q12h)

Cefepime, 1-2 g q8h

Ciprofloxacin, 400 mg q12h, Moxifloxacin 400 mg qd

Ceftriaxone, 2.0 g q24h (if meningitis, 2 g q12h)

Levofloxacin, 750 mg q24h

Gentamicin, 7 mg/kg q24h

Ampicillin, 2 g q4h

Cefotaxime, 2 g q4h

*Carbapenems are less susceptible to extended-spectrum beta-lactamases (ESBL); base choice on local resistance pattern.

MRSA, methicillin-resistant *S. aureus*.

Format adapted from Simon D, Trenholme G. Antibiotic selection for patients with septic shock. *Crit Care Clin.* 2000;16:215-230.

در عفونت هایی که اندیکاسیون استفاده از کارباپنم وجود ندارد و یا نیاز به استفاده empirical از کارباپنم ها نیست استفاده از پپراسیلین/تازوباکتام گزینه بسیار مناسبی است به خصوص در عفونت های اکتسابی از جامعه. حتی استفاده از آمپی سیلین/سولباکتام در برخی مواد عفونت های خفیف و حساس توصیه شده است.



The End

