



**Central Administration of Pharmaceutical Care  
General Administration For Drug Utilization and Pharmacy Practice Administration**

# **National Guidelines for Preauthorization of Restricted Antimicrobials in Hospitals National Antimicrobial Rational Use Committee 2022**

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## Table of Contents

<b>Content</b>	<b>Page</b>
<b>Antimicrobial Preauthorization Policy</b>	2
<b>Introduction</b>	2
<b>Purpose of The Guidance</b>	3
<b>Policy Statement</b>	3
<b>Procedures for Antimicrobial Preauthorization Policy</b>	4
<b>Performance Metrics (KPIs)</b>	5
<b>The Criteria of Antimicrobials Needed to Be Preauthorized</b>	6
<b>Guide for Antimicrobial Preauthorization List</b>	6
<b>Annex I: List of Watch and Reserve Antibiotics</b>	7
<b>Annex II: Sample Preauthorization Prescribing Form</b>	9
<b>Preauthorized Antibiotic Criteria Stickers</b>	10
<b>Contributors</b>	14
<b>References</b>	15

## Antimicrobial Preauthorization Policy

### 1. Introduction

Hospital formularies include a variety of anti-infective agents that are ideally selected based on the needs of the facility. A number of these agents require some sort or extent of restriction or specific guidance for use to avoid the emergence of resistance. Antimicrobial restriction is restricted dispensing of targeted antimicrobial agent(s) on the hospital's formulary, according to approved criteria. Implementing antimicrobial restriction is one of the antimicrobial stewardship interventions for improving the prescribing practices. Antimicrobials preauthorization (expert approval prior to prescription) is considered one of the restrictive interventions (limitations to prescribing targeted antibiotics). Other restrictive interventions include: compulsory order forms for targeted agents, automatic stop orders (e.g. after a single dose of surgical prophylaxis), selective susceptibility reporting from the laboratory (report susceptible first-line narrow spectrum antibiotics to regular wards).

In terms of the rationale for restricting individual antimicrobial agents, recommendations are made based on patient safety, disease state complexity, risk of antimicrobial resistance, best practice prescribing and public health interests. Pharmacoeconomic considerations may also be taken into account, however expenditure should rarely be the sole factor involved in a recommendation for antimicrobial restriction.

Carbapenems are considered as an example of class of the antibiotics which needed to be restricted at many healthcare facilities. As per the national database of Egypt and the antimicrobial consumption data, carbapenem antibiotics are among the most antibiotics irrationally used and this leads to emergence of carbapenem resistance. Carbapenems are a class of beta-lactam antibiotics that are active against many aerobic and anaerobic gram-positive and gram-negative organisms.

The emergence of Carbapenem-resistance has become a serious public worldwide health threat. This type of antimicrobial resistance is spreading at an alarming rate, resulting in major outbreaks and treatment failure of community-acquired and nosocomial infections caused by the clinically relevant carbapenem-resistant Enterobacteriaceae.

#### List of Carbapenems available in Egypt:

- Ertapenem
- Imipenem / Cilastatin
- Meropenem

National Office for Handling and Reduction of Medication Errors (NO HARME)/EDA received 24522 reports related to antimicrobial use from 86 hospitals of different sectors all over the different cities of Egypt during the period from March 2020 to July 2021, the reports were analyzed and given the following results regarding Carbapenems:

- Carbapenems are among the most common antimicrobial classes to be associated with Drug Therapy Problems (DTPs).
- The most common Carbapenems DTP/ME (medication error) are inappropriate dose/interval (66% of Carbapenems DTPs/MEs) and Carbapenems prescribing with no valid indication.

## 2. Purpose of The Guidance

- Ensures the prudent use of antimicrobials (help control use of antimicrobials).
- Reduces rates of emerging resistance.
- Avoids overuse or misuse of broad spectrum antibiotics (e.g., use of broad-spectrum agents where narrower spectrum agents are more appropriate).
- Reserves antimicrobials for treatment of multi-drug-resistant organisms.
- Improves the quality of antimicrobial prescribing.
- Delivers a range of positive outcomes across many areas of patient care.
- Decreases risk for serious adverse effects and the medication errors.
- Pharmacoeconomic considerations (reduce medicine costs for hospitals over time).

## 3. Policy Statement

- Each hospital's antimicrobial prescribing guidelines must include a list that stipulates which antimicrobials are preauthorized (approval of authorized person/a specialist is required).
- Preauthorization policy is typically developed by the hospital antimicrobial stewardship program or antimicrobial subcommittee/team, and should be periodically reviewed and updated.
- Preauthorization prescribing form should be completed by the prescriber and approved by the antimicrobial stewardship team member (authorized person).

#### 4. Procedures for Antimicrobial Preauthorization Policy

- 1- Head of pharmacy assigns a team of clinical pharmacists and supply chain pharmacists to prepare the list of available antimicrobials and the antimicrobials suggested to be preauthorized (antimicrobial preauthorization list) N.B., The criteria of antimicrobials needed to be preauthorized and the antimicrobial preauthorization list which mentioned in this document can be used as a guidance.
- 2- Head of pharmacy presents a list of hospital available antimicrobials and the suggested antimicrobial preauthorization list in the relevant committee meeting e.g., antimicrobial stewardship committee / rational antimicrobial use committee / drug and therapeutics committee.
- 3- The responsible committee members discuss the suggested antimicrobial preauthorization list, approve it and assign an authorized person/s (e.g., clinical pharmacist who is a member of the antimicrobial stewardship team (AMS clinical pharmacist)) to approve the use of preauthorized antimicrobials.
- 4- The antimicrobial preauthorization list and the policy should be available to all healthcare providers.
- 5- The antimicrobial preauthorization list must be reviewed on a regular basis, in light of the hospital's antimicrobial usage data and rates of antimicrobial resistance.
- 6- An authorized person will be available promptly through a clearly defined pathway 24 hours/seven days a week OR the antimicrobials can be dispensed during weekends, holidays and nightshifts till the authorization next day (N.B., any antimicrobial agent proven to be prescribed and dispensed inappropriately should be stopped).
- 7- If a physician is considering prescribing one or more preauthorized antimicrobial agents, he/she should contact the authorized person on-call.
- 8- The prescriber should complete a preauthorization prescribing form with clinical details to justify use of the preauthorized antimicrobial agents, including the following information:
  - a. Patient identification(s), patient`s demographics (gender, age, weight and height of the patient).
  - b. Diagnosis, indication and site of infection.
  - c. Clinical indication for antimicrobial therapy.
  - d. Drug allergies.
  - e. Microbiology culture results, if available.
  - f. Renal and hepatic function.
  - g. Patient`s clinical notes.
  - h. Prescriber`s signature.

#### 4. Procedures for Antimicrobial Preauthorization Policy, Cont.,

- 9- Authorized person will respond to the prescriber in a timely manner, and on the basis of the above information, may:
- Recommend the use of the restricted agent(s)
  - Recommend an alternative therapeutic option
  - Recommend further investigations or clinical follow-up
- 10- The authorized person should complete a preauthorization prescribing form:
- Specify whether approve or not.
  - Rationale if not approved.
  - Comments and recommendations.
  - Signature and date.
- 11- Dispensing pharmacist should confirm approval before dispensing preauthorized antimicrobials, sign and date the preauthorization prescribing form upon dispensing.
- 12- Preauthorization approval should be updated every 7 days to assess whether the preauthorized antimicrobial agent should be continued or stopped or replaced by other antimicrobials (depending on the culture results and patient response), dispensing pharmacist should confirm that the approval gets updated by the authorized person before dispensing.
- 13- ASP team should train healthcare professionals about the policy, distributing educational materials; auditing and giving feedback to ensure applying the policy (to determine if preauthorization policy is followed).

#### 5. Performance Metrics (KPIs)

The following KPIs can be used to assess the performance and impact of the antimicrobial preauthorization policy:

- Consumption of preauthorized antimicrobials e.g., carbapenems.
- Compliance with preauthorization policy.
- Resistance to the preauthorized antimicrobials.

## 6. The Criteria of Antimicrobials Needed to Be Preauthorized

A number of agents require restrictions or specific guidance for use because of one or more of the following:

- Potential to promote resistance e.g., some antibiotics in watch category of AWaRe. (Annex I)
- Need to reserve for treatment of multi-drug-resistant organisms e.g., Reserve category of AWaRe. (Annex I)
- Potential for/documented overuse or misuse (e.g., use of broad-spectrum agents where narrower spectrum agents are more appropriate).
- Pharmacoeconomic considerations.
- Threaten the patient safety.

## 7. Guide for Antimicrobial Preauthorization List

The following antimicrobial classes, or specific agents, should be considered for inclusion in preauthorization lists, following local agreement from the responsible Committee (N.B., The agents in this table fulfil the criteria of antimicrobials needed to be preauthorized):

Antimicrobial	Rational for preauthorization
Colistin IV Tigecycline Aztreonam Ceftaroline fosamil Ceftazidime and Avibactam Ceftolozone and Tazobactam Tedizolid Antivirals: Ganciclovir, Valganciclovir Antifungals: Micafungin, Anidulafungin, Caspofungin, Voriconazole & Amphotericin B	<ul style="list-style-type: none"> <li>• Used in the management of complex, resistant or emerging infectious diseases</li> <li>• Rarely indicated as a first line agent</li> </ul>
Carbapenems (e.g., meropenem, imipenem, ertapenem)	<ul style="list-style-type: none"> <li>• Broad-spectrum agent; linked with emergence of multi-resistant Gram-negative organisms.</li> </ul>
Azithromycin	<ul style="list-style-type: none"> <li>• It is one of the most frequently used antibiotic substances (according to WHO GLASS report for the Egyptian antimicrobial consumption data).</li> </ul>
Fluoroquinolones (e.g., ciprofloxacin, ofloxacin, levofloxacin)	<ul style="list-style-type: none"> <li>• linked with emergence of multi-resistant</li> </ul>
Teicoplanin Linezolid	<ul style="list-style-type: none"> <li>• Rarely indicated as a first line agent.</li> </ul>
Aminoglycosides (e.g., gentamicin)	<ul style="list-style-type: none"> <li>• High toxicity</li> <li>• It is one of the most frequently used antibiotic substances (according to WHO GLASS report for the Egyptian antimicrobial consumption data).</li> </ul>

**N.B.**, the above list is not exhaustive, and hospitals may choose to include other agents on preauthorization lists such as anti-fungal agents, anti-viral agents, or other antibiotics.

**N.B.**, National policies issued by the National Committee for the Rational Antimicrobial Use at the Egyptian Drug Authority include guiding procedures can be tailored according to the different circumstances within each entity.

## Annex I: List of Antibiotics (Watch and Reserve according to WHO AWaRe list) Registered in Egyptian Drug Authority Database

Watch	Reserve
<p>This group includes:</p> <ul style="list-style-type: none"> <li>• Antibiotic classes that have higher resistance potential.</li> <li>• Selected Watch group antibiotics are recommended as essential first or second choice empiric treatment options for a limited number of specific infectious syndromes.</li> </ul>	<p>This group includes:</p> <ul style="list-style-type: none"> <li>• Antibiotics and antibiotic classes that should be reserved for treatment of confirmed or suspected infections due to multi- drug-resistant organisms.</li> <li>• Reserve group antibiotics should be treated as “last resort” options.</li> <li>• These antibiotics should be tailored to highly specific patients and settings, when all alternatives have failed or are not suitable.</li> </ul>
Azithromycin	Aztreonam
Cefaclor	Ceftaroline fosamil
Cefdinir	Ceftazidime and Avibactam
Cefepime	Ceftolozane and Tazobactam
Cefixime	Colistimethate IV/Oral
Cefoperazone	Linezolid
Cefotaxime	Tedizolid
Cefoxitin	Tigecycline
Cefpodoxime	Dalbavancin
Cefprozil	Daptomycin
Ceftazidime	
Ceftriaxone	
Cefuroxime	
Ciprofloxacin	



Clarithromycin	
Cefodizime	
Cefonicid	
Cefotetan	
Cefotiam	
Cefpirome	
Ceftibuten	
Fusidic-acid	
Josamycin	
Midecamycin	
Miocamycin	
Pefloxacin	

## Annex II: Sample pre-authorization prescribing form

Date: -----

<b>Patient information &amp; Diagnosis</b>				
Patient name:	Medical record number:	Department:	Ward:	
Age:	Weight:      Height:	Sex: Male <input type="checkbox"/> Female <input type="checkbox"/>		Allergies:
Serum Creatinine:	Cr Cl/GFR:      ALT:      AST:	<b>Diagnosis:</b>		
<b>Indication for antimicrobial treatment</b>				
<input type="checkbox"/> Community acquired <input type="checkbox"/> Hospital acquired/health care associated (number of days after admission:      )				
<b>Site of infection e.g., respiratory, urinary tract:</b>				
<b>Request for pre-authorized/restricted antimicrobials</b>				
Antimicrobial (s) requested	Dose and duration	Administration route	Interval	Reason for request
<b>Has the patient already received antimicrobial(s)?      Yes <input type="checkbox"/>      No <input type="checkbox"/>      If yes, what?</b>				
Antimicrobial (s) prescribed	Dose and duration	Administration route	Interval	Why is the treatment not adequate?
<b>Requesting physician's name/contact number:</b> _____				
<b>Patient Response</b>				
	Baseline (1 <sup>st</sup> approval)	2 <sup>nd</sup> approval	3 <sup>rd</sup> approval	
Clinical Parameters				
Lab Parameters				
Culture (date, specimen, pathogen sensitivity)				
Others				
<b>Decision</b>				
1st approval      Yes      No Reasons for not approving		Name/signature of approver: Dispensing pharmacist:		Date:
2nd approval (after 7 days)      Yes      No Reasons for not approving		Name/signature of approver: Dispensing pharmacist:		Date:
3rd approval (after 14 days)      Yes      No Reasons for not approving		Name/signature of approver: Dispensing pharmacist:		Date:

Reference: WHO practical toolkit Antimicrobial Stewardship Programs in health-care facilities in low- and middle-income countries

## 8. Preauthorized Antibiotic Criteria Stickers

Antibacterial Agents	
Amikacin / Gentamicin	Azithromycin
<p>Treatment of serious infections due to gram-negative organisms.</p> <ul style="list-style-type: none"> <li>• bloodstream infection</li> <li>• bone infection</li> <li>• respiratory tract infection</li> <li>• endocarditis</li> </ul>	<ul style="list-style-type: none"> <li>• Pneumonia, community-acquired</li> <li>• Pelvic inflammatory disease</li> <li>• Uncomplicated Skin and skin structure infection</li> </ul>
Aztreonam	Colistin IV
<p>Treatment of the following infections caused by gram-negative bacilli</p> <ul style="list-style-type: none"> <li>• Urinary tract infections</li> <li>• lower respiratory tract infections</li> <li>• septicaemia</li> <li>• skin/skin structure infections</li> <li>• Intra-abdominal infections</li> <li>• Gynaecological infections</li> </ul>	<ul style="list-style-type: none"> <li>• Highly resistant gram-negative infections for which alternative therapies are not injectable appropriate</li> </ul>
Ceftazidime and Avibactam Ceftolozone and Tazobactam	Ceftaroline fosamil
<ul style="list-style-type: none"> <li>• Intra-abdominal infections</li> <li>• Pneumonia, hospital-acquired and ventilator-associated</li> <li>• Urinary tract infection, complicated (pyelonephritis or urinary tract infection with systemic signs/symptoms)</li> </ul>	<ul style="list-style-type: none"> <li>• Pneumonia</li> <li>• Skin and soft tissue infections</li> <li>• Bloodstream infection (pathogen-directed therapy for methicillin-resistant <i>S. aureus</i>)</li> </ul>
Ciprofloxacin IV	Ertapenem
<ul style="list-style-type: none"> <li>• Treatment of complicated urinary tract infections and pyelonephritis</li> <li>• complicated intra-abdominal infections (in combination with metronidazole)</li> <li>• Surgical site infection</li> <li>• Peritoneal dialysis catheter-related infection</li> <li>• Neutropenic fever, low-risk cancer patients (empiric therapy)</li> <li>• Diabetic foot</li> </ul>	<ul style="list-style-type: none"> <li>• Intra-abdominal infection, mild to moderate, community-acquired in patients</li> <li>• Pelvic infection</li> <li>• Pneumonia, community acquired</li> <li>• Skin and skin structure infection, complicated</li> <li>• Urinary tract infection, complicated</li> </ul> <p>N.B., not used for Pseudomonas caused infections</p>

Imipenem	Levofloxacin
<ul style="list-style-type: none"> <li>Bloodstream infection (gram-negative bacteraemia)</li> <li>Bone and joint infections</li> <li>Gynaecologic infections</li> <li>Intra-abdominal infection</li> <li>Pneumonia</li> <li>Skin and soft tissue infection, moderate to severe</li> <li>Urinary tract infection (complicated and uncomplicated)</li> </ul>	<ul style="list-style-type: none"> <li>Community-acquired pneumonia</li> <li>Nosocomial pneumonia</li> <li>Acute exacerbation of Chronic obstructive pulmonary disease</li> <li>Urinary tract infection</li> <li>Skin or skin structure infections</li> </ul>
Linezolid	Meropenem
<ul style="list-style-type: none"> <li>Enterococcal infections (vancomycin-resistant)</li> <li>Pneumonia</li> <li>Skin and skin structure infections</li> </ul> <p>N.B., Linezolid is not indicated for treatment of gram-negative infections; if a concomitant gram-negative pathogen is documented or suspected, initiate specific therapy immediately.</p>	<ul style="list-style-type: none"> <li>Intra-abdominal infection</li> <li>Meningitis, bacterial</li> <li>Skin and skin structure infection</li> <li>Bloodstream infection (gram-negative bacteraemia)</li> </ul>
Tigecycline	Tedizolid
<ul style="list-style-type: none"> <li>Intra-abdominal infection</li> <li>Pneumonia, community acquired</li> <li>Skin and skin structure infections, complicated</li> </ul> <p>N.B., Not indicated for treatment of diabetic foot infections, hospital-acquired or ventilator-associated pneumonia</p>	<ul style="list-style-type: none"> <li>Acute bacterial skin and soft tissue infections caused by susceptible isolates of gram-positive microorganisms</li> </ul>
Teicoplanin	
<ul style="list-style-type: none"> <li>Serious infections (e.g., endocarditis, bacteraemia).</li> <li>Severe sepsis or septic shock and bone/joint infections.</li> </ul>	
Antiviral Agents	
Ganciclovir / Valganciclovir	
<ul style="list-style-type: none"> <li>Cytomegalovirus disease, prophylaxis (transplant recipients).</li> <li>Cytomegalovirus retinitis (immunocompromised patients)</li> </ul>	

Antifungal Agents	
<b>Amphotericin B</b>	<b>Liposomal Amphotericin B</b>
<ul style="list-style-type: none"> <li>• Treatment of patients with progressive and potentially life-threatening fungal infections N.B., it should not be used to treat non-invasive forms of fungal disease such as oral thrush, vaginal candidiasis, and oesophageal candidiasis in patients with normal neutrophil counts.</li> </ul>	<ul style="list-style-type: none"> <li>• Treatment of cryptococcal meningitis in patients with HIV.</li> <li>• Empiric treatment in febrile neutropenic patients with presumed fungal infection.</li> <li>• Systemic fungal infections in patient's refractory to conventional amphotericin B deoxycholate therapy or when renal impairment or unacceptable toxicity precludes the use of the deoxycholate formulation.</li> <li>• Leishmaniasis (visceral)</li> </ul>
<b>Anidulafungin</b>	<b>Caspofungin</b>
<ul style="list-style-type: none"> <li>• Treatment of candidemia and other forms of Candida infections (intra-abdominal abscess and peritonitis).</li> <li>• Candidiasis, oesophageal, refractory disease</li> </ul>	<ul style="list-style-type: none"> <li>• Aspergillosis, invasive in patients who are refractory to or intolerant of other therapies (e.g., amphotericin B, lipid formulations of amphotericin B).</li> <li>• Treatment of candidemia and the following Candida infections Intra-abdominal abscesses, peritonitis, and pleural space infections.</li> <li>• Candidiasis, oesophageal</li> <li>• Empiric therapy for presumed fungal infections in febrile neutropenia.</li> </ul> <p><b>Limitations of use:</b></p> <ul style="list-style-type: none"> <li>▪ Has not been studied in endocarditis, osteomyelitis, and meningitis due to Candida.</li> <li>▪ Not approved for the treatment of oropharyngeal candidiasis.</li> </ul>
<b>Micafungin</b>	<b>Voriconazole</b>
<ul style="list-style-type: none"> <li>• Candidemia, acute disseminated candidiasis, and Candida peritonitis and abscesses</li> <li>• Oesophageal candidiasis</li> <li>• Prophylaxis against invasive fungal infections (hematopoietic cell transplant recipients)</li> </ul>	<ul style="list-style-type: none"> <li>• Treatment of fungal infections in patient's intolerant of, or refractory to, other therapy</li> </ul>

## 9. Contributors

Editorial board	
<p><b>Dr. Shima Nasr</b> Head of Rational Drug Use unit at Drug Utilization &amp; Pharmacy Practice General Administration – EDA Member of National AMC/AMU Team</p>	<p><b>Dr. Lobna Samy</b> Head of Pharmaceutical Care Initiatives Unit at Drug Utilization &amp; Pharmacy Practice General Administration – EDA Member of National AMC/AMU Team</p>
<p><b>Dr. Mohammed Elsayed Eldesokey</b> Drug Utilization &amp; Pharmacy Practice General Administration member (GA DU&amp;PP) - EDA</p>	<p><b>Dr. Nesma Atef</b> Head of Drug Formulary Unit at Drug Utilization &amp; Pharmacy Practice General Administration – EDA</p>
Members of the National Rational Antimicrobial Use Committee (Ordered Alphabetically)	
<p><b>Dr. Ahmed Motawea</b> Chief of medical supply dept. Armed Forces Medical Services Authority</p>	<p><b>Dr. Hema Soliman</b> Quality and Patient Safety Consultant (GAHAR Representative)</p>
<p><b>Dr. Asaad Sadek</b> Acting Director IPC General Directorate (MOHP Representative)</p>	<p><b>Prof. Dr. Maha Abdel Aziz El-touny</b> Prof. Internal medicine ASU. IPC consultant Ministry of Interior</p>
<p><b>Dr. Elizabeth Tayler</b> Senior Technical Officer Antimicrobial Resistance and Communicable Disease WHO Egypt country office</p>	<p><b>Dr. Mohammed Abdelfattah</b> Head of the central administration for preventive affairs (MOHP Representative)</p>
<p><b>Dr. Eman Nadim</b> Clinical pharmacist at the central administration for unified procurement (The Egyptian Authority for unified procurement Representative)</p>	<p><b>Prof. Dr. Nirmeen Ahmed Sabry</b> Professor of clinical pharmacy Cairo University Medication management consultant</p>
<p><b>Prof. Dr. Ghada Esmail</b> Prof. of Clinical Pathology (Microbiology) at Faculty of Medicine Ain Shams University. Head of IPC University Hospitals (University Hospitals representative)</p>	<p><b>Dr. Sherif Kamal</b> Consultant of the General Authority of Healthcare (GAH Representative) Clinical pharmacy programs director Children cancer hospital Egypt</p>
<p><b>Dr. Ghada Ali Younis</b> Former General Manager of the Drug Utilization and Pharmacy Practice General Administration at the Egyptian Drug Authority</p>	<p><b>Dr. Shereen Abdel Gawad</b> Head of the Pharmaceutical Care Central Administration – Head of National Rational Antimicrobial Use Committee-EDA.</p>

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