

Republic of the Philippines PHILIPPINE HEALTH INSURANCE CORPORATION

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PhilHealth Circular No. <u>023-2015</u>

TO

ALL ACCREDITED HEALTH CARE PROVIDERS,

PHILHEALTH MEMBERS, PHILHEALTH REGIONAL

OFFICES AND ALL OTHER CONCERNED

SUBJECT

Policy statements on the Diagnosis, Empiric Management, and Prevention of Community-acquired Pneumonia (CAP) in Immunocompetent Adults as reference by the Corporation in

ensuring quality of care

I. RATIONALE

The revised Implementing Rules and Regulations of the National Health Insurance Act of 2013 (RA 7875 as amended by RA 9241 and RA 10606) under Title V (Quality Assurance and Accreditation) Rule 1 (Quality Assurance) Section 51 provides the implementation of quality assurance standards as reference for ensuring quality of care services.

Compliance to clinical practice guidelines (CPGs) shall be one of the strategies in the implementation of quality assurance standards. The CPG recommendations based on best available evidence shall be translated into policy statements and shall be used primarily to provide guidance to doctors, hospitals and patients as to what tests, medicines, and procedures are strongly recommended if benefits clearly outweigh the harms. It shall be used by the Corporation as one of its references in assessing the quality of care rendered by PhilHealth-accredited health care providers to members through performance monitoring and other activities when necessary. Moreover, this Circular shall focus on moderate-risk and high-risk CAP which are being reimbursed by the Corporation.

Community acquired pneumonia (CAP) is considered as one of the top illnesses in claims reimbursement. Moderate-risk and high-risk CAP require inpatient care because of the need for intravenous treatment and close observation due to risk of developing complications. The recommendations in this document incorporate updated information from the CAP Policy Statements published by the Corporation in The HTA Forum 2006. Furthermore, these evidence-based policy recommendations were approved by the PhilHealth Quality Assurance Chemittee (QAC) as reference for ensuring quality of care.

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A. DEFINITION

CAP is commonly defined as an acute infection of the pulmonary parenchyma with symptoms of acute illness accompanied by abnormal chest findings. Patients who acquire the infection in hospitals or long-term facilities are typically not part of the definition.

B. DIAGNOSIS

Clinical judgment is needed to make a diagnosis of community-acquired pneumonia (CAP). Patients usually presents with:

- 1. A history of cough within the past 24 hours or less than 2 weeks;
- 2. Abnormal vital signs of tachypnea (respiratory rate >20 breaths per minute), tachycardia (cardiac rate >100 per minute) and fever (temperature >37.8 C); and
- 3. With at least 1 abnormal chest finding of diminished breath sounds, rhonchi, crackles, or wheeze.

C. INITIAL CHEST RADIOGRAPHY

- 1. Chest x-ray (standing posteroanterior and lateral views) should be done for all patients suspected of pneumonia.
- 2. Chest CT scan should not be done routinely in the evaluation of pneumonia.

D. HOSPITAL ADMISSION

- 1. Only moderate-risk and high-risk CAP should be admitted [Grade A recommendation]. Refer to Annex A for the clinical features of patients with moderate risk and high risk CAP.
- 2. Chest x-ray may be repeated for hospitalized patients suspected of pneumonia but have initial "normal" chest radiographic findings.

E. MICROBIOLOGIC STUDIES

For moderate-risk and high-risk CAP, blood cultures AND gram stain and culture with antibiotic sensitivity tests of respiratory specimens should be done prior to starting any antibiotic treatment. [Grade A recommendation]

F. TREATMENT

- 1. Among patients with moderate-risk and high-risk CAP, initial empiric antibiotic therapy based on initial risk stratification is recommended (refer to Annex B). [Grade B]
- 2. Routine use of mucolytics is NOT recommended in treatment of troublesome cough associated with pneumonia.

G. MONITORING RESPONSE TO INITIAL THERAPY

- 1. Patients with CAP should be monitored within 72 hours after initial therapy for clinical response based on improvement of temperature, respiratory rate, blood pressure, sensorium, oxygen saturation, and inspired oxygen concentration.
- 2. If there is no improvement after 72 hours of treatment, patient should be reassessed for possible resistance to the antibiotics or "for presence of other pathogens such as M. tuberculosis, viruses, parasites or fungi." [Grade B recommendation]
- 3. Measurement of arterial oxygenation is important in the initial evaluation of patients with CAP. The use of pulse oximetry may complement rather than replaces clinical severity scoring tools.
- 4. A follow-up chest x-ray is recommended only for patients who are not clinically improving. [Grade B recommendation]
- 5. Follow-up cultures of blood and sputum are not indicated for patients who are responding to treatment. [Grade A recommendation]



H. STREAMLINING EMPIRIC ANTIBIOTIC THERAPY

Patients started on parenteral antibiotics can be switched to oral therapy (refer to Annex C) once the patient is clinically improving, is hemodynamically stable and has a functioning gastrointestinal tract. [Grade B recommendation]

I. HOSPITAL DISCHARGE

Patients diagnosed with moderate-risk and high-risk CAP can be discharged based on the following criteria:

- 1. Absence of unstable co-existing illness or other life-threatening complication;
- 2. Stable vital signs; and
- 3. Ability to maintain oral intake

[Grade A recommendation]

J. LENGTH OF STAY

- 1. Patients with moderate-risk and high-risk CAP should be confined for a minimum of 4 days to provide sufficient time for proper evaluation of patient's response to therapy. Moreover, IV antibiotics should be administered for at least 3 days.
- 2. Hospital stay can be extended for longer period in high-risk CAP patients due to clinical instability of the condition.

K. PREVENTION

The following are recommended for the prevention of CAP:

- 1. Pneumococcal and influenza vaccinations
- 2. Smoking cessation

[Grade A recommendation]

II. Monitoring and Evaluation

The health care provider shall be bound by the provisions of the performance commitment and subject to the rules on monitoring and evaluation of performance as provided for in PhilHealth Circular No. 54 s. 2012 re: Provider Engagement through Accreditation and Contracting for Health Services (PEACHeS) and PhilHealth Circular No. 031-2014 re: Health Cate Provider Performance Assessment System (HC-PAS).

III. Repealing Clause

All provisions of previous issuances, circulars and directives that are inconsistent with any of the -provisions of this Circular for this particular circumstance wherein the same is exclusively applicable, are hereby amended, modified and repealed accordingly.

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IV. Separability Clause

In the event that a part or provision of this Circular is declared unauthorized or rendered invalid by any Court of Law or competent authority, those provisions not affected by such declaration shall remain valid and effective.

V. Effectivity

This Circular shall take effect for all admissions starting September 15, 2015. It shall be published in any newspaper of general circulation and shall be deposited thereafter with the National Administrative Register at the University of the Philippines Law Center

President and CEO.

Date Signed: 8/25/

Title:

PhilHealth Circular on Policy Statements on the Diagnosis, Empiric Management, and Prevention of Community-Acquired Pneumonia (CAP) in Immunocompetent Adults as Reference by the Corporation in Ensuring Quality of Care

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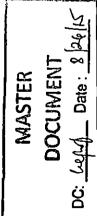
Annex A. Clinical Features of patients with CAP according to risk categories (adapted from CAP Guidelines 2010)

Moderate-risk CAP	High-risk CAP		
Any of the following:	Any of the criteria under moderate risk		
	CAP category		
Unstable vital signs	Plus		
• RR ≥30 breaths/min	Severe sepsis and septic shock		
• PR ≥125 beats/min	- ·		
• Temp ≥40 C or =36 C</td <td></td>			
• SBP <90 mmHg			
• DBP = 60 mmHg</td <td>Need for mechanical ventilation</td>	Need for mechanical ventilation		
_	}		
Altered mental state of acute onset			
Suspected aspiration			
Decompensated co-morbid condition			
Chest X-ray:			
- multilobar infiltrates			
- pleural effusion or abscess			
These patients need to be hospitalized for	These patients warrant admission in the		
closer monitoring and/or parenteral	intensive care unit. [Grade A]		
therapy. [Grade A]			



Annex B. Usual Recommended Dosages of Formulary Antibiotics in 50-60 kg adults with normal liver and renal functions

Risk Category	Empiric Therapy	Antibiotic/Dosage
Moderate Risk	IV non-antipseudomonal B-lactam	Amoxicillin + Clavulanic acid 1.2 gm q8h
CAP	•	Cefoxitin 1-2 gm q8h
		Cefuroxime Na 1.5 gm q8h
		Cefotaxime 1-2 gm q8h
		Ceftriaxone 1-2 gm q24h
]		Ertapenem 1 gm q24h
		22
	PLUS	Azithromycin dehydrate PO/IV 500 mg q24h
	Oral extended macrolide	Clarithromycin PO/IV 500 mg q12h
		Erythromycin PO/IV 0.5 – 1 gm q6h
}	OR .	I flow DO /TV 500 75024h
		Levofloxacin PO/IV 500-750 mg q24h
	Oral respiratory fluoroquinolones	Moxifloxacin PO/IV 400 mg q24h
High Risk CAP	No risk for Pseudomonas	
(all antibiotics	aeruginosa:	
are given		
intravenously)	IV non-antipseudomonal B-lactam	Amoxicillin + Clavulanic acid 1.2 gm q6-8h
		Cefotaxime 1-2 gm q8h
		Ceftriaxone 1-2 gm q24h
		Ertapenem 1 gm q24h
}	DITIC	A sick as a second dilumbates DO /IV/ 500 as a second
	PLUS	Azithromycin dihydrate PO/IV 500 mg q24
	IV extended macrolide	Clarithromycin PO/IV 500 mg q12h
		Erythromycin PO/IV 0.5 - 1 gm q6h
	OR.	Levofloxacin PO/IV 500-750 mg q24h
}	IV respiratory fluoroquinolone	5.
	With risk for Pseudomonas	
	aeruginosa:	
J	IV antipseudomonal B-lactam	Cefepime 2 gm q8-12h
	1 vanispoudomonar 2 monari	Gotopano 2 gm do 1211
	PLUS	Azithromycin dihydrate 500 mg q24h
	IV extended macrolide	Clarithromycin 500 mg q12h
1	PLUS	Gentamicin 3 mg/kg q24h
	Aminoglycoside	Netilmicin 7 mg/kg q24h
		Amikacin 15 mg/kg q24h
	OR	Ciprofloxacin 400 mg q12h
المر	IV anti-pseudomonal	Levofloxacin 750 mg q24h
1 (fluoroquinolones (high dose)	
11	Others	Oxacillin (Staphylococcus) 1-2 gm q4-6h
		Clindamycin (Staphylococcus and anaerobes)
		600 mg q6-8h
11		Metronidazole (anaerobes) 50 mg q6-8h
		Vancomycin (MRSA) 1 gm q12h



Annex C. Antibiotics dosage agents for streamlining or switch therapy*

Antibiotic	Dosage	Antibiotic	Dosage
Amoxicillin + clavulanic	625 mg TID or 1 gm BID	Cefuroxime axetil	500 mg BID
acid	500 mg OD	Cefdinir	300 mg BID
Azithromycin dihydrate	500 mg BID	Cefixime	200 mg BID
Clarithromycin		Levofloxacin	200 mg BID

^{*}for adults weighing 50 to 60 kg with normal liver and renal funsction

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