Management of suspected/confirmed Influenza (Adults)



In Canada seasonal influenza generally begins in October, peaks in December-January and ends in late April (Quebec surveillance data can be accessed at https://www.inspq.qc.ca/influenza).

Most cases of influenza result in mild illness, but viral pneumonias with/without bacterial superinfection as well as extrapulmonary complications (myositis, myocarditis, encephalitis) can occur in patients with high-risk conditions.

Based on observational studies, **early initiation** of antiviral therapy (ideally within 12 hours of onset of symptoms) may provide mortality benefit in hospitalized patients at high-risk of complications.

Suspected influenza: Fever and/or new onset/exacerbation of respiratory symptoms **Confirmed influenza**: Laboratory detection of Influenza A or B in respiratory sample

PATIENTS AT HIGH RISK OF COMPLICATIONS:

- ≥ 65 years
- Pregnant or postpartum (within 2 weeks of delivery)
- Underlying comorbidies:
 - Pulmonary disease (COPD, asthma, cystic fibrosis)
 - o Diabetes
 - Cardiovascular disease excluding isolated hypertension
 - o Chronic kidney disease
 - o Cirrhosis
- Immunocompromise: hematological malignancy, HSCT recipients, solid organ transplant, HIV with CD4 < 200, immunosuppressive medication (high dose corticosteroids, anti-TNF therapy, chemotherapy, etc.)
- Morbid obesity
- Nursing home residents

ADMISSION CRITERIA

- Respiratory criteria:
 - Dyspnea at rest or minimal activity
 - Respiratory rate > 22/min
 - PaO₂ < 65 mm Hg or O₂Sat
 < 90%
- Non-respiratory criteria
 - Altered mental status
 - Signs of sepsis/shock
 - Other considerations as per treating team

DIAGNOSTIC CONSIDERATIONS

- Nasopharyngeal swab or aspirate (NPS or NPA), or BAL for testing by RT-PCR
- If moderate-severe illness requiring admission:
 - Chest X-ray
 - Blood and sputum/BAL cultures before starting any antibiotics



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PHARMACOLOGIC MANAGEMENT

Mild disease	No risk factors for complicated disease:	
(no supplemental O ₂)	No antimicrobials	
	 <u>Pregnant:</u> <u>Oseltamivir¹ 75 mg PO BID x 5 days</u> (regardless of when symptoms started) <u>Any risk factor for complicated disease and symptom onset < 48 h:</u> <u>Oseltamivir¹ 75 mg PO BID x 5 days</u> 	
Moderate disease (Supplemental O ₂)	• Oseltamivir ¹ 75 mg PO BID x 5 days (regardless of when symptoms started)	
Severe disease	 Oseltamivir¹ 75 mg PO BID x 5 days AND 	
(extensive pneumonia,	Ceftriaxone 2 g IV q24h + Azithromycin 500 mg IV/PO q24h	
respiratory failure, septic shock)	Reassess in 48 h based on culture results; max. duration 5 days	
Clinical deterioration after initial improvement on antiviral	• Ceftriaxone 2 g IV q24h Reassess in 48 h based on culture results	
If MRSA colonized	 Add vancomycin² 15-25 mg/kg IV q8h-12h 	
If hospital-associated pneumonia	• Piperacillin-tazobactam 4.5 g IV q8h (extended-infusion over 3-4 hours) instead of ceftriaxone	
¹ Adjustment for renal dysfunction		
Creatinine clearance (use Cockroft	Gault Dose oseltamivir	

Creatinine clearance (use Cockroft-Gault equation)	Dose oseltamivir
31-60 mL/min	30 mg PO/PT BID
≤ 30 mL/min	30 mg PO/PT daily
Intermittent hemodialysis	75 mg PO/PT after each HD session

²See Vancomycin Therapeutic Drug Monitoring guideline; consult pharmacy for dosing adjustments

ADDITIONAL CONSIDERATIONS

- Offer annual vaccination to patients at high-risk of complications

REFERENCES

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Drafted by G. Moussa (Pharmacy Department), F. Bourdeau (Pharmacy Department) and Q. Li (Pharmacy Department) Reviewed by M. Semret (ID)

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