



# REGIONAL PEDIATRIC PARENTERAL DRUG MONOGRAPH

GENERIC NAME

# acetylcysteine



Effective Date: Dec 2011 CLASSIFICATION OTHER NAMES PAGE **Antidote for** Mucomyst

Revised Date: July 2024 acetaminophen 1 of 2

## **ADMINISTRATION POLICY:**

IV Infusion May be administered by a nurse IV Intermittent – May be administered by a nurse

IM Injection – DO NOT administer

#### RECONSTITUTION/DILUTION/ADMINISTRATION:

**Available as:** 200 mg/mL – 10 mL, 30 mL vial.

- Contact the Manitoba/Ontario Poison Control Centre at 1-855-776-4766.
- WRHA/HSC Sites use the acetylcysteine (3 bag protocol) monograph. Dosing and administration in this monograph will not be congruent with WRHA/HSC Toxicology dosing and administration guidelines.

## Patient is less than or equal to 20 kg:

Remove 37.5 mL from a 250 mL bag of D5W Add 37.5 mL (7 500 mg) acetylcysteine 200 mg/mL

Final volume: 250 mL Final concentration: 30 mg/mL (3%)

## Patient is 21 to 40 kg:

Remove 75 mL from a 500 mL bag of D5W. Add 75 mL (15 000 mg) acetylcysteine 200 mg/mL Final volume: 500 mL Final concentration: 30 mg/mL (3%)

## Patient is greater than 40 kg:

Remove 150 mL from a 1000 mL bag of D5W. Add 150 mL (30 000 mg) acetylcysteine 200 mg/mL Final volume: 1000 mL Final concentration: 30 mg/mL (3%)

## IV intermittent (Loading Dose): Pump Library: (acetylLp)

Acetylcysteine loading

~ .					
Concentration	Total Dose	VTBI	Administration Time		
(mg/mL)	(mg)	(mL)	(hours)		
	(max 24 000)				
30	Calculated dose	variable	4		
	(per dosage section)				
Clinical Advisory: High Alert					
Soft Low Dose Limit: 1920 mg		Soft High Dose Limit: 24 000 mg			
Care unit: Pediatric					

**LOADING DOSE:** 60 mg/kg/hr (maximum of 6000 mg/hr) of 3% *N-A*cetylcysteine X 4 hours

Calculate N-Acetylcysteine dose using total body weight up to a maximum of 100 kg.

Eg. Patient weight=120kg (above max weight of 100kg) 60mg x 100kg (use max weight, not 120kg) =

6000mg/hr x 4 hours = **24 000** mg total dose

Eg. Patient weight= 30 kg

60mg x 30kg = 1800mg/hr x 4 hours = **7 200 mg total dose** 

**Note**: Discard the rest of the bag once loading dose is complete. Use a new bag for the maintenance dose that follows.





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Effective Date: Dec 2011	CLASSIFICATION	OTHER NAMES	PAGE
D . 1D . 11 2024	Antidote for	Mucomyst	2 62
Revised Date: July 2024	acetaminophen		2 of 2

**IV infusion: Pump Library:** 

Acetylcysteine Maintenance

Drug Library	Dose Rate	Short Name	Care Unit
Yes	mg/kg/h	acetylMp	Pediatric
Drug	Diluent	Final Volume (VTBI)	Final Concentration
variable	variable	250, 500, or 1000 mL	30 mg/mL

Patient weight: enter patient weight to a MAX of 100kg in the pump (eg. if patient weighs 120kg enter 100kg)

Clinical Advisory: High Alert and maximum 600mg/hour

Soft Low Dose Limit: 6 mg/kg/h Soft High Dose Limit: 6 mg/kg/h

**DOSAGE:** 

MAINTENANCE DOSE: 6 mg/kg/hr (maximum of 600 mg/hr) of 3% N-Acetylcysteine continuously until

advised to STOP by the Poison Control Centre

Calculate N-Acetylcysteine dose using total body weight up to a maximum of 100 kg.

Eg. Patient weight=120kg (above max weight of 100kg)

6 mg x 100 kg (use max weight, not 120 kg) = 600 mg/hr

Eg. Patient weight= 30kg

6mg x 30kg= **180mg/hr** 

### STABILITY/COMPATIBILITY

**Stability of Final Admixture:** 24 hours at room temperature **Compatibility:** D5W preferred, normal saline

## PRECAUTIONS, POTENTIAL ADVERSE REACTIONS:

- Nausea, vomiting, hypotension may occur (asthmatics especially at risk)
- Hypersensitivity reactions skin rash, hives, flushing and urticarial
- Patients with asthma may develop bronchospasm

## ADDITIONAL NOTES AND NURSING CONSIDERATIONS:

- *Obesity:* In patients who weigh greater than 100 kg, the calculation of the IV acetylcysteine dose should be capped at 100 kg of body weight
- Do not order intravenous *N*-Acetylcysteine to run over a fixed duration (e.g. 21 hours) or a fixed dose (e.g. 100mg/kg over 16 hours), but instead order as an open-ended hourly infusion, with reassessment at least q12 hours based on serial laboratory testing as recommended by the Poison Centre.