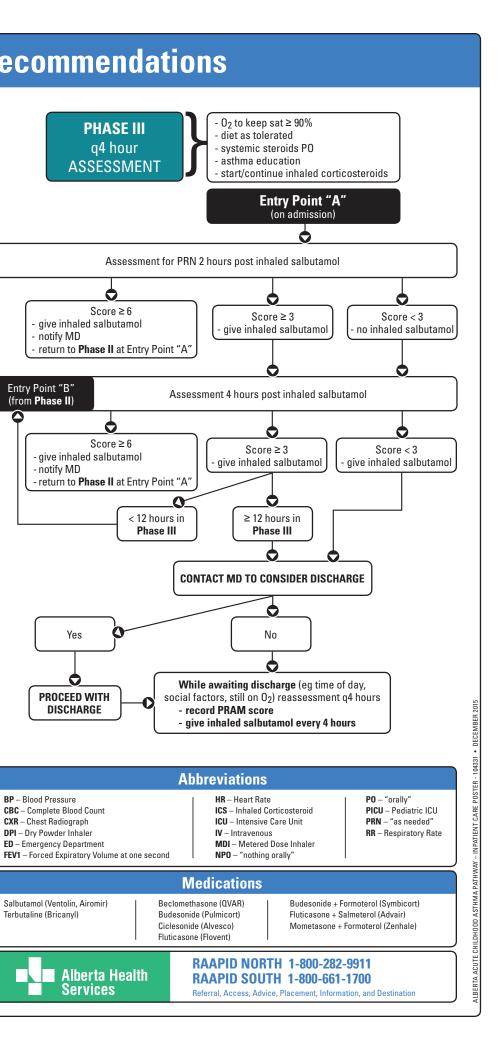


In regional centers, as patient transport to a tertiary center often requires intubation and intubation should be avoided when possible, patient care at the regional site directed by the pediatrician may be preferred. Consultation support without transport OR discussion of potential transport can be obtained via the tertiary care PICU (or via RAAPID).

* To view online pathway, continuing education module, and supporting evidence go to www.albertachildhoodpathways.com



Inpatient Care: Tertiary and Regional Centres

Summary of Orders and Inpatient Pathway

- Diet / Fluids / Electrolytes
 <u>Diet:</u> Phase I: consider clear fluids until in Phase II. NPO if not tolerating PO intake or if deteriorating (possible ICU). Resume PO intake as soon as possible. Phase II and III: Diet as tolerated. · Fluids: If vomiting, dehydration, poor intake or prolonged need for q1 hour aersolized salbutamol: IV fluid as needed.
- D5/0.45 with 20mEq KCI/L (30-40mEq KCI/L if K+ is low). Reduce and discontinue IV as soon as oral intake improves. • Electrolytes: If frequent inhaled salbutamol in ED or on ward, consider labs to check K+. If needing at least maintenance
- IV fluid, electrolyte check a24 hours recommended.

2. Oxygen

- Phase I and II: Suggest to keep sats ≥ 93%. Periodic saturation checks q2 hours and PRN before aersolized salbutamol. • Once in Phase III Suggest to keep sats ≥ 90% as long as there is no increased work of breathing.
- 3. Prednisone/Prednisolone 1-2 mg/kg (max 60mg) PO for 5 days total (Alternate Option (ie. for emesis): Dexamethasone 0.15-0.3mg/kg/dose (max 10mg)). Three to five days of dexamethasone suggested although literature is insufficient to support a particular length of treatment. Evidence regarding equivalency of prednisone and dexamethasone is weak. There are no published studies of dexamethasone use in inpatients.
- · Consider IV steroid if unable to tolerate PO, more severe cases or if started in ED due to severity: IV methylprednisolone 2mg/kg load and then start 1-2 mg/kg/day (max 80mg/day) divided q6 hours. Discontinue once oral tolerated or patient is improving and start oral steroids.
- · A longer course of therapy may be indicated for those on oral steroids recently prior to admission or if response to therapy has been slow.

4. Pathway Entry/Assessment of Clinical Status

- Modified PRAM score is used to determine if inhaled salbutamol treatment is needed.
- · Vital signs: RR. HR to be done with each assessment, BP routine.
- Phase II and III: FEV1 recommended for patients ≥ age 6 for those capable of spirometry, suggest once or twice per day and prior to discharge. May not be possible at all sites due to spirometry accessibility.
- · As per ED pathway, assessment for admission occurs at least 4 hours after administration of oral steroids; prior to this interval, ED pathway is most appropriate.
- · Admit to Phase I if patient on q1 hourly salbutamol prior to admission.
- · Admit to Phase II if patient on q2 hourly salbutamol prior to admission
- Phase III rarely indicated at admission (usually discharge from ED when on q4 hourly salbutamol).

PHASE I

- If patient is being admitted into Phase I Entry Point "A" first assessment is to be done 30 minutes after last inhaled salbutamol then every 30 minutes or 1 hour as per pathway (see algorithm).
- Repeat assessment after inhaled salbutamol (15-30 minutes post), noting response to treatment on assessment form. On post assessment, PRAM may be unchanged and score may be ≥ 3 but do not repeat salbutamol unless clinically indicated
- If on assessment 1 hour after inhaled salbutamol, score < 3, do not give inhaled salbutamol. Patient is ready to move to Phase II Entry Point "B" (needs MD order).
- · If on assessment patient is requiring inhaled salbutamol every 30 minutes on 3 subsequent assessments or if on assessment score is ≥ 6, MD involvement is needed to decide course of therapy which will vary depending on the clinical situation (see algorithm for considerations for MD assessment and when patient is deteriorating) After 6 hours in Phase I, MD reassessment re: stay in Phase I or move to Phase II.

PHASE II

- If patient is being admitted into Phase II Entry Point "A" first assessment is one hour after last inhaled salbutamol (PRN assessment) then 2 hours after last inhaled salbutamol then every 2 hours as long as inhaled salbutamol required (score > 3)
- If patient is being moved from Phase I to Phase II Entry Point "B" first assessment is done 2 hours after last inhaled salbutamol then every 2 hours as long inhaled salbutamol required (score \geq 3).
- · Repeat assessment after inhaled salbutamol (15-30 minutes post), noting response to treatment on assessment form. On post assessment, PRAM may be unchanged and score may be ≥ 3 but do not repeat salbutamol unless clinically indicated
- If on assessment 2 hours after inhaled salbutamol, score is < 3, do not give inhaled salbutamol. Patient is ready to move to Phase III Entry Point "B".
- Nurse or RT directed transfer can occur if score < 3, patient has been ≥ 4 hours in Phase II and there has been no increased O₂ needs or increased respiratory rate.
- If greater than 24 hours in Phase II or if score < 3 but criteria for nurse or RT directed transfer are not met, continue salbutamol at minimum of q2 hours and contact MD to consider transfer to Phase III.
- If score ≥ 6, notify MD and return to Phase I Entry Point "A".

PHASE III

- If patient is being admitted into Phase III Entry Point "A" first assessment is 2 hours after last inhaled salbutamol and then 4 hours after the last inhaled salbutamol then every 4 hours.

 If patient is being moved from Phase II to Phase III Entry Point "B" first assessment is 4 hours after last inhaled
- salbutamol then every 4 hours.
- · Repeat assessment after inhaled salbutamol (15-30 minutes post), noting response to treatment on assessment form. On post assessment, PRAM may be unchanged and score may be ≥ 3 but do not repeat salbutamol unless clinically indicated.
- If on assessment 4 hours after last inhaled salbutamol score is < 3, give inhaled salbutamol. Patient is ready for potential discharge (see below). If there is a delay in discharge - assess every 4 hours and inhaled salbutamol to be given every 4 hours as a minimum
- If score ≥ 6, notify MD and return to Phase II Entry Point "A"
- 5. Salbutamol Therapy by MDI/Spacer is strongly recommended
- Dose: 100mcg/puff weight < 20kg 5 puffs/dose; ≥ 20kg 10 puffs/dose. Once in Phase III reduce to 5 puffs/dose for all weights. If less effective, increase by 1-2 puff/dose; if increased side effects (HR, jittery), decrease by 1-2 puff/dose.
- Max MDI dose 10 puffs
- Alternate: Nebulization dose 2.5mg/dose for < 20kg and 5mg/dose for ≥ 20kg; nebulization should be considered when patient requires high flow oxygen by face mask, when patient is deteriorating in Phase I and/or becoming fatiqued or when PRAM score is ≥ 8
- Once in Phase III, can switch to home inhaled β2 Agonist and ICS device if not being discharged with MDI and Spacer. Note: Ventolin Diskus and Bricanyl Turbuhaler 1 puff = 2 puffs inhaled β_2 Agonist by MDI/Spacer.

* To view online pathway, continuing education module, and supporting evidence go to www.albertachildhoodpathways.com

Alberta Acute Childhood Asthma Pathway: Evidence based* recommendations

Inpatient Care: Tertiary and Regional Centres

- 6. Ipratropium
 Not recommended routinely for inpatient therapy BUT may be used in asthmatic patient who is severe or deteriorating after admission. Use only in first 24 hours of admission
- Dose: MDI (20mcg/puff) 4 puffs/dose OR nebulizer 250mcg/dose for all weights, x3 doses given along with each inhaled salbutamol treatment

7. Long Acting β_2 Agonists or Leukotriene Receptor Antagonists Continue usual maintenance therapy.

- 8. Inhaled Corticosteroid
- · Usual therapy should continue in hospital. If no maintenance therapy, begin as soon as possible
- Suggested dosing in hospital if not previously using daily inhaled cortcosteroid (ICS): Alvesco MDI (200 mcg) 1 puff OD-BID OR Flovent MDI (125mcg) 1-2 puffs BID OR Flovent Diskus (100mcg) 1-2 puffs BID OR Pulmicort Turbuhaler (200mcg) 1-2 puffs BID OR QVAR MDI (100mcg) 1-2 puffs BID.
- · In general ICS are of similar effectiveness. However, caution should be exercised when using all inhaled corticosteroids at higher doses because they pose a risk for significant adverse effects such as adrenal axis suppression or inhibition of growth (see online pathway for details*).
- Consideration for DPI is recommended for those age 6 and over; consideration should be given for child preference, parent preference, cost and drug coverage.

9. Magnesium Sulphate

- In Phase I consider Magnesium sulphate if deterioration unresponsive to treatment
- When considering use of Magnesium sulphate, in a tertiary center, PICU consult should be initiated or if in a regional center, consider seeking advice from PICU.
- Dosing: 40 mg/kg IV bolus over 20 minutes (max 2 grams)
- · Monitoring requirements: magnesium sulphate can cause hypotension, respiratory depression HR and BP should be closely followed: cardiorespiratory monitoring recommended

10. Investigations/Antibiotics

- CXR only if atypical presentation; deterioration after admission; suspected pneumonia.
- · Capillary (or arterial or venous) blood gas if deterioration; altered mental status; underlying chronic lung disease.
- CBC, cultures if high fever; toxic appearance; clinical deterioration. • Antibiotics – if definite pneumonia, sinusitis, otitis media.
- 11. Asthma Education

- Should be completed for all inpatients, best done in Phase II or Phase III. • Further outpatient asthma education is highly recommended.

12. At tertiary care sites Respirology Consultation should occur when: ICU admission

- Regularly followed by Respiratory Service or Asthma Clinic.
- Respirology Consultation can be considered when:
- Severe exacerbation.
- · Historical features suggestive of poor outpatient management.

13. Discharge Criteria in Phase III:

- Score < 3 on assessment 4 hours after last treatment or 12 hours in Phase III
- Room air, saturations > 90%.
- · Asthma education completed · Family able to continue treatment at home.
- · Follow-up arranged Asthma outpatient care is essential; if no family physician able to be obtained, consider outpatient nediatric consultation · Continue ICS at discharge until seen by the community care provider.
- Discharge action plan completed and communicated to family and community physician (\$
- Discharge instructions given to family (\$\$\chickleftarrow\$)
- Prescriptions given (\$\$)
- For above 🛠 discharge items: use triplicate "Pediatric Asthma Discharge Prescription and Short Term Plan" for all purposes (action plan, discharge instructions and prescription) OR use site mandated medication/reconcilliation process and use the duplicate Pediatric Asthma Short Term Plan for action plan and discharge instructions.

Device Recommendations

• 0-4 years: MDI/Spacer with mask • ≥ 4 years: MDI/Spacer with <u>mouthpiece</u>

Abbreviations						
BP – Blood Pressure CBC – Complete Blood Count CXR – Chest Radiograph DPI – Dry Powder Inhaler ED – Emergency Department FEVI – Forced Expiratory Volume at one second		HR – Heart Rate ICS – Inhaled Corticosteroid ICU – Intensive Care Unit IV – Intravenous MDI – Metered Dose Inhaler NPO – "nothing orally"		PO – "orally" PICU – Pediatric ICU PRN – "as needed" RR – Respiratory Rate		
Medications						
Salbutamol (Ventolin, Airomir) Terbutaline (Bricanyl)	Beclomethasone (QVAR) Budesonide (Pulmicort) Ciclesonide (Alvesco)		Budesonide + Formoterol (Symbicort) Fluticasone + Salmeterol (Advair) Mometasone + Formoterol (Zenhale)			



≥ 6 years: DPI preferred



not include 02 saturation.

that same patient.

4 hours prior to discharge.

Signs Suprasternal Indrawing Scalene Retractions Wheezing

Air Entry

Phase I or Phase II.

† Excludes O₂ saturation

Alberta Acute Childhood **Asthma Pathway: Evidence based* recommendations**

Inpatient Care: Tertiary and Regional Centres

Pathway Inclusions

Age 1-18 years with asthma; 1st time wheeze if diagnosis is likely asthma; **NOT** bronchiolitis; **NOT** pneumonia unless the pneumonia is felt to be a more minor issue compared to the asthma.

Pathway Entry on Admission

MD to determine Phase to enter on admission based on response to treatment prior to admission. · As per ED pathway, assessment for admission occurs at least 4 hours after administration of oral steroids; prior to this interval, ED pathway is most appropriate.

Admit to Phase I if patient on g1 hourly salbutamol prior to admission.

· Admit to Phase II if patient on q2 hourly salbutamol prior to admission.

• Phase III rarely indicated at admission (usually discharge from ED when on g4 hourly salbutamol.

Inpatient Assessment

In ED/urgent care, the PRAM score is used for assessment of severity of exacerbation at triage and following respiratory status.

The inpatient pathway uses a modified PRAM score (see below). The modified PRAM score does

When reviewing PRAM scores in ED prior to admission, most patients are on oxygen such that their PRAM score will be 1-2 points higher than the inpatient modified PRAM score would be for

In the inpatient pathway, the modified PRAM score is used to assess if salbutamol treatment is indicated and to extend the intervals of assessment. The patient moves from Phase I to Phase I to Phase III as their assessment intervals extend from q30-60 minutes to q2 hours and then every

Inpatient Assessment Score (Modified PRAM[†])

0	1	2	3
absent		present	
absent		present	
absent	expiratory only	inspiratory & expiratory	audible without stethoscope/silent chest
normal	decreased at bases	widespread decrease	absent/minimal

Phase Change Criteria: SCORE of < 3 at routine assessment or MD order on a reassessment in

For salbutamol assessment: if SCORE \geq 3, give salbutamol, if < 3 no salbutamol.

Repeat PRAM Score 15-30 minutes post any salbutamol treatment.

For any assessment SCORE ≥ 6, give salbutamol and notify MD. If in Phase II or Phase III move back to previous phase. If in Phase I consider further investigations, reassess therapy salbutamol frequency, IV, oxygen, etc.) and consider PICU consultation if not responding to treatment.

