



MEDICAL DIRECTIVE

Critical Care Outreach Team (CCOT) – Respiratory Distress

Approved by/Date: Medical Advisory Committee – October 27, 2015

Authorizing physician(s)

Intensivists who are part of the Critical Care Physician Section

Authorized to who

CCOT Responders (RRTs and RNs) that have the knowledge, skill and judgment and who have successfully attained certification by a course of self-study supplied by the Intensivist – Educators appointed by the Authorizing Physicians, participated in Didactic and Simulation education, completed orientation with Critical Care Outreach Team, and have successfully passed examinations.

Patient Description / Population

Adult Patients over 18. Patients who present with symptoms of shortness of breath and exhibit respiratory distress including but not restricted to symptoms of airway compromise, tachypnea, dyspnea, orthopnea, cyanosis, shortness of breath, accessory muscle use, stridor, crackles, wheezes and /or cyanosis.

Medical Directive Description/Physician's Order

1. Manage airway including support of oxygenation and ventilation
2. Oxygen therapy as required to maintain oxygen saturation above or equal to 92%; patients who have COPD 88-92%
3. Initiate monitoring including cardiac, blood pressure and pulse oximetry
4. Vital signs including temperature Q 5-30 minutes and prn
5. Review patient history and diagnosis
6. Stat POC or i-STAT ABG (pH, pO₂ and pCO₂)
7. STAT Portable Chest X-ray – upright if possible
8. Stat Blood Work (CBC, electrolytes, glucose, urea, creatinine, Magnesium, Phosphorus, Corrected calcium)
9. Insert a large (18 if possible) gauge IV
10. Intraosseous access may be attained when it is a very unstable, life threatening situation and when IV access has not been successful after 2 attempts or 90 seconds of searching for a suitable vein
11. **If on auscultation CRACKLES** - Position patient in Semi to high Fowlers

Originating Committee: Critical Care – June 18, 2015

Medical Advisory Committee: October 27, 2015

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12. **If on auscultation wheeze or silent chest** – Salbutamol 4-9 puffs q15 min up to 3 times and Ipratropium 4-9 puffs once
13. Continuous Positive Airway Pressure (CPAP)/Bi-level Positive Airway Pressure (Bipap) or CPAP mask initiation:
 - **NOTE: patients who require ongoing CPAP/Bipap must be transferred to a Critical Care Unit**
 - If severe respiratory distress (respiratory rate greater than 30) and/or oxygen saturation less than 90% on 100% oxygen by non-rebreather or high-flow/Optiflow
 - Contraindications: systolic pressure less than 90 mmHg systolic, suspected pneumothorax, decreased level of consciousness leading to compromise of airway patency, inability to sit upright, unable to cooperate with CPAP/BiPAP, abdominal surgery within 7 days, active nausea and vomiting.
 - Start CPAP at 5 cm H₂O and increased by 5 cm H₂O to maximum 15 cm H₂O as required to relieve symptoms/hypoxia and as tolerated by patient
 - May initiate Bipap if ongoing respiratory distress despite CPAP or evidence of hypercapnea (PaCO₂ greater than 50 and pH less than 7.3 OR EtCO₂ greater than 50 and no ABG available)
 - Initiate inspiratory pressure 5 cm H₂O above CPAP setting and titrate up as required to maximum 25 cm H₂O total inspiratory pressure.

Specific conditions/circumstances that must be met before the Directive can be implemented

- Patient must be assessed and placed into a severity category by the CCOT responder.
- The patient must have symptoms of crackles, wheezing, shortness of breath, cyanosis, dyspnea, tachypnea, orthopnea, and/or accessory muscle use.
- Each intervention will be explained to the patient and/or family when possible.
- Medications given by hospital personnel or taken by the patient just prior to the event must be included in the calculation of maximum doses.
- Personal protective equipment (mask, protective eyewear or face shield) must be used by staff when within two metres of any patient on CPAP/Bipap, with or without symptoms of an acute respiratory infection.

Contraindications to the implementation of the Directive

- Patient refuses therapy.
- Allergy to salbutamol, ipratropium or MDI propellant will prevent use of the specific agent.
- Patient is in pre-morbid state (silent chest, cyanosis, confusion) – code blue must be activated.



- Patient is incapable of cooperating with the procedures.
- Patients exhibiting signs and symptoms of acute respiratory infection must be managed with Contact/Droplet isolation in a private room to have CPAP/Bipap initiated on the floor.

Identify relevant Delegated Control Act or Added Skill associated with this Directive

Administering a substance by injection or inhalation.

Performing a procedure below the dermis:

- IV Insertion Certification
- IO Insertion Certification

Documentation requirements

- Implementation of the Medical Directive must be documented on the chart under physician orders.
- Vital signs pre and q15 to 30 minutes post medication.
- Response to bronchodilator medication must be documented in the CCOT note.

Review/Evaluation Process (how often/by who)

Every 2 years by Medical Department - Emergency Medicine and Critical Care Council.

Related Documents

Canadian Asthma Consensus Report – Canadian Medical Association Journal 1999; 161:S1- S12.

ORNGE Medical Directives and Standing Orders. Airway-Airway Protocol – Crash. April. 2005. Pg. 33-34.

Ontario Provincial Primary Care Paramedic Medical Directives - Bronchoconstriction Medical Directive. Waterloo Region, Ontario, Canada. Base Hospital Program Jan 1, 2007 pg. 8-9

Ontario Provincial Primary Care Paramedic Medical Directives – Sever Asthma Exacerbation Medical Directive. Waterloo Region, Ontario, Canada. Base Hospital Program Jan 1, 2007 pg. 10.

Ontario Provincial Primary Care Paramedic Medical Directives – Pulmonary Edema Medical Directive. Waterloo Region, Ontario, Canada. Base Hospital Program Jan 1, 2007 pg. 13.



Lakeridge Health Corporation. Emergency Medical Directives: Adult Asthma. Sept 2009.

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20. Faculty of Medicine, Liverpool Health Science, Liverpool, Australia. Medical Emergency Team, 2005, pg. 1-3.
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