

Inclusion Criteria

All patients undergoing endotracheal tube intubation (ETI) using rapid sequence intubation

Exclusion Criteria

Intubations not using standard RSI approach

Concern for respiratory failure or inability to protect airway

[Definition & Diagnosis](#)

Consider

[Signs and Symptoms of Hypoxia & Hypercapnia](#)

Consider

[Diagnostic Testing](#)

Off Pathway

No

Clinical indications for endotracheal intubation?

Yes

- NPO
- Cardiac monitor
- Continuous pulse oximetry
- Capnography

[Rapid Sequence Intubation Agents for Medical Patients](#)

[Rapid Sequence Intubation Agents for Trauma Patients](#)

[Post Intubation Medications](#)

Prepare for intubation and post intubation sedation and analgesia by using
Intubation Order Set

Optimize
hemodynamics prior to
intubation

Utilize

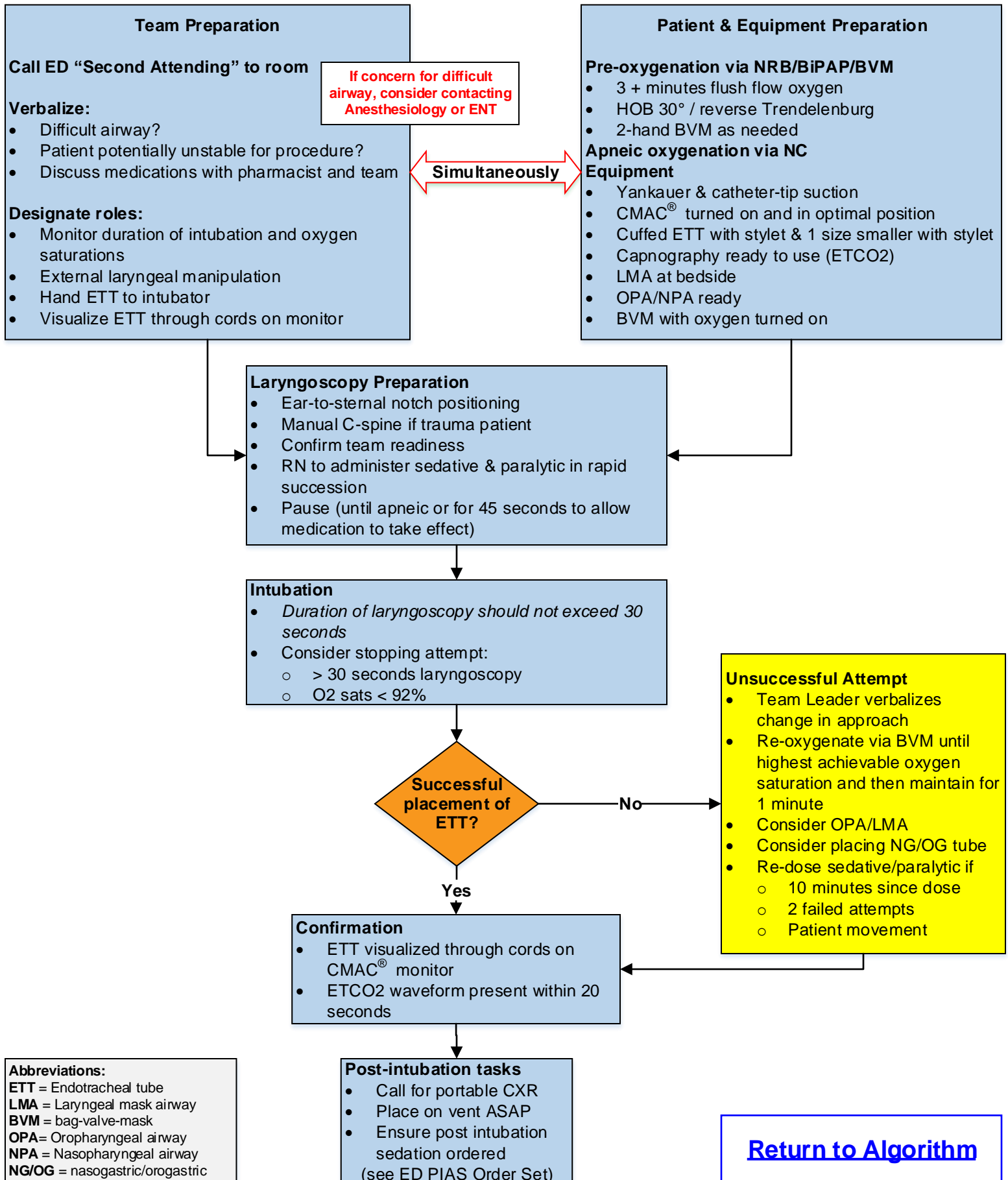
[Rapid Sequence Intubation Checklist](#)

After successful intubation

- Portable Chest Xray
- NG/OG placement (as needed for GI decompression)
- Mechanical Ventilation

Admit to ICU

ED Rapid Sequence Intubation Checklist



Signs & Symptoms of Hypoxia and Hypercapnia

Severity Level	Hypoxia	Hypercapnia ^a
Mild	<ul style="list-style-type: none"> • None or depressed efficiency 	<ul style="list-style-type: none"> • Flushed skin • Headaches
Moderate	<ul style="list-style-type: none"> • Dyspnea • Headaches, dizziness • Fatigue • Pallor • Tachycardia, cardiac arrhythmias • Hypertension • Mood swings: euphoria, disorientation, or depression • Ataxia, tingling 	<ul style="list-style-type: none"> • Tachypnea • Tachycardia • Dyspnea • Muscle twitches, depressed tendon reflexes • Drowsiness, confusion • Hypertension
Severe	<ul style="list-style-type: none"> • Cyanosis • Hypotension • Bradycardia • Visual impairment • Loss of consciousness, seizures, coma 	<ul style="list-style-type: none"> • Papilledema • Coma
^a In chronic hypercapnia, signs and symptoms of hypercapnia are observed when PCO ₂ increases above baseline level. Vo P, Kharasch K. <i>Respiratory Failure. Pediatrics in Review.</i> 2014;35(11):476-486.		

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Definition & Diagnosis

Respiratory failure occurs when the respiratory system fails in oxygenation or carbon dioxide elimination or both.

2 types of respiratory failure:

- Hypoxemic = $P_{aO_2} < 60$ mm hg with normal or low P_{aCO_2}
- Hypercapnic = $P_{aCO_2} > 50$ mm hg

Vo P, Kharasch K. Respiratory Failure. Pediatrics in Review. 2014;35(11):476-486.

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Indications for Intubation

- Failure to ventilate
- Failure to oxygenate
- Inability to protect airway
 - The absence of a gag reflex is not sensitive or specific as an indicator of a loss of airway reflexes.
 - Spontaneous or volitional swallowing better illustrates the patient's ability to protect the airway.
 - Pooled secretions in the posterior oropharynx suggest an inability to protect the airway.
- Anticipated clinical deterioration
- Airway protection needed for further evaluation or transport

Brown CA, Sackles JC, Mick NW. The Walls Manual of Emergency Airway Management. 5th Edition. Wolters Kluwer, 2018.

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Diagnostic Testing

Interpretation of Blood Gas Results^a

Condition	pH	Paco ₂	Base Excess
Acute respiratory acidosis or acute hypoventilation	↓	↑	↔
Acute respiratory alkalosis or acute hyperventilation	↑	↓	↔
Acute or chronic respiratory acidosis	↓	–/↑	↑
Acute metabolic acidosis with respiratory compensation	↓	↓	↓
Chronic respiratory acidosis with metabolic compensation	Normal/slightly ↓	↑	↑

^a*In chronic hypercapnia, signs and symptoms of hypercapnia are observed when Pco₂ increases above baseline level.*

↓=decrease; ↑=increase; ↔=no change

Vo P, Kharasch K. *Respiratory Failure. Pediatrics in Review.* 2014;35(11):476-486.

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Rapid Sequence Intubation Agents for Medical Patients

DRUG	DOSE	ROUTE	ONSET	DURATION	USE/BENEFIT	WARNING/COMMENTS
Premedication Agents						
Atropine	0.01-0.02 mg/kg No minimum dose; max 0.5 mg	IV, IM	1-5 min	> 30 min	Infants <1 year of age; bradycardia; copious secretions	
Lidocaine	1 mg/kg	IV	1-2 min	10-20 min		No longer recommended to prevent ICP increase due to conflicting/limited data
Analgesic Agents						
Fentanyl	1-2 mcg/kg (max 100 mcg)	IV, IM	Immediate	30-60 min	Desire blunting of the sympathetic response	Chest wall rigidity with rapid administration
Sedation Agents						
Etomidate	0.3 mg/kg (max 30 mg)	IV	30 sec	5 min	Hemodynamic instability	Suppresses cortisol – caution in sepsis; myoclonus – avoid in seizure disorders Hemodynamically neutral
Ketamine	1-2 mg/kg IV (max 100 mg) 4-10 mg/kg IM	IV, IM	IV: 30 sec IM: 3-4 min	5-10 min 12-25 min	Hemodynamic instability, sepsis, asthma	Bronchodilator; increases secretions, BP, HR
Midazolam	0.1-0.2 mg/kg (max 5-10 mg)	IV, IM	IV: 1.5-2 min IM: 15min	10-30 min		Monitor for hypotension
Propofol	1-2 mg/kg (max 100 mg)	IV	30-45 sec	3-10 min	Bronchospasm, head injuries, elevated ICP	Monitor for hypotension
Neuromuscular Blocking Agents						
Succinylcholine	IV: 1-2 mg/kg IM: 4 mg/kg (max 150 mg)	IV, IM	IV: 30-60 sec IM: ~3 min	IV: 4-10 min IM: 10-30 min		Increases ICP, hyperkalemia, fasciculation, ↑IOP, ↓HR Monitor for malignant hyperthermia
Rocuronium	1.2 mg/kg (max 100 mg)	IV	60 sec	25-40 min		
Vecuronium	0.1 mg/kg (max 10 mg)	IV	2-3 min	45-60 min		

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Rapid Sequence Intubation Agents for Trauma Patients

Patients < 1 year of age

- Premedication: Atropine (0.02 mg/kg, max 0.5 mg) to prevent reflex bradycardia
- Sedation : Ketamine (2 mg/kg*, max 100 mg)
- Paralytic: Rocuronium (1.2 mg/kg, max 100 mg)

Patients greater than 1 year of age

- Sedation: Ketamine (2 mg/kg*, max 100mg)
- Paralytic: Rocuronium (1.2 mg/kg, max 100mg)

*Ketamine can worsen hypotension and exacerbate myocardial depression in patients who are catecholamine depleted. This would include patients who have had prolonged hypotension; a maximum dose of 1.5 mg/kg, up to 100mg, is recommended in these patients

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Post Intubation Medications

Recommendations:

- Use PRN IV medications prior to continuous infusion arriving
- Initiate **ONE** continuous infusion
 - Fentanyl infusion is preferred at MCED
 - If at LCED, Dexmedetomidine is preferred
- Fentanyl and Versed:
 - Start the infusion by giving a bolus off the drip, then start the infusion rate
 - Bolus dosing is the same as the continuous infusion dosing
- Dexmedetomidine
 - Bolus 0.2 mg/kg over 2-3 minutes
 - Only bolus once

If unable to maintain appropriate sedation, recommend using PRN IV medications before ordering a second continuous infusion medication

Patients ≤ 50 kg				
Medication	Starting Dose (Initial Dosing Range)	Titrate by Q30 minutes PRN	Max Dosing	Side Effects & Other Information
Fentanyl	1 mcg/kg/hr (0.5-1 mcg/kg/hr)	0.5 mcg/kg/hr	3-4 mcg/kg/hr	Rapid, onset. lipophilic with adipose deposition, withdrawal symptoms after prolonged infusion
Dexmedetomidine (Precedex)	0.4 mcg/kg/hr (0.3-0.7 mcg/kg/hr)	0.2 mcg/kg/hr	1.2 mcg/kg/hr	Bradycardia, hypotension, loss of airway reflexes
Midazolam (Versed)	0.1 mg/kg/hr (0.05-0.1 mg/kg/hr)	0.05 mg/kg/hr	0.15 mg/kg/hr	Respiratory depression, hypotension

**For patients > 50 kg, dosing depends on Ideal Body Weight and/or Adjusted Body Weight.
Please consult pharmacy for dosing**

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Quality Metrics

Goal:

- To improve, standardize, and optimize post-intubation care for all patients requiring intubation in the Pediatric Emergency Department.

Process Measures:

- Pathway Visualization
- Post Intubation Medication Order Set utilization

Outcome measures:

- Patients receiving post intubation sedation within 10 minutes of intubation

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Pathway Team & Process

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Clinical Pathway Development

This clinical pathway was developed using the process described in the NCH Clinical Pathway Development Manual Version 6, 2022. Clinical Pathways at Nationwide Children's Hospital (NCH) are standards which provide general guidance to clinicians. Patient choice, clinician judgment, and other relevant factors in diagnosing and treating patients remain central to the selection of diagnostic tests and therapy. The ordering provider assumes all risks associated with care decisions. NCH assumes no responsibility for any adverse consequences, errors, or omissions that may arise from the use or reliance on these guidelines. NCH's clinical pathways are reviewed periodically for consistency with new evidence; however, new developments may not be represented, and NCH makes no guarantees, representations, or warranties with respect to the information provided in this clinical pathway.

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