

GOVERNMENT OF WEST BENGAL  
HEALTH & FAMILY WELFARE DEPARTMENT  
SWASTHYA BHABAN, BLOCK GN-29, SECTOR V  
SALT LAKE CITY, KOLKATA -91

Memo No. M/6AI

Date: 17.04.2020

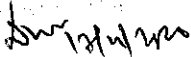
To  
The Principal (All Medical Colleges)  
The Medical Superintendents cum Vice Principal (All Medical Colleges)  
The Chief Medical Officer of Health (All Districts and Health Districts)  
The Superintendents (All DH/ SSH/ SDH/ SGH)  
The Block Medical Officer of Health (All blocks)


**State Protocol for Clinical Management of COVID-19 Cases, West Bengal**

In partial modification of the Memo No. HPH/9M-21/2020/77 dated 31.03.2020, the following final guideline in order to streamline the management protocol of COVID-19 affected patient across the State, prepared by the Expert Committee is hereby being circulated.

All concerned are hereby instructed to adhere to the guidelines outlined in the enclosed protocol.

All concerned are further instructed to share the guidelines to all faculties, specialists, Medical Officers under their control.

  
Director of Medical Education  
Government of West Bengal

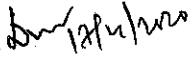
  
Director of Health Services  
Government of West Bengal


Memo No. M/6AI/1CA

Dated: 17.04.2020

Copy forwarded for information and necessary action please to:

1. DDHS (PH) and SSO, IDSP, West Bengal.
2. SNO, IDSP, West Bengal.
3. Dy. CMOH-II, all Districts and Health Districts.
4. Guard File

  
Director of Medical Education  
Government of West Bengal

  
Director of Health Services  
Government of West Bengal

**SALIENT POINTS IN MANAGEMENT OF SEVERE COVID-19 IN ADULTS**


<p><b>Oxygen therapy</b></p>	<ul style="list-style-type: none"> <li>• Administer oxygen to all Severe Acute Respiratory Illness (SARI) patients with respiratory distress / hypoxemia / shock</li> <li>• Start with nasal prongs @ 5L/min or simple face mask / venturi mask / nonrebreathing mask @ 10-15L/min, as needed</li> <li>• Titrate for target SpO<sub>2</sub></li> <li>• Initial target SpO<sub>2</sub> ≥ 94%</li> <li>• Target SpO<sub>2</sub> after initial stabilization: 90-96%</li> </ul>
<p><b>Initial fluid management</b></p>	<ul style="list-style-type: none"> <li>• Conservative fluid strategy if no evidence of shock</li> <li>• Cautious IV fluids</li> <li>• Monitor for worsening of oxygenation during fluid therapy</li> </ul>
<p><b>Empiric antimicrobials</b></p>	<ul style="list-style-type: none"> <li>• To all patients as early as possible, preferably within the first hour</li> <li>• Choose drugs to cover all suspected bacteria and influenza</li> <li>• Try to send blood cultures before starting antimicrobials; do not delay antimicrobials waiting to send cultures</li> <li>• De-escalate based on culture results or clinical judgment</li> </ul>
<p><b>Monitor closely for signs of clinical deterioration (rapidly progressive respiratory failure)</b></p>	<ul style="list-style-type: none"> <li>• Monitor vital signs and gas exchange at regular intervals</li> <li>• Check whether tolerating oxygen therapy</li> <li>• Do not delay intubation if worsening</li> <li>• If High Flow Nasal Cannula (HFNC) is available, consider a short HFNC trial in selected patients under close monitoring → Do not delay intubation if worsening</li> <li>• If HFNC not available, consider a short Noninvasive Positive Pressure Ventilation (NIPPV) trial in selected patients under close monitoring → Do not delay intubation if worsening</li> </ul>


	<ul style="list-style-type: none"> <li>• Airborne precautions must during HFNC /NIPPV / Endotracheal intubation</li> <li>• MDI with spacer preferred to nebulizers, if possible</li> </ul> <p>NB: Aerosol generating procedures: Intubation, Extubation, HFNC, NIPPV, Bag masking, Open suctioning, T piece / any open circuit, Bronchoscopy, CPR, Nebulisation, Tracheostomy</p>
<b>Monitor closely for signs of clinical deterioration (shock or sepsis)</b>	<ul style="list-style-type: none"> <li>• Monitor vital signs at regular intervals</li> <li>• Appropriate investigations at regular intervals</li> </ul>
<b>Investigations</b>	<ul style="list-style-type: none"> <li>• <b>Baseline investigations:</b> Complete haemogram with neutrophil-lymphocytic ratio, LFT, Urea, Creatinine, CRP, ECG, CXR, ABG with lactate (if SpO<sub>2</sub>&lt;94%), Ferritin(if available),Urine RE</li> <li>• <b>Follow up investigations:</b> Investigations to monitor organ functions regularly; Other investigations as decided by treating team; Early detection of myocardial involvement by troponins, NT-pro BNP and echocardiography; USG Abdomen</li> </ul>
<b>Address comorbidities</b>	<ul style="list-style-type: none"> <li>• Tailor management according to comorbidities</li> </ul>
<b>Criteria of CRITICAL CARE UNIT (CCU) admission</b>	<ul style="list-style-type: none"> <li>• Requiring mechanical ventilation</li> <li>• Hypotension requiring vasopressor support</li> <li>• Worsening mental status</li> <li>• Multi-organ dysfunction syndrome</li> </ul>
<b>When to intubate</b>	<ul style="list-style-type: none"> <li>• Features of respiratory fatigue with increased work of breathing and worsening respiratory parameters indicating respiratory failure</li> <li>• Haemodynamic instability</li> <li>• Altered sensorium with a threatened airway</li> </ul> <p>NB: Although intubation decision should be individualized, <b>keep a low threshold for intubation</b></p>
<b>How to intubate</b>	<ul style="list-style-type: none"> <li>• Full complement of PPE with face shield</li> <li>• Ensure scene safety &amp; check readiness of all</li> </ul>

	<p>essential drugs &amp; equipments prior to intubation</p> <ul style="list-style-type: none"> <li>• Most experienced team member to intubate</li> <li>• Complete airway assessment prior to intubation</li> <li>• Hemodynamic evaluation &amp; optimization, if needed, prior to intubation</li> <li>• Use Heat Moisture Exchanger (HME) with bacterial-viral filter in every oxygenation interface (face mask, circuit, endotracheal tube (ETT), catheter mount, LMA)</li> <li>• Use closed system suctioning</li> <li>• Preoxygenation with 100% oxygen</li> <li>• Rapid sequence intubation using induction agent (propofol or etomidate) and muscle relaxant (succinylcholine or rocuronium)</li> <li>• Limit bag mask ventilation unless unavoidable</li> <li>• Apply cricoid pressure only in case of ongoing regurgitation</li> <li>• Use videolaryngoscope with separate screen, if available</li> <li>• In anticipated difficult airway, anaesthesiologist may be called to intubate</li> <li>• In unanticipated difficult airway, use LMA and simultaneously call for expert help</li> <li>• Clamp ETT during unavoidable disconnections &amp; connections</li> <li>• Use end-tidal CO<sub>2</sub> and CXR to confirm correct position of ETT</li> <li>• After intubation, appropriate cleaning/disinfection of equipment and environment is mandatory</li> </ul>
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**COVID-19 & ARDS: Invasive mechanical ventilation**

<b>Initial mode</b>	<p>Volume Control (VC)</p> <p>NB: Can use Pressure Control (PC) if tidal volume goals are met. But mostly PC is tried when initial VC mode fails</p>
<b>Initial Settings</b>	<ul style="list-style-type: none"> <li>• Tidal volume (VT): 6ml/kg predicted body weight (PBW)</li> <li>• Rate: to match baseline minute ventilation (not &gt; 35)</li> </ul> <p>NB: PBW= Males: 50 + 2.3 (Height in inches – 60)  Females: 45.5 + 2.3 (Height in inches – 60)</p>

	<b>Tidal volume adjustment:</b>									
	<ul style="list-style-type: none"> <li>• Check Plateau Pressure (Pplat)</li> <li>• Plateau Pressure goal <math>\leq 30</math> cm H<sub>2</sub>O</li> <li>• If Pplat&gt;30: decrease VT by 1ml/kg steps to minimum 4ml/kg</li> <li>• If breath stacking(auto PEEP) or severe dyspnea occurs, may increase VT to 7-8 ml/kg if Pplat remains<math>\leq 30</math></li> </ul>									
	<b>Set PEEP according to the PEEP-FiO<sub>2</sub> tables to achieve oxygenation goal (PaO<sub>2</sub> 55-80 mm Hg / preferably SpO<sub>2</sub> 90-96%):</b>									
	<i>Lower PEEP-Higher FiO<sub>2</sub> combinations: (start with minimum value for a given FiO<sub>2</sub>)</i>									
	FiO <sub>2</sub>	0.3	0.4	0.5	0.6	0.7	0.8	0.9	1.0	
	PEEP	5	5-8	8-10	10	10-14	14	14-18	18-24	
	<i>Higher PEEP- Lower FiO<sub>2</sub> combinations:</i>									
	FiO <sub>2</sub>	0.3	0.3	0.3	0.3	0.3	0.4	0.4	0.5	0.5
	PEEP	5	8	10	12	14	14	16	16	18
	FiO <sub>2</sub>	0.5	0.6	0.7	0.8	0.8	0.9	1.0	1.0	
	PEEP	20	20	20	20	22	22	22	24	
	<ul style="list-style-type: none"> <li>• Higher PEEP (&gt;10) in moderate to severe ARDS</li> <li>• Lower PEEP (<math>\leq 10</math>) in mild ARDS and "non-ARDS like" severe pneumonia</li> <li>• Continue with higher PEEP if PEEP responsive (recruiters) and lower PEEP if PEEP non-responsive (non-recruiters)</li> <li>• PEEP responsive (Recruiters): Keeping FiO<sub>2</sub> unchanged, <i>usually</i> oxygenation improves with increase in PEEP with minimal / no drop in mean arterial pressure, minimal / no rise in PaCO<sub>2</sub> and minimal / no rise in driving pressure</li> <li>• Try to keep Pplat<math>\leq 30</math> and driving pressure (Pplat-PEEP) &lt;15</li> </ul>									
<b>Fluid management</b>	<ul style="list-style-type: none"> <li>• Conservative strategy in absence of tissue hypoperfusion</li> <li>• Try to avoid hypervolemia</li> </ul>									
										
<b>Oxygenation improving</b>	<ul style="list-style-type: none"> <li>• Reduce PEEP &amp; FiO<sub>2</sub> gradually</li> <li>• Shift to a partial assist / spontaneous mode, if tolerated</li> <li>• Plan for protocolised liberation from ventilation</li> </ul>									
<b>Oxygenation not improving</b>	Search for and address reasons of failure: <ul style="list-style-type: none"> <li>• Ensure conservative fluid management</li> </ul>									

	<ul style="list-style-type: none"> <li>• Treat patient-ventilator dyssynchrony, if present</li> <li>• Shift mode (volume limited to pressure limited)</li> <li>• Search for complications of disease or ventilation</li> </ul>								
									
<b>Oxygenation improving</b>	Optimize and persist with above-mentioned approaches till patient is ready for liberation from ventilation								
<b>Oxygenation not improving</b>	<b>If acceptable gas exchange not achievable without incurring Pplat &gt; 30, consider rescue therapies:</b>								
	<table border="1"> <tr> <td><b>Prone ventilation</b></td> <td> <ul style="list-style-type: none"> <li>• Most preferred rescue therapy</li> <li>• Consider in <math>PaO_2/FiO_2 &lt; 150</math> with a <math>FiO_2 \geq 0.6</math> and <math>PEEP \geq 5</math> or <math>PaO_2 / FiO_2 \leq 100</math> with a <math>PaO_2 \leq 60</math> despite optimization of ventilator settings at <math>FiO_2</math> of 1</li> <li>• Consider early proning (within first 36 hours)</li> <li>• 12-16 hours / day</li> <li>• Always check for contraindications and complications</li> </ul> </td> </tr> <tr> <td><b>Recruitment maneuvers</b></td> <td> <ul style="list-style-type: none"> <li>• Consider in PEEP responsive patients</li> <li>• Preferred method: Sustained high pressure inflation (35-40 cm H<sub>2</sub>O of CPAP for 40 seconds)</li> <li>• Avoid staircase maneuvers (Incremental PEEP)</li> <li>• Avoid routine use of recruitment maneuvers</li> </ul> </td> </tr> <tr> <td><b>Neuromuscular blockers</b></td> <td> <ul style="list-style-type: none"> <li>• Consider continuous infusion for up to 48 hrs in case of persistently high plateau pressures or severe dyssynchrony</li> <li>• Can use intermittent boluses to facilitate lung protective ventilation, if needed</li> </ul> </td> </tr> <tr> <td><b>Pulmonary vasodilators</b></td> <td> <ul style="list-style-type: none"> <li>• If available, a trial of inhaled prostacyclin or nitric oxide may be considered, if other rescue strategies have failed</li> </ul> </td> </tr> </table>	<b>Prone ventilation</b>	<ul style="list-style-type: none"> <li>• Most preferred rescue therapy</li> <li>• Consider in <math>PaO_2/FiO_2 &lt; 150</math> with a <math>FiO_2 \geq 0.6</math> and <math>PEEP \geq 5</math> or <math>PaO_2 / FiO_2 \leq 100</math> with a <math>PaO_2 \leq 60</math> despite optimization of ventilator settings at <math>FiO_2</math> of 1</li> <li>• Consider early proning (within first 36 hours)</li> <li>• 12-16 hours / day</li> <li>• Always check for contraindications and complications</li> </ul>	<b>Recruitment maneuvers</b>	<ul style="list-style-type: none"> <li>• Consider in PEEP responsive patients</li> <li>• Preferred method: Sustained high pressure inflation (35-40 cm H<sub>2</sub>O of CPAP for 40 seconds)</li> <li>• Avoid staircase maneuvers (Incremental PEEP)</li> <li>• Avoid routine use of recruitment maneuvers</li> </ul>	<b>Neuromuscular blockers</b>	<ul style="list-style-type: none"> <li>• Consider continuous infusion for up to 48 hrs in case of persistently high plateau pressures or severe dyssynchrony</li> <li>• Can use intermittent boluses to facilitate lung protective ventilation, if needed</li> </ul>	<b>Pulmonary vasodilators</b>	<ul style="list-style-type: none"> <li>• If available, a trial of inhaled prostacyclin or nitric oxide may be considered, if other rescue strategies have failed</li> </ul>
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<b>Ventilator precautions</b>	<ul style="list-style-type: none"> <li>• Fresh ventilator circuit for every new patient</li> <li>• HME with bacterial-viral filters must be fitted</li> <li>• Tubings and HME with bacterial-viral filters to be changed every 48 hours / when visibly soiled</li> <li>• Use closed system suctioning</li> <li>• Avoid routine suctioning</li> <li>• Avoid unnecessary disconnections</li> <li>• Clamp ETT for unavoidable disconnections &amp; connections</li> </ul>
<b>Sample collection in severe disease</b>	<ul style="list-style-type: none"> <li>• If upper respiratory tract samples (nasopharyngeal &amp; oropharyngeal) test negative, but still the clinical suspicion remains high specially in presence of pneumonia or severe disease, send lower respiratory tract samples also (expectorated sputum or tracheal aspirates / mini BAL / BAL in intubated patients)</li> <li>• Can send only lower respiratory tract samples in intubated patients</li> </ul>
<b>COVID-19 &amp; Shock: Haemodynamic support</b>	
<b>Fluid therapy</b>	<b>Strategy of acute resuscitation:</b> <ul style="list-style-type: none"> <li>• Individualize, monitoring tissue perfusion</li> <li>• Conservative strategy preferred to liberal</li> <li>• Try to avoid hypervolemia</li> <li>• Follow lactate</li> </ul>
	<b>Choice of fluids:</b> <ul style="list-style-type: none"> <li>• Buffered / balanced crystalloids</li> <li>• Avoid HES / Dextran / Gelatin / routine use of albumin</li> </ul>
	<b>Assess fluid responsiveness, whenever possible:</b> <ul style="list-style-type: none"> <li>• Use dynamic parameters for assessing preload responsiveness (e.g. Passive Leg Raising), as feasible</li> </ul>
<b>Vasoactive agents</b>	<ul style="list-style-type: none"> <li>• First line vasopressor: Noradrenaline</li> </ul>

	<p>(Vasopressin / Adrenaline if Noradrenaline not available)</p> <ul style="list-style-type: none"> <li>• Second line vasopressor: Add Vasopressin</li> <li>• MAP target: 60-65 mm Hg</li> <li>• Presence of cardiac dysfunction &amp; persistent hypoperfusion despite fluids &amp; Nordadrenaline: Add Dobutamine</li> <li>• Avoid Dopamine</li> <li>• Refractory shock despite fluids &amp; vasopressors: Add IV Hydrocortisone (200mg / day as continuous infusion / intermittent doses)</li> </ul>
<b>Renal replacement therapy</b>	
<b>When to dialyze in Acute Kidney Injury (AKI)</b>	<ul style="list-style-type: none"> <li>• Volume overload</li> <li>• Severe metabolic acidosis</li> <li>• Refractory hyperkalemia</li> <li>• Uremic encephalopathy</li> <li>• Uremic pericarditis</li> </ul>
<b>Strategy</b>	<ul style="list-style-type: none"> <li>• All modalities of renal replacement therapy can be used depending on clinical status</li> <li>• Preferably bedside dialysis. Portable reverse osmosis water in a tank may be used, if needed.</li> <li>• Acute peritoneal dialysis can be tried in selected patients where hemodialysis facility is not available</li> <li>• Cytokine removal therapies not recommended</li> </ul>
<b>Other critical care management</b>	
<ul style="list-style-type: none"> <li>• Protocolised light sedation</li> <li>• Enteral nutrition</li> <li>• Glycemic control</li> <li>• Prevention of hospital acquired infections</li> <li>• Deep vein thrombosis prophylaxis</li> <li>• Stress ulcer prophylaxis</li> <li>• Pressure ulcer prevention by two hourly turning</li> <li>• Protocolised liberation from ventilation</li> <li>• Caution about premature extubation (especially without facilitative HFNC / NIPPV) and subsequent reintubation</li> <li>• Early physical therapy</li> <li>• Use point-of-care ultrasound as much as possible to avoid transfers out of CCU for</li> </ul>	



investigations (e.g. CT scans)

- De-isolate after clinical recovery and two RT-PCR negative samples taken 24 hours apart