The novel coronavirus disease 2019 (COVID-19) pandemic caused by severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) is causing significant morbidity and mortality worldwide. The common presentations in children include involvement of respiratory system leading to pneumonia and acute respiratory distress syndrome, as well as multiorgan dysfunction syndrome and multisytem inflammatory syndrome in children (MIS-C). Pediatric COVID-19 is a milder disease as compared with the adults. Also, there is rise in MIS-C cases which is a hyperinflammatory condition temporally associated with SARS-CoV-2. Since respiratory system is predominantly involved, few of these critically ill children often require respiratory support which can range from simple oxygen delivery devices, high-flow nasal cannula (HFNC), noninvasive ventilation (NIV), invasive mechanical ventilation, and extracorporeal membrane oxygenation (ECMO). Most of the oxygen delivery devices and respiratory interventions generate aerosols and pose risk of transmission of virus to health care providers (HCPs). The use of HFNC and NIV should be limited to children with mild respiratory distress preferably in negative pressure rooms and with adequate personnel protective equipments (PPEs). However, there should be low thresholds for intubation and invasive mechanical ventilation in the event of clinical deterioration while on any respiratory support. The principle of providing respiratory support requires special droplet and air-borne precautions to limit exposure or transmission of virus to HCPs and at the same time ensuring safety of the patient.
COVID-19.1 Since lungs are the predominant organs involved in COVID-19, and respiratory support is the most common intervention, it should be provided taking all precautions to limit transmission of virus to health care providers (HCPs) and ensuring safety of the patients.

Pediatric COVID-19

Pediatric COVID-19 is relatively mild disease when compared with adults and accounts for 1 to 5% of the case load.4 Data suggest that Pediatric COVID-19 cases might be less severe than adults and that children might experience different symptoms than adults.4–8 According to previous studies, the involvement of children is as follows: asymptomatic (4–21%), mild (51–58%), moderate (19–39%), severe (1–5.2%), and critical disease (0.6–1.7%). The children with oxygen saturation <92% were in range of 1 to 2.3% and very few required assisted-mechanical ventilation. The mortality rate in children ranged from 0 to 0.6%.5,6,8,9

In a retrospective Chinese study on the epidemiological characteristics of 2143 Pediatric patients with COVID-19, it was demonstrated that 34.1% (n = 731) cases were laboratory-confirmed and young children, particularly infants, were vulnerable to SARS CoV-2 infection. The proportion of severe and critical cases was 10.6, 7.3, 4.2, 4.1, and 3.0% for the age group of <1, 1 to 5, 6 to 10, 11 to 15, and 16 years, respectively. Only one 14-year-old boy died.6 Among 149,082 cases of COVID-19 reported from the United States, 2,572 (1.7%) were aged <18 years.10 Hospitalization status was available for 745 (29%) cases in children aged <18 years and 35,061 (31%) in adults aged 18 to 64 years. Among children with COVID-19, 147 (5.7–20%) were reported to be hospitalized and 15 (0.58–2%) were admitted to an ICU. Children aged <1 year accounted for the highest percentage (15–62%) of hospitalization among pediatric patients with COVID-19. There were three deaths in this report.10 Similarly, in Madrid, Spain, during the first 2 weeks of the epidemic, 60% (25/41) children with confirmed COVID-19 were hospitalized, 9.7% (4/41) were admitted to ICU, and 9.7% (4/41) needed respiratory support beyond nasal prongs. Only one kid required mechanical ventilation and no death was reported.11

This fact is further reiterated in one of the recently published systematic review of Pediatric cases of COVID-19, which demonstrated that most children and adolescents presented with mild symptoms.12 Out of 1,065 children with SARS-CoV-2 infection, respiratory symptoms were mild except in one 13-month-old infant who presented with severe lower respiratory tract infection (COVID-19 pneumonia), complicated by shock and kidney failure, required ICU and invasive ventilation.13 No Pediatric death from COVID-19 was reported in the age range of 0 to 9 years. As compared with adults, children with COVID-19 showed a better prognosis and recovered within 1 to 2 weeks.12 Children with chronic diseases, like bronchopulmonary dysplasia and cystic fibrosis, were also less infected than adults.14

In late April 2020, clinicians from the United Kingdom (UK) reported a cluster of eight previously healthy children who presented with hyperinflammatory shock syndrome temporarily associated with COVID-19.15 Thereafter, multiple reports of pediatric inflammatory multisystem syndrome temporally associated with SARS-CoV-2 (PIMS-TS), multisystem inflammatory syndrome in children (MIS-C), Kawasaki’s disease (KD), and Kawasaki’s-like syndrome were published from the countries with high case load of COVID-19 like the UK, France, Italy, and the United States.16–19

The peak in the cases with PIMS-TS or MIS-C followed 2 to 6 weeks after the peak of COVID-19 pandemic.16–19 The age group involved was of older children and the most common presenting features were fever, gastrointestinal symptoms, rash, conjunctival injection, and respiratory symptoms. Majority had positive SARS-CoV-2 antibody test and one-third had positive reverse-transcription polymerase chain reaction (RT-PCR). The commonest laboratory abnormalities were elevated C-reactive protein (CRP), D-dimer, procalcitonin, brain natriuretic peptide (BNP), fibrinogen, ferritin, troponin, and interleukin-6 (IL-6); lymphopenia, hypoalbuminemia, and thrombocytopenia. The cardiovascular manifestations noted were shock (60–65%), myocardial dysfunction (30–60%), myocarditis (65%), and coronary artery abnormalities (30–40%). About 65 to 70% required pediatric intensive care unit (PICU) admission, 40 to 60% vasoactive drugs, and 15 to 25% required mechanical ventilation. The common treatment provided was intravenous immunoglobulin (IVIg; 65–80%), steroids (50–60%), antiplatelet drugs (64%), and anticoagulation (50–55%). The overall mortality was low (1–2%).18,20–25

Radiological Assessment in Children

The chest computed tomography (CT) manifestations of COVID-19 in pediatric cases are diverse and lack specificity. As compared with adults, ground glass opacities (GGOs) in pediatric cases with COVID-19 are more localized with lower attenuation, and less lobular involvement. Other CT manifestations noted are consolidation, GGOs with consolidation, or interlobular septa thickening.26 However, the radiological picture often gets complicated by the underlying coinfection in children.27 Moreover, there are some pediatric cases with positive RT-PCR presenting with normal CT scans. Therefore, CT provides an unnecessary risk of exposure to radiation, therefore must be chosen with extreme caution in children. It is also unclear whether CT scanning has any additional value as a screening tool to rule out COVID-19 infections in children with little or no respiratory symptoms or with negative or missing RT-PCR test results.28 Even in adult population, the sensitivity of a chest CT to detect abnormalities in proven COVID-19 patients ranges from 44 to 97% (median, 69%).28

The evidence for point of care lung ultrasound (POCUS) in COVID-19 has grown in this pandemic, although it mainly includes case reports, opinion pieces, and tutorials.29 POCUS has a high sensitivity for the pulmonary manifestations of COVID-19, such as ARDS and consolidation.29 POCUS could also be utilized to monitor treatment response. Furthermore it’s an easy available bed-side tool and minimizes nosocomial spread of the disease.29,30 In a meta-analysis of seven studies (122 patients), almost all patients had abnormal lung ultrasound.31
The common abnormalities were interstitial involvement/B-pattern (97%), pleural line abnormalities (70%), pleural thickening (54%), consolidation (39%), and pleural effusion (14%). Thus there is a potential role of this modality in the triage, diagnosis, and follow-up of COVID-19 patients.

Why COVID-19 is Less Frequent and Less Severe in Children?

The answer to this relevant question is not clear and several hypotheses have been postulated related to both exposure and host factors. Children are having less exposure of COVID-19 cases, as they are usually well cared for at home, the schools are closed, and outdoor activity is limited. Children may be less susceptible to SARS-CoV-2 as a result of possible age-related differences in the expression of angiotensin-converting enzyme 2 (ACE2) receptors. Adults who required ICU care had higher levels of cytokines, whereas children with COVID-19 showed a lower degree of immune dysregulation in contrast to adults. Lymphopenia is a common finding in adults, whereas it was noted in only 3.5% of pediatric cases in the Wuhan Children's Hospital Series. Furthermore, as the children's immune system is evolving, it might respond to different viral antigens differently. Also, frequent intercurrent viral infections and vaccination could possibly enhance the immunity and play a protective role against developing COVID-19 in children.

Transmission from Children to Adults and Caregivers

The current literature is limited regarding risk of SARS-CoV-2 transmission from children to other children, adults, and caregivers. Children usually have mild or asymptomatic disease; transmission has been demonstrated to occur from asymptomatic, as well as symptomatic individuals, including transmission from children to adults within family clusters. Although SARS-CoV-2 is thought to be predominantly transmitted by respiratory droplets, fecal–oral transmission is also possible. It is unclear whether fecal RT-PCR represents active viral replication or residual noninfectious viral genomic material and the potential risk for transmission. However, there are contradictory reports which suggested that children have not played a significant role in household transmission. The data on school-related transmission is also very limited. In a systematic review based on 69 pediatric cases, the mean duration of viral shedding through the respiratory tract was 11.1 ± 5.8 days and from gastrointestinal tract was 23.6 ± 8.8 days from symptom onset. In 89% of cases, viral shedding via the gastrointestinal tract persisted for 4 weeks after nasopharyngeal or throat swabs became negative. Further studies are required to determine whether children are less or more infectious than adults. Till then, standard prevention measures should be diligently followed.

Supplemental Oxygen

In adults with COVID-19, the guidelines suggest starting supplemental oxygen if the peripheral oxygen saturation (SpO2) is <92% on room air, and recommend starting supplemental oxygen if SpO2 is <90%. It is recommended that in adult patients with acute hypoxemic respiratory failure on oxygen, the saturation should be maintained no higher than 96%.

In children with COVID-19 with respiratory distress and/or hypoxia, oxygen can be started with low flow devices according to tolerability by the patient, familiarity by the treating team, availability, amount of oxygen needed (FiO2), and cost. The nasal prongs (1–5 L/min), nasal cannula, face-mask, venturi mask, non-rebreathing mask, or bubble continuous positive airway pressure (bCPAP) can be used based on above-mentioned parameters. High-flow oxygen delivery devices have potential for aerosol generation. If tolerated, children can be advised to wear triple layer surgical mask over the nasal prongs or nasal cannula.

Noninvasive Ventilation and Heated Humidified High-Flow Nasal Cannula

In critically ill adult patients with COVID-19 in China, the use of NIV and HFNC was about one-third and two-thirds, respectively. The main concern in using these devices was the increased risk of exposure to HCPs due to aerosol generation, if negative pressure rooms are not available. The literature is ambiguous regarding the safety concerns as regards the aerosol generation. Though previous data suggest that NIV was associated with nosocomial transmission...
in cases of SARS, on the contrary human laboratory data suggest that NIV does not generate aerosols.

There are several pediatric masks available with multiple sizes, facilitating NIV therapy in children. Commonly used mask interface for NIV in children includes nasal mask, oronasal mask, nasal pillows, and total-faced masks. Multiple factors require consideration in the selection of an interface including age and developmental stage, facial anatomy, a child’s personal mask choice, interface availability, and available NIV machine. Careful mask selection, a well-fitting headgear, and time investment for mask desensitization are some important recommendations for adequate mask adaptation in children. Data in children comparing performance among various mask interfaces is scarce. In one retrospective study involving 62 children (>2 years of age), no difference was seen in NIV adherence, correction of abnormalities in nocturnal gas exchange, and leak values between nasal pillows, nasal, and oronasal masks.

Similarly, there might be safety issues with the use of HFNC. Studies that evaluated the safety of HFNC were not designed to show whether or not HFNC is AGP and did not examined the spread of viruses. HFNC does not seem to confer an increased risk of transmission of disease. In studies, evaluating bacterial environmental contamination, HFNC presented as a contamination risk similar to that of conventional oxygen. If we take evidence from studies during the SARS epidemic, HCPs exposed to HFNC were not at increased risk of developing disease. Also, HFNC was more comfortable than the conventional oxygen therapy. Therefore, in adults with COVID-19 and acute hypoxic respiratory failure, guidelines prefer HFNC over conventional oxygen therapy.

However, in children with COVID-19 who persist to have increased work of breathing and hypoxemia on supplemental oxygen should receive HFNC if available. Patients with progressive respiratory distress, or where HFNC is unavailable, can be escalated to NIV, bCPAP, or bilevel positive airway pressure (BiPAP).

A systematic review which included 12 randomized controlled trials (RCTs; 1,989 patients), provided low-certainty evidence that HFNC may reduce invasive ventilation (relative risk \( RR = 0.85; 95\% \) confidence interval [CI]: 0.74–0.99) and escalation of oxygen therapy (RR = 0.71; 95\% CI: 0.51–0.98) in patients with respiratory failure. No difference in mortality was seen between patients receiving HFNC versus conventional oxygen therapy.

Although NIV might reduce intubation and mortality in mild ARDS, it is associated with higher mortality in moderate-to-severe ARDS from multiple causes. In one trial, using NIV, failure was reported in 49\% of patients with hypoxic respiratory failure requiring escalation to intubation. In addition, patients with hypoxic respiratory failure randomized to NIV had higher mortality (28\%; 95\% CI: 21–37\%) as compared with the conventional oxygen therapy (23\%; 95\% CI: 16–33\%) or HFNC (13\%; 95\% CI: 7–20\%; \( p = 0.02 \)). In another cohort of Middle East Respiratory Syndrome (MERS) adult patients, NIV was associated with a high failure rate (92.4\%) leading to intubation.

Moreover, NIV may generate large tidal volumes and greater transpulmonary pressures, further aggravating the already injured lung. Also, the delayed initiation of invasive mechanical ventilation may lead to emergency or more unstable intubation that increases the risk of aerosol transmission.

In children, HFNC or NIV are safe and efficacious modes of respiratory support may provide adequate respiratory support to prevent the need for invasive mechanical ventilation. However, patient on noninvasive mode of ventilation (HFNC or NIV) should be monitored closely for a possible deterioration. In case of nonimprovement or deterioration, early intubation should be planned in a controlled setting rather than waiting for long time and then performing high-risk intubation in emergency and uncontrolled setting which may increase the risk of nosocomial infection to HCPs.

Thus, HFNC and NIV may be reserved for children with mild ARDS without hemodynamic instability, with strict close monitoring, airborne precautions, and in single patient rooms with negative pressure. To limit environmental contamination, the nonvented masks as interface, double lumen tubings, and viral filter at expiratory limb should be used while delivering NIV. There should be low thresholds for intubation and invasive mechanical ventilation in the event of deterioration while on HFNC or NIV.

**Endotracheal Intubation**

Children with worsening clinical status, respiratory fatigue, hemodynamic instability, \( \text{PaO}_2/\text{FiO}_2 < 300 \), or altered mental status should be considered for early intubation and mechanical ventilation. Since intubation is an AGP, previous studies on SARS and MERS showed that intubation poses a risk of viral transmission to HCPs. Therefore, it is of utmost importance to maintain safety of patient, as well as of HCPs, while performing endotracheal intubation.

The endotracheal intubation should be performed in negative pressure room or in single-patient well-ventilated room. Use full PPE along with N95 or equivalent respirators and perform hand hygiene. The checklist for drugs and equipments, and clear roles should be in place, and ventilator should be ready and on stand-by mode with disposable tubing with viral filter between expiratory limb of the circuit and machine. Inside the room, limit the number of staff to three-four (intubator, airway assistant, nurse for administering medication, and team leader). The most skilled or experienced operator should perform the endotracheal intubation to minimize the number of intubations attempts and risk of transmission. The necessary plan should be communicated and alternative plan for the difficult airways should be ready. It is preferable to perform endotracheal intubation using video-guided laryngoscopy to reduce the distance between patient’s airway and the HCP. The transparent aerosol entrainment box or plastic sheets can be used to cover the patients head and upper body while intubation to limit exposure to aerosols. These can also be used during extubation.
Since bag and mask ventilation generates aerosols, the preoxygenation can be done with nonrebreathing mask or tight-fitted face mask and bag with high-efficiency particulate air (HEPA) filter between face mask and bag but without providing positive pressure breaths. If positive pressure breaths are needed, create a tight mask seal (two-hand technique by one HCP and bagging by other) and provide positive pressure breaths at minimal rate and with small tidal volume. The cuffed or microcuffed endotracheal tubes should be used to limit peritubal air leak and risk of transmission. Use of rapid sequence induction with muscle relaxants will reduce coughing and prevent transmission to the HCPs. Once intubated, immediately inflate the cuff, connect the endotracheal tube to the already set ventilator with closed in-line suction and viral filter between endotracheal tube and circuit, and turn on the ventilator. Observation of chest rise, end-tidal carbon dioxide detection, or lung ultrasound should be used to confirm endotracheal tube placement. Use in-line closed suctioning systems postintubation to reduce aerosol generation. The unnecessary disconnections from the ventilator should be avoided. In case of circuit disconnection, the tube should be immediately clamped and connection should be reestablished. After intubation, all surfaces should be cleaned with 1% sodium hypochlorite, equipment’s should be properly cleaned or disposed, and doffing which should be monitored by trained observer or buddy. Since the process is complex and needs a lot of coordination, it is suggested to run regular simulation session in the unit.

Mechanical Ventilation

The mechanical ventilation strategies in children with COVID-19 and ARDS are more or less similar to any other child with ARDS due to other cause except for the aerosol precautions and infection control precautions to limit spread of infection to HCPs. The major focus of mechanical ventilation for COVID-19 is the avoidance of ventilator-induced lung injury while facilitating gas exchange via lung-protective ventilation. It is recommended to use low tidal volume (VT) ventilation (VT, 4–8 mL/kg of predicted body weight), over higher tidal volumes (VT > 8 mL/kg). Target plateau pressures (Pplat) should be kept <30 cm of H2O when ventilating patients with ARDS. There are no clinical trials examining the effect of positive end-expiratory pressure (PEEP) on coronavirus-induced ARDS. However, for mechanically ventilated adults with COVID-19 with moderate-to-severe ARDS, guidelines suggest using a higher PEEP strategy over a lower PEEP strategy, keeping check on barotrauma. There is no defined limit of lower or higher PEEP; however, any PEEP levels >10 cm of H2O constitutes a higher PEEP strategy. Start with PEEP of 7 to 10 cm of H2O and increase to 10 to 15 cm of H2O.

Other strategies include adequate sedation and analgesia, neuromuscular blockade, restrictive fluid strategy, prone positioning, permissive hypoxemia to limit FiO2 <60%, and permissive hypercapnia. Judicious use of neuromuscular blocking agents (NMBA), either intermittent or continuous infusion till 48 hours should be used to facilitate lung protective ventilation. The optimal fluid strategy in COVID-19 is not known; however, the strategy should remain the same as in any case of ARDS and use a conservative fluid strategy over a liberal fluid strategy.

Prone positioning should be applied early for 12 to 16 hours/day when PaO2/FiO2 <150, given its association with reduced mortality in moderate to severe ARDS. Every unit should have a protocol for prone positioning. Even cardiopulmonary resuscitation is advocated in prone position to minimize aerosolization. Clinicians should be aware of its complications such as pressure sores, vascular line and endotracheal tube displacement, facial edema, transient hemodynamic instability, corneal abrasions, brachial plexus injury, and hemodialysis vascular access flow issues. The surviving sepsis campaign guidelines suggests to use corticosteroids in mechanically ventilated adults with COVID-19 and ARDS (weak recommendation). This decision needs to be taken on case-to-case basis in pediatric population and is not routinely recommended.

In patients with severe ARDS due to COVID-19 not improving on optimizing ventilation and rescue strategies, a trial of inhaled bronchodilator (nitric oxide), recruitment maneuver, and if available, venovenous extracorporeal membrane oxygenator (VV-ECMO) can be used. There are no clinical trials of ECMO in COVID-19 patients. Since the availability of ECMO is questionable in resource-limited centers, and it requires an experienced center with full infrastructure, ECMO should only be considered in carefully selected patients with COVID-19 and severe ARDS.

Nebulization

Since nebulization also generates aerosols, it should be avoided. If bronchodilators are required, they can be administered by a metered dose inhaler with spacer.

Weaning and Extubation

In children recovering from the disease, the routine weaning and extubation protocols should be followed. During extubation, the transparent aerosol entrainment box or plastic sheets can be used to cover the patients’ head and upper body to limit exposure to HCPs. The patient should be extubated to nasal prongs with continuous monitoring to assess any need for HFNC or NIV.

Conclusion

Children with COVID-19 often need respiratory support which can be provided by simple measures like nasal prongs, nasal cannula, face mask, bCPAP, venturi mask, and nonrebreathing mask. The use of HFNC and NIV should be limited to children with mild ARDS preferably in negative pressure rooms and with adequate PPEs. Early intubation and invasive mechanical ventilation in children with moderate-to-severe ARDS, and those with hemodynamic compromise, altered sensorium, rapid deterioration, and multiple organ dysfunction syndrome are needed. While performing intubation and mechanical
ventilation, every attempt should be made to limit exposure to aerosols. The lung-protective mechanical ventilation strategies should be used with adequate sedation, analgesia, and neuromuscular blockers.

**Authors’ Contributions**

S.G. and S.K.A. reviewed the literature and prepared the initial draft. V.K. critically evaluated and modified the manuscript. S.G. will act as guarantor of the paper.

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