IV. Application of Bronchoscopy in the Diagnosis and Management of COVID-19 Patients

Flexible bronchoscopy is versatile, easy to use, and well tolerated in mechanically ventilated COVID-19 patients. Its applications include:

(1) Collection of respiratory specimens from the lower respiratory tract (i.e. sputum, endotracheal aspirate, bronchoalveolar lavage) for SARS-CoV-2 or other pathogens guides the selection of appropriate antimicrobials, which may lead to clinical benefits. Our experience indicates that lower respiratory specimens are more likely to be positive for SAR-CoV-2 than upper respiratory specimens.

(2) Can be used for localization of the site of bleeding, cessation of hemoptysis, sputum or blood clots removal; if the site of bleeding is identified by bronchoscopy, local injection of cold saline, epinephrine, vasopressin, or fibrin as well as laser treatment can be performed via the bronchoscope.

(3) Assist in the establishment of artificial airways; guide tracheal intubation or percutaneous tracheotomy.

(4) Drugs such as infusion of α-interferon and N-acetylcysteine can be administrated via the bronchoscope.

Bronchoscopic views of extensive bronchial mucosal hyperemia, swelling, mucus-like secretions in the lumen and jelly-like sputum blocking the airway in critically ill patients. (Figure 7).

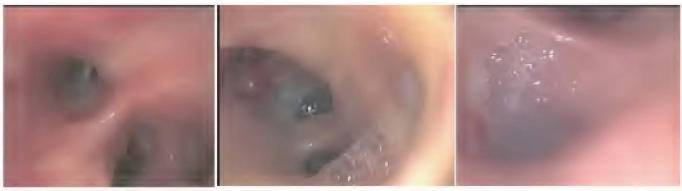


Figure 7: Bronchoscopic manifestations of COVID-19: bronchial mucosa swelling and congestion; large amounts of mucus secretions in the lumen

V. Diagnosis and Clinical Classification of COVID-19

Early diagnosis, treatment and isolation should be carried out whenever possible. Dynamic monitoring of lung imaging, oxygenation index and cytokine levels are helpful for early identification of patients who may develop into severe and critical cases. A positive result of the nucleic acid of SARS-CoV-2 is the gold standard for the diagnosis of COVID-19. However, considering the possibility of false negatives in nucleic acid detection, suspected cases characteristic manifestations in CT scans can be treated as confirmed cases even if the nucleic acid test is negative. Isolation and continuous tests of multiple specimens should be carried out in such cases.

The diagnostic criteria follow Protocols for the Diagnosis and Treatment of COVID-2019. A confirmed case is based on epidemiological history (including cluster transmission), clinical manifestations (fever and respiratory symptoms), lung imaging, and results of SARS-CoV-2 nucleic acid detection and serum-specific antibodies.

Clinical Classifications:

1. Mild Cases

   The clinical symptoms are mild and no pneumonia manifestations can be found in imaging.

2. Moderate Cases

   Patients have symptoms such as fever and respiratory tract symptoms, etc. and pneumonia manifestations can be seen in imaging.

3. Severe Cases

   Adults who meet any of the following criteria: respiratory rate ≥ 30 breaths/min; oxygen saturation ≤ 93% at a rest state; arterial partial pressure of oxygen (PaO_2)/oxygen concentration (FiO_2) ≤ 300 mmHg. Patients with > 50% lesions progression within 24 to 48 hours in lung imaging should be treated as severe cases.

4. Critical Cases

   Meeting any of the following criteria: occurrence of respiratory failure requiring mechanical ventilation; presence of shock; other organ failure that requires monitoring and treatment in the ICU.

   Critical cases are further divided into early, middle and late stages according to the oxygenation index and compliance of respiratory system.

   - Early stage: 100 mmHg <oxygenation index ≤150 mmHg; compliance of respiratory system ≥30 mL / cmH_2O; without organ failure other than the lungs. The patient has a great chance of recovery through active antiviral, anti-cytokine storm, and supportive treatment.

   - Middle stage: 60 mmHg < oxygenation index ≤100 mmHg; 30 mL/cmH_2O > compliance of respiratory system ≥15 mL/cmH_2O; may be complicated by other mild or moderate dysfunction of other organs.

   - Late stage: oxygenation index ≤ 60 mmHg; compliance of respiratory system <15 mL/cmH_2O; diffuse consolidation of both lungs that requires the use of ECMO; or failure of other vital organs. The mortality risk is significantly increased.

VI. Antiviral Treatment for Timely Elimination of Pathogens

An early antiviral treatment can reduce the incidence of severe and critical cases. Although there is no clinical evidence for effective antiviral drugs, currently the antiviral strategies based on the characteristics of SAR-CoV-2 are adopted according to Protocols for Diagnosis and Treatment of COVID-19: Prevention, Control, Diagnosis and Management.

1 Antiviral Treatment

At FAHZU, lopinavir/ritonavir (2 capsules, po q12h) combined with arbidol (200 mg po q12h) were applied as the basic regimen. From the treatment experience of 49 patients in our hospital, the average time to achieve negative viral nucleic acid test for the first time was 12 days (95% CI: 8-15 days). The duration of negative nucleic acid test result (negative for more than 2 times consecutively with interval ≥ 24h) was 13.5 days (95% CI: 9.5 - 17.5 days).

If the basic regimen is not effective, chloroquine phosphate can be used on adults between 18-65 years old (weight ≥ 50 kg: 500 mg bid; weight ≤50 kg: 500 mg bid for first two days, 500 mg qd for following five days).

Interferon nebulization is recommended in Protocols for Diagnosis and Treatment of COVID-19. We recommend that it should be performed in negative-pressure wards rather than general wards due to the possibility of aerosol transmission.

Darunavir/cobicistat has some degree of antiviral activity in viral suppression test in vitro, based on the treatment experience of AIDS patients, and the adverse events are relatively mild. For patients who are intolerant to lopinavir/ritonavir, darunavir/ cobicistat (1 tablet qd) or favipiravir (starting dose of 1600 mg followed by 600 mg tid) is an alternative option after the ethical review. Simultaneous use of three or more antiviral drugs is not recommended.

2 Course of Treatment

The treatment course of chloroquine phosphate should be no more than 7 days. The treatment course of other regimens has not been determined and are usually around 2 weeks. Antiviral drugs should be stopped if nucleic acid test results from sputum specimens remain negative for more than 3 times.

VII. Anti-shock and Anti-hypoxemia Treatment

During the progression from the severe to critically ill stage, patients may develop severe hypoxemia, cytokine cascade and severe infections that might develop into shock, tissue perfusion disorders, and even multiple organ failure. Treatment is aimed at incentive removal and fluid recovery. The artificial liver support system (ALSS) and blood purification can effectively diminish inflammatory mediators and cytokine cascade and prevent the incidence of shock, hypoxemia and respiratory distress syndrome.

1 Usage of Glucocorticoids when Necessary

Appropriate and short-term use of corticosteroids to inhibit cytokine cascade and to prevent disease progression should be considered for patients with severe COVID-19 pneumonia as early as possible. However, a high dose of glucocorticoids should be avoided due to adverse events and complications.

1.1 Indication for Corticosteroids

① for those in severe and critically ill stage;

② for those with persistent high fever (temperature above 39°C);

③ for those whose computerized tomography (CT) demonstrated patchy ground-glass attenuation or more than 30% area of the lungs are involved;

④ for those whose CT demonstrated rapid progression (more than 50% area involved in pulmonary CT images within 48 hours);

⑤ for those whose IL-6 is above ≥ 5 ULN.

1.2 Application of Corticosteroids

Initial routine methylprednisolone at a dose of 0.75~1.5 mg/kg intravenously once a day (nearly 40 mg once or twice a day) is recommended. However, methylprednisolone at a dose of 40 mg q12h can be considered for patients with falling body temperature or for patients with significantly increased cytokines under routine doses of steroid. Even methylprednisolone at a dose of 40 mg-80 mg q12h can be considered for critical cases. Closely monitor body temperature, blood oxygen saturation, blood routine, C-reactive protein, cytokines, biochemical profile and lung CT every 2 to 3 days during the treatment as necessary. The dosage of methylprednisolone should be halved every 3 to 5 days if medical conditions of patients are improved, the body temperature normalizes, or involved lesions on CT are significantly absorbed. Oral methylprednisolone (Medrol) once a day is recommended when the intravenous dose is reduced to 20 mg per day. The course of corticosteroids in not defined; some experts have suggesting ceasing corticosteroids treatment when patients are nearly recovered.

1.3 Special Consideration during Treatment

① screening of TB by T-SPOT assay, HBV and HCV by antibody assay should be performed before corticosteroid therapy;

② proton pump inhibitors could be considered to prevent complications;

③ blood glucose should be monitored. High blood glucose should be treated with insulin when necessary;

④ low serum potassium should be corrected;

⑤ liver function should be monitored closely;

⑥ traditional Chinese herbal medicine may be considered for patients who are sweating;

⑦ sedative-hypnotics can be administered temporarily for patients with sleep disorder.

2 Artificial Liver Treatment for Suppression of Cytokine Cascade

The artificial liver support system (ALSS) can conduct plasma exchange, adsorption, perfusion, and filtration of inflammatory mediators such as endotoxins and harmful metabolic substances of small or medium molecular weight. It can also provide serum albumin, coagulation factors, balance fluid volume, electrolytes and acid-base ratio, and manifest anti-cytokine storms, shock, lung inflammation, et al. In doing so it can also help to improve multiple organ functions including the liver and kidney. Thus, it can increase treatment success and reduce the mortality of severe patients.

2.1 Indication for ALSS

① serum inflammatory indicator (such as IL-6) level rises to ≥ 5 ULN, or rising rate is ≥1 time per day;

② involved area of pulmonary CT or X-ray images ≥10% progression per day;

③ artificial liver support system is required for the treatment of underlying diseases.

Patients meeting ① + ②, or patients meeting ③.

2.2 Contraindications

There is no absolute contraindication in the treatment of critically ill patients. However, ALSS should be avoided in the following situations:

① Severe bleeding disease or disseminated intravascular coagulation;

② Those who are highly allergic to blood components or drugs used in the treatment process such as plasma, heparin and protamine;

③ Acute cerebrovascular diseases or severe head injury;

④ Chronic cardiac failure, cardiac functional classification ≥ grade III;

⑤ Uncontrolled hypotension and shock;

⑥ Severe arrhythmia.

Plasma exchange combined with plasma adsorption or dual plasma molecular adsorption, perfusion, and filtration is recommended according to the patients' situation. 2000 mL of plasma should be exchanged when ALSS is performed. Detailed operating procedures can be found in the Expert Consensus on the Application of Artificial Liver Blood Purification System in the Treatment of Severe and Critical Novel Coronavirus Pneumonia.

ALSS significantly reduces the time that critically ill patients stay in the ICU in our hospital. Typically, the levels of serum cytokines such as IL-2/IL-4/IL-6/TNF-α are remarkably decreased, and oxygen saturation is significantly improved after ALSS.

3 Oxygen Therapy for Hypoxemia

Hypoxemia can present due to impaired respiratory functions by COVID-19. Oxygen supplementation treatment can correct hypoxemia, relieving secondary organ damage caused by respiratory distress and hypoxemia.

3.1 Oxygen therapy

(1) Continual oxygen saturation monitoring during oxygen therapy

Some patients do not necessarily have impaired oxygenation functions at the onset of infection but may manifest rapid deterioration in oxygenation over time. Therefore, continual monitoring of oxygen saturation is recommended, before and during oxygen therapy.

(2) Oxygen therapy as soon as possible

Oxygen therapy is not necessary for patients with oxygen saturation (SpO_2) of more than 93% or for patients without obvious symptoms of respiratory distress without oxygen treatment. Oxygen therapy is strongly recommended to the patients with symptoms of respiratory distress. It should be noted that some severe patients with $PaO_2/FiO_2 < 300$ had no obvious symptoms of respiratory distress.

(3) Treatment goal of oxygen therapy

The treatment goal of oxygen therapy is to maintain the oxygen saturation (SpO_2) at 93%-96% for patients without chronic pulmonary disease and at 88%-92% for patients with chronic type II respiratory failure. Specially, the oxygen concentration should be increased to 92%-95% for patients whose SpO_2 drops below 85% frequently during daily activities.

(4) Control oxygen therapy

PaO_2/FiO_2 is a sensitive and accurate indicator of oxygenation function. The stability and monitorability of FiO_2 are very important for patients with disease progression and PaO_2/FiO_2 below 300 mmHg. Controlled oxygen therapy is the preferred treatment.

High-flow nasal cannula (HFNC) oxygen therapy is recommended for patients with the following conditions: SpO_2 < 93%; PaO_2/FiO_2 < 300 mmHg (1 mmHg = 0.133 kPa); respiratory rate > 25 times per min at bed; or remarkable progression on X-ray imaging. Patients should wear a surgical mask during HFNC treatment. The airflow of HFNC oxygen therapy should start at a low level and gradually increased up to 40-60 L/min when PaO_2/FiO_2 is between 200-300 mmHg so that patients do not feel obvious chest tightness and shortness of breath. An initial flow of at least 60 L/min should be given immediately for patients with obvious respiratory distress.

Tracheal intubation for patients is dependent on disease progression, systemic status and complication of patients for those with stable situation but with a low oxygenation index (<100 mmHg). Thus, detailed evaluations of the clinical condition of patients is very important before decision making. Tracheal intubation should be performed as early as possible for patients with an oxygenation index less than 150 mmHg, worsening symptoms of respiratory distress or multiple organ dysfunction within 1-2 hours after high-flow (60 L/min) and high-concentration (> 60%) HFNC oxygen therapy.

Older patients (> 60 years old) with more complications or PaO_2/FiO_2 less than 200 mmHg should be treated in ICU.

3.2 Mechanical Ventilation

(1) Noninvasive Ventilation (NIV)

NIV is not strongly recommended in COVID-19 patients who fail HFNC treatment. Some severe patients progress to ARDS rapidly. Excessive inflation pressure may cause gastric distension and intolerance which contribute to aspiration and worsen lung injury. A short-term (less than 2 hours) use of NIV can be closely monitored if the patient has acute left heart failure, chronic obstructive pulmonary disease or is immunocompromised. Intubation should be performed as early as possible if improvement of respiratory distress symptoms or PaO_2/FiO_2 is not observed.

NIV with a double circuit is recommended. A virus filter should be installed between the mask and the exhalation valve when applying NIV with a single tube. Suitable masks should be chosen to reduce the risk of virus spread through air leakage.

(2) Invasive Mechanical Ventilation

① Principles of invasive mechanical ventilation in critically ill patients

It is important to balance the ventilation and oxygenation demands and the risk of mechanical ventilation-related lung injury in the treatment of COVID-19 .

• Strictly set the tidal volume to 4 - 8 mL/kg. In general, the lower the lung compliance, the smaller the preset tidal volume should be.

• Maintain the platform pressure < 30 cmH_2O (1 cmH_2O = 0.098 kPa) and driving pressure <15 cmH_2O.

• Set PEEP according to the ARDS's protocol.

• Ventilation frequency: 18-25 times per minute. Moderate hypercapnia is allowed.

• Administer sedation, analgesia, or muscle relaxant if the tidal volume, platform pressure and driving pressure are too high.

② Lung Recruitment

Lung recruitment improves the heterogeneous distribution of lesions in patients with ARDS. However, it may result in severe respiratory and circulatory complications and therefore, the lung recruitment maneuver is not routinely recommended. The assessment of lung expandability should be performed prior to the application.

(3) Prone Position Ventilation

Most critically ill patients with COVID-19 respond well to prone ventilation, with a rapid improvement of oxygenation and lung mechanics. Prone ventilation is recommended as a routine strategy for patients with PaO_2/FiO_2 < 150 mmHg or with obvious imaging manifestations without contraindications. Time course recommended for prone ventilation is more than 16 hours each time. The prone ventilation can be ceased once PaO_2/FiO_2 is greater than 150 mmHg for more than 4 hours in the supine position.

Prone ventilation while awake may be attempted for patients who have not been intubated or have no obvious respiratory distress but with impaired oxygenation or have consolidation in gravity-dependent lung zones on lung images. Procedures for at least 4 hours each time is recommended. Prone position can be considered several times per day depending on the effects and tolerance.

(4) Prevention of Regurgitation and Aspiration

Gastric residual volume and gastrointestinal function should be routinely evaluated. Appropriate enteral nutrition is recommended to be given as earlier as possible. Nasointestinal feeding and continuous nasogastric decompression are recommended. Enteral nutrition should be suspended and aspiration with 50 mL syringe be done before transfer. If no contraindication exists, a 30° semi-sitting position is recommended.

(5) Fluid Management

Excessive fluid burden worsens hypoxemia in COVID-19 patients. To reduce pulmonary exudation and improve oxygenation, the amount of fluid should be strictly controlled while ensuring the patient's perfusion.

(6) Strategies to Prevent Ventilator-Associated Pneumonia (VAP)

VAP bundled strategies should be strictly implemented:

① Select appropriate type of endotracheal tube;

② Use a endotracheal tube with subglottic suction (once every 2 hours, aspirated with 20 mL empty syringe each time);

③ Place the endotracheal tube at the right position and correct depth, fix properly and avoid pulling;

④ Maintain the airbag pressure at 30 - 35 cmH_2O (1 cmH_2O = 0.098 kPa) and monitor every 4 hours;

⑤ Monitor the airbag pressure and deal with water condensates when the position changes (two people cooperate in dumping and pouring the water condensates into a capped container containing a pre-made disinfectant chlorine solution); deal with secretions accumulated in the airbag;

⑥ Clean up secretions from the mouth and nose timely.

(7) Weaning of Ventilation

Sedatives is reduced and discontinued before awakening when the patient's PaO_2/FiO_2 is more than 150 mmHg. Intubation withdrawal should be performed as earlier as possible if permitted. HFNC or NIV is used for sequential respiratory support after withdrawal.

VIII. The Rational Use of Antibiotics to Prevent Secondary Infection

COVID-19 is a disease of viral infection, therefore antibiotics are not recommended to prevent bacterial infection in mild or ordinary patients; it should be used carefully in severe patients based on their conditions. Antibiotics can be used with discretion in patients who have the following conditions: extensive lung lesions; excess bronchial secretions; chronic airway diseases with a history of pathogen colonization in the lower respiratory tract; taking glucocorticoids with a dosage ≥ 20 mg × 7d (in terms of prednisone). The options of antibiotics

include quinolones, the second or third generation cephalothins, β-lactamase inhibitor compounds, etc. The antibiotics should be used for the prevention of bacterial infection in critically severe patients, especially those with invasive mechanical ventilation. The antibiotics such as carbapenems, β-lactamase inhibitor compounds, linezolid and vancomycin can be used in critically ill patients according to the individual risk factors.

The patient's symptoms, signs and indicators such as blood routine, C-reactive protein, and procalcitonin, need to be closely monitored during the treatment. When the change of a patient's condition is detected, a comprehensive clinical judgment needs to be made. When the secondary infection cannot be ruled out, qualified specimen need to be collected for testing by smear preparation, cultivation, nucleic acid, antigen and antibody, in order to determine the infectious agent as early as possible. Antibiotics can be empirically used in the following conditions: ① more expectoration, darker sputum color, especially yellow pus sputum; ② the rise of body temperature which is not due to exacerbation of the original disease; ③ the marked increase of white blood cells and/or neutrophils; ④ procalcitonin ≥ 0.5 ng/mL; ⑤ Exacerbation of oxygenation index or circulatory disturbance that are not caused by the viral infection; and the other conditions suspiciously caused by bacteria infections.

Some COVID-19 patients are at the risk of secondary fungal infections due to weakened cellular immunity caused by viral infections, the use of glucocorticoid and/or broad-spectrum antibiotics. It is necessary to do respiratory secretions microbiological detections such as smear preparation and cultivation for critically ill patients; and provide timely D-Glucose (G-test) and galactomannan (GM-test) of blood or bronchoalveolar lavage fluid for suspected patients.

It is necessary to be vigilant with possible invasive candidiasis infection and anti-fungal therapy. Fluconazole or echinocandin can be used in the following conditions: ① patients are given broad-spectrum antibiotics for seven days or more; ② patients have parenteral nutrition; ③ patients have invasive examination or treatment; ④ patients have positive candida culture in the specimen obtained from two body parts or more; ⑤ patients have significantly increased results of G-test.

It is necessary to be vigilant with possible invasive pulmonary aspergillosis. Anti-fungal therapy such as voriconazole, posaconazole, or echinocandin are considered to be used in the following conditions: ① patients are given glucocorticoid for seven days or more; ② patients have agranulocytosis; ③ patients have chronic obstructive pulmonary disease and aspergillus culture are tested positive in the specimen obtained from the airway; ④ patients have significantly increased results of GM-test.

IX. The Balance of Intestinal Microecology and Nutritional Support

Some COVID-19 patients have gastrointestinal symptoms (such as abdominal pain and diarrhea) due to direct viral infection of the intestinal mucosa or antiviral and anti-infective drugs. There has been report that the intestinal microecological balance is broken in COVID-19 patients, manifesting a significant reduction of the intestinal probiotics such as lactobacillus and bifidobacterium. Intestinal microecological imbalance may lead to bacterial translocation and secondary infection, so it is important to maintain the balance of intestinal microecology by microecological modulator and nutritional support.

1 Microecologics Intervention

(1) Microecologics can reduce bacterial translocation and secondary infection. It can increase dominant gut bacteria, inhibit intestinal harmful bacteria, reduce toxin production and reduce infection caused by gut microflora dysbiosis.

(2) Microecologics can improve the gastrointestinal symptoms of patients. It can reduce water in feces, improve fecal character and defecation frequency, and reduce diarrhea by inhibiting intestinal mucosal atrophy.

(3) The hospital with relevant resources can perform intestinal flora analysis. Therefore, the intestinal flora disturbance can be discovered early according to the results. Antibiotics can be adjusted timely and probiotics can be prescribed. These can reduce the chances of intestinal bacterial translocation and gut-derived infection.

(4) Nutrition support is an important means to maintain intestinal microecological balance.Intestinal nutrition support should be applied timely on the basis of effective evaluations of nutritional risks, gastroenteric functions, and aspiration risks.

2 Nutrition Support

The severe and critically ill COVID-19 patients who are in a state of severe stress are at high nutritional risks. Early evaluations of nutrition risk, gastrointestinal functions and aspiration risks, and timely enteral nutritional support are important to the patient's prognosis.

(1) Oral feeding is preferred. The early intestinal nutrition can provide nutritional support, nourish intestines, improve intestinal mucosal barrier and intestinal immunity, and maintain intestinal microecology.

(2) Enteral nutrition pathway. Severe and critically ill patients often harbor acute gastrointestinal damages, manifested as abdominal distension, diarrhea, and gastroparesis. For patients with tracheal intubation, intestinal nutrition tube indwelling is recommended for post-pyloric feeding.

(3) Selection of nutrient solution. For patients with intestinal damage, predigested short peptide preparations, which are easy for intestinal absorption and utilization, are recommended. For patients with good intestinal functions, whole-protein preparations with relatively high calories can be selected. For hyperglycemia patients, nutritional preparations which are beneficial to glycemic controlling are recommended.

(4) Energy supply. 25-30 kcal per kg body weight, the target protein content is 1.2-2.0 g/kg daily.

(5) Means of nutritional supply. Pump infusion of nutrients can be used at a uniform speed, starting with a low dosage and gradually increasing. When possible, the nutrients can be heated before feeding to reduce intolerance.

(6) The elderly patients who are at high aspiration risks or patients with apparent abdominal distention can be supported by parenteral nutrition temporarily. It can be gradually replaced by independent diet or enteral nutrition after their condition improves.

X. ECMO Support for COVID-19 Patients

COVID-19 is a novel, highly infectious disease primarily targeting pulmonary alveoli, which damages primarily the lungs of critically ill patients and leads to severe respiratory failure. For the application of extracorporeal membrane oxygenation (ECMO) in COVID-19 treatment, medical professionals need to pay close attention to the following: the time and means of intervention, anticoagulant and bleeding, coordination with mechanical ventilation, awake ECMO and the early rehabilitation training, strategy of handling for complications.

1 ECMO Intervention Timing

1.1 Salvage ECMO

In the state of mechanical ventilation support, measures such as lung protective ventilation strategy and prone position ventilation have been taken for 72 h. With the onset of one of the following conditions, salvage ECMO intervention needs to be considered.

(1) $PaO_2/FiO_2 < 80$ mmHg (regardless of what the PEEP level is);

(2) Pplat ≤ 30 mmHg, $PaCO_2 > 55$ mmHg;

(3) The onset of pneumothorax, air leakage > 1/3 tidal volume, duration > 48 h;

(4) Circulation deterioration, the dosage of norepinephrine > 1 μg/(kg×min);

(5) Cardio-pulmonary resuscitation in vitro life support ECPR.

1.2 Replacement ECMO

When the patient is not suitable for long-term mechanical ventilation support, i.e., the patient is not able to obtain the expected results, ECMO replacement needs to be adopted immediately. With the onset of one of the following conditions, ECMO replacement needs to be considered.

(1) Decreased lung compliance. After the pulmonary recruitment maneuver, the compliance of the respiratory system < 10 mL/cmH_2O;

(2) Persistent exacerbation of pneumomediastinum or subcutaneous emphysema. And the parameters of mechanical ventilation support cannot be reduced within 48 h, according to the estimation;

(3) $PaO_2/FiO_2 < 100$ mmHg. And it cannot be improved by routine methods in 72 h.

1.3 Early Awake ECMO

Early awake ECMO can be applied to patients who have been supported by mechanical ventilation with the expected high parameters for more than 7 days and who meet the necessary conditions of awake ECMO. They might benefit from it. All the following conditions must be met:

(1) The patient is in a clear state of consciousness and is fully compliant. He or she understands how ECMO works and its maintenance requirements;

(2) The patient is not complicated with neuromuscular diseases;

(3) Pulmonary damage score Murry > 2.5;

(4) Few pulmonary secretions. The time interval between the two airway suction procedures > 4 h;

(5) Stable hemodynamics. Vasoactive agents are not required for assistance.

2 Cathetering Methods

Because the ECMO supporting time for most COVID-19 patients is greater than 7 days, the seldinger method should be used as much as possible for the ultrasound guided peripheral catheter insertion, which reduces the bleeding damages and infection risks brought about by intravascular cathterization by venous angiotomy, especially for the early awake ECMO patients. Intravascular catheterization by venous angiotomy may be considered only for the patients with bad blood vessel conditions, or the patients whose catheterization cannot be identified and selected by ultrasound, or the patients whose seldinger technique failed.

3 Mode Selection

(1) The first choice for the patients of respiratory impairment is the V-V mode. The V-A mode should not be the first option just because of the possible circulation problems.

(2) For the respiratory failure patients complicated with cardiac impairment, PaO_2/FiO_2 < 100 mmHg, the V-A-V mode ought to be selected with the total flux > 6 L/min and V/A = 0.5/0.5 is maintained by current limiting.

(3) For the COVID-19 patients without severe respiratory failure but complicated with serious cardiovascular outcomes leading to cardiogenic shock, the V-A assisted by ECMO mode ought to be selected. But IPPV support is still needed and the awake ECMO should be avoided.the awake ECMO should be avoided.

4 Flux Set-value and Target Oxygen Supply

(1) The initial flux > 80% cardiac output (CO) with a self-cycling ratio < 30%.

(2) SPO_2 > 90% is to be maintained. FiO_2 < 0.5 is supported by mechanical ventilation or the other oxygen therapy.

(3) To ensure the target flux, 22 Fr (24 Fr) vein access canula is the first choice for the patient with a body weight below (above) 80 kg.

5 Ventilation Setting

Normal ventilation maintenance by adjusting the sweep gas level:

(1) The initial air flow is set to be Flow: sweep gas = 1:1. The basic target is to maintain $PaCO_2$ < 45mmHg. For the patients complicated with COPD, $PaCO_2$ < 80% basal level.

(2) The patient's spontaneous respiratory strength and respiratory rate (RR) should be maintained, with 10 < RR < 20 and without chief complaint of breathing difficulty from the patient.

(3) The sweep gas setup of the V-A mode needs to ensure the 7.35-7.45 PH value of the bloodstream out of the oxygenator membrane.

6 Anti-Coagulation and Bleeding Prevention

(1) For the patients without active bleeding, without visceral bleeding, and with platelet count > 50×109/L, the recommended initial heparin dosage is 50 U/kg.

(2) For the patients complicated with bleeding or with platelet count < 50×109/L, the recommended initial heparin dosage is 25 U/kg.

(3) The activated partial thromboplastin time (aPPT) being 40—60 sec is proposed to be the target of anticoagulation maintenance dosage. The trend of D-dimer change should be considered at the same time.

(4) Heparin-free operation may be performed in the following circumstances: the ECMO support must continue but there is fatal bleeding or active bleeding that has to be controlled; whole heparin coated loop and catheterization with blood flow > 3 L/min. The recommend operation time < 24 hour. Replacement devices and consumables need to be prepared.

(5) Heparin resistance. Under some conditions of heparin usage, aPTT is not able to reach the standard and blood coagulation happens. In this case, the activity of plasma antithrombin III (ATIII) needs to be monitored. If the activity reduces, fresh frozen plasma needs to be supplemented to restore heparin sensitivity.

(6) Heparin induced thrombopenia (HIT). When HIT happens, we recommend to perform plasma exchange therapy, or to replace heparin with argatroban.

7 Weaning from ECMO and Mechanical Ventilation

(1) If a patient treated by V-V ECMO combined with mechanical ventilation satisfies the awake ECMO condition, we suggest to first try to remove the artificial airway, unless the patient has ECMO related complications, or the expected time of removal of all the assisting machines is less than 48 h.

(2) For a patient who has too much airway secretions that frequent artificial suction clearance is needed, who is expected to have a long-term mechanical ventilation support, who satisfies the conditions $PaO_2/FiO_2 > 150$ mmHg and time > 48 h, whose lung image changes for the better, and whose damages related to mechanical ventilation pressure have been controlled, the ECMO assistance may be removed. It is not recommended to keep ECMO intubation.

XI. Convalescent Plasma Therapy for COVID-19 Patients

Since Behring and Kitasato reported the therapeutic effects of diphtheria antitoxin plasma in 1891, plasma therapy has become an important means of pathogen immunotherapy for acute infectious diseases. The disease progression is rapid for severe and critically ill patients of an emerging infectious disease. In the early phase, the pathogens damage the target organs directly and then lead to severe immuno-pathological damage. The passive immune antibodies can effectively and directly neutralize the pathogens, which reduces the damage of the target organs and then block the subsequent immune-pathological damages. During multiple global pandemic outbreaks, WHO also emphasized that "convalescent plasma the rapy is one of the most recommended potential therapies, and it has been used during other epidemic outbreaks". Since the outbreak of COVID-19, the initial mortality rate was rather high due to the lack of specific and effective treatments. As mortality rate is an important metric that the public concerns, clinic treatments which can reduce the fatality rate of critical cases effectively are key to avoid public panic. As a provincial-level hospital in Zhejiang province, we have been responsible to treat the patients from Hangzhou and the critically ill patients of the province. There are abundant potential convalescent plasma donors and critically ill patients who need convalescent plasma treatment in our hospital.

1 Plasma collection

In addition to the common requirements of blood donation and procedures, the following details should be noted.

1.1 Donors

At least two weeks after recovery and being discharged (the nucleic acid test of the sample taken from the lower respiratory tract remains negative≥14 days). 18 ≤ Age ≤ 55. The body weight>50 kg (for male) or >45 kg (for female). At least one week since last glucocorticoid usage. More than two weeks since last blood donation.

1.2 Collection Method

Plasmapheresis, 200-400 mL each time (based on medical consultation).

1.3 Post-Collection Testing

In addition to the general quality test and the test of blood-borne disease, the blood samples need to be tested for:

(1) Nucleic acid testing for SARS-CoV-2;

(2) 160-fold dilution for the qualitative test of SARS-CoV-2 specific IgG and IgM detection; or 320-fold dilution for the qualitative test of whole antibody detection. If possible, keep > 3 mL plasma for the viral neutralization experiments.

The following should be noted. During the comparison of virus neutralization titer and luminescent IgG antibody quantitative detection, we found that the present SARS-CoV-2 specific IgG antibody detection does not fully demonstrate the actual virus neutralization capability of the plasma. Therefore, we suggested the virus neutralization test as the first choice, or test the overall antibody level with the 320-fold dilution of the plasma.

2 Clinical Use of the Convalescent Plasma

2.1 Indication

(1) Severe or critically ill COVID-19 patients tested positive in respiratory tract test;

(2) The COVID-19 patients who are not severe or critically ill, but in a state of immunity suppression; or have low CT values in the virus nucleic acid testing but with a rapid disease progression in the lungs.

Note: In principle, the convalescent plasma should not be used on COVID-19 patients with disease course exceeding three weeks. But in clinical applications, we found that the convalescent plasma therapy is effective for patients with a disease course exceeding three weeks and whose virus nucleic acid tests continuously to show positive from respiratory tracts specimen. It can speed up virus clearance, increase the numbers of the plasma lymphocytes and NK cells, reduce the level of plasma lactic acid, and improve renal functions.

2.2 Contraindication

(1) Allergy history of plasma, sodium citrate and methylene blue;

(2) For patients with history of autoimmune system diseases or selective IgA deficiency, the application of convalescent plasma should be evaluated cautiously by clinicians.

2.3 Infusion plan In general, the dosage of convalescent plasma therapy is ≥400 mL for one infusion, or ≥ 200 mL per infusion for multiple infusions.

XII. TCM Classification Therapy to Improve Curative Efficacy

1 Classification and Stage

COVID-19 can be divided into early, middle, critical and recovery stages. At the early

stage, the disease has two main types: "wet lungs" and "external cold and internal heat." The middle stage is characterized by "intermittent cold and heat." The critical stage is characterized by "internal block of epidemic toxin." The recovery stage is characterized by "qi deficiency in lung-spleen." The disease initially belongs to wet lung syndrome. Due to fever, both intermittent cold and heat treatments are recommended. In the middle stage, cold, dampness, and heat coexist, belonging to "cold-heat mixture" in terms of TCM. Both cold and heat therapy should be considered. According to the theory of TCM, heat should be treated with cold drugs. But cold drugs impair Yang and lead to a cold spleen and stomach and cold-heat mixture in the middle-Jiao. Therefore, in this stage both cold and heat therapies should be considered. Because cold-heat symptoms are commonly seen in COVID-19 patients, the cold-heat therapy is better than other approaches.

2 Therapy Based on Classification

(1) Wet lungs Ephedra Herb 6 g, Semen Armeniacae Amarumg 10 g, Coix Seed 30 g, Liquoric Root 6 g, Baical Skullcap Root 15 g, Huoxiang 10 g, Reed Rhizome 30 g, Cyrtomium Rhizome 15 g, Indian Buead 20 g, Chinese Atractylodes Rhizome 12 g, Officinal Magnolia Bark 12 g.

(2) External cold and internal heat

Herba Ephedrae 9 g, Raw Gypsum Fibrosum 30 g, Semen Armeniacae Amarumg 10 g, Liquoric Root 6 g, Baical Skullcap Root 15 g, Pericarpium Trichosanthis 20 g, Fructus Aurantii 15 g, Officinal Magnolia Bark 12 g, Tripterospermum Cordifolium 20 g, White Mulberry Root-bark 15 g, Pinellia Tuber 12 g, Indian Buead 20 g, Platycodon Root 9 g.

(3) Intermittent cold-heat

Pinellia Tuber 12 g, Baical Skullcap Root 15 g, Golden Thread 6 g, Dried Ginger 6 g, Chinese Date 15 g, Kudzuvine Root 30 g, Costustoot 10 g, Indian Buead 20 g, Thunberg Fritillary Bulb 15 g, Coix Seed 30 g, Liquoric Root 6 g.

(4) Internal block of epidemic toxin

Use cheongsimhwan for treatment.

(5) Qi deficiency of lung and spleen

Membranous Milkvetch Root 30 g, Pilose Asiabell Root 20 g, Roasted Largehead Atractylodes Rhizome 15 g, Indian Buead 20 g, Fructus Amomi 6 g, Siberian Solomonseal Rhizome 15 g, Pinellia Tuber 10 g, Tangerine Peel 6 g, Wingde Yan Rhizome 20 g, Semen Nelumbinis 15 g, Chinese Date 15 g.

Patients in different stages should take different approaches. One dose per day. Boil the medicine in water. Take it every morning and evening.

XIII. Drug Use Management of COVID-19 Patients

COVID-19 patients are often complicated with underlying diseases receiving multiple types of drugs. Therefore, we should pay more attention to the adverse drug reactions and drug interactions so as to avoid drug-induced organ damage and improve the success rate of treatment.

1 Identification of adverse drug reactions

It has been demonstrated that the incidence of abnormal liver function is 51.9% in COVID-19 patients who have received lopinavir/ritonavir combined arbidol antiviral treatment. Multivariate analysis revealed that antiviral agents and more concomitant medications are two independent risk factors of abnormal liver function. Therefore, monitoring of the adverse drug reactions should be strengthened; the unnecessary drug combinations should be reduced. The main adverse reactions of antiviral agents include:

(1) Lopinavir /ritonavir and darunavir/cobicistat: diarrhea, nausea, vomit, the increase of serum aminotransferase, jaundice, dyslipidemia, the increase of lactic acid. Symptoms will recover after drug withdrawal.

(2) Arbidol: the increase of serum aminotransferase and jaundice. When combined with lopinavir, the incidence rate is even higher. The symptoms will recover after drug withdrawal. Sometimes a slowdown of the heart could be induced; thus it is necessary to avoid the combination of arbidol with β-receptor inhibitors such as metoprolol and propranolol. We suggest to stop taking the drugs when the heart rate drops below 60/min.

(3) Fapilavir: elevation of plasma uric acid, diarrhea, neutropenia, shock, fulminant hepatitis, acute kidney injury. The adverse reactions were commonly seen in elderly patients or patients complicated with cytokine storm.

(4) Chloroquine phosphate: dizziness, headache, nausea, vomit, diarrhea, different kinds of skin rash. The most severe adverse reaction is cardiac arrest. The main adverse reaction is the ocular toxicity. An electrocardiogram needs to be examined before taking the drug. The drug should be prohibited for patients with arrhythmia (e.g., conduction block), retinal disease, or hearing loss.

2 Therapeutic Drug Monitoring

Some antiviral and antibacterial drugs need therapeutic drug monitoring (TDM). Table 1 presents the plasma concentrations of such drugs and their dosage adjustment. Upon the onset of aberrations of plasma drug concentration, the treatment regimens need to be adjusted by considering the clinical symptoms and concomitant drugs.

Table 1 The range of concentrations and points for attention of the common TDM drugs for the COVID-19 patients

Drug names	Time points of blood collection	The range of concentrations	Principles of dosage adjustment
lopinavir/ ritonavir	(peak) 30 min after drug administration (trough) 30 min before drug administration	lopinavir: (trough) > 1 μg/mL (peak) < 8.2 μg/mL	Correlated with drug efficacy and side effects.
imipenem	10 min before the drug administration	1~8 μg/mL	Interpretation and adjust the plasma drug concentration based on MIC of the pathogen testing
meropenem	10 min before the drug administration	1~16 μg/mL	
vancomycin	30 min before the drug administration	10~20 mg/L (15~20 mg/L for the severe MRSA infection)	The trough concentration correlates with the failure rate of anti-infective therapy and renal toxicity. When the concentration is overly high, reduction of drug frequency or single dose is required.
linezolid	30 min before the drug administration	2~7 μg/mL	The trough concentration correlates with myelosuppression adverse reactions. The blood routine test needs to be closely monitored.
voriconazol	30 min before the drug administration	1~5.5 μg/mL	The trough concentration correlates with the therapeutic efficacy and adverse reactions such as impaired liver function.

3 Paying attention to the potential drug interactions

Antiviral drugs such as lopinavir/ritonavir are metabolized through the enzyme CYP3A in the liver. When patients receiving concomitant medications, the potential drug interactions need to be carefully screened. Table 2 shows interactions between antiviral drugs and common drugs for underlying diseases.

Table 2 Interactions between antiviral drugs and common drugs for underlying

Drug names	Potential interactions	Contraindication in combined medication
lopinavir/ ritonavir	When combined with drugs associated with CYP3A metabolism (e.g., statins, immunosuppressors such as tacrolimus, voriconazole), the plasma concentration of the combined drug may increase; leading to 153%, 5.9 folds, 13 folds increase of the AUC of rivaroxaban, atrovastatin, midazolam, respectively. Pay attention to clinical symptoms and apply the TDM.	Combined use with amiodarone (fatal arrhythmia), quetiapine (severe coma), simvastati (rhabdomyolysis) is prohibited.
darunavir/ cobicistat	When combined with drugs associated with CYP3A and/or CYP2D6 metabolism, the plasma concentration of the combined drugs may increase. See lopinavir/ ritonavir.	See lopinavir/ritonavir.
arbidol	It interacts with CYP3A4, UGT1A9 substrates, inhibitors, and inducers.	——
fapilavir	① Theophyllinum increases the bioavailability of fapilavir. ② It increases the bioavailability of acetaminophen by 1.79 folds. ③ Its combination with pyrazinamide increases the plasma uric acid level. ④ Its combination with repaglinide increases the plasma repaglinide level.	——
chloroquine phosphate	——	Prohibit to combine with the drugs that may lead to the prolonged Q-T interval (such as moxifloxacin, azithromycin, amiodarone, etc.).

Note: "—" : no relevant data; TDM: therapeutic drug monitoring; AUC: area under the curve; UGT1A9: uridine diphosphate glucosidase 1A9.

4 Avoiding medical damage in special populations

Special populations include pregnant women, patients with hepatic and renal insufficiency, patients supported by mechanical ventilation, patients under continuous renal replacement therapy (CRRT) or, extracorporeal membrane oxygenation (ECMO), etc. The following aspects need to be noted during drug administration.

(1) Pregnant women

Lopinavir/ritonavir tablets could be used. Favipiravir and chloroquine phosphate are prohibited.

(2) Patients with hepatic insufficiency Drugs that are excreted unchanged through the kidney are preferred, such as penicillin and cephalosporins, etc.

(3) Patients with renal insufficiency (including those on hemodialysis)

Drugs that are metabolized through the liver or excreted through the liver-kidney double channels are preferred, such as linezolid, moxifloxacin, ceftriaxone, etc.

(4) Patients under CRRT for 24h For vancomycin, the recommended regimen is: loading dose 1 g and maintenance dose 0.5 g, q12h. For imipenem, the maximum daily dosage should not exceed 2 g.

XIV. Psychological Intervention with COVID-19 Patients

1 The psychological stress and symptoms of COVID-19 patients

Confirmed COVID-19 patients often have symptoms such as regret and resentment, loneliness and helplessness, depression, anxiety and phobia, irritation and sleep deprivation. Some patients may have panic attacks. Psychological evaluations in the isolated wards demonstrated that, about 48% of confirmed COVID-19 patients manifested psychological stress during early admission, most of which were from their emotional response to stress. The percentage of delirium is high among the critically ill patients. There is even a report of encephalitis induced by the SARS-CoV-2 leading to psychological symptoms such as unconsciousness and irritability.

2 Establishing a dynamic mechanism for evaluation and warning of psychological crisis

Patients' mental states (individual psychological stress, mood, sleep quality, and pressure) should be monitored every week after admission and before discharge. The self-rating tools include: Self-Reporting Questionnaire 20 (SRQ-20), Patient Health Questionnaire 9 (PHQ-9) and Generalized Anxiety Disorder 7 (GAD-7). The peer-rating tools include: Hamilton Depression Rating Scale (HAMD), Hamilton Anxiety Rating Scale (HAMA), Positive and Negative Syndrome Scale (PANSS). In such a special environment as the isolated wards, we suggest that patients should be guided to complete the questionnaires through their cell phones. The doctors can interview and perform scale assessing through face-to-face or online discussion.

3 Intervention and treatment based on the assessment

3.1 Principles of intervention and treatment

For mild patients, psychological intervention is suggested. Psychological self-adjustment includes breathing relaxation training and mindfulness training. For moderate to severe patients, intervention and treatment by combining medication and psychotherapy are suggested. New antidepressants, anxiolytics, and benzodiazepines can be prescribed to improve the patients' mood and sleep quality. The second generation antipsychotics such as olanzapine and quetiapine can be used to improve psychotic symptoms such as illusion and delusion.

3.2 The recommendation of psychotropic medications in elderly patients

Middle-aged or elderly COVID-19 patients' medical situations are often complicated by physical diseases such as hypertension and diabetes. Therefore, when selecting psychotropic medications, the drug interactions and their effects on respiration must be fully considered. We recommend using citalopram, escitalopram, etc. to improve depression and anxiety symptoms; benzodiazepines such as estazolam, alprazolam, etc. to improve anxiety and sleep quality; olanzapine, quetiapine, etc. to improve psychotic symptoms.

XV. Rehabilitation Therapy for COVID-19 Patients

Severe and critically ill patients suffer from different degrees of dysfunction, especially respiratory insufficiency, dyskinesia and cognitive impairment, during both acute and recovery stages.

1 Rehabilitation therapy for severe and critically ill patients

The goal of early rehabilitation intervention is to reduce breathing difficulties, relieve symptoms, ease anxiety and depression and lower the incidence of complications. The process of early rehabilitation intervention is: rehabilitation assessment - therapy - reassessment.

1.1 Rehabilitation assessment

Based on general clinical assessment, especially functional evaluation, including respiration, cardiac status, motion and ADL should be emphasized. Focus on respiratory rehabilitation assessment, which includes the evaluation of thoracic activity, diaphragm activity amplitude, respiratory pattern and frequency, etc.

1.2 Rehabilitation therapy

The rehabilitation therapy of severe or critically ill COVID-19 patients mainly includes position management, respiratory training, and physical therapy.

(1) Position management. Postural drainage may reduce the influence of sputum on the respiratory tract, which is especially important to improve the patient's V/Q. Patients must learn to tip themselves into a position which allows gravity to assist in draining excretion from lung lobes or lung segments. For patients using sedatives and suffering from consciousness disturbance, a standing-up bed or the bed head elevation (30°-45°-60°) may be applied if the patient's condition permits. Standing is the best body position for breathing in a resting state, which can effectively increase the patient's respiratory efficiency and maintain lung volume. As long as the patient feels good, let the patient take a standing position and gradually increase the time standing.

(2) Respiratory exercise. Exercise can fully expand the lungs, help the excretions from pulmonary alveoli and airway expel into the large airway so that sputum would not accumulate at the bottom of the lungs. It increases the vital capacity and enhances lung function. Deep-slow breathing and chest expansion breathing combined with shoulder expansion are the two major techniques of respiratory exercises.

① Deep-slow breathing: while inhaling, the patient should try his/her best to move the diaphragm actively. The breathing should be as deep and slow as possible to avoid the reduction of respiratory efficiency caused by fast-shallow breathing. Compared with thoracic breathing, this kind of breathing needs less muscle strength but has better tidal volume and V/Q value, which can be used to adjust breathing when experiencing short of breath.

② Chest expansion breathing combined with shoulder expansion: Increase pulmonary ventilation. When taking a deep-slow breath, one expands his/her chest and shoulders while inhaling; and moves back his/her chest and shoulders while exhaling. Due to the special pathological factors of viral pneumonia, suspending breathing for a long time should be avoided in order not to increase the burden of respiratory function, and the heart, as well as oxygen consumption. Meanwhile, avoid moving too fast. Adjust the respiratory rate at 12-15 times/min.

(3) Active cycle of breathing techniques. It can effectively remove bronchus excretion and improve lung function without exacerbation of hypoxemia and airflow obstruction. It consists of three stages (breathing control, thoracic expansion and exhalation). How to form a cycle of breathing should be developed according to the patient's condition.

(4) Positive expiratory pressure trainer. The pulmonary interstitium of COVID-19 patients has been severely damaged. In mechanical ventilation, low pressure and low tidal volume are required to avoid damages to the pulmonary interstitium. Therefore, after the removal of mechanical ventilation, positive expiratory pressure trainer can be used to help the movement of excretions from the low volume lung segments to the high-volume segments, lowering the difficulty of expectoration. Expiratory positive pressure can be generated through air flow vibration, which vibrates the airway to achieve airway supporting. The excretions can then be removed as the high-speed expiratory flow moves the excretions.

(5) Physical therapy. This includes ultrashort wave, oscillators, external diaphragm pacemaker, electrical muscle stimulation, etc.

XVI. Lung Transplantation in Patients with COVID-19

Lung transplantation is an effective treatment approach for final-stage chronic lung diseases. However, it is rarely reported that lung transplantation has been performed to treating acute infectious lung diseases. Based on current clinical practice and results, FAHZU summarized this chapter as a reference for medical workers. In general, following the principles of exploration, doing the best to save life, highly selective and high protection, if lung lesions are not significantly improved after adequate and reasonable medical treatment, and the patient is in critical condition, lung transplantation could be considered with other evaluations.

1 Pre-transplantation assessment

(1) Age: It is recommended that the recipients are not older than 70. Patients over 70 years old are subject to careful evaluation of other organ functions and postoperative recovery capability.

(2) The course of the disease: There is no direct correlation between the length of the disease course and the severity of the disease. However, for patients with short disease courses (fewer than 4-6 weeks), a full medical assessment is recommended to evaluate whether adequate medication, ventilator assistance, and ECMO support have been provided.

(3) Lung function status: Based on the parameters collected from lung CT, ventilator, and ECMO, it is necessary to evaluate whether there is any chance of recovery.

(4) Functional assessment of other major organs: a. Evaluation of the consciousness status of patients in critical condition using brain CT scan and electroencephalography is crucial, as most of them would have been sedated for an extended period; b. Cardiac assessments, including electrocardiogram and echocardiography that focus on right heart size, pulmonary artery pressure and left heart function, are highly recommended; c. The levels of serum creatinine and bilirubin should also be monitored; for patients with liver failure and renal failure, they should not be subjected to lung transplantation until the functions of the liver and kidney are recovered.

(5) The nucleic acid test of COVID-19: The patient should be tested negative for at least two consecutive nucleic acid tests with a time interval longer than 24 hours. Given the increased incidents of COVID-19 test result returning from negative to positive after treatment, it is recommended to revise the standard to three consecutive negative results. Ideally, negative results should be observed in all body fluid samples, including blood, sputum, nasopharynx, broncho-alveolar lavage, urine, and feces. Considering the difficulty in operation, however, at least the testing of sputum and broncho-alveolar lavage samples should be negative.

(6) Assessment of infection status: With the extended in-patient treatment, some COVID-19 patients may have multiple bacterial infections, and thus a full medical assessment is recommended to evaluate the situation of infection control, especially for multidrug-resistant bacterial infection. Moreover, post-procedure antibacterial treatment plans should be formed to estimate the risk of post-procedure infections.

(7) The preoperative medical assessment process for lung transplantation in COVID-19 patients: a treatment plan proposed by the ICU team → multidisciplinary discussion → comprehensive medical evaluation → analysis and treatment of relative contraindications → pre-habilitation before lung transplantation.

2 Contraindications

Please refer to The 2014 ISHLT Consensus: A consensus document for the selection of lung transplantation candidates issued by the International Society for Heart and Lung Transplantation (updated in 2014).

XVII. Discharge Standards and Follow-up Plan for COVID-19 Patients

1 Discharge standards

(1) Body temperature remains normal for at least 3 days (ear temperature is lower than 37.5 ℃);

(2) Respiratory symptoms are significantly improved;

(3) The nucleic acid is tested negative for respiratory tract pathogen twice consecutively (sampling interval more than 24 hours); the nucleic acid test of stool samples can be performed at the same time if possible;

(4) Lung imaging shows obvious improvement in lesions;

(5) There is no comorbidities or complications which require hospitalization;

(6) SpO_2 > 93% without assisted oxygen inhalation;

(7) Discharge approved by multi-disciplinary medical team.

2 Medication after discharge

Generally, antiviral drugs are not necessary after discharge. Treatments for symptoms can be applied if patients have mild cough, poor appetite, thick tongue coating, etc. Antiviral drugs can be used after discharge for patients with multiple lung lesions in the first 3 days after their nucleic acid are tested negative.

3 Home isolation

Patients must continue two weeks of isolation after discharge. Recommended home isolation conditions are:

① Independent living area with frequent ventilation and disinfection;

② Avoid contacting with infants, the elderly and people with weak immune functions at home;

③ Patients and their family members must wear masks and wash hands frequently;

④ Body temperature are taken twice a day (in the morning and evening) and pay close attention to any changes in the patient's condition.

4 Follow-up

A specialized doctor should be arranged for each discharged patient's follow-ups. The first follow-up call should be made within 48 hours after discharge. The outpatient follow-up will be carried out 1 week, 2 weeks, and 1 month after discharge. Examinations include liver and kidney functions, blood test, nucleic acid test of sputum and stool samples, and pulmonary function test or lung CT scan should be reviewed according to the patient's condition. Follow-up phone calls should be made 3 and 6 months after discharge.

5 Management of patients tested positive again after discharge

Strict discharge standards have been implemented in our hospital. There is no discharged case in our hospital whose sputum and stool samples are tested positive again in our follow-ups. However, there are some reported cases that patients are tested positive again, after being discharged based on the standards of national guidelines (negative results from at least 2 consecutive throat swabs collected at an interval of 24 hours; body temperature remaining normal for 3 days, symptoms significantly improved; obvious absorption of inflammation on lung images). It is mainly due to sample collection errors and false negative testing results. For these patients, the following strategies are recommended:

(1) Isolation according to the standards for COVID-19 patients.

(2) Continuing to provide antiviral treatment which has been proved to be effective during prior hospitalization.

(3) Discharge only when improvement is observed on lung imaging and the sputum and stool are tested negative for 3 consecutive times (with an interval of 24 hours).

(4) Home isolation and follow-up visits after discharge in accordance with the requirements mentioned above.

Part Three Nursing

I. Nursing Care for Patients Receiving High-Flow Nasal Cannula (HFNC) Oxygen Therapy

1 Assessing

Provide detailed information of the HFNC oxygen therapy to get the patient's cooperation before implementation. Use low dose sedative with close monitoring if necessary. Choose a proper nasal catheter based on the diameter of the patient's nasal cavity. Adjust the head strap tightness and use decompression plaster to prevent device-related pressure injuries on the facial skin. Maintain the water level in the humidifier chamber. Titrate the flow rate, the fraction of inspired oxygen (FiO_2), and the water temperature based on the patient's respiratory demands and tolerance.

2 Monitoring

Report to the attending physician to seek medical decision of replacing HFNC by mechanical ventilation if any of the followings occur: hemodynamic instability, respiratory distress evidenced by obvious contraction of accessory muscles, hypoxemia persists despite oxygen therapy, deterioration of consciousness, the respiratory rate > 40 breaths per minute continuously, significant amount of sputum.

3 Treatment of Secretions

Patients' drool, snot, and sputum should be wiped with tissue paper, be disposed in a sealed container with chlorine-containing disinfectant (2500 mg/L). Alternatively, secretions can be removed by oral mucus extractor or suctioning tube and be disposed in a sputum collector with chlorine-containing disinfectant (2500 mg/L).

II. Nursing Care for Patients with Mechanical Ventilation

1 Intubation Procedures

The number of the medical staff should be limited to the minimum number that can ensure the patient's safety. Wear powered air-purifying respirator as PPE. Before intubation, perform administration of sufficient analgesia and sedative, and use muscle relaxant if necessary. Closely monitor the hemodynamic response during intubation. Reduce movement of staff in the ward, continuous purify and disinfect the room with plasma air purification technology for 30 min after completion of intubation.

2 Analgesia, Sedation and Delirium Management

Determine the target pain management goal every day. Assess pain with every 4 hours (Critical-Care Pain Observation Tool, CPOT), measure sedation with every 2 hours (RASS/BISS). Titrate the infusion rate of analgesics and sedatives to achieve pain management goals. For the known painful procedures, preemptive analgesia is administered. Perform CAM-ICU delirium screening in every shift to ensure an early diagnosis of COVID-19 patients. Apply centralization strategy for delirium prevention, including pain relief, sedation, communication, quality sleep, and early mobilization are used.

3 Prevention of Ventilator-Associated Pneumonia (VAP)

The ventilator bundle is used to reduce VAP, which includes hand washing; raising the tilt angle of the patient's bed by 30-45° if no contradiction is presented; oral care every 4 to 6 hours by using a disposable oral mucus extractor; maintain endotracheal tube (ETT) cuff pressure at 30-35 cmH_2O every 4 hours; enteral nutrition support and monitor gastric residual volume every 4 hours; evaluating daily for ventilator removal; using washable tracheal tubes for continuous subglottic suctioning combined with 10 mL syringe suctioning every 1 to 2 hours, and adjusting the suctioning frequency according to the actual amount of secretions. Dispose retentate below the glottis: the syringe containing the subglottic secretions is immediately used to aspirate an appropriate amount of chlorine-containing disinfectant (2500 mg/L), then be re-capped and disposed of in a sharp container.

4 Sputum Suction

(1) Use a closed sputum suction system, including closed suction catheter and closed disposable collection bag, to reduce the formation of aerosol and droplets.

(2) Collection of sputum specimen: use a closed suction catheter and a matching collection bag to reduce exposure to droplets.

5 Disposal of Condensation from Ventilators

Use disposable ventilator tubing with dual-loop heating wire and automatic humidifier to reduce the formation of condensation. Two nurses should cooperate to dump the condensation promptly into a capped container with chlorine-containing disinfectant (2500 mg/L). The container can then be directly put in a washing machine, which can be heated up to 90 °C, for automatic cleaning and disinfection.

6 Nursing Care for the Prone Position Ventilation (PPV)

Before changing the position, secure the position of tubing and check all the joints to reduce the risk of disconnection. Change the patient's position every 2 hours .

III. Daily Management and Monitoring of ECMO (Extra Corporeal Membrane Oxygenation)

1. **ECMO equipment should be managed by ECMO perfusionists and the following items should be checked and recorded every hour:** Pump flow rate/rotation speed; blood flow; oxygen flow; oxygen concentration; ensuring that the temperature controller is flowing; temperature setting and actual temperature; preventing clots in circuit; no pressure to the cannulae and the circuit tubing is not kinked, or no "shaking" of ECMO tubes; patient's urine color with special attention to red or dark brown urine; pre & post membrane pressure as required by the doctor.

2. **The following items during every shift should be monitored and recorded:** Check the depth and fixation of cannula to ensure that the ECMO circuit interfaces are firm, the water level line of the temperature controller, the power supply of the machine and the connection of the oxygen, the cannula site for any bleeding and swelling; measure leg circumference and observe whether the lower limb on the operation side is swollen; observe lower limbs, such as dorsalis pedis artery pulse, skin temperature, color, etc.

3. **Daily monitoring:** Post membrane blood gas analysis.

4. **Anticoagulation management:** The basic goal of ECMO anticoagulation management is to achieve a moderate anticoagulation effect, which ensures that certain coagulation activity under the premise of avoiding excessive activation of coagulation. That is to maintain the balance among anticoagulation, coagulation and fibrinolysis. The patients should be injected with heparin sodium (25-50 IU/kg) at the time of intubation and maintained with heparin sodium (7.5-20 IU/kg/h) during the pump flow period. The dosage of heparin sodium should be adjusted according to APTT results which should be held between 40-60 seconds. During the anticoagulation period, the number of skin punctures should be reduced as less as possible. Operations should be taken gently. The status of bleeding should be observed carefully.

5. Implement the "ultra-protective lung ventilation" strategy to avoid or reduce the occurrence of ventilator-related lung injury. It is recommended that the initial tidal volume is < 6 mL/kg and the intensity of spontaneous breathing is retained (breathing frequency should be between 10-20 times/min).

6. Closely observe the vital signs of patients, maintain MAP between 60-65 mmHg, CVP < 8 mmHg, SpO_2 > 90%, and monitor the status of urine volume and blood electrolytes.

7. Transfuse through the post membrane, avoiding infusion of fat emulsion and propofol.

8. According to the monitoring records, evaluate the ECMO oxygenator function during every shift.

IV. Nursing Care of ALSS (Artificial Liver Support System)

ALSS nursing care is mainly divided into two different periods: nursing care during treatment and intermittent care. Nursing staff should closely observe the conditions of patients, standardize the operating procedures, focus on key points and deal with complications timely in order to successfully complete ALSS treatment.

1 Nursing Care during Treatment

It refers to nursing during each stage of ALSS treatment. The overall operation process can be summarized as follows: operator's own preparation, patient evaluation, installation, pre-flushing, running, parameter adjustment, weaning and recording. The following are the key points of nursing care during each stage:

(1) Operator's own preparation

Fully adhere to Level III or even more strict protective measures.

(2) Patient assessment

Assess the patient's basic conditions, especially allergy history, blood glucose, coagulation function, oxygen therapy, sedation (for sober patients, pay attention to their psychological state) and catheter function status.

(3) Installation and pre-flushing

Use consumables with closed-loop management while avoiding the exposure to patient's blood and body fluids. The corresponding instruments, pipelines and other consumables should be selected according to the planned treatment mode. All basic functions and characteristics of the consumables should be familiarized.

(4) Running

It is recommended that the initial blood draw speed is ≤ 35 mL/min to avoid low blood pressure which might be caused by high speed. Vital signs should be monitored as well.

(5) Parameter Adjustment

When the patient's extracorporeal circulation is stable, all treatment parameters and alarm parameters should be adjusted according to the treatment mode. A sufficient amount of anticoagulant is recommended in the early stage and the anticoagulant dose should be adjusted during the maintenance period according to different treatment pressure.

(6) Weaning

Adopt "liquid gravity combined recovery method"; the recovery speed ≤ 35 mL/min; after weaning, medical waste should be treated in accordance to the SARS-Cov-2 infection prevention and control requirements and the treatment room and instruments should be cleaned and disinfected as well.

(7) Recording

Make accurate records of the patient's vital signs, medication and treatment parameters for ALSS and take notes on special conditions.

2 Intermittent Care

(1) Observation and treatment of delayed complications:

Allergic reactions, imbalance syndromes, etc.;

(2) ALSS Intubation Care:

Medical staff during each shift should observe the patient's conditions and make records; prevent catheter-related thrombosis; carry out professional maintenance of the catheter every 48 hours;

(3) ALSS Intubation and Extubation Care:

Vascular ultrasonography should be performed before extubation. After extubation, the lower limb with the intubation side of patients should not be moved in 6 hours and the patient should rest in bed for 24 hours. After extubation, the surface of the would to be observed.

V. Continuous Renal Replacement Treatment (CRRT) Care

1 Preparation before CRRT

Preparation for patient: establish effective vascular access. Generally, central vein catheterization is performed for CRRT, with the internal jugular vein preferred. A CRRT device can be integrated into the ECMO circuit if the two are applied at the same time. Prepare equipment, consumables, and ultrafiltration medication before CRRT.

2 In-treatment Care

(1) Vascular Access Care:

Perform professional catheter care every 24 hours for patients with central venous catheterization to properly fix access to avoid distortion and compression. When CRRT is integrated into ECMO treatment, the sequence and the tightness of the catheter connection should be confirmed by two nurses. Both the outflow and the inflow CRRT lines are suggested to be connected behind the oxygenator.

(2) Closely monitor consciousness and the vital signs of patients; accurately calculate the fluid inflow and outflow. Closely observe blood clotting within the cardiopulmonary bypass circuit, respond effectively to any alarms, and ensure that the machine is operating properly. Assess the electrolyte and acid-base balance in the internal environment through blood gas analysis every 4 hours. The replacement liquid should be prepared freshly and labeled clearly under strict sterile conditions.

3 Postoperative Care

(1) Monitor blood routine, liver and kidney function and coagulation function.

(2) Wipe the CRRT machine every 24 hours if continuous treatment is applied. Consumables and wasted liquid should be disposed in accordance with hospital requirements to avoid nosocomial infection.

VI. General Care

1 Monitoring

Patient vital signs should be continuously monitored, especially changes in consciousness, respiration rate and the oxygen saturation. Observe symptoms such as cough, sputum, chest tightness, dyspnea, and cyanosis. Monitor arterial blood gas anlysis closely. Timely recognition of any deterioration to adjust strategies of oxygen therapy or to take urgent response measures. Pay attention to ventilator associated lung injury (VALI) when under high positive end-expiratory pressure (PEEP) and high-pressure support. Closely monitor changes in airway pressure, tidal volume and respiratory rate.

2 Aspiration Prevention

(1) Gastric retention monitor: perform continuous post-pyloric feeding with a nutrition pump to reduce gastroesophageal reflux. Evaluate gastric motility and gastric retention with ultrasound if possible. Patient with normal gastric emptying are not recommended for routine assessment;

(2) Evaluate gastric retention every 4 hours. Re-infuse the aspirate if the gastric residual volume is < 100 mL; otherwise, report to the attending physician;

(3) Aspiration prevention during patient transportation: before transportation, stop nasal feeding, aspirate the gastric residues and connect the gastric tube to a negative pressure bag. During transportation, raise the patient's head up to 30°;

(4) Aspiration prevention during HFNC: Check the humidifier every 4 hours to avoid excessive or insufficient humidification. Remove any water accumulated in the tubing promptly to prevent cough and aspiration caused by the accidental entry of condensation into the airway. Keep the position of the nasal cannula higher than the machine and tubes. Promptly remove condensation in the system.

3 Implement strategies to prevent catheter-related bloodstream infection and catheter-related urinary tract infection.

4 Prevent pressure-induced skin injuries, including device-related pressure-induced injuries, incontinence-associated dermatitis and medical adhesive-related skin injuries. Identify patients at a high risk with the Risk Assessment Scale and implement preventive strategies.

5 Assess all patients upon admission and when their clinical conditions change with the VTE risk assessment model to identify those who are at a high risk and implement preventive strategies. Monitor coagulation function, D-dimer levels and VTE-related clinical manifestations.

6 Assist eating for patients who are weak, short of breath or those with an obvious fluctuating oxygenation index. Intensify oxygenation index monitoring on these patients during meals. Provide enteral nutrition at early stages for those who are unable to eat by mouth. During each shift, adjust the enteral nutrition rate and quantity according to the tolerance of enteral nutrition.

Appendix

I. Medical Advice Example for COVID-19 Patients

1 Medical Advice of Mild COVID-19 Cases

1.1 Ordinary

- Air isolation, blood oxygen saturation monitoring, oxygen therapy with nasal cannula

1.2 Examinations

- 2019 Novel Coronavirus RNA Detection (Three Sites) (Sputum) qd
- 2019 Novel Coronavirus RNA Detection (Three Sites) (Feces) qd
- Blood routine, biochemical profile, urine routine, stool routine + OB, coagulation function + D dimer, blood gas analysis + lactic acid, ASO + RF + CPR + CCP, ESR, PCT, ABO + RH blood type, thyroid function, cardiac enzymes + quantitative assay of serum troponin, four routine items, respiratory virus test, cytokines, G/GM test, angiotensin converting enzyme
- Liver, gallbladder, pancreas and spleen ultrasound, echocardiography and lung CT scan

1.3 Medication

- Arbidol tablets 200 mg po tid
- Lopinavir/Ritonavir 2 tablets po q12h
- Interferon spray 1 spray pr. tid

2 Medical Advice of Moderate COVID-19 Cases

2.1 Ordinary

- Air isolation, blood oxygen saturation monitoring, oxygen therapy with nasal cannula

2.2 Examinations

- 2019 Novel Coronavirus RNA Detection (Three Sites) (Sputum) qd
- 2019 Novel Coronavirus RNA Detection (Three Sites) (Feces) qd
- Blood routine, biochemical profile, urine routine, stool routine + OB, coagulation function + D dimer, blood gas analysis + lactic acid, ASO + RF + CPR + CCP, ESR, PCT, ABO + RH blood type, thyroid function, cardiac enzymes + quantitative assay of serum troponin, four routine items, respiratory virus test, cytokines, G/GM test, angiotensin converting enzyme
- Liver, gallbladder, pancreas and spleen ultrasound, echocardiography and lung CT scan

2.3 Medication

- Arbidol tablets 200 mg po tid
- Lopinavir/Ritonavir 2 tablets po q12h
- Interferon spray 1 spray pr.nar tid
- NS 100 mL + Ambroxol 30mg ivgtt bid

3 Medical Advice of Severe COVID-19 Cases

3.1 Ordinary

- Air isolation, blood oxygen saturation monitoring, oxygen therapy with nasal cannula

3.2 Examinations

- 2019 Novel Coronavirus RNA Detection (Three Sites) (Sputum) qd
- 2019 Novel Coronavirus RNA Detection (Three Sites) (Feces) qd
- Blood routine, biochemical profile, urine routine, stool routine + OB, coagulation function + D dimer, blood gas analysis + lactic acid, ASO + RF + CPR + CCP, ESR, PCT, ABO + RH blood type, thyroid function, cardiac enzymes + quantitative assay of serum troponin, four routine items, respiratory virus test, cytokines, G/GM test, angiotensin converting enzyme
- Liver, gallbladder, pancreas and spleen ultrasound, echocardiography and lung CT scan

3.3 Medication

- Arbidol tablets 200 mg tid
- Lopinavir/Ritonavir 2 tablets po q12h
- Interferon spray 1 spray pr.nar tid
- NS 100 mL + methylprednisolone 40 mg ivgtt qd
- NS 100 mL + pantoprazole 40 mg ivgtt qd
- Caltrate 1 tablet qd
- Immunoglobulin 20 g ivgtt qd
- NS 100 mL + Ambroxol 30 mg ivgtt bid

4 Medical Advice of Critical COVID-19 Cases

4.1 Ordinary

Air isolation, blood oxygen saturation monitoring, oxygen therapy with nasal cannula

4.2 Examinations

- 2019 Novel Coronavirus RNA Detection (Three Sites) (Sputum) qd
- 2019 Novel Coronavirus RNA Detection (Three Sites) (Feces) qd
- Blood routine, ABO + RH blood type, urine routine, stool routine + OB, four routine items, respiratory virus test, thyroid function, electrocardiogram, blood gas analysis + electrolyte + lactic acid + GS, G/GM test, blood culture ONCE
- Blood routine, biochemical profile, coagulation function + D dimer, blood gas analysis + lactic acid, natriuretic peptide, cardiac enzyme, quantitative assay of serum troponin, immunoglobulin + complement, cytokine, sputum culture, CRP, PCT qd
- Blood glucose measurement q6h
- Liver, gallbladder, pancreas and spleen ultrasound, echocardiography and lung CT scan

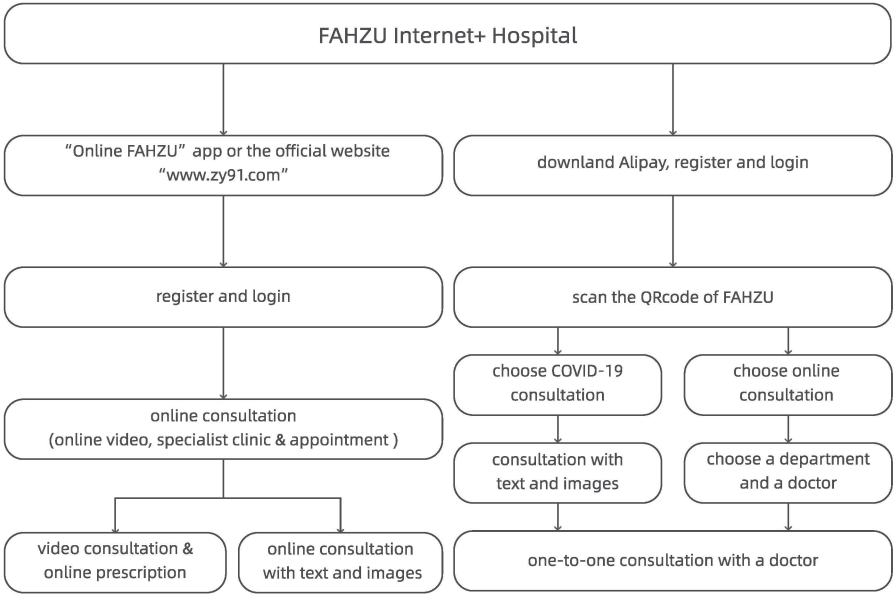
4.3 Medication

- Arbidol tablets 200 mg po. tid
- Lopinavir/Ritonavir 2 tablets q12h (or darunavir 1 tablet qd)
- NS 10 mL + methylprednisolone 40 mg iv q12h
- NS 100 mL + pantoprazole 40 mg ivgtt qd
- Immunoglobulin 20 g ivgtt qd
- Thymic peptides 1.6 mg ih biw
- NS 10 mL + Ambroxol 30 mg iv bid
- NS 50 mL + isoproterenol 2 mg iv-vp once
- Human serum albumin 10 g ivgtt qd
- NS100 mL + piperacillin/tazobactam 4.5 ivgtt q8h
- Enteral nutrition suspension (Peptisorb liquid) 500 mL nasogastric feeding bid

II. Online Consultation Process for Diagnosis and Treatment

2.1 Online Consultation for Diagnosis and Treatment

Instructions on FAHZU Internet+ Hospital



"Online FAHZU" app or the official website "www.zy91.com"

FAHZU Internet+ Hospital

Please feel free to contact us:
Email: zdyy6616@126.com, zyinternational@163.com

2.2 Online Doctors' Communication Platform

Instructions on the International Medical Expert Communication Platform of The First Affiliated Hospital, Zhejiang University School of Medicine

Visit www.dingtalk.com/en or scan the QR code (Figure1) to download DingTalk app

Sign up for a DingTalk account with your name and phone number (not open to public).

Method 1: Select "Contacts" > "Join Team" > "Join by Team Code" > Enter Input ID: 'YQDK1170'.

Method 2: Scan the QR code (Figure 2) of the International Medical Expert Exchange Platform of FAHZU.

Fill out your information to apply to join. Enter your name, country, and medical institution.

Become the member of the team after the admin's approval.

Communicate with experts in group chat by using your mobile or PC.

Instant picture and text communication assisted with AI translation

Remote video conference

Access to medical treatment guideliness

Figure1: Scan to Download DingTalk App

Figure 2: QR Code of FAHZU Communication Platform

Figure 3: User Guide

Note: Scan the QR code of Figure 3 to download user guide

Editorial Board

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浙大一院

Overview of FAHZU

Founded in 1947, The First Affiliated Hospital, Zhejiang University School of Medicine (FAHZU), is the earliest affiliated hospital of Zhejiang University. With six campuses, it has now evolved into a medical center integrating health care, medical education, scientific research and preventative care. In terms of overall strength, FAHZU is ranked 14th in China.

As a large-size general hospital, it currently has over 6,500 employees, including academicians of the Chinese Academy of Engineering, National Distinguished Young Scholars and other outstanding talents. There is a total of 4,000 beds available to patients in FAHZU. Its main campus handled 5 million emergency and outpatient visits in 2019.

Over the years, FAHZU has successfully developed a number of renowned programs in organ transplantation, pancreatic diseases, infectious diseases, hematology, nephrology, urology, clinical pharmacy, etc. FAHZU helps many realize the radical resection of cancer and enjoy long-term survival. FAHZU is also an integrated provider of liver, pancreas, lung, kidney, intestine and heart transplantation. In the fight against SARS, H7N9 avian flu and COVID-19, it has gained rich experience and fruitful results. As a result, its medical professionals have published many articles in journals such as ***New England Journal of Medicine, the Lancet, Nature*** and ***Science.***

FAHZU has been extensively engaged into overseas exchanges and collaboration. It has established partnerships with over 30 prestigious universities around the world. Productive achievements have also been accomplished through exchange of our medical experts and technologies with Indonesia, Malaysia and other countries.

Adhering to the core value of seeking truth with prudence, FAHZU is here to offer quality health care to everyone in need.


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