Pulmonary vascular disease

ORIGINAL ARTICLE

Pulmonary hypertension in the intensive care unit. Expert consensus statement on the diagnosis and treatment of paediatric pulmonary hypertension. The European Paediatric Pulmonary Vascular Disease Network, endorsed by ISHLT and DGPK

Michael Kaestner,1 Dietmar Schranz,2 Gregor Warnecke,3,4 Christian Apitz,1 Georg Hansmann,5 Oliver Miera5

ABSTRACT

Acute pulmonary hypertension (PH) complicates the course of several cardiovascular, pulmonary and other systemic diseases in children. An acute rise of RV afterload, either as exacerbating chronic PH of different aetiologies (eg, idiopathic pulmonary arterial hypertension (PAH), chronic lung or congenital heart disease), or pulmonary hypertensive crisis after corrective surgery for congenital heart disease, may lead to severe circulatory compromise. Only few clinical studies provide evidence on how to best treat children with acute severe PH and decompensated RV function, that is, acute RV failure. The specific treatment in the intensive care unit should be based on the underlying pathophysiology and not only be focused on so-called ‘specific’ or ‘tailored’ drug therapy to lower RV afterload. In addition therapeutic efforts should aim to optimise RV preload, and to achieve adequate myocardial perfusion, and cardiac output. Early recognition of patients at high risk and timely initiation of appropriate therapeutic measures may prevent the development of severe cardiac dysfunction and low cardiac output. In patients not responding adequately to pharmacotherapy, (1) novel surgical and interventional techniques, temporary mechanical circulatory support with extracorporeal membrane oxygenation, (2) pumpless lung assist devices and (3) or lung or heart-lung transplantation should be timely considered. The invasive therapeutic measures can be applied in a bridge-to-recovery or bridge-to-lung transplant strategy. This consensus statement focuses on the management of acute severe PH in the paediatric intensive care unit and provides an according treatment algorithm for clinical practice.

INTRODUCTION

Despite recent advances in the specific treatment of pulmonary hypertension (PH), RV failure following or in the context of severe rise of pulmonary vascular resistance (PVR) is a challenging complication of PH and is associated with substantial morbidity and mortality.

PH leading to a decompensation of the cardiovascular system can be considered a syndrome with non-specific signs and symptoms presenting late in the disease process and may be strongly associated with, or directly caused by several, very heterogeneous underlying conditions.1 Distinction between precapillary and postcapillary aetiologies (or establishment of a combination of the two) is important to initiate specific individual therapy.

Pathophysiology of acute PH and RV failure

Chronic PH causes adaptation and remodelling of the RV to increased loading conditions. Pulmonary hypertensive crisis (PHC) occurs when compensatory mechanisms fail, RV systemic function decompensates and RV preload acutely decreases resulting in abolished cardiac output and coronary perfusion.2 3 Acute elevation of afterload (pulmonary artery pressure (PAP)) is poorly tolerated by the unprepared RV. In healthy adult individuals, the RV cannot acutely generate a mean PAP >40 mm Hg.4 Adaptive changes of the RV microstructure and function do not work in the setting of acute rise of RV afterload. Thus, although myocardial contractility may initially rise with RV concentric hypertrophy and preserved systolic and diastolic function, excessive pressure overload results in maladaptive remodelling with RV dilatation and failure.5 Among many others, arrhythmia, myocardial ischaemia and/or pulmonary disease such as infections or pulmonary arterial embolism may trigger an acute rise of PAP and PVR (figure 1). With subsequent RV dilatation, the contractile sarcomere apparatus is damaged, resulting in RV failure. The combination of right-to-left septal shift with subsequent LV compression, related to limited space within the pericardial sac, results in low LV output, systemic arterial hypotension and elevated IV and RV end-diastolic pressure. With or without RV hypertrophy, these factors decrease coronary perfusion leading to myocardial ischaemia that perpetuates RV failure.6 The resulting metabolic acidosis further increases PVR and PAP Moreover, a sudden increase in PAP causes airway obstruction due to arterial dissection of the smaller intrapulmonary arteries and lung oedema. The resulting dead space ventilation and ventilation/perfusion mismatch causes hypoxia and respiratory acidosis, eventually resulting in even more elevated PAP and PVR (figure 1).2 3 7

In children with congenital heart disease (CHD) and systemic-to-pulmonary shunt, PH crisis may occur following corrective surgery. Factors that promote the development of postoperative PH are
not yet completely understood, however, endothelial cell dysfunction was found to be a contributing factor preoperatively and postoperatively.8–10 Inflammatory response to cardiopulmonary bypass, and ischaemia-reperfusion injury and its impact on heart-lung function contribute to the rise in PVR.11 Pain, awakening reactions in mechanically ventilated children, tracheal secretions or tracheal suctioning may trigger acute severe elevation of PVR presenting as PHC and potentially leading to cardiocirculatory collapse and death (figure 1).

In this consensus statement, we distinguish between and elaborate on three common scenarios: (1) acute PH crisis after cardiac surgery for CHD, with different management strategies subdivided for patients with univentricular and biventricular physiology (2) acute deterioration in a child with previously known chronic PH (eg, a child with chronic parenchymal lung disease and acute viral chest infection and acute deterioration in a child with new diagnosis of group 1 PH (pulmonary arterial hypertension (PAH), eg, idiopathic PAH), and (3) secondary PH.12 We finally focus on the rapid assessment and efficient management of acute severe PH in the paediatric intensive care unit (PICU), including concise recommendations and a treatment algorithm for clinical practice.

METHODS

The recommendations given in table 2 relate to the grading system currently suggested by the European Society of Cardiology (ESC) and the American Heart Association (AHA), and was based on paediatric data only (class of recommendation, level of evidence). The grading and voting process within the writing group is outlined in the executive summary13 of this online supplement. Computerised searches of the PubMed/MEDLINE bibliographical database were conducted between 1990 and June 2015. Clinical trials, guidelines and reviews limited to paediatric data were searched using the terms ‘pulmonary hypertension’ and ‘intensive care’, ‘heart failure’, ‘congenital heart’, ‘postoperative’, ‘surgery’, ‘ECMO’, ‘lung assist’, ‘ventricular assist’, ‘mechanical circulatory support’, ‘cardiopulmonary bypass’, and ‘pulmonary hypertensive crisis’. The primary focus of this manuscript is on group 1 PH according to the World Symposium on Pulmonary Hypertension (WSPH).
and transthoracic echocardiography are mandatory to guide
that is beyond the scope of this article.

Kaestner M, et al

Assessment, the following questions have to be addressed:
rapidly lead to cardiorespiratory collapse. During the initial
Acute elevation of PVR is a life-threatening event that may
Initial assessment and monitoring of children with acute PH
▸ Are there any other causes that could explain the symptoms
▸ Are there any precipitating factors that might be responsible
▸ Are there any other causes that could explain the symptoms
▸ High output and tissue hypoxia?

After the initial clinical assessment of vital signs, chest X-ray
Chest X-ray helps to differentiate between PH with and
without ventilation-perfusion mismatch and may show involve-
ment of one or both lungs. In addition, in patients with
chronic PH.14 15 In patients who have undergone surgery
for CHD, the ECG may still display changes that occur with the
underlying heart defect, but sometimes it may be normal. Loss
of sinus rhythm (eg, in atrial flutter or junctional tachycardia) is
an important diagnosis as it aggravates RV failure.

ECG
Signs of right atrial dilatation, right axis deviation, right bundle
branch block or RV hypertrophy are typically seen in patients
with chronic PH.14 15 In patients who have undergone surgery
for CHD, the ECG may still display changes that occur with the
underlying heart defect, but sometimes it may be normal. Loss
of sinus rhythm (eg, in atrial flutter or junctional tachycardia) is
an important diagnosis as it aggravates RV failure.

Chest X-ray
Chest X-ray helps to differentiate between PH with and
without ventilation-perfusion mismatch and may show involve-
ment of one or both lungs. In addition, in patients with
chronic or newly diagnosed PH, signs of enlargement of the
eric or newly diagnosed PH, signs of enlargement of the
therapy. Monitoring of haemodynamics and organ function
(brain, liver, kidney, coagulation system, etc) is an essential part
of the routine set-up and procedures in the PICU.1

Table 1  Recommendations on the therapy of acute pulmonary hypertension in the paediatric ICU

<table>
<thead>
<tr>
<th>Recommendations</th>
<th>COR</th>
<th>LOE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oxygen should be given when transcutaneous oxygen saturation &lt;95% in children with PH and normal cardiac anatomy (^2)</td>
<td>I</td>
<td>C</td>
</tr>
<tr>
<td>Intravenous prostanoids should be considered to treat children with severe PH (^3)</td>
<td>Ia</td>
<td>B</td>
</tr>
<tr>
<td>iNO may be considered for treatment of postoperative PH in mechanically ventilated patients to improve oxygenation and reduce the risk of pulmonary hypertensive crisis (^24)</td>
<td>Iib</td>
<td>B</td>
</tr>
<tr>
<td>Concomitant sildenafil should be administered to prevent rebound PH in patients who have signs of increased PAP on withdrawal of iNO, and require re-start of iNO despite preceding gradual weaning of iNO.</td>
<td>I</td>
<td>B</td>
</tr>
<tr>
<td>Intravenous sildenafil may be considered for treatment of PH in critically ill patients, especially in those with an unsatisfactory response to iNO (^31)</td>
<td>Iib</td>
<td>B</td>
</tr>
<tr>
<td>Inhaled iloprost may be as effective as iNO in children with postoperative PH (^25-27)</td>
<td>Iib</td>
<td>B</td>
</tr>
<tr>
<td>In children who develop signs of low cardiac output or profound pulmonary failure despite optimal medical therapy, extracorporeal life support may be considered (^7-36)</td>
<td>Iib</td>
<td>C</td>
</tr>
</tbody>
</table>

COR, class of recommendation; ICU, intensive care unit; iNO, inhaled nitric oxide; LOE, level of evidence; PH, pulmonary hypertension.

Table 2  Medications used for treatment of pulmonary hypertension in the intensive care unit

<table>
<thead>
<tr>
<th>Drug</th>
<th>Dose</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Epoprostenol intravenous</td>
<td>Start with 1–3 ng/kg/min, increase gradually to 60 (and more) ng/kg/min intravenous</td>
<td>Caution: systemic arterial hypotension. Need to change drug vial/delivery system every 12–24 h</td>
</tr>
<tr>
<td>Iloprost inhal ed, intravenous</td>
<td>0.25 μg/kg inhal, max. 10 μg; 6x/day or 1–5 μg/kg/min intravenous</td>
<td>Caution: systemic arterial hypotension</td>
</tr>
<tr>
<td>iNO</td>
<td>2–40 ppm inhal</td>
<td></td>
</tr>
<tr>
<td>Sildenafil intravenous, oral</td>
<td>0.4 mg/kg bolus over 3hrs IV (optional), then 1.6–2.4 mg/kg/day continuous infusion oral 8–20 kg: 3×10 mg oral &gt;20 kg: 3×20 mg oral</td>
<td>do not exceed 30mg/d. Higher sildenafil doses up to 7.2mg/kg/day IV have been used in newborn infants with PPHN associated with congenital diaphragmatic hernia</td>
</tr>
<tr>
<td>Epinephrine</td>
<td>0.01–1 μg/kg/min intravenous</td>
<td>Positive inotropy. Increases myocardial oxygen consumption, tachycardia. Moderate effects on PVR and SVR</td>
</tr>
<tr>
<td>Norepinephrine</td>
<td>0.01–1 μg/kg/min intravenous</td>
<td>Increases SVR and PVR</td>
</tr>
<tr>
<td>Vasopressin</td>
<td>0.0003–0.002 IU/kg/min intravenous</td>
<td>Probably does not increase PVR (advantage vs norepinephrine)</td>
</tr>
<tr>
<td>Terlipressin</td>
<td>5–10 ng/kg/min intravenous</td>
<td>Probably does not increase PVR (advantage vs norepinephrine)</td>
</tr>
<tr>
<td>Dobutamine</td>
<td>5–20 μg/kg/min intravenous</td>
<td>Increases myocardial oxygen consumption, tachycardia. Probably does not increase PVR</td>
</tr>
<tr>
<td>Milrinone</td>
<td>0.375–1.0 μg/kg/min intravenous</td>
<td>Lowers PVR. Caution: systemic arterial hypotension</td>
</tr>
<tr>
<td>Levosimendan</td>
<td>0.1 μg/kg/min intravenous</td>
<td>Lowers PVR. Caution: systemic arterial hypotension</td>
</tr>
</tbody>
</table>

COR, class recommendation; ICU, intensive care unit; iNO, inhaled nitric oxide; PPHN, persistent pulmonary hypertension in the newborn; PVR, pulmonary vascular resistance; SVR, systemic vascular resistance.
right heart and central pulmonary arteries are typically present. In more severe cases, one may recognise peripheral rarefaction of the pulmonary vasculature (increased radiotranc-
sluency of the lung fields). However, right heart enlargement
may be best seen in the lateral plane (radiopaque retrosternal
space that is filled by the dilated RV). In patients with under-
lying CHD, changes on the chest X-ray can be related to the
cardiac lesion itself.15 16

Echocardiography
Echocardiography is the most important tool for the assessment
of ventricular function and RV-LV interaction.17 Doppler ana-
lysis of tricuspid valve and pulmonary valve regurgitation to
estimate PAP and diastolic inflow characteristics of both ventri-
cles may help in guiding therapy in the ICU. The estimated PAP
and right ventricular systolic pressure (RVSP) need to be inter-
preted in the context of the degree of RV dysfunction. Low
pressure gradients across the tricuspid and pulmonary valves
may be due to RV failure; they should be seen in relation to
other variables (eg, tricuspid annular plane systolic excursion
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absence of pericardial
and pleural effusions, or ascites. If pulmonary vein obstruction
(TAPSE), and in light of the presence or absent

Invasive monitoring
Invasive monitoring with an arterial and a central venous line
should be established in all patients with cardiopulmonary com-
promise in whom vasopressor or inotropic therapy may be
necessary. Insertion of invasive lines should be done with suffi-
cient local anaesthesia and adequate analgesedation. Additional
general anaesthesia might be hazardous (see anaesthesia and
ventilation). Invasive monitoring is also indicated in patients at
risk for systemic hypotension secondary to PH targeted therapy,
even if this does not include vasopressor/inotropic therapy.
There is a broad controversy on the most useful way to haemo-
dynamically monitor a child with PH, in particular, we cannot
make a clear recommendation for or against invasive PAP mon-
toring/Swan-Ganz catheters, in the PICU. Measurement of
filling pressures of both ventricles may be of value to guide fluid
management and escalation of therapy in distinct scenarios.

THERAPY OF ACUTE PH IN THE ICU
Basic measures in the ICU
Oxygen
As a potent pulmonary vasodilator and a weak systemic vaso-
constrictor, oxygen is indicated in children with ventilation-
perfusion mismatch based on arterial oxygen saturations of less
than 93% (figure 2). Sufficient supply prevents anaerobic metab-
olism in peripheral organs.2 In children with systemic-to-pulmonary
shunts, supplemental oxygen augments pulmonary overcirculation
with the risk of worsening cardiac
and pulmonary function. In cyanotic heart disease with
pulmonary-to-systemic (right-to-left) shunt flow, a higher
haemoglobin level and shunt flow guarantees adequate systemic
oxygen delivery. In these patients, oxygen is indicated in con-
comitant parenchymal lung disease or in deep cyanosis; arterial
oxygen saturations of 75–85% are generally accepted as
sufficient.

Alkalisation
Alkalisation is effective for immediate treatment of PH crisis.18 19 Acidosis elevates PVR and impairs the effect of ino-
tropic and vasopressor drugs. Therefore, acidosis, as measured
by negative base excess, should be abolished. Alkalisation with
sodium bicarbonate to achieve a pH of 7.44 resulted in signifi-
cantly reduced PVR. Of note, neurodevelopmental outcome
might be negatively affected after prolonged hypocapnic alkali-
osis in newborns.20

Sedation
Anxiety and agitation increase PVR and oxygen consumption
and should be avoided. Sedation of a critically ill child has to be
done with caution: in spontaneously breathing patients, hypo-
ventilation and apnoea have to be avoided. In ventilated
patients, loss of sufficient LV preload together with substantial
decrease of systemic vascular resistance (SVR) can lead to circu-
lar arrest due to loss of coronary perfusion pressure.

Anaesthesia and ventilation
Anaesthesia, intubation and insertion of invasive lines are among
the most crucial steps in the management of a child with immin-
ent deterioration. Mechanical ventilation is indicated in severe
PH with profound cyanosis, respiratory or metabolic acidosis not
responding to initial therapy, respiratory failure or in cardiac cir-
culatory arrest. In patients responding to medical therapy mecha-
nical ventilation should be avoided. Anaesthesia should be
performed by the most experienced person available. Induction
is usually started with a rapidly acting sedative and muscle relax-
ant and may be followed by an opioid. Data regarding ketamine
are ambiguous because its effect on PVR depends on comedica-
tion.21 Induction of anaesthesia for intubation may cause a pro-
nounced fall in SVR, and circulatory collapse. To overcome the
fall in SVR vasopressor support may become necessary. Nursing
care and respiratory therapy of ventilated patients requires aware-
ness of cardiopulmonary interactions. Maneouvers triggering pul-
monary hypertensive crises such as insufficient sedation, rise of
pCO2 or suctioning should be avoided. Moreover, positive pres-
sure ventilation impairs cardiac filling and output,22 especially
in the failing RV. Normoventilation (pCO2 levels 35–40 mm Hg)
and long expiratory times are recommended. Hyperventilation
reduces cardiac output, increases SVR19 and induces lung injury.
In patients with failing RV or in univentricular circulation, the
pulmonary perfusion pressure (flow) in relation to mean airway
pressure has to be monitored to guarantee sufficient pulmonary flow.

Fluid management
RV function in neonates and patients with significant PH is
preload dependent. Volume loss is poorly tolerated and in an
acute PH crisis, volume challenge may be useful, but haemo-
dynamic monitoring is mandatory. On the other hand, chronic
RV failure is associated with fluid overload and systemic venous
congestion. Nevertheless, rapid fluid removal with diuretic
therapy or haemofiltration has to be used with caution because
RV unloading can induce low cardiac output.

‘Specific’ pharmacotherapy to decrease RV afterload
Prostanoids and nitric oxide
PH targeted therapy improves pulmonary blood flow and decreases RV afterload. Intravenous prostanoids (epoprostenol,
iloprost, treprostinil) should be considered in the critically ill
patient, if a severe ventilation-perfusion mismatch has been

excluded. Since prostanoids lower SVR, concomitant systemic vasopressor therapy may become necessary.

Inhalative therapy with prostanoids or inhaled nitric oxide (iNO) has less impact on SVR and should be considered early, especially if the systemic blood pressure is low. In addition, inhaled or aerosolised application does not worsen the ventilation-perfusion mismatch in the same way as the intravenous route may do. INO is the first choice in mechanically ventilated patients. After cardiopulmonary bypass, iNO reduces PVR and may lower the risk of PH crisis and shorten the postoperative course. However, in a recent meta-analysis of iNO in patients with CHD, its effect on survival and several haemodynamic secondary end points has been questioned. Inhaled iloprost has been proposed as an alternative to iNO with comparable effects on PVR, but well designed prospective clinical trials are lacking. Acute haemodynamic effects of other inhaled drugs, such as nitroglycerine or milrinone, have been described in small case series, but their usefulness is not well established.

PDE5-inhibitors (sildenafil)
Oral sildenafil is reasonable to facilitate weaning from iNO. However, in the intensive care setting oral medications have the

Figure 2  Algorithm for management of acute pulmonary hypertension in the ICU. CPP, coronary perfusion pressure; CPR, cardiopulmonary resuscitation; CVP, central venous pressure; FBC, full blood count; ICU, intensive care unit; LFT, lung function test; / indicates "or"; SBP, systemic blood pressure; SvO₂, systemic venous oxygen saturation, Lact: lactate, SVR, systemic vascular resistance; TR, tricuspid regurgitation; VA-ECMO, venoarterial extracorporeal membrane oxygenation.

disadvantage of unpredictable absorption. Prophylactic use of sildenafil before surgical correction of CHD has been proven to be useful. The administration of intravenous sildenafil has been described but its effectiveness has not been clearly determined. Systemic arterial hypotension and impairment of oxygenation have been described as adverse events.

Pharmacotherapy to increase myocardial perfusion and/or counteract right-to-left interventricular septal shift (LV compression and low LV output)

Inotropes
In the case of severe RV failure in PH, inotropic support may be necessary. Milrinone and levosimendan can be useful as positive inotropic drugs since they have little or no effect on heart rate but lower PVR. Single centre studies have shown positive effects of levosimendan on PVR in children after cardiac surgery. Levosimendan’s major effect is thought to be improvement of myocardial contractility. Additionally, there is a vasodilator effect due to inhibition of PDE3 activity. Dobutamine and epinephrine improve RV contractility but may induce tachycardia which impairs diastolic filling and coronary perfusion subsequently lowering cardiac output.

Non-pharmacological therapy
Surgical or interventional intra-atrial communication
On the basis of clinical experience in adults, patients with systemic hypertensive pulmonary hypertension, and to treat low SVR caused by PH targeted therapy. Elevation of systemic blood pressure may become necessary to maintain coronary perfusion pressure and to reduce leftward septal shift. Vasopressin and terlipressin have been shown in small case series to lower PVR while increasing SVR. Patients on vasopressor therapy should be monitored closely for adequacy of cardiac output.

Vasopressors
Vasopressor therapy with norepinephrine, vasopressin or terlipressin may be indicated to induce reshifting of the interventricular septum from left to right, to improve tissue perfusion in patients with systemic hypertension, and to treat low SVR caused by PH targeted therapy. Elevation of systemic blood pressure may become necessary to maintain coronary perfusion pressure and to reduce leftward septal shift. Vasopressin and terlipressin have been shown in small case series to lower PVR while increasing SVR. Patients on vasopressor therapy should be monitored closely for adequacy of cardiac output.

Potts shunt
As an alternative to lung transplantation, creation of an aorto-pulmonary shunt (Potts shunt) in children with suprasystemic idiopathic PH has been described in a small case series. Although these first long-term results are promising, no recommendation can be made because of the limited experience at this stage.

Extracorporeal membrane oxygenation
Depending on RV function veno-venous (VV) or venoarterial (VA) extracorporeal membrane oxygenation (ECMO) may be considered as a bridge to recovery or bridge to transplantation. In patients with large atrial communications, circulatory support with VV-ECMO may be feasible even in the failing RV by providing oxygenated blood shunting right-to-left through the defect. The indication for mechanical support with ECMO depends on aetiology of RV or lung failure. Its short-term use in post cardiac surgery PH is generally accepted. However, longer support times had been described with new therapeutic strategies (e.g. awake-ECMO) making it feasible as bridge to transplantation in selected patients with other aetiologies. Timing of mechanical support in children is less well established compared with adults. The Interagency Registry for Mechanically Assisted Circulatory Support has defined clinical profiles of patients failing on optimal therapy for heart failure. Levels 1 and 2 are generally accepted indications for mechanical support in acute heart failure. In children after cardiac surgery, implantation of ECMO with metabolic acidosis or during cardiopulmonary resuscitation (CPR) is a risk factor for increased mortality and brain injury.

PH crisis after cardiac surgery may cause circulatory collapse requiring CPR. If spontaneous return of circulation cannot be achieved by conventional CPR measures, VA-ECMO implanted during CPR is an option (reversibility of disease). Overall survival in VA-ECMO implanted during CPR is 38%, with better outcome in the absence of severe metabolic acidosis before support; 11% of survivors suffered from cerebral seizures and 6% had evidence of brain injury in a CT scan.

In patients with idiopathic PH who deteriorate rapidly due to progressive RV failure, eligibility for transplantation has to be evaluated. If lung transplantation is deemed feasible, ECMO should be considered as bridging treatment. Acute decompression in chronically ill children due to reversible disease (eg, in chronic lung disease and pneumonia) or in patients in whom targeted therapy for PH is suboptimal may require a bridge to recovery strategy. Although described in the literature, implantation of ECMO in children with idiopathic PH is controversial due to its irreversible nature and in consideration of the shortage of donor organs. If cannulated peripherally, patients can be awake and extubated, which allows longer support times and reduces the risks of ventilator associated complications. In severe RV compromise, VA-ECMO is needed to guarantee adequate cardiac output. Alternatively, a pumpless paracorporeal lung assist that is connected to the pulmonary artery and left atrium after sternotomy may be indicated in patients with hypoxaemia but sustained RV and LV functions.

This device decompresses the RV sufficiently, provides oxygenation and removes carbon dioxide.

With normalisation of cardiac output after lung transplantation the chronically unloaded LV may develop substantially elevated filling pressures leading to pulmonary oedema and lung failure. Successful, scheduled postoperative VA-ECMO as a bridging strategy to LV reverse remodelling and recovery has been described in adults and children.

Ventricular assist device
Ventricular assist device treatment has not been proven to be effective in children with RV failure due to PH with preserved LV function. Pulsatile devices may lead to pulmonary haemorrhage due to high pulse pressures; while afterload dependent continuous flow devices may not provide sufficient pump flow in patients with high PVR. Therefore ventricular assist device implantation in RV failure due to PH with normal LV filling pressures is not recommended.

Lung transplantation
Bilateral lung transplantation should be considered in children with inadequate clinical response on maximal combination therapy who remain in functional class III or IV. Due to long waiting times and influence on outcome, transplantation should be considered before cardiopulmonary decompression has occurred. Mortality of children who are mechanically ventilated before transplantation is significantly increased (HR 2.6, CI
1.72 to 4.07). The median survival rate after lung transplantation in children is between 5.6 years and 6.1 years.

**Therapeutic strategies in different clinical scenarios**

**CPR of children with acute PH crisis and cardiopulmonary arrest**

Resuscitating a patient with severely elevated PVR, acute heart failure and cardiopulmonary arrest is particularly difficult (figures 2 and 3). Resuscitation is initiated and conducted following published guidelines. From a pathophysiological perspective, it is important to proceed with tailored, imminent treatment as mentioned above. Special attention should be given to coronary perfusion (pressure, flow), especially in the setting of severe RV hypertrophy, elevated RV filling pressures and/or tachycardia. RV hypertrophy, decreased aortic to RV end diastolic pressure gradient, and shortening of diastole contribute to RV myocardial ischaemia, RV dilation and subsequent LV compression, causing a rise in LV filling pressures and—together with low cardiac output and coronary perfusion.

Figure 3  BAS, balloon atrioseptostomy; Clinical algorithm for the management of the scenarios for acute PH in the ICU. CHD, congenital heart disease; E-CPR, extracorporeal membrane oxygenation with cardiopulmonary resuscitation; ICU, intensive care unit; IPAH, idiopathic pulmonary arterial hypertension; PDE5, phosphodiesterase 5; PH, pulmonary hypertension; PHC, pulmonary hypertensive crisis; SaO2, arterial oxygen saturation; VA-ECMO, venoarterial extracorporeal membrane oxygenation; VAD, ventricular assist device.
pressure—to additional LV myocardial ischaemia. Studies comparing outcome of standard CPR and CPR plus targeted therapy are lacking. In appropriate patients with acute PH crisis, early implementation of mechanical support (VA-ECMO) should be considered. Clinical data on the neurological outcome of children with pulmonary hypertension, who acutely arrested, are lacking.

Therapy of acute postoperative pulmonary hypertensive crisis (PHC)

Acute PHC is a life-threatening emergency and has to be treated aggressively since the risk of cardiocirculatory collapse necessitating cardiopulmonary resuscitation is high. Providing normal ventilation and normalising pCO2 (if necessary with a manual ventilation bag), boluses of norepinephrine, sedative drugs and muscle relaxants restore circulation in the majority of cases. Medication lowering RV afterload, augmenting RV preload and contractility and maintaining adequate coronary perfusion pressure (as described above) preserve LV function and oxygen delivery in patients at risk. Adequate postoperative analgesia and sedation, minimal handling and minimising tracheal suctioning help to prevent acute PHC. Preoperative administration of PDE5 inhibitors such as sildenafil has been tried, but due to the paucity of clinical data, no corresponding recommendation can be made. VA-ECMO should be considered early on when pharmacological and ventilator strategies fail.

Patients with univentricular heart and partial (‘Glenn’) or total cavopulmonary anastomosis/’Fontan-like’ circulation (ie, no subpulmonary ventricle).

Pulmonary vessels of newborns are highly reactive to stimuli which elevate PVR. In children with univentricular heart after Glenn operation, an increase of PVR may provoke deep cyanosis, stasis of blood in the shunt and shunt thrombosis. Management of suspected shunt thrombosis must be aggressive. No clinical studies comparing different treatment strategies of acute shunt thrombosis are available. Management generally includes: optimisation of ventilation, sedation, muscle relaxation if necessary, iNO, bolus of heparin and prompt treatment of arterial hypotension (bolus of volume if filling pressures are low; norepinephrine to increase afterload, epinephrine if systolic function is impaired). If immediate recovery of arterial oxygen saturations fails to occur, opening of the chest to massage the shunt or implantation of VA-ECMO may become necessary.

In patients with univentricular physiology and total cavopulmonary connection, a low PVR is a prerequisite for adequate ventricular filling and output. iNO,52 milrinone53 and ventilation with low mean airway pressure or better spontaneous breathing54 all contribute to improved outcomes in the early postoperative course. Negative pressure ventilation has been described in the failing Fontan circulation.55 In children who are mechanically ventilated long term, mild permissive hypercapnia may allow lower ventilator settings without elevation of PVR as long as there is no respiratory acidosis (figure 3).

Therapy of PH in a child with acute deterioration of previously known chronic PH

Initial assessment, oxygen supplementation and establishment of monitoring are the first steps in a child with acute deterioration of chronic PH. Decisions on how to treat the child are based on an individualised strategy since scientific evidence is insufficient. As long as the child’s condition is not substantially improved (ie, normalisation of lactate, blood gases, vital parameters, urine output) frequent re-evaluations are mandatory. Elevated or increasing lactate levels, development of metabolic or respiratory acidosis, hypotension or other signs of low cardiac output should prompt escalation of therapy including anaesthesia and mechanical ventilation (see Anaesthesia and ventilation) or—if indicated—mechanical support (see Extracorporeal membrane oxygenation). Usually, escalation of specific medical treatment to decrease RV afterload in a child that is not responding to current drugs, means adding a second or third drug rather than exchanging the substances.56 Choice of treatment options is guided by the clinical condition. Initiation of therapy in a spontaneously breathing child with an inhaled drug has the advantage that no venous access is needed and it does not increase ventilation-perfusion mismatch. With respiratory fatigue or in small children, inhalation may become impossible. Intravenous prostanoids should be considered in the critically ill child; blood pressure monitoring is mandatory. If a child is mechanically ventilated iNO may be strongly considered. Clinicians may decide to add sildenafil orally or intravenously, especially in the child who is not responding to therapy; due to the paucity of data, no recommendation can be made in this regard. Inotropic and vasopressor therapy is adapted to haemodynamic parameters. Chronic specific therapy should be continued. If liver function tests are above three times the upper limits, bosentan must be withheld.

Therapy of PH in a child with acute deterioration due to new diagnosis of PH

The principles of treatment are comparable to those described above (see Therapy of PH in a child with acute deterioration due to previously known chronic PH). The clinical condition of a critically ill child initially often precludes a complete diagnostic workup. However, diagnoses and conditions which change the therapeutic regimen such as acute pulmonary embolism must not be missed.

Conclusions

Acute PH is a serious complication in children at risk, including those after surgery for CHD, and has major impact on clinical outcome in patients with and without chronic PH. Treatment in the ICU should be based on the underlying pathophysiology but ultimately needs to focus on the basic goals of lowering RV afterload and augmenting RV preload and contractility. Furthermore, maintaining adequate coronary perfusion pressure and flow will help to preserve myocardial oxygen and energy supply, and thus RV and LV systolic function and oxygen delivery. Early recognition of patients at particular risk, and timely establishment of efficient therapeutic actions may prevent the development of severe cardiac dysfunction, low cardiac output and death.

Author affiliations

1Department of Paediatric Cardiology, University Children’s Hospital Ulm, Ulm, Germany
2Paediatric Heart Centre, University Hospital of Giessen and Marburg, Giessen, Germany
3Department of Cardiothoracic, Transplantation and Vascular Surgery, Hannover Medical School, Hannover, Germany
4German Centre for Lung Research, BREATH, Hannover, Germany
5Department of Paediatric Cardiology and Critical Care, Hannover Medical School, Hannover, Germany
6Department of Congenital Heart Disease and Paediatric Cardiology, Deutsches Herzzentrum Berlin, Berlin, Germany

Correction notice

In table 2 the Sildenafil intravenous dosage has been updated from 0.4mg to 0.4mg/kg since this paper was first published online.
Pulmonary vascular disease

Funding CA currently receives grant funding from Stiftung KinderZentrum. GW currently receives grant support from the German Research Foundation (DFG; SFB 738). GH currently receives grant support from the German Research Foundation (DFG; HA 4348/2-1), Fördereingangsdeutsche Kinderherzerzten (WH-001/2014) and Stiftung KinderZentrum (2511-6-13). This supplement was produced with support from an unrestricted educational grant from Actelion Pharmaceuticals Germany GmbH, Bayer Pharma AG, and Pfizer Inc. None of these organisations had any influence on the composition of the writing group or the content of the articles published in this supplement. Open Access publication of this article was sponsored by Actelion Pharmaceuticals Germany GmbH.

Competing interests None declared.

Provenance and peer review Commissioned; externally peer reviewed.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) licence, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/


