Pulmonary manifestations of sickle cell disease
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Pulmonary complications account for significant morbidity and mortality in patients with sickle cell disease. Clinical lung involvement manifests in two major forms: the acute chest syndrome and sickle cell chronic lung disease. Acute chest syndrome is characterised by fever, chest pain, and appearance of a new infiltrate on chest radiograph. Sickle cell chronic lung disease, on the other hand, manifests as radiographic interstitial abnormalities, impaired pulmonary function, and, in its most severe form, by the evidence of pulmonary hypertension. Progress has been made in understanding the pathophysiology and management of these complications. In this review the current knowledge of the mechanism, diagnosis, and treatment of pulmonary complications of sickle cell disease are discussed.

Sickle cell disease (SCD) is one of the most prevalent genetic disorders. There are more than 200 million carriers of sickle cell trait worldwide, and 200,000 to 300,000 people are born annually with major haemoglobinopathies. Approximately 0.14% of African Americans children are homozygous for the sickle cell gene and 8% have sickle cell trait, making SCD the most prevalent inherited disorder in African Americans. The protein manifestations of SCD are caused by substitution of glutamic acid by valine in the beta subunits of the haemoglobin molecule. Upon exposure to low oxygen tension the mutant haemoglobin S becomes less soluble and aggregates into large polymers. This results in a distorted erythrocyte with marked decrease in its deformability contributing to the vaso-occlusive and haemolytic aspects of the disease. With improved supportive care the median age of survival has risen to 42 years for men and 48 years for women. As survival into adulthood has become more common in patients with SCD, there has been an increased incidence of chronic organ failure. The lung is among the major organs involved in SCD. The pulmonary manifestations of SCD are both acute and chronic (see box 1). They remain the leading cause of morbidity and mortality in patients with SCD. Clinical lung involvement commonly takes two major forms: the acute chest syndrome (ACS) and sickle cell chronic lung disease (SCCLD). More than 20% of adults with SCD have fatal pulmonary complications. Although pulmonary manifestations of SCD are common, they remain under-diagnosed by physicians. This review summarises the current knowledge and management of pulmonary manifestations associated with SCD.

AIRWAY HYPER-REACTIVITY
Airway hyper-reactivity is a common pulmonary function test abnormality among young subjects with SCD. The reported incidence of obstructive lung defects in children with SCD is 35% to 37%. A prospective controlled trial demonstrated airway reactivity to cold air in 60% patients with SCD compared with none in the control population, suggesting an association between SCD and airway reactivity. The pathogenesis of increased airways reactivity and its relationship to ACS and SCCLD, however, is unknown. In the absence of data on long term benefit, routine use of bronchodilators is not recommended in patients with SCD.

NOCTURNAL OXYHAEMOGLOBIN DESATURATION
Nocturnal oxyhaemoglobin desaturation has been reported in SCD subjects, with a prevalence of up to 40% in children and adolescents. Proposed mechanisms are obstructive sleep apnoea, intrinsic lung disease, and an abnormality in oxyhaemoglobin affinity. Several investigators documented obstructive sleep apnoea in SCD with or without nocturnal oxyhaemoglobin desaturation. Although it has been shown that low nocturnal oxygen saturation is associated with higher rate of painful crisis in childhood, the relationship between nocturnal oxyhaemoglobin desaturation and vaso-occlusive crisis or ACS is poorly understood.

THROMBOEMBOLISM
Patients with SCD are known to be hypercoagulable. A variety of mechanisms are postulated from low levels of protein S and C, fibrinogen, and factor V Leiden.

Box 1: Pulmonary manifestations of sickle cell disease
- Increased airway reactivity.
- Nocturnal oxyhaemoglobin desaturation.
- Thromboembolism.
- Acute chest syndrome.
- SCCLD: radiographic interstitial lung abnormalities; pulmonary function test abnormalities; pulmonary hypertension.

Abbreviations: ACS, acute chest syndrome; MACSS, multicentre acute chest syndrome study; NO, nitric oxide; NYHA, New York Heart Association; SCCLD, sickle cell chronic lung disease; SCD, sickle cell disease; VCAM-1, vascular cell adhesion molecule-1.
Figure 1 Chest radiograph of a 34 year old man admitted with ACS showing infiltration of right lower and middle lobes. Patient improved after receiving antibiotic therapy and blood transfusion.

Box 2: Risk factors associated with acute chest syndrome
- Younger age.
- Homozygous haemoglobin SS.
- Low haemoglobin F concentration.
- High steady state leucocyte counts.
- High haemoglobin levels.
- Previous history of acute chest syndrome.
- Avascular necrosis of bone.

Infection
Patients with SCD are at an increased risk of infection due to abnormalities in host defences including impairment of the complement system and functional asplenia. In the MACSS study, infection was documented in 30% of cases with ACS, and 27 different pathogens were identified. *Chlamydia pneumoniae* was the most frequent pathogen followed by *Mycoplasma pneumoniae* and respiratory syncytial virus.

Pulmonary fat embolism
The occurrence of pulmonary emboli containing fat and necrotic bone marrow in patients with SCD is well known. Fat embolism associated ACS was identified in 9% cases of ACS in the MACSS report. The pulmonary signs and symptoms typically are preceded by bone pain, with laboratory evidence of a significant decrease in haemoglobin and platelet count and an increased plasma level of free fatty acids and phospholipase A2. The diagnosis of pulmonary fat embolism is supported by the presence of lipid-laden macrophages in the bronchoalveolar lavage fluid. The Corwin index is used to quantify the amount of lipid in the pulmonary macrophages. ACS due to pulmonary fat embolus is associated with severe haematological and clinical abnormalities and has a fourfold higher mortality rate in adults compared to children.

In situ thrombosis
Pulmonary infarction secondary to in situ thrombosis is a potential cause of ACS, though is uncommonly documented pathologically.

Hypoventilation from pain and/or narcotic analgesics
There is a high correlation between thoracic bone infarction on bone scan and the presence of pulmonary infiltrates. It has been suggested that bone pain can cause splinting and atelectasis and can present as ACS. Similarly, postoperative thoracic and upper abdominal pain may predispose to ACS. Incentive spirometry and adequate pain control therefore becomes necessary under such situations. On the other hand, excessive use of narcotics may lead to hypoventilation and ACS as a complication of their use.

Iatrogenic pulmonary oedema
Although it has been suggested that aggressive hydration may lead to ACS in patients with SCD, this association is not established.

Pathogenesis
The mechanism responsible for ACS is not fully understood. It is believed to be a specific form of acute lung injury that can progress to acute respiratory distress syndrome. The lung...
injury is caused by various insults superimposed on the genetically based pathophysiology of SCD. In addition to haemoglobin S polymerisation and red cell sickling, increased expression of adhesion molecules on sickle erythrocytes and endothelium, release of inflammatory mediators, interaction of sickle red cell with leucocytes, microvascular thrombosis, and endothelial damage may all contribute to microvasculature occlusion and tissue infarction.

More recent data suggest that abnormalities in endothelial cell nitric oxide (NO) production and metabolism as well as oxidant status may contribute to the development of ACS. NO regulates vascular tone and endothelial function, and maintains tissue oxygenation by reducing shunt physiology. Hypoxia induces cytokine and endothelin-1 release which up-regulates expression of the endothelial adhesion receptor vascular cell adhesion molecule-1 (VCAM-1), thereby enhancing adhesion of red cells to endothelial cells. Additionally, fat embolism and release of free fatty acids also enhances the adhesion process by VCAM-1 up-regulation. Under normal conditions NO inhibits both endothelin-1 production and endothelial VCAM-1 expression. During ACS, however, hypoxia and red cell sickling inhibit NO production while free radical species released by leucocytes and endothelium, contribute to microvasculature occlusion and tissue infarction.

Clinical features
The Cooperative Study of Sickle Cell Disease, which prospectively followed up 3751 patients, reported data on 1722 ACS episodes in 939 patients. Patients with ACS presented with fever (80%), cough (74%), chest pain (57%), dyspnoea (28%), productive cough (24%), wheezing (11%), and haemoptysis (2%) accompanied by hypoxia, leucocytosis, and infiltrates on chest radiographs that often progressed to multilobar pulmonary disease indistinguishable from acute respiratory distress syndrome.

The symptoms of ACS are age dependent; ACS is often preceded by febrile episodes in children and by vaso-occlusive crisis in adults. Children are more likely to have fever and cough, whereas adults more commonly experience chest pain and dyspnoea.

Management of acute chest syndrome
General measures
SCD patients admitted for painful crisis should be considered in the prodromal phase of ACS. Prophylactic manoeuvres like close pulmonary monitoring, cautious hydration, optimal pain control, and incentive spirometry remain essential components of therapy. Physicians should have a low threshold for ordering a chest radiograph and arterial blood gas analysis. Vigorous hydration may result in pulmonary oedema and worsening respiratory distress and should be avoided. Judicious use of analgesics is necessary to relieve bone pain and splinting to prevent lung atelectasis. If there is no contraindication, non-steroidal agents should be the part of all pain control regimens as they have a narcotic-sparing effect. Efficacy of incentive spirometry in preventing the pulmonary complications (atelectasis and infiltrates) associated with the ACS has been proven in a randomised controlled trial and should be routinely used in patients with SCD.

Oxygen therapy
Oxygen should be administered for the correction of hypoxia, which if untreated creates a risk of multiorgan failure. Pulse oximeter correlation with arterial oxygen tension in patients with SCD may at times be poor and arterial blood gas confirmation is required in hypoxic patients.

Box 3: Diagnostic testing and laboratory monitoring in acute chest syndrome
- Blood and sputum/tracheal secretion culture.
- Daily blood counts and appropriate metabolic profile.
- Serial measurement of arterial blood gases as necessary.
- Chest radiographs.
- Flexible bronchoscopy with bronchoalveolar lavage as appropriate.

Antibiotics
Broad spectrum antibiotics, including a macrolide or quinolone in view of atypical infections such as chlamydia and mycoplasma, should be administered. Penicillin prophylaxis and use of Haemophilus influenzae vaccine has reduced bacterial infection significantly and should be given to all patients with SCD.

Bronchodilators
Bronchodilators are given to patients when airflow obstruction is present, though some investigators recommend their routine use in all patients. Many patients with SCD have airway hyper-reactivity and this fact should be considered when managing these patients on the ventilator to prevent autotpeous end expiratory pressure and dynamic hyperinflation.

Bronchoscopy
Bronchoscopy may be considered in patients not responding to initial therapy. It can be utilised both as an airway clearance technique by removing the thick and tenacious airway secretions often found in patients with ACS and for more accurate diagnosis of the episode.

Transfusion
Simple and exchange blood transfusion may be beneficial by lowering the fraction of sickle haemoglobin and also by improving the oxygen carrying capacity of blood. Both simple and exchange transfusion have resulted in similar improvement in oxygenation in patients with ACS. Transfusion is indicated in individuals with severe disease, multilobe involvement, persistent or worsening hypoxaemia, neurological abnormalities, multiorgan failure, or those with a history of cardiac disease. In patients with severe anaemia, simple transfusion appears to be the modality of choice. However, a post-transfusion increment in haemoglobin levels above 110 g/l may be dangerous due to increased blood viscosity and therefore in individuals with a relatively high haemoglobin exchange transfusion should be performed. Severe episodes may not respond to transfusion therapy and may require support with mechanical ventilation or extracorporeal membrane oxygenation. Efficacy of chronic transfusion has been shown in a randomised controlled trial in the prevention of future episodes of ACS.

Corticosteroids
A short course of corticosteroids has been shown to reduce the need for blood transfusion and the length of hospitalisation in children with ACS. However, its use in vaso-occlusive pain crisis is associated with more rebound attack. Corticosteroids therapy, therefore, should be considered experimental in ACS.

Nitric oxide
NO may be beneficial in severe cases that are refractory to standard therapy. There are isolated reports of significant reduction in pulmonary artery pressure, pulmonary vascular resistance, and alveolar-arterial gradient with concomitant improvement in cardiac output in patients with ACS after the inhalation of NO.
SICKLE CELL CHRONIC LUNG DISEASE

The exact incidence, prevalence, natural history, and methods of diagnosis of SCCLD have not been established due to the lack of detailed epidemiological studies. It is suggested that SCCLD has a prevalence of approximately 4% in patients with SCD.10–12 SCCLD is presumably related to recurring episodes of infarction and infection and is characterised by a decrease in radiolucency of the lungs, moderate to severe impairment of pulmonary function, and in its most severe form by evidence of pulmonary hypertension.1,10–12

Radiographic abnormalities

Significant radiographic interstitial lung disease has been seen in patients with SCD.13 In a prospective study, 41% patients with SCD who had at least one prior episode of ACS were found to have significant multifocal interstitial lung abnormalities on thin section computed tomography scans of the chest.14 A correlation was found between the severity and extent of interstitial abnormalities on computed tomography and the number of prior episodes of ACS.15

Pulmonary function abnormalities in sickle cell disease

Pulmonary function abnormalities in SCD are frequent and are characterised by airway obstruction, restrictive lung disease, abnormal diffusion capacity, and hypoxaemia.16–20 However, a restrictive airways abnormality is typically seen in patients with SCCLD.16–20

Pulmonary hypertension

Prevalence

Secondary pulmonary hypertension has become an increasingly recognised complication in patients with SCD. It is associated with impaired exercise tolerance, progressive heart failure, and a high mortality.21,22–24 Retrospective studies of echocardiograms of patients with SCD suggested a prevalence of pulmonary hypertension from 30% to 56%.25–27 A recent prospective study based on echocardiograms in 154 patients with SCD, however, revealed a prevalence of 34% and approximately 27% of these patients found to have severe pulmonary hypertension.28

Box 4: Treatment of acute chest syndrome

- Identify and treat all underlying precipitating factors.
- Supplemental oxygen to treat hypoxia and maintain arterial oxygen tension above 70 mm Hg.
- Optimal pain control and fluid management.
- Ongoing respiratory therapy by using incentive spirometry and chest physical therapy as necessary.
- Provide empirical antibiotic with coverage for atypical organisms.
- Bronchodilators for patients with reactive airway disease.
- Simple or exchange transfusion to reduce haemoglobin S concentration and to enhance oxygen carrying capacity.
- Miscellaneous: NO inhalation, systemic steroids, mechanical ventilation, and extracorporeal membrane oxygenation.

Hydroxyurea

The incidence of ACS is significantly reduced in adult patients treated with hydroxyurea.29–31 A double blind, placebo controlled trial of 299 adults with SCD was terminated early, because hydroxyurea reduced the frequency of ACS, hospitalisation, and need for transfusion.32 Hydroxyurea reduces sickling by increasing fetal haemoglobin level and hence decreasing the relative concentration of haemoglobin S in the erythrocytes.33 It should, therefore, be considered in patients with recurrent episodes of ACS.

Right sided cardiac catheterisation is the gold standard test for the diagnosis of pulmonary hypertension.34–36 Haemodynamic studies of patients with cardiopulmonary symptoms by right sided heart catheterisation have also demonstrated a raised pulmonary systolic and diastolic pressure.37–39 Although SCD patients with echocardiographic evidence of pulmonary hypertension may not have the symptoms of cardiac dysfunction, mortality is significantly increased in these patients compared with patients without pulmonary hypertension.40–42

Pathophysiology

Pulmonary hypertension is characterised by progressive obliteration of the pulmonary vasculature. The mechanism of pulmonary hypertension complicating SCD is unknown and likely to be multifactorial. Possible causes include: sickle cell related vasculopathy due to sequestration of sickle erythrocytes, fat embolism, or recurrent infection;43–44; chronic hypoxic stress causing irreversible remodelling of the vasculature with smooth muscle proliferation and fibrosis;45,46; recurrent pulmonary thromboembolism;47,48; increased blood viscosity with consequent right ventricular volume or pressure overload; and pulmonary scarring from repeated episodes of ACS.49–51 Regardless of the pathophysiology of pulmonary hypertension, once it becomes established, the patients are at risk for right sided heart failure.

Clinical features

Patients tend to be asymptomatic in the early stage, though with moderate to severe pulmonary hypertension they experience chest pain, dyspnoea, and hypoxaemia at rest. With disease progression they are at risk for right heart failure and sudden death from pulmonary thromboembolism, systemic hypotension, and cardiac arrhythmia.52

Management

Limited data are available in the literature regarding management of pulmonary hypertension related to SCD. The efficacy of continuous intravenous epoprostenol (prostacyclin) and calcium channel blockers, which are effective therapy in primary pulmonary hypertension, are of unknown efficacy in pulmonary hypertension secondary to SCD.53–56 Patients with significant parenchymal lung disease may develop considerable shunt and increased oxygen requirement while receiving epoprostenol.57,58 Likewise, systemic blood pressure is lower than normal in patients with SCD and it is not known whether SCD patients can tolerate vasodilators.59 Various other agents including prostaglandin analogues such as subcutaneous
treprostinil, inhaled iloprost, and oral beraprost as well as oral bosentan, an endothelin receptor antagonist, have shown efficacy in both primary and secondary pulmonary hypertension in randomised controlled trials. However, the therapeutic role of these agents in pulmonary hypertension related to SCD is not known. Recent clinical and experimental data have suggested that altered metabolism of NO may play a part in the pathogenesis of pulmonary hypertension. Therapeutic NO inhalation may prove efficacious in patients with SCD and secondary pulmonary hypertension due to its ability to selectively dilate the pulmonary vasculature which reduces pulmonary pressure and increases oxygenation. Studies are underway to determine if inhaled NO will reduce morbidity and mortality in SCD.

In addition to vasodilators, a number of adjunctive therapies can be useful in pulmonary hypertension secondary to SCD. Since pulmonary hypertension may be related to polymerisation of haemoglobin S, hydroxyurea treatment should be considered. Furthermore, hydroxyurea decreases the incidence of ACS, multiple episodes of which may contribute to pulmonary hypertension and may also have a protective effect on the pulmonary vasculature. Long term oxygen therapy should be considered if resting hypoxaemia is present. Results with simple or exchange transfusion are disappointing. However, the combination of exchange transfusion with long term oxygen therapy may be beneficial in pulmonary hypertension. A retrospective and a non-randomised prospective study suggested that anticoagulation increases survival in patients with primary pulmonary hypertension. Chronic anticoagulation therapy is, therefore, recommended in SCD with pulmonary hypertension unless a contraindication is present. Caution is required in the use of diuretics and cardiac glycosides may be helpful in relieving symptoms in patients with right heart failure. Pulmonary thromboendarterectomy in surgically accessible chronic thromboembolic disease with pulmonary hypertension has been performed successfully in patients with SCD and should be considered in patients with chronic macrovascular occlusion of pulmonary arteries. The role of atrial septostomy or single or bilateral atrial septostomy may be considered. Lung transplantation in patients with pulmonary hypertension has been performed successfully in patients with SCD and should be considered in cases of deterioration or more advanced disease, intravenous epoprostenol should be started and atrial septostomy may be considered. Lung transplantation is less likely to be beneficial in patients with SCD due to an expected high procedure related mortality in these patients.

CONCLUSION
Pulmonary complications are common and account for a large proportion of deaths among adults with sickle cell disease. They result from the complex pathophysiology of sickle cell disease. New therapies are being developed that may prove to be beneficial in the various complications. Clinicians should be aware of acute and chronic complications of sickle cell disease, as they have emerged as major threats to the health and longevity of patients with this condition.

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Sickle cell disease


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