Advanced sickle cell associated interstitial lung disease presenting with cor pulmonale in a Nigerian

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INTRODUCTION

The chronic complications of sickle cell disease (SCD) are usually collectively referred to as sickle cell chronic lung disease (SCLD). It is defined by radiologic and clinical features of ventilatory dysfunction and pulmonary hypertension, which may later progress to cor pulmonale.1 SCLD was first reported as a cause of cor pulmonale by Yater and Hansmann in 1936.2 The precise prevalence remains largely unknown due to lack of detailed epidemiological studies. A prevalence rate of 4% was reported by Powars et al.3 We reported a prevalence rate of 18.4% among young adult SCD patients in our center.4 Previous studies have reported abnormal pulmonary function among Nigerians with SCD5,6 but we are not aware of any previous report with computed tomography (CT) evidence of extensive interstitial lung disease. We report a SCD patient who presented with CT proven interstitial lung disease complicated by pulmonary hypertension and cor pulmonale.

CASE REPORT

Mrs. A.O., a 40-year-old known SCD patient with hemoglobin SS genotype was referred to the pulmonary division on account of dry cough of 3 weeks, difficulty in breathing of 2 weeks, and bilateral leg swelling of 4 days. There was low-grade fever, no weight loss, and no contact with chronic cough index. Difficulty in breathing initially occurred on exertion but later occurred even at rest. There was orthopnea but no paroxysmal nocturnal dyspnea. There was bilateral leg swelling, which worsened progressively. No swelling in other parts of the body. No reduction in her urine volume. She was not a diagnosed asthmatic, hypertensive, or diabetic and she was not previously treated for tuberculosis or any other chronic lung diseases. She was only admitted for childbirth in her adulthood years. She had a single episode of blood transfusion when she was 6 years old. Afterwards, she has been crisis free for more than 20 years. She neither smokes cigarettes nor consumes alcoholic drinks. No history of exposure to cigarette smoke from others. She is married in a monogamous family with two girls and one boy. Her last childbirth was 15 years prior to presentation. She works as a secretary in the registration office of a state polytechnic. No history of exposure to occupational dust or fumes. No history of chronic drug use and no history of allergy.

On general examination, she was a middle aged looking woman, conscious, dyspneic with central cyanosis, afebrile (Temperature=36.5), not pale, anicteric, not dehydrated, grade 3 digital clubbing, bilateral pitting pedal edema up to the lower third of the leg, weight=84 kg, height=1.65 m, and BMI=30.9 kg/m². Oxygen saturation in room air was 78% and this increased to 92% on 5 L/min of oxygen via nasal prong.

Respiratory examination revealed that she was in respiratory distress as evidenced by respiratory rate of 42/min, flaring of alae nasi, and use of accessory muscles of respiration, and the trachea was central with symmetrical chest movement. Percussion note was resonant with coarse
crepitations over the entire right lung and posterior lower zone of the left lung.

Cardiovascular examination: pulse=100 beats/min, regular with premature beats, full volume; arterial wall was not thickened. No locomotor brachialis. BP was 105/60 mmHg. JVP was raised to the angle of the jaw. She had a right parasternal heave and apex beat was in the 5th left intercostal just lateral to the mid-clavicular line. Heart sounds were S1, S2, S3 gallop rhythm with loud pulmonary component of second heart sound.

Abdomen – full, moves with respiration, liver was 4 cm below the costal margin, mildly tender, smooth surface, and span of 16 cm. Spleen and kidneys were not enlarged. No demonstrable ascites.

Other systems were within normal limits.

**Investigations**

Her packed cell volume was 32% with a white blood cell count of 13.5 ×10^9/L (neutrophil=52%, lymphocyte=38%, other=9%) and erythrocyte sedimentation rate of 3 mm/h. Sputum examination for acid and alcohol fast bacilli was negative in three samples; sputum culture did not yield any growth. Chest X-ray showed widespread reticulonodular opacities involving the whole of the right lung field and left upper zone with an area of hyperluscency in the left mid and lower zones. Electrocardiography revealed sinus tachycardia of 120 beats/min, P pulmonale, and right ventricular hypertrophy with ventricular extrasystole. Echocardiography demonstrated tricuspid regurgitation with pulmonary hypertension. Spirometry showed severe restrictive pattern {FVC=1.13 L (36% of predicted); FEV\(_1\)=0.84 L (35% of predicted) FEV\(_1\)/FVC ratio=74.5% (92% of predicted)}. Chest CT demonstrated diffuse ground glass opacity with thickening of interlobar septa in the entire right lung field as well as left apex and part of the left lower region. There were multiple bullae with reduced attenuation (suggestive of compensatory emphysema of the remaining lung tissue) in the left lung field, sparing the apex. The pulmonary trunk was prominent and there was cardiomegaly [Figures 1-3].

She was admitted and managed initially in the intensive care unit with intranasal oxygen, cardiopulmonary monitoring, tablet levofloxacin 500 mg daily, intravenous frusemide 60 mg 8 hourly, tablet spironolactone 25 mg twice daily, and tablet aspirin 75 mg daily.

The heart failure gradually improved but she continued to desaturate in room air. She was advised on domiciliary oxygen, which she initially rejected but later purchased a cylinder of oxygen (did not purchase oxygen concentrator because of poor power supply) and pulse oximeter. She was discharged home, on 1–3 L/min of domiciliary oxygen (to be titrated with the pulse oximeter readings), aspirin,
and oral diuretics, after 1 month of hospitalization. She was re-admitted 1 week later with productive cough and features of right heart failure. She was managed with oxygen, diuretics, antibiotics, and aspirin. She improved and was discharged after 3 weeks. She has remained relatively stable at subsequent clinic visits. Her clinic oxygen saturation was 91–94% (off oxygen) during clinic visits.

DISCUSSION

We report a sickle cell patient with CT evidence of extensive interstitial lung disease, abnormal oxygen saturation, and severe restrictive pattern on spirometry as well as clinical, electrocardiographic, and echocardiographic features of pulmonary hypertension and cor pulmonale. These features are in keeping with advanced SCLD in the patient despite being relatively crisis free and without sickle cell-related hospital admissions for over 20 years prior to presentation. This is likely due to chronic occult pulmonary insults by similar mechanisms to acute chest syndrome. The occult insults, which are too tiny to be discernible at the active stage, accumulate over the years and may later become visible radiologically. As far back as 1970, Lagundoye reported generalized loss of translucency of the lung fields on chest radiographs of stable patients with sickle cell anemia and related hemoglobinopathies in Nigeria. Our patient has generalized loss of translucency of the entire right lung field with some involvement of the left lung field. Similarly, the chest CT shows diffuse ground glass opacity with thickening of interlobar septa in the entire right lung field and some involvement of the left lung. There was no diffuse interstitial fibrosis. This is similar to the report of a study on CT scan findings of 29 patients who were 5 to 54 years old in which none of the patients had diffuse interstitial fibrosis or honeycombing. Even autopsy studies show little fibrosis.

Although SCD has been reported as a cause of pulmonary hypertension/cor pulmonale in Nigeria, we are not aware of any previous report associating it with CT proven interstitial lung disease in sickle cell patients. SCLD is a major cause of death in adult sickle cell patients. In a longitudinal study, the mean time to death after diagnosis of patients in advanced SCLD (Stage 3 to Stage 4) was 2.5 years. There are no data available to support any specific treatment of interstitial lung disease in sickle cell patients. Little is also known about the factors that determine the development and extent of progression of SCLD. Many of the recent studies on chronic pulmonary complications of SCD are focusing mainly on pulmonary hypertension in sickle cell patients. However, studies are needed to address the pathogenesis and treatment of the interstitial lung disease component of SCLD. While we await further studies and in the absence of proven treatments for SCLD, symptomatic treatment is indicated as we did in our patient.

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