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Predicting risk factors for thromboembolic complications in patients with sickle cell anaemia – lessons learned for prophylaxis

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Abstract

Objective: To assess the clinical and laboratory predictors of venous thromboembolism (VTE) in patients with sickle cell anaemia (SCA) and its relationship to morbidity and mortality.

Methods: This retrospective case–control study analysed data from patients with SCA that experienced VTE compared with matched control patients with SCA but no VTE (2:1 ratio).

Results: A total of 102 patients with SCA were enrolled (68 cases with VTE and 34 controls). Amongst the 68 cases (median age, 29.5 years), 26 (38.2%) presented with isolated pulmonary embolism (PE). A higher prevalence of splenectomy (73.5% versus 35.3%) was observed in the cases compared with the controls. A significantly higher prevalence of central venous catheter (CVC) insertion (42.6% versus 8.8%) was observed in the cases compared with the controls. High white blood cell counts, serum lactic dehydrogenase (LDH), bilirubin and C-reactive protein (CRP) and low haemoglobin (Hb) and HbF were significant risk factors for VTE. Forty-two cases (61.8%) developed acute chest syndrome, 10 (14.7%) had a stroke and seven (10.3%) died.

Conclusions: VTE in patients with SCA has a high impact on morbidity and mortality. PE was the leading presentation of VTE, with CVC insertion, high LDH, bilirubin, CRP and white blood cell counts along with low Hb and HbF constituting other significant risk factors.

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Introduction

Sickle cell anaemia (SCA) is a common and life-threatening haematological disorder that affects millions of people worldwide.^{1,2} In Oman, SCA is a major public concern, where 5.7% of the population carry the sickle cell gene and 0.3% Omani people have SCA.^{3,4} It is a monogenic recessively inherited disorder with abnormal beta-globin alleles, where glutamic acid is replaced by valine in the haemoglobin subunit beta (*HBB*) gene.^{5,6} Upon deoxygenation, this genetic mutation causes erythrocytes to undergo intracellular polymerization in which flexible biconcave shaped red cells change into an elongated rigid form (sickled cells). As the process of sickling and unsickling continues, the erythrocyte membrane loses the flexibility to change and the cell remains permanently sickled.⁷ Sickled red blood cells are rigid with a non-deformable membrane and are prematurely destroyed leading to a chronic haemolytic anaemia.^{1,7} The intravascular haemolysis of red blood cells results in elevated free plasma haemoglobin with depletion of nitric oxide in the microcirculation contributing to the high prevalence of morbidity and mortality in adults with SCA.⁸

Sickled erythrocytes occlude blood vessels and disrupt blood flow, causing repeated ischaemia and inflammation. This vessel occlusion is a complex process involving an interaction between the sickle cell, vessel endothelial cells, platelets and other blood components.^{9,10} The adherence of the sickle cell to the vessel endothelium results in activation of the endothelial cells and secretion

of inflammatory mediators and biomarkers that increase platelet activation and induce blood coagulation.¹⁰

Patients with SCA frequently develop acute pulmonary complications, including acute chest syndrome (ACS), asthma and a higher risk for thromboembolism.^{11–13} Several mechanisms have been suggested to explain this hypercoagulability including platelet reactivity, with elevated plasma concentrations of platelet-derived microparticles, hydroxyl radical formation, fibronectin and thrombospondin elevation, deficient proteins S and C, and elevated levels of factor VIII and tissue factor.^{14,15}

Several studies in SCA patients have recorded a high prevalence of venous thromboembolism (VTE).^{16–18} These studies found that the incidence of VTE events in SCA patients was higher than in the non-SCA population.^{16–18} Also, several laboratory parameters were abnormal in the patients with VTE such as heterozygosity for factor V Leiden, elevated platelet counts as well as high factor VIII: C levels.¹⁸ A previous study that investigated the prevalence of pulmonary embolism (PE) and deep vein thrombosis (DVT) in SCA patients reported that 0.44% of patients had a discharge diagnosis of PE and DVT, as compared with 0.12% of non-SCA patients, indicating a significantly high prevalence.¹⁸ A previous study reported that the cumulative incidence of VTE was 17.1% for severe sickle cell disease (SCD) patients (hospitalized ≥ 3 times a year) versus 8.0% for the matched asthma controls and was associated with increased

mortality.¹⁹ The Cooperative Study of Sick Cell Disease on 1523 SCD patients aged ≥ 15 years with 8862 years of follow-up reported an incidence rate for first VTE of 5.2 events/1000 person-years (95% confidence interval [CI] 3.8, 6.9) with a cumulative incidence of 11.3% (95% CI 8.3, 15.3) by age 40 years.²⁰ Thus, it is apparent that SCA is a risk factor for DVT and PE.

This current study aimed to assess the clinical and pathological characteristics of VTE amongst patients with SCA in order to determine the predisposing factors and determine its impact on morbidity and mortality.

Patients and methods

Patient population

This retrospective case-control study enrolled consecutive patients presenting with thromboembolic complications including unprovoked PE, upper and lower extremity DVT, portal vein thrombosis (PVT), superior vena cava thrombosis, right atrial thrombosis and cerebrovascular accident (CVA) in the Department of Haematology, Sultan Qaboos University Hospital, Muscat, Oman between January 2007 and January 2018. The inclusion criterion for the case group was patients with SCA that were admitted with venous thrombosis or thromboembolic complications. Patients that had SCA without thrombosis that attended the outpatient clinic of the Department of Haematology, Sultan Qaboos University Hospital for routine appointments and prescription medication or were admitted for pregnancy formed the control group in a ratio of 2:1 due to the consecutive enrolment during the study period. None of the patients included in the study were on a regular transfusion protocol for the prevention of SCA complications. All the patients in this study were confirmed to have SCA by high

performance liquid chromatography. All patient data were collected from the hospital information system at the Sultan Qaboos University Hospital.

The Medical Research Ethics Committee of the Sultan Qaboos University Hospital approved the study (no. MREC 1491). Patient consent was not required due to the retrospective design of the study. The reporting of this study conforms to STROBE guidelines.²¹

Data collection

The data collected from the hospital information system included age, sex and SCA-associated complications, such as frequency of vaso-occlusive crisis (VOC), ACS, splenic sequestration and febrile episodes. The VOC frequency was categorized as high (>6 VOC/year), moderate (3–6 VOC/year) and low frequency (1–2 VOC/year). ACS was defined as patients presenting with fever and/or respiratory symptoms, accompanied by new infiltrates on the chest radiograph. Steady state was defined as the absence of fever or VOC in the previous ≥ 4 weeks in the patient. Amongst the cases and controls, asplenia was defined as splenic atrophy as confirmed by ultrasound/computed tomography (CT) scan and/or past surgical splenectomy. All central venous catheter (CVC) lines used in these patients included peripherally inserted central lines, PORT-A-CATH®, regular central line in the neck and femoral lines. The type of thrombosis, CVC insertion, hydroxyurea (HU) use and pregnancy details were recorded as clinical parameters. Laboratory data were recorded at baseline and during the thromboembolic episode in the cases. Haematological parameters included haemoglobin (Hb) level, white blood cell (WBC) and platelet counts, reticulocyte %, HbS% and HbF%. Biochemical parameters included serum total bilirubin, serum lactic dehydrogenase

(LDH), serum vitamin B12, serum fibrinogen, D-dimer and C-reactive protein (CRP). In addition, the radiological results of magnetic resonance imaging, CT angiography and abdominal ultrasound were recorded. The impact of VTE on morbidity and mortality in all patients was determined. VTEs were managed using a standard international protocol based on the American College of Chest Physicians guidelines.²⁰

Statistical analyses

All statistical analyses were performed using IBM SPSS Statistics for Windows, Version 23.0 (IBM Corp., Armonk, NY, USA). Normally distributed continuous data are presented as mean $\pm$ SD. Continuous data that were not normally distributed are presented as median with interquartile range (IQR). Mann–Whitney

U-test was used to compare the continuous data between the case and control groups; and between baseline and the VTE episode in the case group. Categorical data are presented as frequency of patients. Independent *t*-test was used to compare the continuous data between the case and control groups. χ^2 -test was used to compare the categorical data between the case and control groups. A *P*-value <0.05 was considered statistically significant.

Results

This retrospective case–control study enrolled 102 patients with SCA. Table 1 shows the demographic and clinical characteristics of the cases ($n=68$) and controls ($n=34$). The median age of the cases was 29.5 years (IQR 24.3, 35.0; range, 12–80 years) compared with 33.0 years (IQR

Table 1. Demographic and clinical characteristics of patients with sickle cell anaemia with (case group, $n=68$) and without (control group, $n=34$) venous thromboembolism (VTE).

Characteristic	Case group $n=68$	Control group $n=34$	Statistical analyses ^a
Age, years	29.5 (24.3, 35.0)	33.0 (24.0, 41.3)	NS
Sex, male:female	24:44	16:18	NS
Age in males, years	27.5 (24.0, 30.0)	39.5 (32.0, 45.3)	NS
Age in females, years	33.0 (25.0, 36.0)	26.5 (24.0, 31.3)	NS
Haemoglobin genotype			
HbSS homozygotes	64 (94.1)	30 (88.2)	NS
Hb β -Thal/HbS	4 (5.9)	2 (5.9)	NS
History of ACS	56 (82.4)	22 (64.7)	NS
History of splenomegaly	7 (10.3)	6 (17.6)	NS
History of splenectomy	50 (73.5)	12 (35.3)	NS
VOC/year, 1–2	25 (36.8)	17 (50.0)	NS
VOC/year, 3–6	18 (26.5)	12 (35.3)	NS
VOC/year, >6	28 (41.2)	5 (14.7)	$P=0.04$
History of hydroxyurea therapy	36 (52.9)	21 (61.8)	NS
History of previous pregnancy	12 (27.3)	3 (16.7)	NS
History of pulmonary embolism in pregnancy or puerperium	16 (36.4)	0 (0.0)	$P<0.05$

Data presented as median (interquartile range) or *n* of patients (%).

^a χ^2 -test was used to compare the categorical data between the case and control groups; NS, no significant between-group difference ($P \geq 0.05$).

ACS, acute chest syndrome; VOC, vaso-occlusive crisis.

24.0, 41.3; range, 12–55 years) for the control group. There were no significant differences in age and sex distribution between the two groups. Asplenia (including autosplenectomy) was observed in 50 (73.5%) cases compared with 12 (35.3%) controls, but the difference did not reach significance (Table 1). There was a significantly higher proportion of patients in the case group with high VOC frequency ($>6/\text{year}$) compared with the control group ($P=0.04$). There were no significant differences between the two groups in terms of Hb genotype, history of ACS, history of splenomegaly, history of HU therapy or the rate of pregnancy. A history of PE during pregnancy or puerperium (up to 6 weeks after delivery) was observed significantly more often in the case group than the control group ($P<0.05$).

Amongst the haematological parameters (Table 2), there were significant differences between the two groups at baseline in Hb, WBC count, reticulocytes, HbS% and HbF

% ($P<0.05$ for all comparisons). CRP, LDH and total bilirubin were all significantly higher in the cases compared with the controls ($P<0.05$ for all comparisons). There were no significant differences between the two groups in terms of kidney function, liver function, tobacco use or ferritin levels (data not shown).

The pattern of thromboembolism in the study population is shown in Table 3. Isolated PE and upper and lower limb DVT were the predominant presentations in the case group. Among the risk factors for thromboembolic complications, the rate of CVC insertion was significantly higher in the case group compared with the control group ($P<0.05$). There were no significant differences in the rate of ACS or stroke between the two groups. The mortality rate was significantly higher in the case group compared with the control group ($P<0.05$). Mortality in the case group was 10.3% ($n=7$), of which five died after

Table 2. Laboratory parameters in patients with sickle cell anaemia with (case group, $n=68$) and without (control group, $n=34$) venous thromboembolism (VTE).

Parameter	Case group $n=68$			Control group $n=34$	Statistical analyses ^a
	At VTE episode	At baseline versus at VTE episode		At baseline	Cases versus controls
Haemoglobin, g/dl	8.5	9.7	$P<0.05$	9.9	$P<0.05$
White blood cell count, $\times 10^9/\text{l}$	12.7	10.1	$P<0.05$	8.9	$P<0.05$
Reticulocytes, %	5.3	5.4	NS	4.2	$P<0.05$
Platelet count, $\times 10^9/\text{l}$	392	400	NS	343	NS
Haemoglobin S, %	78.2			84.5	$P<0.05$
Haemoglobin F, %	5			8.5	$P<0.05$
Fibrinogen, g/l	4			3.2	NS
Vitamin B12, pmol/l	222.5			253.5	NS
Serum lactic dehydrogenase, u/l	483	292	$P<0.05$	334	$P<0.05$
C-reactive protein, mg/l	45			7	$P<0.05$
Serum total bilirubin, mg/dl	39.5	25.0	$P=0.03$	28.5	$P<0.05$
Elevated D-Dimers	37 (54.4)			0 (0.0)	—

Data presented as median or n of patients (%).

^aMann–Whitney U -test was used to compare the continuous data between the case and control groups; and between baseline and the VTE episode in the case group; NS, no significant between-group difference ($P\geq 0.05$).

Table 3. Comparison of events and outcomes in patients with sickle cell anaemia with (case group, $n = 68$) and without (control group, $n = 34$) venous thromboembolism (VTE).

Outcomes	Case group $n = 68$	Control group $n = 34$	Statistical analyses ^a
Isolated PE	26 (38.2)	0 (0.0)	$P < 0.0005$
DVT ^b	25 (36.8)		
Upper extremity DVT	13 (52.0)	0 (0.0)	$P = 0.002$
Lower extremity DVT	12 (48.0)	0 (0.0)	$P = 0.002$
Other VTEs	17 (25.0)	0 (0.0)	$P = 0.005$
Ischaemic cerebral event	10 (58.8)	0 (0.0)	$P = 0.005$
Right atrial and SVC thrombosis	2 (11.8)	0 (0.0)	NS
Abdominal vessel thrombosis	5 (29.4)	0 (0.0)	$P = 0.04$
Combined PE + VTE/DVTs	6 (8.8)	0 (0.0)	$P = 0.03$
CVC insertion	29 (42.6)	3 (8.8)	$P < 0.05$
PE with CVC	7 (24.1)	0 (0.0)	$P = 0.02$
Upper limb DVT with CVC	18 (62.1)	0 (0.0)	$P = 0.0002$
Lower limb DVT with CVC	4 (13.8)	0 (0.0)	NS
ACS	42 (61.8)	22 (64.7)	NS
Stroke	10 (14.7)	2 (5.9)	NS
Death	7 (10.3)	0 (0.0)	$P < 0.05$

Data presented as n of patients (%).

^a χ^2 -test was used to compare the categorical data between the case and control groups; NS, no significant between-group difference ($P \geq 0.05$).

^bUpper extremity DVT: axillary, subclavian, internal jugular; Lower extremity DVT: all lower limb up to the iliac veins. PE, pulmonary embolism; DVT, deep vein thrombosis; SVC, superior vena cava; CVC, central venous catheter; ACS, acute chest syndrome.

a PE episode and two died after having an ischaemic stroke.

There were moderate correlations between VTE and PE and CVC insertion, with correlation coefficients of $r = 0.35$ and $r = 0.3$, respectively.

Discussion

Several studies have shown that VTE is one of the common complications of SCA with a cumulative incidence that increases with the age, with the mean age ranging from 24 to 37 years.^{18,19,22} Researchers have also compared SCA with inherited thrombophilia with a thrombosis prevalence around 25%, which is similar to those seen in a family with strong thrombophilic defects.²³ In this current retrospective study, the median age of SCA patients

that developed VTE was 29.5 years (range, 12–80 years), which was not significantly different from the median age of the control group (33.0 years) and similar to previous research.¹⁹ However, it was lower than the median age of 68 years for thrombosis seen in the general population.²⁴ In addition, the current study found that sex was not a risk factor for VTE in patients with SCA, but the prevalence data indicated a preponderance for females, probably driven by pregnancy, although it did not reach statistical significance. These data were similar to another study from the US in which females were more at risk for VTE than males, again, not in keeping with the observations in the general population.¹⁹

The spectrum of thrombosis in the current study cohort revealed that DVT/VTEs ($n = 42$) were seen in more than half of the

cases as compared with isolated PEs ($n=26$). This has been reported in the previous studies.^{17,19} A previous study observed that the prevalence of DVT in SCA and non-SCA patients was similar, whereas, the prevalence of PE in SCA patients was twice that of non-SCA patients.¹⁸ It has been suggested that thrombosis-in-situ could be responsible for this striking difference and that PE may be an aetiological factor in SCA patients that develop respiratory symptom.²⁵ Pulmonary hypertension with tricuspid regurgitation jet velocity ≥ 2.5 m/s is linked to VTE.²⁶ Although, SCD is a hypercoagulable state, the contribution of VTE to an elevated tricuspid regurgitation velocity is unknown.²⁶ This raises the possible pathogenic link between large-vessel venous thrombosis and the small-vessel thrombi observed in patients with sickle cell with pulmonary hypertension.²⁷ Although this current study found a high rate of PE and VTE, arterial thrombosis was rarely seen (one case each, for cerebral arterial and portal arterial thrombosis, but excluded from current data analysis).

One of the aims of this current study was to identify the predisposing factors that increase the risk of developing VTE in patients with SCA. There are a number of potential predisposing factors to thrombosis in SCA, including frequent hospitalizations, raised inflammation and hypercoagulable state associated with this disease, however, those factors are not well studied. Although the increased frequency of VOC is not a direct risk factor, it could be considered as a surrogate factor, since it leads to frequent and prolonged hospitalization of SCA patients, which is a risk factor for thromboembolism. This current study found that patients with more frequent VOCs had a significantly higher prevalence of VTE. This may again reflect disease severity and has been seen in other research.¹⁹ This current study found that a

high percentage of the VTE patients had a history of ACS (82.4%), implying that it did predispose to the occurrence of VTE, and such patients suffered from ACS as a complication of PE. Furthermore, previous research has demonstrated that PE is considered to significantly predispose for ACS, with PE episodes developing during ACS,²⁵ which is again similar to these current observations. However, it is difficult to conclude (from the current data), whether PE is a cause or a consequence of ACS.²⁸ It is believed that patients with SCA are more prone to infection as a result of immune impairment because of asplenia.²⁹ Infection, by promoting an increase in inflammatory cytokines, can provoke thrombosis, especially in asplenic patients.³⁰ Furthermore, asplenia in haemolytic diseases including SCA was reported to be associated with an increased risk of thrombosis, as seen in this current study.¹⁶

Central venous catheter insertion has a significant correlation with VTE episodes.¹⁵ CVC is generally required for many patients with SCA and it has been shown to be an independent risk factor for VTE.^{15,31} Furthermore, prolonged use of catheter implantation increases the risk of thrombosis,^{17,31} as seen in the current study cohort. Among the 68 patients with VTE in the case group, 29 patients had CVC at the time of VTE, including 18 with upper limb DVT, seven with PE and four had lower limb DVT.

Pregnancy is significantly correlated with VTE as the rate of VTE during pregnancy and puerperium was 1.5–2.5-times greater than in non-pregnant SCA women.^{32,33} It is reported that pregnancy-related VTE in females with SCA is 1.5–5-times greater than pregnancy-related VTE in the general population.³² The current study demonstrated that 27.3% of cases compared with 16.7% of controls had a history of pregnancy. Interestingly, PE was reported in 16 (36.4%) of females in the case group,

whereas none of the three pregnant females in the control group developed PE ($P < 0.05$). There are several possible explanations for this, such as impaired venous return from lower limbs, high level of steroid hormones and changes in the haemostatic system, in addition to SCA, which together contribute to elevate the risk of thrombosis during the pregnancy period.^{32,33} This has led to the recommendation of prophylactic anticoagulation in all pregnant patients with SCA.³⁴

The use of HU therapy over the last 30 years has been shown to increase HbF in SCA patients leading to improved clinical efficacy.³⁵ It is associated with a reduction in the levels of biomarkers of hypercoagulability in SCA patients and is likely to reduce the prevalence of VTE.³⁶ In this current study, the median HbF levels were significantly higher in the control group compared with the case group. Furthermore, a higher percentage of control patients (61.8%) were treated with HU compared with 52.9% in the case group, which might explain the higher median HbF in the controls. Although there are contradicting results on the impact of HU on VTE prevention, these current results were inconclusive with respect to a positive role of HU in VTE prevention, knowing that HbF inhibits intracellular HbS polymerization.^{37,38}

In the current study population, there were elevations in the median WBC count, LDH level, bilirubin and CRP in the case group from baseline to VTE episode; and the baseline levels were significantly different compared with the control group. Higher WBC count and elevated LDH level are reported to be risk factors for VTE in SCA.³⁹ Biomarkers are helpful in identifying different pathophysiological mechanisms in SCA, including hypercoagulability, haemolysis, nitric oxide depletion, inflammation, oxidative stress, vasculopathy, reperfusion injury and tissue

necrosis with infarction.^{7-10,19,25} Over half of the patients in the case group in the current study had elevated D-dimer (54.4%) at the VTE episode and the CRP levels were significantly higher at the VTE episode compared with the controls at baseline, which were presumably due to an acute phase reaction, often seen in patients with thromboembolism and in sickle cell crisis.³³

Among the clinical complications of VTE in this current group of patients, a large majority of the patients in the case group developed ACS (61.8%), which was consequential of the development of PE; and 10 patients (14.7%) developed stroke as a manifestation of venous and arterial thrombosis within cerebral vessels. In 306 autopsies of SCA patients, a previous study reported that 9.8% died due to stroke and 4.9% died due to pulmonary emboli/thrombi.⁴⁰ However, another autopsy study from Nigeria reported that the mortality rate from fatal thromboembolic complications was 37%.⁴¹ Thus, although infections and multiorgan failure are major causes of death in SCA, SCA with VTE is still associated with a significant mortality.^{40,41} These data support the regular use of prophylactic low molecular weight heparin for the purpose of VTE prevention and improved VOC management.^{42,43}

In conclusion, acknowledging the limitation of the retrospective nature of this current study, VTE was one of the common complications of SCA and it occurred at a younger age than the general population. PE showed a higher prevalence in patients with SCA. Several risk factors contributed to the development of VTE in patients with SCA such as CVC insertion and pregnancy. In addition, coagulation activity in patients with SCA was enhanced with abnormal activation of the fibrinolytic system and increased levels of WBCs, LDH, bilirubin and CRP. This current study demonstrated that thromboembolic complications in patients with SCA had a high impact on

morbidity and mortality and calls for the use of prophylactic anticoagulation in all patients with SCA during their hospital stay.

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Author contributions

All authors made substantial contributions. They have all seen and approved the final version of the manuscript. S.A., A.R.A., S.R.A., S. Y.A., N.F., S.K.B. and A.V.P. were fully involved in the conception and design of the study, recruitment and care of patients, acquisition of data, or analysis and interpretation of the data. S.A. and A.V.P. were instrumental in the drafting the article and critical appraisal before submission.

Declaration of conflicting interest

The authors declare that there are no conflicts of interest.

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