Asymptomatic bacteriuria in sickle cell disease: a cross-sectional study

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Background: It is known that there is significant morbidity associated with urinary tract infection and with renal dysfunction in sickle cell disease (SCD). However, it is not known if there are potential adverse outcomes associated with asymptomatic bacteriuria (ASB) infections in sickle cell disease if left untreated. This study was undertaken to determine the prevalence of ASB, in a cohort of patients with SCD.

Methods: This is a cross-sectional study of patients in the Jamaican Sickle Cell Cohort. Aseptically collected mid-stream urine (MSU) samples were obtained from 266 patients for urinalysis, culture and sensitivity analysis. Proteinuria was measured by urine dipsticks. Individuals with abnormal urine culture results had repeat urine culture. Serum creatinine was measured and steady state haematology and uric acid concentrations were obtained from clinical records. This was completed at a primary care health clinic dedicated to sickle cell diseases in Kingston, Jamaica. There were 133 males and 133 females in the sample studied. The mean age (mean ± sd) of participants was 26.6 ± 2.5 years. The main outcome measures were the culture of ≥ 10⁵ colony forming units of a urinary tract pathogen per milliliter of urine from a MSU specimen on a single occasion (probable ASB) or on consecutive occasions (confirmed ASB).

Results: Of the 266 urines collected, 234 were sterile and 29 had significant bacteriuria yielding a prevalence of probable ASB of 10.9% (29/266). Fourteen patients had confirmed ASB (prevalence 5.3%) of which 13 had pyuria. Controlling for genotype, females were 14.7 times more likely to have confirmed ASB compared to males (95% CI 1.8 to 121.0). The number of recorded visits for symptomatic UTI was increased by a factor of 2.5 (95% CI 1.4 to 4.5, p < 0.005) but serum creatinine, uric acid and haematology values were not different in patients with confirmed ASB compared with those with sterile urine. There was no association with history of gram negative sepsis.

Conclusion: ASB is a significant problem in individuals with SCD and may be the source of pathogens in UTI. However, further research is needed to determine the clinical significance of ASB in SCD.
Background
The definition of asymptomatic bacteriuria (ASB) is controversial as some have defined it as the quantitative growth of bacteria, greater than or equal to $10^5$ colony forming units per millilitre urine of the same organism, on aseptically collected midstream urine specimens, in the absence of symptoms of urinary tract infection on two or more consecutive occasions [1,2] while for others a single occasion is sufficient [3-5]. The prognostic significance of ASB resides in the observation that persons with ASB in certain medical conditions, such as diabetes mellitus and pregnancy, are at increased risk of pyelonephritis and renal impairment [5-7].

In sickle cell disease renal disease and dysfunction are common and increase morbidity and mortality [8,9]. Additionally, symptomatic urinary tract infection (UTI) is associated with painful crisis, bacteraemia, pneumonia, and osteomyelitis in SCD [10-13]. Whether ASB is a prelude to more serious renal disease and dysfunction in SCD is unclear. This present study was undertaken to estimate the prevalence and possible risk factors of ASB in the Jamaican sickle cell cohort.

Methods
Patients
The sample comprised a group of patients in the Jamaican Sickle Cell Cohort attending the 2004 Annual Sickle Cell Cohort Review conducted at the Sickle Cell Unit Clinic at the University of the West Indies, Kingston, Jamaica. The Jamaican Sickle Cell Cohort includes patients with sickle cell diseases detected using standard criteria during screening of 100,000 consecutive non-operative deliveries at a large maternity hospital (the Victoria Jubilee Hospital) between 1973 and 1981. This screening yielded 580 patients with different sickle genotypes, of which 315 had HbSS disease, 201 Sickle cell-haemoglobin C, 33 Sickle cell-β-thalassaemia, 14 Sickle cell-β thalassaemia and 17 sickle variants. They have been followed clinically since birth [14]. At the 2004 cohort review 267 of 302 patients eligible to attend the annual cohort review were seen. The reasons for being absent were: Twenty-two of the patients did not attend the review that year, 11 were lost to follow up, or incarcerated, and 2 were ill at the time of review. One patient refused to leave a sterile urine sample during the cohort review period, thus the final participating patient sample for this study was 266 patients. Ethical approval was granted for this study by the University Hospital of the West Indies, University of the West Indies Faculty of Medical Sciences Ethics Committee.

Measurements
Aseptically collected midstream urines (MSUs) were obtained from the symptom free attendees of 2004 cohort review and sent on the same day for microbiological culture. Urinalysis was performed on aliquots of the MSU specimen using the QuickVue UrinChek™10+SG test strips. Bacterial culture was performed by streaking 0.002 mL of midstream collected urine with a calibrated loop on MacConkey and 5% sheep blood agar plates. These agar plates were incubated at 35°C for 24 hours under aerobic conditions. Isolates were considered significant if there were $\geq 10^5$ colony forming unit/mL (CFU/mL) with 2 or less isolates, doubtful significance if $10^4 - 10^5$ CFU/mL, insignificant if $< 10^4$ CFU/mL. Mixed growths, in any count, of more than two organisms were considered to be contaminated. Significant isolates were selected for identification and antimicrobial susceptibility testing using the vitek (biome'rieux' version R06.01, Missouri USA). Proteinuria was defined as a protein reading of trace or greater on dipstick urinalysis.

Patients complaining of any symptoms, whether related or unrelated to the urinary tract, were not included in the study on that day. Rather, these patients were treated for their complaint and studied on another day when they were without symptoms. Subjects whose urine microbiological report was abnormal were asked to return to the unit for a repeat midstream urine collection and culture.

Blood samples were also taken for the measurement of serum creatinine. Serum creatinine were measured in the Tropical Medicine Research Institute laboratory with an Abbott ALCYON 300I, using the alkaline picrate method.

The clinical records of all cohort patients in the sickle cell unit database were examined and the following information extracted: age, gender, steady state haematology, steady state uric acid levels, history of urinary tract infection, and history of gram negative sepsis.

Definition of asymptomatic bacteriuria
Probable ASB was defined as the presence of at least $10^5$ colony forming units of a urinary tract pathogen per millilitre of urine in a culture of a midstream urine specimen obtained from a patient during the cohort review visit. A confirmed episode of asymptomatic bacteriuria was defined as two or more consecutive cultures with evidence of asymptomatic bacteriuria due to the same urinary tract pathogen with the same sensitivity pattern. A sample was deemed grossly contaminated if it grew more than two urinary tract pathogens even if in significant quantities.

Statistical methods
Values are expressed as counts or means ± sd as appropriate. The prevalence of probable ASB and confirmed ASB were determined as the ratio of the number of urines classified as probable ASB and confirmed ASB, to the total number of collected MSU samples. The 95 percent confidence interval (95%CI) for this proportion was computed.
Results
A total of 267 patients attended the 2004 cohort review. One person refused to participate in the study. Thus 266 patients had screening clean-catch MSU (Figure 1), of which 50% were male. One woman was pregnant at the cohort review visit; and another became pregnant during follow-up of the first abnormal MSU.

As expected there were significant differences in the hematological values by genotype with the mean steady state haemoglobin levels of HbSS group being significantly lower than sickle cell-haemoglobin C group, sickle cell-βthalassaemia, and sickle variants but having higher steady state nucleated cell counts (Table 1). The steady state serum creatinine values were significantly lower in the HbSS group compared with other genotypes except the sickle variant group. In addition the mean uric acid values were lower in the HbSS group compared with sickle cell-βthalassaemia group (Table 1).

There were 32 patients with abnormal first MSUs: 29 were positive MSU samples growing identifiable pathogens (Figure 1), 2 were grossly contaminated and one grew bacteria of doubtful significance (growth of a urinary tract pathogen between 104 and 106 per mL). Of the 29 positive cultures 15 re-grew the same organisms in the next culture but one of these grew the same organism (E. coli) but with a different sensitivity pattern and was therefore classified as a probable ASB. Thus 14 samples satisfied the criteria for a confirmed episode of asymptomatic bacteriuria (Figure 1) yielding a prevalence of 5.3% (95% CI 0.8% to 29.4%). Fifteen subjects were classified as probable ASB. The probable ASB group consisted of seven samples which grew organisms that were different from the first MSU sample, 2 grew bacteria in insignificant quantities, four were no growth and two were grossly contaminated specimens (Figure 1). None of the patients were on antibiotic therapy during this study and all of the patients with identified ASB had at least one follow-up visit.

Sixteen of the 29 positive first MSU samples (55%) grew E. coli in significant quantity. This was the commonest organism cultured in this study. Other urinary tract pathogens identified in this study included: group B Streptococcus, Enterobacter aerogenes, coagulase negative Staphylococcus, Enterobacter cloacae, Proteus mirabilis, Microaerophilic streptococcus and Bacteroides (mixed), group B Streptococcus and Streptococcus viridans (mixed), Enterobacter species, group D Streptococcus, Klebsiella pneumoniae, and Enterobacter koseri. All of these organisms were grown in pure cultures with only the two exceptions listed above. Fifteen of the patients with probable ASB (52%) had evidence of a host response, as evidenced by pyuria. Of those with confirmed ASB, 13 of 14 patients (93%) had a host response as evidenced by pyuria in the second midstream urinalysis.

There were no significant differences for mean haemoglobin and nucleated blood cell counts between the con-
The sexual dimorphism in the prevalence of ASB has been reported before in healthy adults [6]. It is thought that this

Table 1: Clinical Characteristics of enrolled subjects by genotype (N = 266).

| Clinical characteristic       | HbSS N = 148 | HbSC N = 85 | HbSβ+ N = 6  | HbSβ+ N = 22 | Sickle variants N = 5 |
|-----------------------------|--------------|-------------|--------------|--------------|-----------------------|
| Age yr                      | 26.4 ± 2.6   | 27 ± 2.3    | 25.9 ± 2.3   | 26.5 ± 2.5   | 25.1 ± 2.5            |
| Gender (M:F)                | 71:77        | 38:47       | 2:4          | 13:9         | 4:1                   |
| #Creatinine μmol/L          | 45.7 ± 9.4*  | 63.3 ± 11.7b| 70.7 ± 23.1b | 61.9 ± 12.4a | 62 ± 20.4a            |
| #Uric Acid mmol/L           | 0.3 ± 0.1*   | 0.3 ± 0.1*  | 0.4 ± 0.1*   | 0.3 ± 0.1*   | 0.3 ± 0.1*            |
| #Haemoglobin g/L            | 80.2 ± 9.9*  | 108.2 ± 8b  | 109.9 ± 12.4b| 124 ± 22.1b  | 124 ± 22.1b           |
| #Nucleated Blood cells × 10⁹/L | 13.9 ± 2.2b  | 13.3 ± 2.2b | 9.2 ± 2.7b   | 9.9 ± 2.5b    |                       |
| †Proteinuria ≥ trace : No proteinuria (n = 182) | 17 | 7 | 0 | 1 | 0 |
| †Positive History of gram negative sepsis | 2 | 2 | 0 | 0 | 0 |
| †Positive recorded history of ever having UTI | 50 | 25 | 2 | 1 | 1 |

Values are means ± sd; † values are counts. Abbreviations – HbSS = Homozygous sickle cell disease, HbSC = Sickle cell-haemoglobin C, HbSβ+ = Sickle cell-β thalassemia, HbSβ+ = Sickle cell-β* thalassemia; UTI= Urinary Tract Infections.
* Sickle variants include sickle cell-hereditary persistence of foetal haemoglobin, sickle cell -haemoglobin O Arab, sickle cell -haemoglobin Lepore-Boston.
# Significant difference in mean values by genotype. Dissimilar superscript to HbSS values are significantly different (p < 0.03)
may be related a relative deficiency of secretory IgA antibody response from the mucosal surface in the urogenital tract of females compared with males [6].

In non pregnant females the presence of confirmed ASB has a high sensitivity and specificity of predicting a future symptomatic episode of UTI [5,6]. However in the absence of anatomical or functional abnormalities of the urinary tract, ASB per se is not associated with renal scarring, renal dysfunction or hypertension [17]. In adults with SCD, anatomical abnormalities such as renal cortical scarring, enlarged glomeruli, vascular disorganization of the medulla and renal functional abnormality such as increased effective renal blood flow and hyperfiltration are common [8]. These pathophysiological changes are thought to contribute to sickle cell nephropathy and the high prevalence of renal failure observed in subjects with sickle cell disease [8]. In this context, one might expect persons with ASB to have more UTI and worse renal function. Thus the finding of an increased number of expected visits for symptomatic UTI in subjects with ASB compared to those without as well a trend for a significant association between ASB status and a history of UTI is not surprising. On the other hand, the absence of significant differences in measures of renal function between those with ASB and those without ASB was unexpected. However, serum creatinine and uric acid are not sensitive measures of renal function as glomerular filtration rate has to fall by up to 50% before there are changes in serum creatinine [18,19]. Additionally variation in creatinine production due to differences in muscle mass may alter serum creatinine levels independent of renal functional status [20,21]. Notwithstanding, further prospective studies are needed to define the renal risk posed by ASB in sickle cell disease and to inform therapeutic strategies.

In sickle cell disease, the increased purine turnover associated with accelerated erythropoesis results in the greater production of uric acid [22]. This is counterbalanced by increased renal clearance of urate. It is possible that the increased renal clearance may play a predisposing role for ASB by promoting the formation of microcrystals which damage the uroepithelium thereby facilitating colonization.

The infecting organisms identified in this study are in keeping with commonly isolated bacteria in other studies [1-3,5,10,11]. The presence of bacteria in the urine in the absence of a major inflammatory response and hence symptoms, suggest that there are alterations in the host-pathogen interaction. For example it has been reported that uropathogenic strains of microbes have evolved mechanisms to promote survival in the urinary space. Features of these adaptive mechanisms include presence of adhesion which promote attachment to the urogenital tract, and the production of factors such as α-haemolysin,

Table 2: Clinical characteristics of enrolled subjects with confirmed ASB and sterile urine (N = 248)*

| Clinical characteristics | Confirmed ASB N = 14 | Sterile urine N = 234 |
|-------------------------|----------------------|----------------------|
| Age yr                  | 27.9 ± 2.3           | 26.4 ± 2.5           |
| † Gender (M:F)          | 1:13                 | 124:110              |
| ‡Genotype frequency (HbSS:HbSC:HbS[β]:HbS[β]+: sickle variants) | 9:2:1:2:0          | 130:75:5:19:5        |
| Creatinine µmol/L       | 49.1 ± 12.4          | 53.7 ± 14.3          |
| Uric Acid mmol/L        | 0.36 ± 0.29          | 0.33 ± 0.08          |
| † Proteinuria ≥ trace : No proteinuria (n = 168) | 1:7                  | 22:138               |
| † Positive History of gram negative sepsis | 1                    | 1                    |
| † Positive recorded history of ever having UTI | 7                    | 25                   |
| ‡ Cumulative number of recorded clinic visits for symptomatic UTI* | 18                   | 49                   |

*Values of subjects with probable ASB excluded from table. † Values are counts. Abbreviations : ASB = asymptomatic bacteriuria; HbSC= Sickle cell-haemoglobin C, HbS[β]= Sickle cell-[β] thalassemia, HbS[β]+ = Sickle cell-[β] thalassemia; UTI = Urinary Tract Infections

#Sickle variants include sickle cell-hereditary persistence of foetal haemoglobin, sickle cell -haemoglobin O Arab, sickle cell -haemoglobin Lepore-Boston;

Table 3: Model predicting asymptomatic bacteriuria in the Jamaica Sickle Cell Cohort.

| Variables         | Odds ratio | P value   | Lower 95% CI | Upper 95% CI |
|-------------------|------------|-----------|--------------|--------------|
| Age years         | 1.37       | 0.057     | 0.99         | 1.64         |
| Female (coded as 1)| 15.3       | 0.010     | 1.94         | 120.12       |
| Uric acid mmol/l  | 21.5       | 0.087     | 0.64         | 722.21       |
and cytotoxic necrotizing factor 1 (CNF1) [6]. On the other hand the host factors in SCD that increases susceptibility to bacterial colonization are unclear but in Diabetes Mellitus altered leukocyte function contribute to the susceptibility to ASB in this disease [23].

Conclusion
In summary ASB is a significant problem in individuals with SCD and is associated with UTI. However, further research is needed to determine the clinical and renal significance of ASB in SCD.

Abbreviations
SCD: sickle cell disease, ASB: asymptomatic bacteriuria, MSU: midstream urine, UTI: urinary tract infection, CFU: colony forming unit.

Competing interests
The author(s) declare that they have no competing interests.

Authors’ contributions
SA and MR coordinated and supervised the cohort review at the Sickle Cell Clinic; VC collected data, managed the patients with ASB, and wrote the paper in collaboration with TF, MR and KRG. All authors read and approved the final manuscript.

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