INTRODUCTION
Sickle cell anemia (SCA) was presented by Linus Pauling in 1949 as an autosomal recessive disease in which multiple organs are affected and ever since it has become a prevalent genetic disease in the world: more than 30 million people are affected worldwide. A meta-analysis published in 2018 estimated the prevalence of patients with SCA to be 111.91 for every 100,000 live births. In Brazil, between 25,000 and 50,000 cases of SCA are estimated, according to the Ministry of Health. Chronic kidney disease (CKD) develops in 40% of individuals, and 15–30% die due to kidney complications. Among individuals who reach the fourth decade of life, half have end-stage kidney damage.

In the last decades, the average life expectancy of patients with SCA has increased thanks to better management of the numerous complications of the disease, including sickle nephropathy (SN). However, the death rate of patients with SN is still high, with little known predictors related to its development. To answer the question “What predictors are associated with the onset of CKD in patients with SCA?”, this article seeks to contribute towards a better understanding of SN, making possible a new look at the SCA and its kidney complications, which will contribute towards early detection, better management and, consequently, an improvement in the prognosis of the disease.
METHODS

A systematic review was developed, using the PRISMA\textsuperscript{9} recommendation, for cohort studies on predictors related to the outcome of SN in patients with SCA.


Two researchers (LVSM and SMSR) independently and blindly analyzed the titles and abstracts of the articles found. The divergences were analyzed by a third researcher (LRS). After that, a full-text reading was done applying the inclusion criteria: articles published in the period between 10/27/2009 and 10/25/2019; cohort original articles; studies in which measures of creatinine clearance have been obtained by previously validated means; studies carried out in humans. The exclusion criteria were: articles which only address patients with sickle cell trait/other hemoglobinopathies and those that do not address predictors in patients with kidney injury.

Table 1. Results of analyzed articles.

<table>
<thead>
<tr>
<th>Article</th>
<th>n</th>
<th>Age (Years/mean)</th>
<th>Risk factors (p&lt;0.05)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lebensburger, 2011\textsuperscript{13}</td>
<td>144</td>
<td>–</td>
<td>Lower Hb levels, lower HbF levels and increased LDH.</td>
</tr>
<tr>
<td>Aygun, 2011\textsuperscript{11}</td>
<td>260</td>
<td>≤30</td>
<td>Increased cystatin C levels, higher systolic blood pressure, decreased leukocytes count and decreased absolute neutrophil counts.</td>
</tr>
<tr>
<td>McPherson Yee, 2011\textsuperscript{15}</td>
<td>261</td>
<td>2–21</td>
<td>Increased age, lower Hb levels.</td>
</tr>
<tr>
<td>Silva Junior, 2012\textsuperscript{16}</td>
<td>98</td>
<td>19–67</td>
<td>Lower Hb levels, lower hematocrit levels, lower platelet levels, leukocyturia, proteinuria and increased age.</td>
</tr>
<tr>
<td>Saraf, 2014\textsuperscript{14}</td>
<td>795</td>
<td>24–46</td>
<td>Higher mean arterial pressure, lower Hb levels, macroalbuminuria, haemoglobinuria, increased age, AST, increased LDH.</td>
</tr>
<tr>
<td>Gosmanova, 2014\textsuperscript{12}</td>
<td>98</td>
<td>21–42</td>
<td>Increased age, higher systolic and diastolic blood pressure, albuminuria.</td>
</tr>
</tbody>
</table>

CKD: chronic kidney disease; AKI: acute kidney injury; Hb: hemoglobin; HbF: Fetal hemoglobin; CRP: C-reactive protein; AST: aspartate transaminase; LDH: lactate dehydrogenase; NGAL: urine neutrophil gelatinase-associated lipocalin; BMI: body mass index.

RESULTS

Initially, 321 studies were identified in PubMed, of which six were selected to compose this systematic review, as shown in Figure 1.

Of the six selected studies, five were developed in the United States\textsuperscript{11-15} and one in Brazil\textsuperscript{16}. Table 1 summarizes the main

For each full text selected, the CASP\textsuperscript{10} questionnaire was applied to assess the methodological quality and possible bias. A manual search was performed on the references of the selected articles, to select an article that had not been included in our search key. Then, the predictors cited by the articles that had a p-value less than 0.05 were extracted and arranged in a table according to the highest frequency.

Figure 1. Flow of phases of the research.
characteristics of the six studies and the respective predictors associated with CKD in patients with SCA.

Lower levels of hemoglobin, albuminuria and increased age were the main predictors shown in the studies. Other important predictors presented were increased LDH (lactate dehydrogenase), higher blood pressure, lower levels of fetal hemoglobin, among others listed in Table 2.

**DISCUSSION**

Lower levels of hemoglobin were suggested as important predictors for CKD. However, CKD caused by other diseases can also lead to anemia, which raises the question: is CKD linked to anemia or is anemia a predictor of SN? Naik et al. described the pathophysiology of SN, correlating with anemia. The red blood cells of patients with AF when in vessels with lower oxygen tension or lower pH polymerize, taking the form of a sickle, occluding small diameter vessels and leading to ischemic lesions in multiple organs, including the kidney. This also leads to glomerular microinfarctions and apoptosis of epithelial cells, which can decrease the glomerular filtration rate [17].

Patients that developed CKD had greater ages when compared to patients without kidney injury. These data are consistent with a 4 decades study developed with 1056 patients, which demonstrated that renal injury tends to occur later, around the third to fourth decades of life [18].

Albuminuria and proteinuria were also prevalent predictors. The literature points to other studies that demonstrate albuminuria as a predictor linked to the pathogenesis of podocyte loss and disruption of the glomerular filtration barrier [19]. Podocytes are the epithelial cells that line the glomerular capillaries through their foot processes and are responsible for the composition of the glomerular filtration membrane, together with the porous endothelial cells and the glomerular basement membrane. They serve as a barrier by holding proteins through size and load-dependent blocking. As podocytes have their own metabolism, ischemic lesions, toxins, complement deposition, among other situations, lead to the production of reactive oxygen species that induce apoptosis and podocyte processes effacement, through the destruction of the actin cytoskeleton, changes in the negative podocyte surface and failure of calcium metabolism. Vasooclusive episodes occur in SCA patient's vessels, culminating with ischemic lesions to the kidneys. All of these result in the failure of the filtration barrier and trigger proteinuria [20,21].

Arterial hypertension was positively correlated with the development of CKD in individuals with SCA, through higher systolic and diastolic blood pressure seen in CKD patients. Corroborating these findings, a cohort of 158 patients with SCA showed a 50% increase in creatinine levels in patients with systolic blood pressure levels above 140 mmHg or diastolic blood pressure above 90 mmHg [22].

Urinary and serum markers such as lactic dehydrogenase, AST, cystatin C, leukocyturia, and hemoglobinuria were related to the development of renal injury, which suggests that hemolysis and inflammation are present in the pathophysiology of the nephropathy. Fetal hemoglobin was shown as a protective factor as it prevents red blood cells from acquiring the sickle shape [17].

A small sample study demonstrated a decrease in lactic dehydrogenase in patients with SCA who were treated with hydroxyurea. Also, the same study reported lower reticulocyte counts and higher levels of fetal hemoglobin, findings that suggest a reduction in hemolysis in patients receiving hydroxyurea and also show that hemolysis is an important feature in patients with SCA. A larger cohort also demonstrated that lactic dehydrogenase is associated with higher mortality in individuals with SCA, reaffirming that hemolysis is a predictor of poor prognosis in these patients [23,24].

Higher levels of AST were pointed in the studies of this review as predictors of SCA, but other articles in the literature have correlated higher levels of AST, as well as bilirubin and alanine aminotransferase, with another organ failure in SCA: the sickle cell hepatopathy, as demonstrated in the cohort.
developed with more than 1100 subjects. On the other hand, this study does not present any parameter related to kidney function, which provides the missing information of whether AST is related or not with SN\textsuperscript{25}. However, Madu et al. correlated AST levels and proteinuria, without statistical significance, which also indicates a poor correlation between this predictor and kidney injury\textsuperscript{26}.

A study published in 1985 with patients with renal dysfunction caused by multiple diseases, associated decreased glomerular filtration rate with increased levels of cystatin C\textsuperscript{27}. Our study also demonstrated an association between this marker and reduced renal function in patients with SCA, but recent studies have conflicted with these findings. Unal et al. for example, showed no difference in cystatin C levels between SCA patients and healthy controls\textsuperscript{28}. On the other hand, a Greece study developed with 87 patients found an inverse correlation between glomerular filtration rate and cystatin C levels, corroborating with the previous findings of the 1985 study\textsuperscript{29}. All this controversy raises the need for further research on this marker.

Regarding urinary markers, leukocyturia and hemoglobinuria did not correlate with proteinuria in a cross-sectional study conducted in Africa, which differs from our findings\textsuperscript{30}. However, other reports have shown that hemoglobinuria is related to SN as a sign of papillary necrosis, congestion of the renal vasculature, and even renal medullary carcinoma\textsuperscript{31-33}.

Our study has some limitations. Few cohort studies evaluating the development of kidney injury in patients with SCA have been found in the databases, which demonstrates the need for further studies on this topic. In addition, there is high heterogeneity among the articles, with several measures of creatinine clearance and different criteria of albuminuria.

CONCLUSION

In conclusion, the main predictors associated with the development of CKD in individuals with SCA were lower hemoglobin levels, albuminuria, and increased age. New studies evaluating the pathophysiology of kidney injury in SCA are needed to better understand its installation and prevent its progression.

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AUTHORS’ CONTRIBUTIONS

LM: Conceptualization, Data Curation, Methodology.
SR: Conceptualization, Data Curation, Methodology, Writing – Original Draft.
LS: Writing – Original Draft, Writing – Review & Editing.
DBM: Methodology, Writing – Original Draft, Writing – Review & Editing.

REFERENCES


