Sickle cell disease (SCD) is characterised by haemolytic anaemia and ubiquitous vessel occlusion by sickled red cells. Besides polymerisation of deoxyhaemoglobin S, adhesion of erythrocytes to endothelial cells contributes to altering vascular flow and to provoking vasoocclusion. Chronic and acute ischaemic complications are hallmarks of the disease. Surprisingly, the heart is not usually considered a high risk target organ, in particular in children. However, as many as 10–30% of adults suffer cardiac impairment, mostly left ventricular hypertrophy or congestive heart failure.1 Pathological studies have revealed associations with degenerative myocardial changes, interstitial fibrosis, and occlusion of intramural coronary arteries by aggregated sickle cells.2–5 A review of postmortem examinations of 70 patients (adults and children) with sickle cell disease performed between 1950 and 1981 at the University of Southern California found evidence of myocardial infarction and fibrosis in 17%.6

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Recent major cardiac complications (fatal cardiac failure, severe arrhythmia, myocardial ischaemia) in three of our patients prompted us to explore cardiac status of our paediatric cohort with sickle cell disease. Besides the classical tools (ECG, echocardiography), we looked for myocardial ischaemia using single photon emission computed tomography (SPECT) in children selected according to clinical, ECG, and echocardiographic criteria. Twenty two patients met these criteria and underwent thallium-201 SPECT. Our initial aims were to assess the prevalence of myocardial ischaemia and to look for possible risk factors. Eight children in whom myocardial ischaemia was demonstrated were treated with hydroxyurea, and we could assess the efficacy of the treatment in three of them.

METHODS
A cohort of 300 children with sickle cell disease is being followed up in Necker Enfants Malades Hospital. All children receive a complete physical examination every three months, and are systematically asked about chest pain. Chest roentgenography is not systematic in our follow up, and is performed only in case of heart murmur or chest pain, and when patients are hospitalised. ECG is performed in case of heart murmur, chest pain, or cardiomegaly. Echocardiography is performed in children with heart murmur and/or cardiomegaly.

In this study, children with chest pain, heart failure, abnormal ECG, left ventricular dilatation, or hypokinetic left ventricle were selected for myocardial perfusion studies. Twenty two children met one or more of these criteria. Twelve were male and 10 female; mean age was 11.24 (1.04) years (range 3–19). Twenty one had homozygous sickle cell disease (SS), one was S/β-thalassemic. Two children (patients 6 and 22 in tables 1 and 2) had a history of stroke. Patient 6, aged 10 years, had been on regular transfusion for five years, and received daily iron chelation (ferritin 1100 ng/ml). Patient 22, aged 11 years, had a stroke six months before SPECT. She has been transfused monthly since the accident (ferritin level at SPECT: 500 ng/ml). She received bone marrow transplantation some weeks after cardiac evaluation. Patients 9, 16, and 18 (aged 5, 4, and 3.5 years, respectively), had had episodes of cardiac failure. Seven patients had had chest pain, typical of angina in a 19 year old girl. Twelve patients had ECG abnormalities. Fourteen had left ventricular enlargement.

These 22 patients underwent 201 Tl SPECT after stress, and again three hours later after reinjection, using a single head gamma camera equipped with a low energy all purpose collimator. A dose of 201 Tl was injected at peak exercise or during the pharmacological or mixed stress testing (dose (MBq) = weight (kg) × 1.5). Pharmacological stress testing (using dipyridamole) was systematically used before the age of 6 years. In older children, pharmacological or exercise

Abbreviations: ECG, electrocardiography; ERNA, equilibrium radionuclide angiography; LVEF, left ventricular ejection fraction; SCD, sickle cell disease; SPECT, single photon emission computed tomography.
testing (using progressive stages of 10 to 20 watts for 2 minutes until heart rate reaches 190/min), or mixed testing was performed. Another dose of $^{201}$Tl was reinjected before the redistribution acquisition (dose (MBq) = weight (kg) × 0.5). No patient required sedation. The matrix size format was 64 × 64. Starting from the left posterior oblique position to right anterior oblique position, 30 projections over 180° were collected, 30 seconds per step.

The left ventricular ejection fraction (LVEF) was assessed by equilibrium radionuclide angiography (ERNA) at rest on the same day, using the gamma camera. The matrix size format was 64 × 64. Left anterior oblique and lateral views were taken, 5 million counts per view.

Haemoglobin (Hb) levels and pulse oxymetry (SpO2) (using a Nellcor N-395) (Nellcor Inc., Hayward, CA, USA) were measured on the same day, or within one week when SPECT was performed (no blood transfusion was given between these measurements and SPECT). The mean number of hospitalised vasoocclusive crises per year was evaluated from the medical records of the past three years for 19 patients, but could not be calculated for the remaining three. For all patients we also recorded the histories of acute chest syndromes (defined as acute respiratory illness with radiographic findings consistent with consolidation of the lung(s)).

Patient characteristics were compared using Student’s t-test.

RESULTS

Tables 1 and 2 summarise the results.

Myocardial perfusion studies

Myocardial perfusion was abnormal in 14 of 22 patients, and normal in eight. In the 14 patients with abnormal perfusion, nine had reversible defects, and five had fixed defects (fig 1). Mean LVEF was 61.7 ± 8.8% and was not correlated with myocardial perfusion, left ventricular dilatation, or Hb level.

Risk factors

Mean age, Hb level, SpO2, and mean number of vasoocclusive crises/year did not differ significantly between normal SPECT and abnormal SPECT groups (table 3). However, children with perfusion defects tended to be older (12.7 ± 9.2 years, p = 0.09), and have more severe diseases.

Course

Data for the first two patients included in this series (patients 9 and 10) have already been reported.8 Patient 9 had her first acute cardiac failure when she was 3 years old.

Table 1 results: children with normal myocardial perfusion

<table>
<thead>
<tr>
<th>Pt</th>
<th>Age (y)</th>
<th>Nb VOC/y</th>
<th>ACS other</th>
<th>Hb (g/l)</th>
<th>Chest pains</th>
<th>ECG</th>
<th>Echocardiography</th>
<th>Stress testing</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>6</td>
<td>0</td>
<td>0</td>
<td>67</td>
<td>–</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>Death</td>
</tr>
<tr>
<td>2</td>
<td>11</td>
<td>ND</td>
<td>0</td>
<td>80</td>
<td>–</td>
<td>Normal Normal</td>
<td>+</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>3</td>
<td>9</td>
<td>0</td>
<td>0</td>
<td>85</td>
<td>+</td>
<td>Abnormal Normal</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>4</td>
<td>9</td>
<td>0</td>
<td>0</td>
<td>71</td>
<td>–</td>
<td>Normal LV dilatation</td>
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<td>–</td>
</tr>
<tr>
<td>5</td>
<td>6</td>
<td>1.5</td>
<td>0</td>
<td>65</td>
<td>+</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>6</td>
<td>10</td>
<td>0 stroke</td>
<td>96</td>
<td>–</td>
<td>Normal Normal</td>
<td>–</td>
<td>–</td>
<td>–</td>
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<tr>
<td>7</td>
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<td>0</td>
<td>94</td>
<td>+</td>
<td>Normal Normal</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>8</td>
<td>10</td>
<td>ND</td>
<td>0</td>
<td>75</td>
<td>–</td>
<td>Normal Normal</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

VOC, vasoocclusive crises; ACS: acute chest syndrome; LV, left ventricular; ND, not determined.

Figure 1 $^{201}$Tl myocardial SPECT: reversible defect in the anterior wall, left ventricular cavity dilated.

Table 2 children with abnormal myocardial perfusion

<table>
<thead>
<tr>
<th>Pt</th>
<th>Age (y)</th>
<th>Nb VOC/y</th>
<th>ACS other</th>
<th>Hb (g/l)</th>
<th>Chest pains</th>
<th>ECG</th>
<th>Echo</th>
<th>Stress testing</th>
<th>Defect</th>
<th>Treatment outcome</th>
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</thead>
<tbody>
<tr>
<td>9</td>
<td>5</td>
<td>0</td>
<td>0</td>
<td>80</td>
<td>Heart failure</td>
<td>Abnormal LV dilatation</td>
<td>+</td>
<td>Reversible</td>
<td>HU partial correction</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>16.5</td>
<td>0</td>
<td>0</td>
<td>75</td>
<td>Arrhythmia</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Fixed</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>16</td>
<td>1.5</td>
<td>1</td>
<td>66</td>
<td>–</td>
<td>Normal LV dilatation</td>
<td>+</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>16</td>
<td>0</td>
<td>0</td>
<td>59</td>
<td>–</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>13</td>
<td>16</td>
<td>1.5</td>
<td>0</td>
<td>77</td>
<td>+</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>14</td>
<td>14</td>
<td>0</td>
<td>0</td>
<td>55</td>
<td>–</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>HU partial correction</td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>16.5</td>
<td>0</td>
<td>0</td>
<td>67</td>
<td>–</td>
<td>Abnormal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>HU partial correction</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>14</td>
<td>0</td>
<td>0</td>
<td>90</td>
<td>Heart failure</td>
<td>Abnormal LV dilatation</td>
<td>+</td>
<td>Reversible</td>
<td>HU partial correction</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>18</td>
<td>6</td>
<td>1</td>
<td>70</td>
<td>–</td>
<td>Abnormal LV dilatation</td>
<td>+</td>
<td>Reversible</td>
<td>HU partial correction</td>
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<td>18</td>
<td>3.5</td>
<td>1</td>
<td>2</td>
<td>75</td>
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<td>Normal LV dilatation</td>
<td>–</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
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<tr>
<td>19</td>
<td>19</td>
<td>3</td>
<td>3</td>
<td>85</td>
<td>+</td>
<td>Abnormal Normal</td>
<td>+</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>14</td>
<td>0</td>
<td>0</td>
<td>89</td>
<td>+</td>
<td>Normal –</td>
<td>–</td>
<td>Reversible</td>
<td>HU</td>
<td></td>
</tr>
<tr>
<td>21</td>
<td>13</td>
<td>ND</td>
<td>0</td>
<td>74</td>
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<td>Abnormal LV dilatation</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td></td>
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<td>22</td>
<td>11</td>
<td>1.5</td>
<td>4 stroke</td>
<td>82</td>
<td>–</td>
<td>Normal normal</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td></td>
</tr>
</tbody>
</table>

VOC, vasoocclusive crises; ACS: acute chest syndrome; LV, left ventricular; ND, not determined; HU, hydroxyurea; BMT, bone marrow transplantation.
Echocardiography showed an enlarged hypokinetic left ventricle (shortening fraction: 20%). She was treated with diuretics and angiotensin converting enzyme inhibitors. Two years later, she had no signs of cardiac failure, but her left ventricle was still enlarged and hypokinetic. Stress testing was positive and $^{201}$TI SPECT showed an anteroseptal reversible perfusion defect. The child was then lost to follow up. She was hospitalised again at 7 years with severe cardiac failure (shortening fraction: 13%) and died within days despite intensive cardiac resuscitation.

Patient 10 had ventricular fibrillation when he was 8 years old, followed the day after by constrictive chest pain, posterior subepicardial ischaemia, and severe heart failure. Q-T prolongation led to insertion of an implantable pace-maker. Eight years later, the patient had no new episode of arrhythmia and no signs of heart failure. $^{201}$TI SPECT showed a fixed inferior perfusion defect, probably related to myocardial necrosis.

Among the 12 other patients with myocardial perfusion defects, patient 22, who had had a cerebrovascular infarct, received a bone marrow transplant in 2002, and patient 21 returned to Africa. Hydroxyurea treatment was begun in eight children who had myocardial perfusion defects and other complications of SCD: more than one hospitalised painful crisis/year (patients 11, 13, 17), severe anaemia (patients 14, 15), past history of acute chest syndromes (patients 18, 19), or a past history of acute cardiac failure (patient 16).

**Course under hydroxyurea treatment**
These eight patients were treated with hydroxyurea at a mean dosage of 20 mg/kg/day. SPECT was repeated after six months of treatment in three. Two patients had reversible defects that had improved, and one had a fixed defect, the size of which had decreased.

**DISCUSSION**
Our study shows myocardial perfusion defects in children with SCD, which can be demonstrated and followed with $^{201}$TI SPECT. This finding, which to our knowledge has never been reported in children with SCD, raises several questions. The first is the prevalence of these perfusion defects. In our paediatric population selected according to strict criteria (chest pain, heart failure, abnormal ECG, left ventricular dilatation, or hypokinetic left ventricle), this prevalence was as high as 14/22 (63.6%). Prevalence depends clearly on the criteria used to perform SPECT. All the children in our series who had had an overt symptom of cardiac disease (heart failure episodes for patients 9, 16, 18; ventricular arrhythmia for patient 10; angina for patient 19) had perfusion defects. The relevance of ECG or echographic abnormalities may be more problematic to interpret in black children and adolescents, since some changes are considered to be non-specific.2–9 However, we noted that more children had an abnormal ECG (9/14) and left cardiac enlargement (11/14) in our series than in adults17–19 and has been found to be related to the severity of anaemia and percentage of HbS.21,22 Our findings are in favour of a specific ischaemic cardiomyopathy that could worsen the consequences of the increased cardiac output. Furthermore, some children receive very frequent transfusions, and are exposed to the cardiac toxicity of iron overload.

For about 10 years, hydroxyurea has been recognised as having a beneficial effect on the prognosis of sickle cell disease in adults23 and in children.24–26 This cytostatic drug probably alleviates the severity of the disease via a number of pathways: induction of fetal haemoglobin synthesis (which has a sparing effect on polymerisation of haemoglobin S), reduction in reticulocyte adhesion to vascular endothelium,27 modulation of inflammatory processes, induction of NO synthesis,28 etc. Although we do not yet have evidence that hydroxyurea acts directly on vascular remodelling, all these indirect arguments led us to hypothesise that hydroxyurea could at least partially correct the hypoperfusion; we therefore decided to administer it to selected children. However, due to our uncertainty about the long term effects of this cytostatic drug, we restricted this treatment to children for whom we also had other grounds for considering hydroxyurea treatment (for example, 1–2 hospitalised painful crises/
In conclusion, we report that myocardial perfusion abnormalities can be demonstrated using $^{201}$TI SPECT in children affected with SCD. Those with myocardial perfusion abnormalities tend to be older, have a lower Hb level, and more frequent vasoocclusive crises. Perfusion defects may be reversible or fixed and may lead to cardiac failure or arrhythmia. Improvement in cardiac perfusion was observed in three children of our series who were treated with hydroxyurea.

Finally, we postulate that cardiac impairment is underestimated in children because it is under-investigated. Chest pain is usually attributed to bone infarcts and cardiomegaly to chronic anaemia. As a result of our findings, we now intend to perform ECG systematically in cases of chest pain and/or cardiac enlargement. For the moment, we plan to keep the same clinical, ECG, and echographic criteria for deciding whether or not to perform SPECT. If we confirm in more cases that hydroxyurea in fact improves cardiac perfusion, we will consider that the hypothetical risk of malignancy will be counterbalanced by a decrease in the risk of myocardial impairment.

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