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## Drugs for preventing red blood cell dehydration in people with sickle cell disease.

Srikanth Nagalla  
*Thomas Jefferson University*

Samir K. Ballas  
*Thomas Jefferson University*

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## Drugs for preventing red blood cell dehydration in people with sickle cell disease (Review)

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Drugs for preventing red blood cell dehydration in people with sickle cell disease.

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[Intervention Review]

# Drugs for preventing red blood cell dehydration in people with sickle cell disease

Srikanth Nagalla<sup>1</sup>, Samir K Ballas<sup>2</sup>

<sup>1</sup>Department of Medicine, Division of Hematology, Thomas Jefferson University, Philadelphia, Pennsylvania, USA. <sup>2</sup>Cardeza Foundation for Hematologic Research, Department of Medicine, Jefferson Medical College, Thomas Jefferson University, Philadelphia, USA

Contact address: Samir K Ballas, Cardeza Foundation for Hematologic Research, Department of Medicine, Jefferson Medical College, Thomas Jefferson University, 1015 Walnut Street, Philadelphia, PA 19107-5099, USA. [Samir.Ballas@jefferson.edu](mailto:Samir.Ballas@jefferson.edu).

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## ABSTRACT

### Background

Sickle cell disease is an inherited disorder of hemoglobin, resulting in abnormal red blood cells. These are rigid and may block blood vessels leading to acute painful crises and other complications. Recent research has focused on therapies to rehydrate the sickled cells by reducing the loss of water and ions from them. Little is known about the effectiveness and safety of such drugs. This is an updated version of a previously published review.

### Objectives

To assess the relative risks and benefits of drugs to rehydrate sickled red blood cells.

### Search methods

We searched the Cochrane Cystic Fibrosis and Genetic Disorders Group's Haemoglobinopathies Trials Register.

Last search of the Group's Trials Register: 28 November 2015.

### Selection criteria

Randomized or quasi-randomized controlled trials of drugs to rehydrate sickled red blood cells compared to placebo or an alternative treatment.

### Data collection and analysis

Both authors independently selected studies for inclusion, assessed study quality and extracted data.

### Main results

Of the 51 studies identified, three met the inclusion criteria. The first study tested the effectiveness of zinc sulphate to prevent sickle cell-related crises in a total of 145 participants and showed a significant reduction in painful crises over one and a half years, mean difference -2.83 (95% confidence interval -3.51 to -2.15). However, analysis was restricted due to limited statistical data. Changes to red cell parameters and blood counts were inconsistent. No serious adverse events were noted in the study.

The second study was a Phase II dose-finding study of senicapoc (a Gardos channel blocker) compared to placebo. Compared to the placebo group the high dose senicapoc showed significant improvement in change in hemoglobin level, number and proportion of

dense red blood cells, red blood cell count and indices and hematocrit. The results with low-dose senicapoc were similar to the high-dose senicapoc group but of lesser magnitude. There was no difference in the frequency of painful crises between the three groups. A subsequent Phase III study of senicapoc was terminated early since there was no difference observed between the treatment and control groups in the primary end point of painful crises.

### **Authors' conclusions**

While the results of zinc for reducing sickle-related crises are encouraging, larger and longer-term multicenter studies are needed to evaluate the effectiveness of this therapy for people with sickle cell disease.

While the Phase II and the prematurely terminated phase III studies of senicapoc showed that the drug improved red cell survival (depending on dose), this did not lead to fewer painful crises.

We will continue to run searches to identify any potentially relevant trials; however, we do not plan to update other sections of the review until new trials are published.

## **PLAIN LANGUAGE SUMMARY**

### **Drugs that aim to reduce the loss of water from red blood cells in people with sickle cell disease**

#### **Review question**

We reviewed the evidence to assess the relative risks and benefits of drugs to rehydrate sickled red blood cells.

#### **Background**

Sickle cell disease is an inherited condition that causes red blood cells to become sickle shaped when they lose water. This leads to a high risk of the blood vessels becoming blocked. Such blockages can cause pain, stroke and damage to organs. Recent therapies aim to stop the cells becoming sickle shaped by preventing them losing water.

#### **Search date**

The evidence is current to: 28 November 2015.

#### **Study characteristics**

The review included three studies with 524 people with sickle cell disease aged between 12 and 65 years of age. The intervention in one study was zinc sulphate and in two studies was senicapoc. Each study was compared to a placebo group (a substance which contains no medication). For each study people were selected for one treatment or the other randomly. The studies lasted from three months to 18 months.

#### **Key results**

The study with zinc sulphate showed that this drug may be able to reduce the number of sickle cell crises without causing toxic effects. There were 145 participants in this study and results showed a significant reduction in the total number of serious sickle-related crises over one and a half years, mean difference -2.83 (95% confidence interval -3.51 to -2.15). However, our analysis was limited since not all data were reported. Changes to red cell measurements and blood counts were not consistent. No serious adverse events were noted in the study. The two studies with senicapoc demonstrated that this drug increases the red blood survival and has a role in the prevention on red blood cell dehydration in people with sickle cell disease. The higher dose of the drug was more effective compared to the lower dose. But these changes in the red blood cells did not translate into positive clinical outcomes in terms of reduction in the number of sickle cell crises. Senicapoc had a favourable safety profile. More longer-term research is needed on these drugs and others that might prevent water loss in red blood cells.

We will continue to run searches to identify any potentially relevant trials; however, we do not plan to update other sections of the review until new trials are published.

#### **Quality of the evidence**

All the studies appeared to be well run and we do not think any factors will influence the results in a negative way.

## BACKGROUND

### Description of the condition

Sickle cell disease is caused by the inheritance from both parents of a mutation in the beta-globin gene. The different types of the disease (e.g. SS, SC, S $\beta$ Thal) are caused by inheritance of different altered beta-globin genes, and this can be diagnosed with blood tests or genetic testing. It is so named because the red blood cells (erythrocytes) become distorted into unusual shapes when they give up the oxygen they carry, due to polymerisation of the abnormal hemoglobin within. This is accelerated by a cycle of red blood cell dehydration, which increases intracellular concentration of the sickle hemoglobin and hence polymerisation. One of the distinguishing features of sickle cell anemia is the presence of these dense, dehydrated red cells, which are easily destroyed and can mesh together to block blood vessels (vaso-occlusion).

The disease is characterised by episodes of anemia, vaso-occlusive crises and organ damage, with symptoms which are unpredictable and differ between individuals (Serjeant 1992). It is most prevalent in populations originating from sub-Saharan Africa and parts of India; although population movement has made it a worldwide problem, with approximately 60,000 African-American and 10,000 British people with the disease (Davies 1997; Hickman 1999). Clinical management centers around prophylaxis against infection, pain relief, hydration and periodic blood transfusion in many people to reduce the concentration of sickle cells in the blood stream. However, most treatment is symptomatic and, as a result, despite vastly improving care services, many people with the disease still die in childhood due to sudden bouts of infection, stroke, acute chest syndrome or splenic sequestration (Leikin 1989). Clearly preventative measures would be preferable.

### How the intervention might work

In an attempt to prevent the increased concentration of sickle hemoglobin (which causes polymerisation and deformation of the red blood cell), recent research has begun to target the physiological process leading to dehydration of the erythrocyte for therapeutic potential. Red cells are thought to undergo dehydration via two kinetically distinct pathways, a “fast track” process in reticulocytes (immature red blood cells), and a slow process involving mature erythrocytes. Dehydration via either pathway is due to the water loss accompanying primary loss from the cell of potassium and chloride ions. The major routes involved in sickle cell dehydration are the calcium-activated potassium channel (Gardos channel) operating in parallel with the conductive chloride pathway and the electroneutral potassium-chloride co-transporter(s) (Brugnara 1995). Potential therapeutic approaches involve the use of drugs that reduce sickle cell dehydration via a block of these mechanisms.

The first route, and perhaps the most promising for therapy, is the Gardos channel. Upon deoxygenation of the sickle cell, potassium and water loss are induced as free intracellular calcium levels rise, possibly due to increased membrane permeability for calcium. The Gardos channel may also be positively modulated by vaso-active molecules such as endothelins (Rivera 2000). Studies both in a mouse model for sickle cell disease (SAD mouse) and in humans have shown that this channel can be selectively inhibited by a group of drugs known as the imidazole antimycotics, including the anti-fungal agent clotrimazole (Brugnara 1993; Brugnara 1996; de Franceschi 1994) and a related compound Senicapoc (ICA-17043) (Stocker 2000). Blocking of the Gardos channel resulted in increased potassium content and improved hydration of erythrocytes. Senicapoc (bis(4-fluorophenyl)phenyl acetamide) is a selective and potent blocker of the Gardos channel (Stocker 2003). Dimethyl adipimate may also have an anti-sickling role via the Gardos channel (Gibson 2000).

The role of the chloride conductance or exchange system in sickle cell dehydration has also been recently explored. In vitro, loss of potassium from deoxygenated human sickle cells could be limited by the reversible anion conductance inhibitor NS1652 (Bennekou 2000). Further, the related compound, NS3623 (Bennekou 2001), which has a greater half-life in vivo, improved hydration in the SAD mouse.

Potassium-chloride co-transport occurs via a different membrane protein, which is activated by acidity within the cell allowing loss of potassium and water. This is seen almost exclusively in reticulocytes. No drugs have been found to be useful in inhibiting the transporter, but it has been shown that divalent cations, such as magnesium ions (Brugnara 1987), can block the pathway, again protecting against dehydration. Magnesium preparations are currently marketed for treatment of constipation, boils, recurrence of seizures in eclampsia, renal failure and are recommended for emergency treatment of serious arrhythmias. Magnesium pidolate has been investigated in combination with hydroxyurea in a Phase I trial (Hankins 2008).

Other strategies have also been suggested for reducing sickle red cell dehydration. These include a transmembrane sodium-potassium pump mechanism which is partly responsible for increased ion permeability on deoxygenation. While this is thought to be a minor contributor to dehydration, dipyridamole has been shown to be a selective inhibitor of passive ion transfer and there is rationale, including the known effects on platelet and endothelial cell function, for testing its efficacy in sickle cell disease. Oxidative damage to the cell membrane, which can occur in sickle cell disease due to abnormal iron deposits, increases loss of potassium and water contributing to red cell dehydration (Brugnara 1995). Oral iron chelators may help in this case, and antioxidants could also be useful (Gibson 1998). Nitric oxide, an endogenous compound which is thought to be an important regulator of smooth muscle tone and vasodilatation, is reduced in people with sickle

cell disease (Qureshi 2000). Preliminary studies in SAD mice have shown that it can reduce red cell dehydration (Adrie 2000), and oral supplementation with arginine, an amino acid which releases nitric oxide, (5% by weight) also induces a decrease in red cell density (Romero 2000), via Gardos channel activity (Romero 2002). Hydroxyurea, an anti-cancer compound proven to reduce sickle-related events (Charache 1995; Jones 2001), improves the hydration status of sickle red blood cells (Ballas 1989), although it is not clear whether this is a direct effect, or a consequence of increased levels of fetal hemoglobin (HbF).

In Brazil a local plant, *Pfaffia paniculata*, has been anecdotally reported to ameliorate clinical symptoms of sickle cell disease. The plant possibly contains a naturally occurring compound, which acts as a sodium ionophore. 'In vitro' studies have shown increased erythrocyte deformability and sodium content of sickle red cells when treated with it, thus potentially rehydrating sickle cells (Ballas 2000).

Piracetam (2-oxo-1-pyrrolidine acetamide), a derivative of the neurotransmitter gamma-aminobutyric acid (GABA), works primarily in neurotransmission, but appears to have a beneficial effect on red cell rheology and thus able to reduce vaso-occlusive crises in people with sickle cell disease. The exact mechanism of this remains unclear. It does not appear to inhibit the Gardos channel as previously reported (Gini 1987; Stone 1988; Stuart 1990; Stuart 1992). The use of piracetam for reducing the incidence of painful sickle cell disease crises is the subject of another Cochrane Review (Al Hajeri 2007).

### Why it is important to do this review

All of these potential therapies look promising for reducing dehydration of red blood cells, and therefore ameliorating the symptoms of sickle cell disease. This review aims to bring together clinical trials in this area to establish the clinical value of this pharmaceutical approach.

This is an update of previously published versions of the review (Nagalla 2010; Nagalla 2012; Riddington 2002; Singh 2007).

## OBJECTIVES

To determine whether pharmaceutical therapies designed to prevent red cell dehydration in sickle cell disease:

1. reduce mortality;
2. reduce sickle cell vaso-occlusive crises including episodes of pain and stroke;
3. reduce other complications associated with sickle cell disease e.g. acute chest syndrome, infection, anemia, splenic sequestration, organ damage;

4. are associated with unacceptable adverse events.

## METHODS

### Criteria for considering studies for this review

#### Types of studies

Randomized or quasi-randomized trials. Trials in which quasi-randomized methods, such as alternation, are used were included if there was sufficient evidence that the treatment and control groups were similar at baseline.

#### Types of participants

People with known sickle cell disease (SS, SC, S $\beta$ + and S $\beta$ 0, proven by electrophoresis and sickle solubility test, with family studies or DNA tests as appropriate) of all ages and both sexes, in any setting.

#### Types of interventions

Pharmaceutical therapies designed to reduce red cell dehydration to prevent vaso-occlusive events in sickle cell disease compared to comparator interventions. Studies which included anti-sickling drugs for which the exact mechanism of action is not known were considered if there was some evidence that they act via the red blood cell membrane to prevent sickling. Piracetam was excluded since there is insufficient evidence that it acts primarily via the red cell membrane to improve red cell rheology. Hydroxyurea was excluded as the primary mechanism for this drug is through increasing foetal hemoglobin. Studies in which the intervention was given only to treat an existing sickle-related event were excluded.

#### Types of outcome measures

While the interventions should, by definition, decrease dense sickled cells by reducing erythrocyte dehydration, such biochemical parameters were examined separately from clinical outcomes, to determine the effectiveness of the interventions in clinical practice. However, the hematological outcomes may also provide an indication of the efficacy, and potential effectiveness, of treatment, so these were also analyzed.

#### Primary outcomes

1. Number of deaths
2. Number of sickle pain crises (requirement for opiate treatment or self-reported patient scales)
3. Number of serious complications of sickle cell disease including stroke, acute chest syndrome and acute splenic sequestration

## Secondary outcomes

1. Red cell dehydration (proportion of dense cells, mean corpuscular hemoglobin concentration (MCHC), mean corpuscular volume (MCV))
2. Full blood count including hemoglobin, white blood cell and platelet counts, sickle cell hemoglobin (HbS) and foetal hemoglobin (HbF) level
3. Quality of life: days off work or school, hospitalisations, mobility, etc.
4. Number of other sickle-related events, including priapism, leg ulceration

## Other outcomes

1. Any reported adverse effects or toxicity of drugs were recorded

## Search methods for identification of studies

### Electronic searches

Relevant studies were identified from the Cystic Fibrosis & Genetic Disorders Review Group's Haemoglobinopathies Trials Register using the terms: (sickle cell OR (haemoglobinopathies AND general)) AND (dehydration OR (anti-sickling OR crisis)).

The Haemoglobinopathies Trials Register is compiled from electronic searches of the Cochrane Central Register of Controlled Trials (CENTRAL) (updated each new issue of *The Cochrane Library*) and weekly searches of MEDLINE. Unpublished work is identified by searching the abstract books of five major conferences: the European Haematology Association conference; the American Society of Hematology conference; the British Society for Haematology Annual Scientific Meeting; the Caribbean Health Research Council Meetings; and the National Sickle Cell Disease Program Annual Meeting. For full details of all searching activities for the register, please see the relevant section of the [Cystic Fibrosis and Genetic Disorders Group Module](#).

Date of the most recent search of the Group's Haemoglobinopathies Trials Register: 28 November 2015.

### Searching other resources

The bibliographic references of all retrieved literature were reviewed for additional reports of studies.

## Data collection and analysis

### Selection of studies

Two authors independently applied the inclusion criteria in order to select studies for inclusion in the review. If disagreement arose on the suitability of a study for inclusion in the review, the authors reached consensus by discussion.

### Data extraction and management

Two authors independently extracted the data. Each author, using standard data acquisition forms, independently extracted data. If disagreement arose on the study quality, the authors reached a consensus by discussion.

We planned to collect data at monthly time-frames. However, if data were reported at other time periods we planned to consider examining these as well.

### Assessment of risk of bias in included studies

Two authors assessed the risk of bias of each study. In particular, authors examined details of the randomization method, allocation concealment, whether the study was blinded, whether intention-to-treat analyses were possible from the available data and if the number of participants lost to follow up or subsequently excluded from the study was recorded.

### Measures of treatment effect

We recorded dichotomous outcomes e.g. life or death, as present or absent, whilst recording continuous data such as organ function tests as either mean change from baseline for each group or mean post treatment values and standard deviation (SD) for each group. We aimed to calculate a pooled estimate of the treatment effect for each outcome across studies, (for binary outcomes the odds of an outcome among treatment allocated participants to the corresponding odds among controls).

### Unit of analysis issues

Cross-over trials will only be included in future versions of this review if we consider there to be a sufficient washout period between the treatment arms. We will analyze any data from such trials according to methods described by Elbourne ([Elbourne 2002](#)).

### Dealing with missing data

We sought full reports from authors where studies have been published in abstract form, presented at meetings or reported to the co-authors. Where information was missing or unclear, we contacted the primary investigator.

In order to allow an intention-to-treat analysis, we grouped data by allocated treatment groups, irrespective of later exclusion (regardless of cause) or loss to follow up.



## Assessment of heterogeneity

We will test for heterogeneity between studies using a standard chi-squared test and  $I^2$  statistic (Higgins 2003). We will use the following ranges to describe heterogeneity:

- 0% to 40%: might not be important;
- 30% to 60%: may represent moderate heterogeneity;
- 50% to 90%: may represent substantial heterogeneity;
- 75% to 100%: considerable heterogeneity.

## Assessment of reporting biases

Comprehensive searches were done by both the authors to minimize publication and reporting biases. We compared the 'Methods' section of the full published paper to the 'Results' section to ensure that all outcomes which were measured, were reported.

## Data synthesis

We have used fixed-effect analysis with the data we have presented. If there is a high or moderate degree of heterogeneity between studies included in any future updates, we will consider using a random-effects analysis.

## Subgroup analysis and investigation of heterogeneity

If we find heterogeneity between studies, examination of subgroups, such as age of participants, type of sickle cell disease or ethnicity, may help to explain the reasons for this. For future updates, where appropriate, we plan to perform subgroup analysis of different drugs or combinations of drugs to examine their relative risks and benefits.

## Sensitivity analysis

If we include a sufficient number of studies where quasi-randomization methods are used, we will analyze this group separately.

# RESULTS

## Description of studies

### Results of the search

Fifty-one studies of anti-sickling agents were identified from the searches. Three studies were included within the review (Ataga 2008; Ataga 2011; Gupta 1995).

## Included studies

### Trial design

All three studies were described as double-blind, randomized placebo-controlled studies (Ataga 2008; Ataga 2011; Gupta 1995). The Phase II Ataga study was a dose-finding study which compared a 10 mg dose of senicapoc to a 6 mg dose of senicapoc or to placebo. The trial was multicenter (19 medical centers) in the United States of America (Ataga 2008). The study treated participants for 12 weeks (Ataga 2008). The recent senicapoc trial was a Phase III trial was conducted across 75 centers in 6 countries (Ataga 2011). The participants were randomized to senicapoc (20 mg twice daily as a loading dose for 4 days followed by 10 mg daily) or placebo in a 1:1 fashion. The treatment phase consisted of 52 weeks followed by a follow-up period of 8 weeks (Ataga 2011). The Gupta trial was from a single center in India (Gupta 1995). The Gupta study lasted for one and a half years; participants were seen weekly but data were analysed at the end of the follow-up period (Gupta 1995).

### Participants

There were 524 participants with sickle cell disease included in the review. The Phase II senicapoc study randomized 90 participants aged between 18 to 60 years (Ataga 2008). The Phase III senicapoc study randomized 297 participants between the ages of 16 and 65 years, but only the 289 participants in the modified intent-to-treat group (participants who received at least one dose of the drug) were included in this review (Ataga 2011). In the remaining study, Gupta randomized 145 participants aged between 12 to 27 years (Gupta 1995).

In the Phase II senicapoc study, participants had at least one episode of sickle cell-related painful crisis four weeks prior to the study screening and participants taking hydroxyurea had to be on a stable dose of the drug for at least three months prior to their enrolment in the study (Ataga 2008). In the Phase III senicapoc study, participants had at least two episodes of painful crisis within the preceding 12 months that required medical attention (Ataga 2011). Participants on hydroxyurea in preceding 12 months were required to be on a stable dose of the drug prior to inclusion in the study (Ataga 2011). Gupta excluded participants under five years of age or those who had a history of chronic persistent infection, evidence of organ failure, exposure to extreme temperatures or were taking other medication (Gupta 1995).

In the Phase II senicapoc study, all the participants who received at least one week of the drug and underwent the initial efficacy analysis were included in the modified intent-to-treat analysis. Eighty-eight participants met this criteria and only two discontinued the study before the initial efficacy analysis (Ataga 2008). In the Phase III senicapoc study 289 participants from the initial group of 297 were included in the modified intent-to-treat analysis based on participants who received at least one dose of the drug (Ataga

2011). However, in the Gupta study, 15 participants out of the original 145 subsequently dropped out or were lost to follow-up, and these were not included in the analysis (Gupta 1995).

### Interventions

The Phase II senicapoc study compared three groups: high-dose senicapoc (single 150 mg loading dose followed by 10 mg daily maintenance dose); or low-dose senicapoc (single 100 mg loading dose followed by 6 mg daily maintenance dose); or placebo (Ataga 2008). The high-dose senicapoc group had 31 participants, the low-dose senicapoc and placebo groups had 29 and 30 participants assigned respectively. The Phase III senicapoc study stratified participants based on concomitant hydroxyurea therapy. In each of these groups participants were randomized to receive senicapoc (loading dose of 20 mg twice daily for four days followed by 10 mg daily) or placebo in a 1:1 fashion. Out of the 289 participants in the modified intent-to treat analysis, there were 163 participants in the hydroxyurea group and 126 in the non-hydroxyurea group. The hydroxyurea group had 84 participants receiving senicapoc and 79 participants on placebo. The non-hydroxyurea group had 61 senicapoc participants and 65 in the placebo group (Ataga 2011). The third study randomized participants to receive either zinc sulphate (220 mg three times a day orally) or an identical placebo (Gupta 1995).

### Outcomes measured

The Ataga Phase II study reported the effect of senicapoc on hemoglobin levels, red blood cell (RBC) indices, markers of hemolysis and painful crises. The primary end point of the study was the change in hemoglobin level with the secondary end points being changes in the number and proportion on the dense red blood cells (RBCs) and reticulocytes, lactate dehydrogenase, indirect bilirubin, RBC count, RBC indices, hematocrit and frequency of painful crises (Ataga 2008). The primary end point in the Ataga Phase III was the frequency of sickle cell pain crises. Time to the first, second and third acute painful crises was a secondary end point of the study (Ataga 2011). The effect of the drug on markers of hemolysis, hemoglobin, hematocrit, reticulocyte count, red blood cell count and dense erythrocytes were also measured. The Gupta study measured the number of sickle-related crises, and the number of days in hospital and working (or school) days lost for each crisis (Gupta 1995).

### Excluded studies

Forty-eight studies were excluded. Reasons for exclusions were a lack of evidence that they were acting primarily on red cell dehydration pathways, or they were being used to treat a sickle-related crisis, rather than preventing it, or the study was not randomized (see [Characteristics of excluded studies](#) table). Studies on piracetam were excluded since there is insufficient evidence that this

drug acts primarily via the red cell membrane to improve red cell rheology (Alvim 2005; Piracetam Study 1998).

### Risk of bias in included studies

#### Allocation

All three studies were judged to have a low risk of bias for the generation of the allocation sequence (Ataga 2008; Ataga 2011; Gupta 1995). In both of the senicapoc studies an integrated voice response system was used in the centralized randomization protocol (Ataga 2008; Ataga 2011). In the zinc study, New Castle software was used to generate randomization (Gupta 1995).

Both the senicapoc studies and the zinc study did not report on allocation concealment and so all studies were judged to have an unclear risk of bias for this criteria (Ataga 2008; Ataga 2011; Gupta 1995).

#### Blinding

Both the participants and treating physicians were blinded in all three studies and so were judged to have a low risk of bias (Ataga 2008; Ataga 2011; Gupta 1995). Additionally, Ataga reported that the review committee were also blinded (Ataga 2008).

#### Incomplete outcome data

In the senicapoc Phase II study, a modified intent-to-treat population was used in the efficacy analysis (Ataga 2008). Two out of the 90 participants dropped out prior to the initial assessment and so 88 participants were included as a part of the modified intent-to-treat population. The safety analysis included all the 90 participants. A total of 10 participants did not complete the study; full reasons for this were given in the published paper and we therefore judged there to be a low risk of bias due to incomplete outcome data (Ataga 2008).

The Phase III senicapoc study was terminated early due to a lack of efficacy as determined by the unblinded Data Monitoring Committee (Ataga 2011). The reasons for the Committee's decision were based on the fact that there was no significant improvement in the rate of sickle cell painful crises in participants treated with senicapoc compared to those on placebo ( $P = 0.38$  versus  $P = 0.31$  respectively) despite improvements in anemia and hemolysis. Moreover, comparisons of the times to first, second and third crises between the senicapoc and placebo groups were not significantly different. The authors suggest that the increase in Hb level caused an increase in blood viscosity and offset the potential benefits of senicapoc with regards to painful crises. Furthermore, due to the early termination of the study, the duration that the drug was administered was different in the participants who stopped treatment early. Given these facts, we have not entered any data

from this trial into the data tables and results are reported narratively.

In the zinc study, a power calculation, which assumed 40% control event rate and 20% response, found that 130 participants were needed. Fifteen, of the 145 participants recruited, dropped out or were lost to follow-up, and these are not accounted for in the study publication (Gupta 1995). We therefore judge there to be an unclear risk of bias from reported outcome data for this study.

### Selective reporting

In both the senicapoc studies and the zinc study, all outcomes stated in 'Methods' section were reported in the 'Results' section of the papers; therefore there is a low risk of bias for selective reporting for all studies (Ataga 2008; Ataga 2011; Gupta 1995).

### Other potential sources of bias

The participants in the Phase II senicapoc study were evenly matched in terms of their baseline characteristics between the three study arms (Ataga 2008) and this was also true for the Phase III senicapoc study (Ataga 2011). There was no significant difference in the mean age, sex and the hospitalizations in the year prior to the enrolment in the study. We therefore judged both studies to have a low risk of bias for this domain.

The zinc study gave little information regarding baseline characteristics, including disease severity and medical history of participants, making it difficult to ascertain if the groups were sufficiently similar at the start of the study (Gupta 1995). Thus currently we judge there to be an unclear risk of bias for this study.

## Effects of interventions

### Primary outcomes

#### 1. Number of deaths

No treatment related deaths were reported in the included studies (Gupta 1995; Ataga 2008; Ataga 2011).

#### 2. Number of pain crises

In the senicapoc Phase II study there were a total of 15 sickle cell crises, with the events being distributed equally between the placebo and the two intervention arms (five in each) (Ataga 2008). So both the lower (6 mg) and higher doses (10 mg) of senicapoc did not help in the reduction of painful crises compared to placebo. The Phase III senicapoc study was terminated prematurely as it was unlikely to meet the primary endpoint (Ataga 2011). There was no significant difference in the rate of the sickle cell painful crises between the senicapoc and placebo groups (0.38 versus 0.31,  $P =$

0.054). There was no difference in the painful crises rate between the two treatment arms in the hydroxyurea group (0.39 versus 0.33,  $P = 0.483$ ) but the rate of painful crises was significantly higher in the senicapoc group compared to the placebo group in participants not on hydroxyurea (0.37 versus 0.29,  $P = 0.037$ ) (Ataga 2011). The study included pain-only crises, acute chest syndrome, hepatic sequestration, priapism and stroke under the primary end point of sickle cell-related pain crises. Given that the study was terminated prematurely, there are very few participants who completed the treatment. The duration of the drug regimen was different in the participants who stopped treatment. Therefore we have not included any data from this trial in the data analysis table.

In the zinc study there were 91 vaso-occlusive crises in the intervention group, compared to 220 in the control group (Gupta 1995). This amounts to, on average, 1.40 and 3.38 painful crises per participant in intervention and control groups respectively. Standard deviations cannot be calculated from the data provided in the publication, so we are attempting to contact the authors, and hope to provide this information in a future update.

### 3. Number of other serious sickle-related complications

In the Phase II senicapoc study, there were two episodes of pneumonia and one episode of bronchitis in the placebo group (Ataga 2008). There was one episode of pneumonia in each of the intervention arms. There was a single episode of acute chest syndrome in the study and it occurred in the 6 mg (low-dose) senicapoc arm. There was one episode of aseptic necrosis of the bone in the low-dose senicapoc arm and one episode of deep venous thrombosis in the high-dose senicapoc arm. Furthermore, there was one episode of staphylococcal sepsis in the low-dose senicapoc group and one urinary tract infection in the high-dose senicapoc group. The only case of deep vein thrombosis was reported in the high-dose senicapoc group (Ataga 2008).

The Phase III senicapoc study included pain-only crises, acute chest syndrome, hepatic sequestration, priapism and stroke under the primary end point of sickle cell-related pain crises (Ataga 2011).

In the zinc study, altogether there were 160 episodes of sickle-related crises in the intervention group, and 344 in the control group (Gupta 1995). Of these, vaso-occlusive crises accounted for 59% in the intervention group and 64% in the control group, mixed crises for 24% and 20% in the intervention and control groups respectively, 9% and 12% were hemolytic, 1% in each group sequestration and 1% in each group had aplastic crises. The mean number of all sickle-related crises in the intervention group was 2.46 (SD 1.04), compared to 5.29 (SD 2.58), in the control group. The MD between groups was -2.83 (95% CI -3.51 to -2.15), showing a statistically significant reduction in the participants treated with zinc (Gupta 1995).

## Secondary outcomes

### 1. Red cell dehydration

Both the high and low doses of senicapoc resulted in a significant decrease in the percentage of dense red blood cells compared to the placebo (Ataga 2008). The magnitude of decrease was higher with the 10 mg senicapoc compared to the 6 mg senicapoc. There was a significant decrease in the MCV and mean corpuscular hemoglobin (MCH) in the high-dose senicapoc group compared to the placebo. The low-dose senicapoc caused a significant reduction in the MCH but not MCV when compared to the placebo. There was no significant difference in MCHC between the two intervention arms and the placebo arm. The biochemical markers of hemolysis like lactate dehydrogenase and indirect bilirubin were significantly lower in the senicapoc groups compared to the placebo group. The magnitude of the change was dependent on the dose of senicapoc.

The findings seen in the Phase II senicapoc study were confirmed in the Phase III study where senicapoc was compared to a placebo (Ataga 2011).

The Gupta paper reported significant improvements in laboratory parameters of red cell dehydration (hemoglobin, reticulocyte count, serum bilirubin and serum zinc level) in the intervention group (Gupta 1995).

### 2. Full blood count

The senicapoc study demonstrated an increase in the hemoglobin level by 0.68 gm/dl in the high-dose senicapoc group compared to the 0.01 gm/dl increase in the placebo group ( $P < 0.001$ ) (Ataga 2008). There was no significant change in the hemoglobin level when 6 mg of senicapoc was compared to the placebo. There was significant increase in the RBC count with senicapoc compared to the placebo in a dose-dependent manner.

Both the high and low doses of senicapoc resulted in a significant decrease in the percentage and absolute number of reticulocytes (Ataga 2008).

The above findings were confirmed in the Phase III senicapoc study. There was no significant change in the HbF levels (Ataga 2011).

Gupta reported that participants in the intervention group had a mean hemoglobin level of 7.9 g/dl compared to 6.8 g/dl in the control group ( $P < 0.01$ ) (Gupta 1995). Other outcomes were not reported, and SDs were not given. Again, the authors will be contacted to clarify this information.

### 3. Quality of life measures

These measures were not assessed in the Phase II study of senicapoc (Ataga 2008).

In the Phase III senicapoc study there was no difference in the FACIT-Fatigue scores between the two treatment arms irrespective

of the hydroxyurea stratification. Similarly, there was no difference in the number of participants transfused, the number of transfusions or units per participant. The participants on senicapoc spent more days in the hospital compared to those on placebo (Ataga 2011).

Gupta reported the mean length of hospital stay per crisis in the intervention and control groups respectively was 4.3 and 3.9 days, MD 0.40 (95% CI -0.28 to 1.06), with no significant difference (Gupta 1995). Days of work lost per crisis were 3.4 in the zinc group and 4.9 in the control group. Although there is a significant reduction in the number of work days lost in the zinc group, MD -1.50 (95% CI -2.24 to -0.76), on average hospital stay was longer in this group.

### 4. Other sickle-related events

In the Phase II senicapoc study, there was one episode of staphylococcal sepsis in the low-dose senicapoc group and one urinary tract infection in the high-dose senicapoc group (Ataga 2008). There was one report of muscle strain in the high-dose senicapoc group and once case of bronchitis in the placebo group (Ataga 2008).

In the Phase III senicapoc study there was no significant difference in other sickle cell-related events (pneumonia, asthma, fever, catheter-related infections) between the senicapoc and the placebo groups (Ataga 2011).

In the zinc study, there were 108 episodes of infection reported in the intervention group compared to 204 in the control group (Gupta 1995).

### 5. Adverse drug reactions

In the Phase II senicapoc study three participants discontinued the study due to adverse events (Ataga 2008). One of the participants in the low-dose senicapoc group had dyspnoea and weakness and two in the high-dose senicapoc group had acute chest syndrome and painful crises. Diarrhea and nausea of mild to moderate intensity were more common in the senicapoc groups compared to the placebo group.

Nausea and urinary tract infections were significantly higher in the senicapoc-treated participants compared to placebo in the Phase III senicapoc study. All other adverse events were similar between the two treatment groups (Ataga 2011).

The Gupta study states that zinc was well-tolerated with no significant toxicity throughout the study, although it was unclear which potential toxic effects were monitored (Gupta 1995).

## DISCUSSION

Zinc deficiency in sickle cell disease has been shown in observational studies to correlate with disease severity (Gupta 1987; Karayalcin 1974). Laboratory studies have shown that zinc can improve the sickle cell membrane status and antagonise intracellular

calcium, and could therefore affect red cell dehydration (Bennekou 2001). The Gupta study included in this review showed a significant reduction in the number of sickle-related events in one and a half years of use in people treated with zinc sulphate (Gupta 1995). However, lack of data in the publication makes it difficult to analyse the effect on other outcomes, on painful crisis frequency in particular. In addition, changes to hematological indices, which could be used as biomarkers of RBC sickling, were inconclusive. Further research is required to evaluate the effect of zinc on the basic mechanisms of sickle cell disease. Previous studies using zinc also suggest a benefit for other problems in sickle cell disease, including leg ulcers (Serjeant 1970), growth (Zemel 2001), infection (Prasad 1999) and androgen deficiency in male participants (Prasad 1981). However, these studies were not included in the review since they do not consider prevention of vaso-occlusive crises.

Senicapoc (ICA-17043) is a selective and potent blocker of the Gardos channel in the RBC (Stocker 2003). The effect of this drug on the markers of hemolysis and hemoglobin level was evaluated in a 12-week, Phase II randomized double-blind study in people with sickle cell disease (Ataga 2008). The findings of this study suggested an increase in the lifespan of the sickle red cells, but there was no difference in the incidence of painful crises. A subsequent Phase III study was terminated early due to the lack of improvement in the vaso-occlusive pain crisis in people with sickle cell disease (Ataga 2011).

A number of other studies did not satisfy the inclusion criteria, but are still relevant to this review. Cetiedil citrate is believed to act at the Gardos channel, antagonising the calcium-mediated potassium efflux from RBCs, thus affecting red cell hydration and preventing sickling. In two randomized controlled trials investigating this treatment of sickle pain crises, both found it to limit the duration of a pain crisis and reduce the number of sites of pain compared to placebo (Benjamin 1986; Cabannes 1983). No studies have tested its use in preventing sickle crises.

Piracetam improves red cell rheology; however, the exact mechanism of this on the red cell membrane is unclear. It was initially believed to inhibit the Gardos channel, but later studies disproved this theory (Stone 1988; Stuart 1992). It has been shown in one randomized controlled trial to decrease the number of pain crises (Piracetam Study 1998) and in another to have no effect in preventing pain crises (Alvim 2005). For further information, please see the relevant Cochrane Review (Al Hajeri 2007).

Magnesium is known to inhibit the potassium-chlorine ion pump, and observational studies have shown benefits in sickle cell disease.

Sodium cromoglicolate can stabilise erythrocyte membranes by blocking chlorine or calcium channels. In an *ex vivo* study, two groups of nine children were given sodium cromoglicolate either by the nasal route or inhalation (Toppet 2000). The percentage of sickled cells in the blood of all children significantly decreased,

regardless of which group they were assigned to. Further studies are now needed to test the clinical benefit of this agent.

Diltiazem also proved useful in a pilot study which measured sickle pain intensity and duration and various hematological parameters (Rubio 1992). It is a calcium channel blocker and may impact on red cell hydration.

It was been hypothesised that hydroxyurea may increase intrinsic levels of nitric oxide, and this may affect red cell hydration. Effectiveness of hydroxyurea is considered in a different Cochrane Review (Jones 2001).

Zinc sulphate appears to be of benefit to people with sickle cell disease (Gupta 1995).

However, further research in the form of large well-designed randomized controlled trials is required to fully elucidate the true value in sickle cell disease of drugs which aim to prevent red cell dehydration.

We will continue to run searches to identify any potentially relevant trials; however, we do not plan to update other sections of the review until new trials are published.

## AUTHORS' CONCLUSIONS

### Implications for practice

There is evidence that zinc sulphate is associated with a reduction in pain crises, and also with reductions in other sickle cell-related crises. This was despite the treatment not being associated with an improvement in any of the hematological outcomes. While these results are encouraging, widespread introduction of this agent in the management of people with sickle cell disease is not indicated at present.

Senicapoc clearly improves the red cell survival in a dose-dependent manner. But the improvement in the laboratory measures did not translate into clinical benefits. There was no reduction in the incidence of sickle cell crises with the use of senicapoc. Therefore, currently this drug cannot be used in clinical practice for the prevention of vaso-occlusive crises.

### Implications for research

More studies are needed to evaluate the effect of zinc sulphate and of piracetam on the basic mechanisms of sickle cell disease. Further multicenter randomized controlled trials of zinc sulphate in people with sickle cell disease should be conducted. To investigate whether the findings previously reported are consistent and sustained, these future studies should be larger and longer term than the one reported in this review. Senicapoc improved red cell survival and decreased hemolysis. Further studies are needed to ascertain if this drug will be beneficial in some of the other complications of sickle cell disease such as pulmonary hypertension.



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\* Indicates the major publication for the study

## CHARACTERISTICS OF STUDIES

### Characteristics of included studies [ordered by study ID]

#### Ataga 2008

Methods	Phase II, randomized, double-blind, placebo-controlled study. There was a screening phase and a 12-week treatment phase	
Participants	90 sickle cell participants were recruited at 19 medical centers in the USA. The participants were adults aged between 18 and 60 years with at least one episode of sickle cell-related painful crisis 4 weeks prior to the study screening	
Interventions	Participants were randomized to either: 1. high-dose senicapoc: a single 150 mg senicapoc loading dose followed by 10 mg daily maintenance dose; 2. low-dose senicapoc: a single 100 mg senicapoc loading dose followed by 6 mg daily maintenance dose; or 3. placebo.	
Outcomes	Efficacy, safety, pharmacokinetic, and pharmacodynamic measures were obtained on day 1 of study, end of week 1 and every 2 weeks until the end of treatment phase	
Notes		
<b><i>Risk of bias</i></b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Low risk	Centralized randomization protocol using an integrated voice response system
Allocation concealment (selection bias)	Unclear risk	Allocation concealment was not reported.
Blinding (performance bias and detection bias) Participants	Low risk	The participants were blinded.
Blinding (performance bias and detection bias) Clinicians	Low risk	The treating physicians were blinded.
Blinding (performance bias and detection bias) Outcome assessors	Low risk	The review committee was blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Modified intent to treat population was used for analysis. Two participants dropped out prior to the first efficacy analysis. A total of ten participants did not complete the

**Ataga 2008** (Continued)

		study; full reasons for this were given in the published paper
Selective reporting (reporting bias)	Low risk	All outcomes stated in 'Methods' section reported in 'Results'
Other bias	Low risk	No significant differences in the baseline characteristics of the study participants

**Ataga 2011**

Methods	Phase III randomized, double-blind, placebo-controlled, parallel group study. There was a screening phase, a 52-week treatment phase, and a follow-up evaluation 8 weeks after the end of the study	
Participants	297 participants were recruited from 75 centers in the USA, Jamaica, Brazil, France, Trinidad and the United Kingdom. Participants aged between 16 and 65 years and had at least 2 episodes of painful crisis within the preceding 12 months that required medical attention were included. Participants on hydroxyurea in preceding 12 months were required to be on a stable dose of the drug prior to inclusion in the study	
Interventions	Participants were randomized to receive (in a 1:1 fashion): 1. senicapoc (loading dose of 20 mg twice daily for 4 days followed by 10 mg daily); or 2. placebo	
Outcomes	The primary end point was the frequency of sickle cell pain crises. Time to the first second and third acute painful crises was a secondary end point of the study (Ataga 2011). The effect of the drug on markers of hemolysis, hemoglobin, hematocrit, reticulocyte count, red blood cell count and dense erythrocytes was also measured. Quality of life measures were also included	
Notes		

***Risk of bias***

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Centralized randomization protocol using an integrated voice response system
Allocation concealment (selection bias)	Unclear risk	Allocation concealment was not reported.
Blinding (performance bias and detection bias) Participants	Low risk	The participants were blinded.

**Ataga 2011** (Continued)

Blinding (performance bias and detection bias) Clinicians	Low risk	The treating physicians were blinded.
Blinding (performance bias and detection bias) Outcome assessors	Low risk	The review committee was blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Modified intent to treat population was used for analysis. The study had to be terminated prematurely because it was unlikely to meet the primary end point
Selective reporting (reporting bias)	Low risk	All outcomes stated in 'Methods' section reported in 'Results'
Other bias	Low risk	No significant differences in the baseline characteristics of the study participants

**Gupta 1995**

Methods	Double-blind, placebo-controlled, randomized controlled trial
Participants	145 participants from India with SS disease, aged over 5 years. 15 participants were lost to follow up
Interventions	Zinc (oral 220 mg tds) or placebo (identical in appearance). Participants were seen weekly, follow up was for 1.5 years and the data were analysed at the end of the follow-up period
Outcomes	Sickle related crises: vaso-occlusive; mixed; hemolytic; sequestration; and aplastic. Days in hospital and working days lost
Notes	

***Risk of bias***

<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Low risk	New Castle software was used to generate randomization.
Allocation concealment (selection bias)	Unclear risk	Allocation concealment was not reported.
Blinding (performance bias and detection bias) Participants	Low risk	The participants were blinded.

**Gupta 1995** (Continued)

Blinding (performance bias and detection bias) Clinicians	Low risk	The treating physicians were blinded.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	Not discussed.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Fifteen, of the 145 participants recruited, dropped out or were lost to follow up, and these are not accounted for in the study publication
Selective reporting (reporting bias)	Low risk	All outcomes stated in 'Methods' section reported in 'Results'
Other bias	Unclear risk	Little information was given regarding baseline characteristics, including disease severity and medical history of participants, making it difficult to ascertain if the groups were sufficiently similar at the start of the study

IV: intravenous  
 SCD: sickle cell disease  
 SS: sickle cell anemia  
 tds: three times daily

**Characteristics of excluded studies** [ordered by study ID]

Study	Reason for exclusion
Adadevoh 1973	RCT of megestrol acetate. There is no good recent evidence to suggest that sex hormones prevent dehydration of the sickle red cells
Adams-Graves 1997	Trial of poloxamer 188 - inappropriate mechanism of action for inclusion in this review
Akinsulie 2005	RCT of Cajanus cajan. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Al-Jam'a 1999	Trial of isoxsurpine in treatment of sickle cell crises - rather than prevention
Alvim 2005	RCT of piracetam. There is a lack of evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the cell membrane

(Continued)

Ataga 2002	Pharmacokinetic study of ICA 17043, with no measure of clinical outcomes of interest
Ayra 1996	RCT of tucaresol. Prevents sickling by shifting the oxygen affinity, not through red cell dehydration
Bartolucci 2009	RCT of ketoprofen for treatment of, rather than prevention of, vaso-occlusive crises. Not a study of a drug involved in the prevention of red blood cell dehydration
Beatty 2007	Not a study of a drug involved in the prevention of red blood cell dehydration and vaso-occlusive crisis
Benjamin 1986	RCT of cetiedil citrate. Used in treatment rather than prevention of crises
Billet 1989	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Cabannes 1983	RCT of cetiedil citrate. Used in treatment rather than prevention of crises
Cabannes 1984	RCT of ticlopidine. Mainly an inhibitor of platelet action, rather than anti-dehydration
De Abood 1997	RCT of Depo-Provera and Microgynon. There is no good evidence to suggest that sex hormones prevent dehydration of the sickle red cells
De Ceulaer 1982	RCT of medroxyprogesterone acetate. There is no good recent evidence to suggest that sex hormones prevent dehydration of the sickle red cells
De Ceulaer 1990	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Gail 1982	RCT of urea. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells
Gladwin 2011	Not a study of a drug involved in the prevention of red blood cell dehydration
Godeau 2003	RCT of ketoprofen for treatment, rather than prevention, of vaso-occlusive crises
Isaacs 1971	RCT of steroids. There is no good recent evidence to suggest that sex hormones prevent dehydration of the sickle red cells
Jacobson 1997	Risk analysis using data from a previous RCT, in which morphine was administered in treatment, rather than prevention, of acute chest syndrome
Lonsdorfer 1984	RCT of ticlopidine. Mainly an inhibitor of platelet action, rather than anti-dehydration
Mahmood 1969	RCT of a phenothiazine. Used in treatment rather than prevention of crises
Manion 2001	Pharmacokinetic study of aspartame, with no measure of outcomes of interest



(Continued)

Manrique 1987	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Orringer 1991	RCT of 12C79, which prevents sickling primarily by increasing oxygen affinity, with possible secondary effects on the potassium-chloride co-transport channels in the red cell membrane
Osamo 1981	Trial of aspirin. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells
Oski 1968	Study not randomized, looks at promazine chloride.
Oyewo 1987	Study of diflusal, anti-sickling effects not due to red cell dehydration
Pace 2003	Phase II RCT of N-Acetylcysteine for prevention of sickle cell related vaso-occlusion and formation of dense cells. There is insufficient evidence that N-Acetylcysteine acts via red cell dehydration to include this study in the review
Pichard 1987	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the cell membrane
Piracetam Study 1998	RCT of piracetam. There is a lack of evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the cell membrane
Poflee 1991	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the cell membrane
Qari 2007	RCT of tinzaparin. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Rubio 1992	Pharmacokinetic study of diltiazem, with no measure of outcomes of interest
Semple 1984	RCT of ticlopidine. Mainly an inhibitor of platelet action, rather than anti-dehydration
Silva-Pinto 2007	Study of hydroxyurea, no measure of clinical outcomes of interest
Teuscher 1988	RCT of pentoxifylline. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the cell membrane
Toppet 2000	Non-randomized ex-vivo study of sodium cromoglicate, an anti-sickling agent, majority of patients also taking hydroxyurea
Urea Trial 1974	RCT of urea. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells
Urea Trial 2 1974	RCT of urea. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells

(Continued)

Urea Trial 3 1974	RCT of urea. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells
Uzun 2010	Study of tramadol or meperidine for treatment of, rather than prevention of, vaso-occlusive crises. Not a study of a drug involved in the prevention of red blood cell dehydration
Wallen 2007	Not a study of a drug involved in the prevention of red blood cell dehydration
Wambebe 2001	RCT of Niprisan, a naturally occurring compound. There is no evidence that any anti-sickling action is due to a reduction in dehydration of red cells
Weiner 2002	Study of nitric oxide. There is no evidence that the anti-sickling effect of the drug is due to a reduction in water loss through the red cell membrane
Wynn 2007	Study of hydroxyurea, no measure of clinical outcomes of interest
Zago 1984	Trial of aspirin. There is no evidence that any anti-sickling action is due to a reduction in dehydration of the sickle red cells

RCT: randomized controlled trial

## DATA AND ANALYSES

### Comparison 1. Anti-sickling drug versus placebo

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Mortality	1		Odds Ratio (M-H, Fixed, 95% CI)	Totals not selected
2 Number of other serious sickle-related complications	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 Overall	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Quality of life measures	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 Hospital stay per crisis (days)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 Loss of work days/crisis	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

#### Analysis 1.1. Comparison 1 Anti-sickling drug versus placebo, Outcome 1 Mortality.

Review: Drugs for preventing red blood cell dehydration in people with sickle cell disease

Comparison: 1 Anti-sickling drug versus placebo

Outcome: 1 Mortality

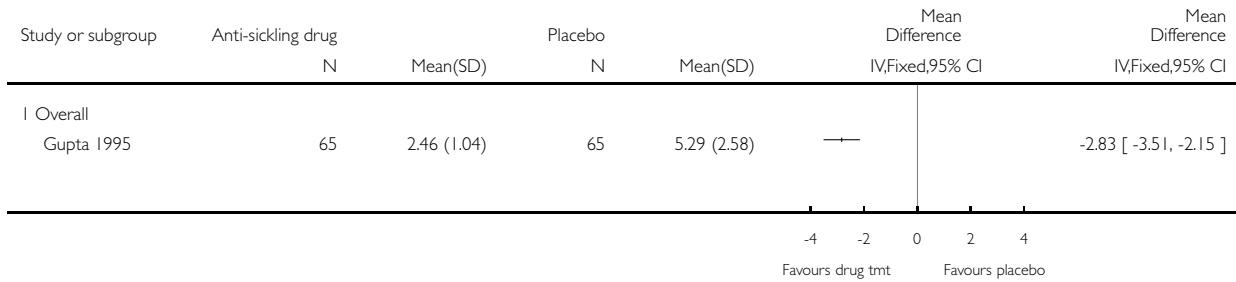
Study or subgroup	Anti-sickling drug	Placebo	Odds Ratio	
	n/N	n/N	M-H,Fixed,95% CI	M-H,Fixed,95% CI
Gupta 1995	0/65	0/65		Not estimable

### Analysis 1.2. Comparison 1 Anti-sickling drug versus placebo, Outcome 2 Number of other serious sickle-related complications.

Review: Drugs for preventing red blood cell dehydration in people with sickle cell disease

Comparison: 1 Anti-sickling drug versus placebo

Outcome: 2 Number of other serious sickle-related complications

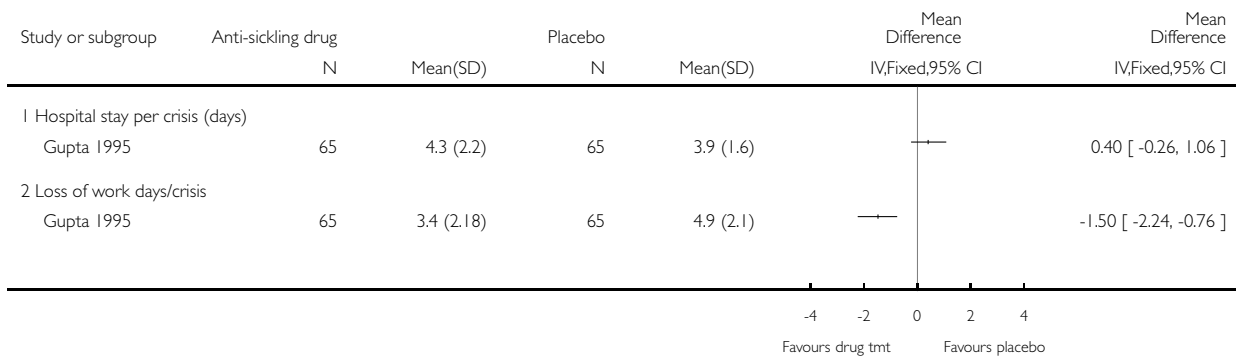


### Analysis 1.3. Comparison 1 Anti-sickling drug versus placebo, Outcome 3 Quality of life measures.

Review: Drugs for preventing red blood cell dehydration in people with sickle cell disease

Comparison: 1 Anti-sickling drug versus placebo

Outcome: 3 Quality of life measures



## WHAT'S NEW

Last assessed as up-to-date: 4 March 2016.

Date	Event	Description
4 March 2016	New citation required but conclusions have not changed	We have made only minor changes to the text throughout the review for this 2016 update
4 March 2016	New search has been performed	We searched the Cochrane Cystic Fibrosis and Genetic Disorders Group's Haemoglobinopathies Trials Register and 28 references were identified; however, none were eligible for inclusion in any section of the review We will continue to run searches to identify any potentially relevant trials; however, we do not plan to update other sections of the review until new trials are published

## HISTORY

Protocol first published: Issue 1, 2002

Review first published: Issue 4, 2002

Date	Event	Description
19 April 2012	New citation required but conclusions have not changed	A new study has been included in the review but did not lead to a change in conclusions ( <a href="#">Ataga 2011</a> ).
19 April 2012	New search has been performed	A search of the Group's Haemoglobinopathies Trials Register identified 34 references of which one has been listed as an included study ( <a href="#">Ataga 2011</a> ) and three as excluded studies ( <a href="#">Bartolucci 2009</a> ; <a href="#">Gladwin 2011</a> ; <a href="#">Uzun 2010</a> ).
20 September 2010	Amended	Contact details updated.
11 November 2009	New citation required but conclusions have not changed	A new lead author (SN) has updated this review with the existing co-author (SB)
11 November 2009	New search has been performed	A search of the Group's Trials Register identified eight new references which were potentially eligible for inclusion in this review One new study was included in the review ( <a href="#">Ataga 2008</a> ). Seven of these have been excluded; four references were additional references to already excluded studies ( <a href="#">Beatty 2007</a> ; <a href="#">Qari 2007</a> ; <a href="#">Weiner 2002</a> ; <a href="#">Wynn 2007</a> )

(Continued)

		<p>; and three were references to new studies (<a href="#">Akinsulie 2005</a>; <a href="#">Silva-Pinto 2007</a>; <a href="#">Wallen 2007</a>).</p> <p>Through personal communication, we have details of a Phase III trial for which publication is pending (<a href="#">Ballas 2009</a>). Full details will be included in a future update</p>
4 September 2008	Amended	Converted to new review format.
19 July 2007	New citation required and conclusions have changed	<p>Substantive amendment.</p> <p>A new review team has updated this review. Dr Priya C. Singh is now lead author with Dr Samir K. Ballas as co-author.</p> <p>The former lead author, Ceri Hirst (née Riddington), and former co-author Dr Lucia de Franceschi are no longer active authors on the review.</p> <p>One trial previously included in this review has been removed (<a href="#">Piracetam Study 1998</a>). The piracetam trial was excluded since the current authors believe there is insufficient evidence that it acts primarily via the red cell membrane to improve red cell rheology.</p> <p>The search of the Group's Haemoglobinopathies Trials Register in November 2006 identified trials possibly eligible for inclusion. Two references were to already excluded studies and these references have been added to the appropriate study IDs (<a href="#">Adams-Graves 1997</a>; <a href="#">Orringer 1991</a>). One reference has been added to <a href="#">Excluded studies</a> (<a href="#">Alvim 2005</a>).</p>
1 June 2004	New search has been performed	<p>The search of the Group's trials register identified eight new references, however, none were found to be eligible for inclusion in the review</p> <p>Five references to three newly identified studies have been excluded. Details of these studies can be found in the <a href="#">Excluded studies</a> section of the review (<a href="#">Adams-Graves 1997</a>; <a href="#">Al-Jam'a 1999</a>; <a href="#">Godeau 2003</a>; <a href="#">Jacobson 1997</a>; <a href="#">Pace 2003</a>).</p> <p>Three references identified were to two already excluded studies (<a href="#">Ataga 2002</a>; <a href="#">Wambebe 2001</a>). These additional references have been added to the appropriate study IDs</p>
1 July 2003	New search has been performed	Five new studies have been identified for inclusion within the <a href="#">Excluded studies</a> section of this review.

## CONTRIBUTIONS OF AUTHORS

### Review from Issue 4, 2009

Dr Singh has stepped down from the review team and Dr Nagalla is now the lead author for this review. Together with Dr Ballas, Dr Nagalla has updated this review and acts as guarantor for this and subsequent versions of the review.

### Review from Issue 4, 2007

Dr Singh is now lead author and updated the review with Dr Samir Ballas (co-author). Dr Singh acts as guarantor of the review. Dr Singh reviewed the current literature on piracetam and revised the manuscript with advice from Dr Ballas. Drs Hirst and de Franceschi have stepped down as active authors on this review.

### Review up to Issue 3, 2007

The review was conceived by the Cochrane Cystic Fibrosis and Genetic Disorders Group and designed by Dr Hirst (née Riddington). Dr Hirst and the Cochrane Cystic Fibrosis and Genetic Disorders Group conducted searches for relevant studies. Dr Hirst and Dr de Franceschi screened, appraised and abstracted data for the review. Additional information from authors was sought by Dr Hirst where necessary. Data entry was performed by Dr Hirst and interpreted by Dr Hirst with advice from the Cochrane Cystic Fibrosis and Genetic Disorders Group.

## DECLARATIONS OF INTEREST

None known.

## SOURCES OF SUPPORT

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- No sources of support supplied

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- National Institute for Health Research, UK.

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## **INDEX TERMS**

### **Medical Subject Headings (MeSH)**

Acetamides [\*therapeutic use]; Anemia, Sickle Cell [\*blood]; Antisickling Agents [\*therapeutic use]; Clinical Trials, Phase II as Topic; Clinical Trials, Phase III as Topic; Dehydration [\*prevention & control]; Early Termination of Clinical Trials; Erythrocyte Aging [drug effects]; Erythrocytes [\*drug effects]; Piracetam [therapeutic use]; Randomized Controlled Trials as Topic; Trityl Compounds [\*therapeutic use]; Zinc Sulfate [\*therapeutic use]

### **MeSH check words**

Humans