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Interventions for treating leg ulcers in people with sickle cell disease

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ABSTRACT

Background
The frequency of skin ulceration makes it an important contributor to the morbidity burden in people with sickle cell disease. Many treatment options are available to the healthcare professional, although it is uncertain which treatments have been assessed for effectiveness in people with sickle cell disease.

Objectives
To assess the clinical effectiveness and safety of interventions for treating leg ulcers in people with sickle cell disease.

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

We searched LILACS (1982 to August 2012), the African Index Medicus (up to August 2012), ISI Web of Knowledge (1985 to August 2012), and the Clinical Trials Search Portal of the World Health Organization (August 2012). We checked the reference lists of all the trials identified. We also contacted those groups or individuals who may have completed relevant randomised trials in this area.

Date of the last search of the Group's Haemoglobinopathies Trials Register: 25 May 2012.

Selection criteria
Randomised controlled trials of interventions for treating leg ulcers in people with sickle cell disease compared to placebo or an alternative treatment.

Data collection and analysis
Two authors independently selected studies for inclusion. All three authors independently assessed the risk of bias of the included studies and extracted data.
Main results

Six studies met the inclusion criteria (198 participants with 250 ulcers). Each trial investigated a different intervention and within this review we have grouped these as systemic pharmaceutical interventions (L-carnitine, arginine butyrate, isoxsuprine) and topical pharmaceutical interventions (Solcoseryl® cream, RGD peptide dressing, topical antibiotics). Three interventions reported on the change in ulcer size (arginine butyrate, RGD peptide, L-carnitine). Of these, RGD peptide matrix significantly reduced ulcer size compared with a control group, mean reduction 6.60cm² (95% CI 5.51 to 7.69). Three trials reported on the incidence of complete closure (isosuprine, arginine butyrate, RGD peptide matrix). None reported a significant effect. No trial reported on: the time to complete ulcer healing; ulcer-free survival following treatment for sickle cell leg ulcers; quality of life measures; or incidence of amputation. There was no reported information on the safety of these interventions.

Authors’ conclusions

There is evidence that a topical intervention (RGD peptide matrix) reduced ulcer size in treated participants compared to controls. This evidence of efficacy is limited by the generally high risk of bias associated with these reports.

We planned to analyse results according to general groups: pharmaceutical interventions (systemic and topical); and non-pharmaceutical interventions (surgical and non-surgical). However, we were unable to pool findings due to the heterogeneity in outcome definitions, and inconsistency between the unit of randomisation and the unit of analysis. This heterogeneity, along with a paucity of identified trials, prevented us performing any meta-analyses.

This Cochrane review provides some evidence for the effectiveness of one topical intervention - RGD peptide matrix. However, this intervention was assessed as having a high risk of bias due to inadequacies in the single trial report. Other included studies were also assessed as having a high risk of bias. We recommend that readers interpret the trial results with caution. The safety profile of the all interventions was inconclusive.

Plain language summary

Treatments for leg ulcers in people with sickle cell disease

Leg ulcers are a chronic complication for people living with sickle cell disease. Ulcers tend to be difficult to treat successfully, healing slowly over months or years. They can severely disrupt quality of life, increase disability, require extended absence from the workplace, and place a high burden of care on healthcare systems. We looked at whether treatments for leg ulcers in people with sickle cell disease were effective and safe.

In this Cochrane review we found six randomised controlled trials including 198 participants with 250 ulcers. Four of the randomised controlled trials were conducted in Jamaica and two in the USA. These trials included medications or dressings applied directly to the ulcer (topical medications) and medications given orally or intravenously (systemic medications). Given the very different modes of action of these two groups, we treated them separately throughout the review. The topical agents included Solcoseryl® cream, RGD peptide matrix dressing and topical antibiotics. Socoseryl aims to improve the use of oxygen by the skin tissue and so promote wound healing. Topical antibiotics are also used to prevent infection. The RGD peptide matrix is a gel that promotes cell growth. The systemic interventions included arginine butyrate, L-carnitine, and isoxsuprine. Aginine butyrate, given intravenously, is thought to accelerate wound healing, L-carnitine, given orally, is thought to improve tissue hypoxia, and isoxsuprine, given orally as isoxsuprine hydrochloride, is thought to widen blood vessels, so increasing blood flow to an affected wound.

One medication, a topical intervention (RGD peptide matrix) reduced ulcer size in treated participants compared to controls. However, this effect should be interpreted with caution given the high risk of bias due to the inadequacies associated with this trial report.

The evidence for the use of interventions to treat people with sickle cell disease and chronic leg ulceration is not strong. All randomised clinical trials that we included in this review were associated with a high risk of bias. This systematic review has shown the need for well-designed, high-quality randomised trials to assess the benefits and harms of interventions to improve the healing of leg ulcers in people with sickle cell disease.
## SUMMARY OF FINDINGS FOR THE MAIN COMPARISON

**Isoxuprine compared to placebo for leg ulcer in people with sickle cell disease**

**Patient or population:** patients with leg ulcer in people with sickle cell disease  
**Settings:**  
**Intervention:** Isoxuprine  
**Comparison:** placebo

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<td>Incidence of complete closure Follow-up: 6 months</td>
<td>See comment</td>
<td>See comment</td>
<td>Not estimable</td>
<td>54 ulcers (1 study; Serjeant 1977)</td>
<td>⊕⊕⊕⊕ very low</td>
<td>This trial shows inconsistency between units of randomisation (30 participants) and unit of analysis (54 ulcers)</td>
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*The basis for the assumed risk (e.g. the median control group risk across studies) is provided in footnotes. The corresponding risk (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).*
BACKGROUND

See Appendix 1 for a glossary of medical terms.

Description of the condition

Sickle cell disease (SCD) is an inherited disease and the most common haemoglobinopathy worldwide (Ballas 2010; Orkin 2010; Rees 2010). Recently, the clinical history and others aspects of SCD have been reviewed (Mousa 2010; Prabhakar 2010; Serjeant 2010). It is common among people with sub-Saharan African, Indian, Middle Eastern or Mediterranean ancestry (Creary 2007; Modell 2008). Sickle cell disease is a major public health problem (Modell 2008) with an estimated 70% of sufferers living in Africa (Makani 2007). Screening programmes for detecting SCD are ongoing worldwide (e.g. Bardakdjian-Michau 2009; Daudt 2007; Henthorn 2004; Mañú 2009; Gulbis 2006; Tshilolo 2008).

The term SCD includes sickle cell anaemia (Hb SS), haemoglobin S combined with haemoglobin C (Hb SC), haemoglobin S associated with β thalassemia (Sβ0 Thal and Sβ+ Thal) and other less prevalent double heterozygous conditions which cause clinical disease (Steinberg 2009; Weatherall 2006). Haemoglobin S combined with normal haemoglobin (A) is known as the sickle cell trait (AS), which is generally asymptomatic and is not part of this review.

One chronic complication of SCD is the sickle cell leg ulcer (SCLU) (Figure 1) (Knox-Macaulay 1983; Minniti 2010; Ramalho 1985). The frequency of skin ulceration makes it an important contributor to the morbidity burden faced by people with SCD, and resistance to therapy makes chronicity an important feature of the condition. Although there is no universally accepted leg ulcer duration that defines the condition as chronic, a research definition from Jamaica has defined a chronic leg ulcer as “an active ulcer recorded at least twice over a minimum period of three months” (Alexander 2004; Serjeant 2005). A SCLU can be a physically disabling complication, with potentially negative psychological and social consequences. It has been considered a marker of disease severity by some authors (Alleyne 1977; Cumming 2008; Eckman 1996; Halabi-Tawil 2008). The rationale behind this severity statement is based on an increasing risk of priapism and pulmonary hypertension in those individuals with SCLU (Halabi-Tawil 2008; Serarslan 2009). People with homozygous SCD and a leg ulcer have a higher prevalence of pulmonary hypertension compared to people with homozygous SCD without a leg ulcer (Serarslan 2009). These vasculopathies have a common denominator: a chronic intravascular hyper-haemolysis which generate a lower level of nitric oxide (Akinsheye 2010; Taylor 2008). Others have used multivariate statistical modelling to explore possible clinical phenotypes in people with homozygous SCD and have suggested a 'leg ulcer phenotype' (Alexander 2004).
Individuals with Hb SS are more likely to experience a leg ulcer than those with other genotypes (Ankra-Badu 1992; Koshy 1989). Geographically, the reported prevalence of this complication varies. In Nigeria, the prevalence of SCLU ranges between 7.5% for people with HbSS and 1.5% for people with HbSC (Akinyanju 1979; Durosinmi 1991). According to the ‘Cooperative Study of Sickle Cell Disease’ in the USA, leg ulcers affected 2.5% of people with SCD, with a higher rate of between 4% and 5% among those with HbSS (Koshy 1989). In Jamaica, a lifetime prevalence of any leg ulceration was reported to be between 70% and 80% (Serjeant 2005). More recently a prevalence of ulcers lasting six months or more was reported as 29.5% and the cumulative incidence as 16.7% (Cumming 2008). These prevalence rate variations are partly due to differing patient age distributions and analysis methodologies. It is generally agreed that leg ulcers are most commonly reported in adolescence and early adulthood in individuals with SCD (Serjeant 2005).

The SCLUs generally occur in areas with less subcutaneous fat, with thin skin, and with decreased blood flow (Trent 2004). The commonest sites are the medial and lateral malleoli (ankles), often becoming circumferential if not controlled early; the medial malleolus is more commonly involved than the lateral malleolus (Serjeant 2005; Trent 2004) (Figure 1). Less common sites are the anterior tibial area, dorsum of the foot, and achilles tendon area (Trent 2004).

The pathogenesis of the SCLU is complex (Aslan 2007; Hagar 2008; Kato 2007; Kato 2009; Mack 2006; Morris 2008; Paladino 2007; Serjeant 2005; Trent 2004; Wood 2008). One or more of the following mechanisms could play a role in the development of leg ulcers in people with SCD:

1. decreased nitric oxide: the haemolysis (breakdown) of the sickle red blood cells releases haemoglobin into the bloodstream which consumes nitric oxide (a powerful vasodilator agent) to the blood, perhaps leading to impaired endothelial function (Mack 2006);

2. infectious process: the role of bacterial infection or colonization, or both, is illustrated by the growth of organisms such as Staphylococcus aureus, Pseudomonas aeruginosa and group A streptococci from ulcer swabs, signs of local inflammation and regional lymphadenopathy (MacFarlane 1986; Mohan 2000; Sehgal 1992);
3. venous incompetence: low oxygen tension in the venous system, inevitable turbulence around venous valves and high white blood cell and platelet counts promote endothelial adhesion and chronic ischemia (Chalchal 2001; Clare 2002; Cumming 2008; Mohan 2000; Serjeant 2005).

4. blood hypercoagulability: an acquired antithrombin III deficiency that has been described in individuals with SCLUs, along with evidence of fibrinolysis (D-dimer fragment, and fibrinogen or fibrin degradation products) (Cacciola 1990a).

5. defective immunity: the complement system contributes to the immune system's defense against infection, and an inability to fix this system has been proposed as a risk factor for developing this chronic complication (Morgan 1981).

6. genetic risk factors: there is a relationship between genes of the TGF-beta/BMP superfamily and endothelial function and nitric oxide biology which may explain the genesis of the SCLU (Nolan 2006; Ofosu 1987; Steinberg 2009).

7. postural vasoconstriction: by using the laser Doppler flowmeter, it had been demonstrated that individuals with SCLU have a low red cell flux at the ulcer or scar site (Mohan 1997).

In brief, the increased susceptibility to leg ulcers in people with SCD is due to a chronic ischemia and defective immunity. Chronic ischemia may be explained by a blood hypercoagulability, venous incompetence, or postural vasoconstriction, with these mechanisms conditioned by a low nitric oxide that may itself be genetically influenced.

**Description of the intervention**

There are a wide range of possible treatments for leg ulcers in people with SCD, and many of these treatments have been considered as interventions among people with leg or foot ulcers resulting from other pathologies, e.g. a silver-based wound dressing and topical agents for treating diabetic foot ulcers (Bergin 2006; O'Meara 2010), the debridement of diabetic foot ulcers (Edwards 2002), oral zinc for arterial and venous leg ulcers (Wilkinson 1998) and compression for venous leg ulcers (O'Meara 2009). Conventional care generally include occlusive dressing (Palfreyman 2006; Vermeulen 2004; Wasiak 2008) and debridement and cleansing (Moore 2005; Nelson 2000; Smith 2011). It is unknown if ulcers in people with SCD might experience differential benefits of these common treatments compared to ulcers from other pathologies. Many possible treatments exist as adjuncts to conventional treatment and we have classified these into two major treatment groups: pharmaceutical interventions; and non-pharmaceutical interventions.

1. Pharmaceutical interventions

   i) **Systemic pharmaceutical interventions:**

      a) vascular drugs (such as pentoxifylline (blood viscosity-reducing agent) (Frost 1990), isosuprime hydrochloride (β-adrenergic receptor stimulant) (Serjeant 1977), xanthinol nicotinate (vasodilator) (Afifi 1979));

      b) antioxidant agents (such as L-carnitine) (Harrel 1990; Serjeant 1997);

      c) recombinant agents and related (such as recombinant human erythropoietin (erythropoiesis-stimulating agent which increases the haemoglobin levels) (al-Momen 1991), antithrombin III concentrate (potent coagulation inhibitor) (Cacciola 1989));

      d) growth factors: such as Bosentan (a receptor endothelin receptor blocker) (Lionet 2008).

      e) minerals (oral zinc sulphate) (Serjeant 1970).

      f) pharmacologic stimulation of HbF synthesis agents (such as arginine butyrate) (Sher 1994);

   ii) **Topical pharmaceutical interventions:**

      a) antibiotics and antiseptics (such as topical antibiotic (Baum 1987), collagen dressing (Reindorf 1989), natural honey (Okany 2004));

      b) growth factors and related (such as topical granulocyte-macrophage colony-stimulating factor (GM-CSF) (Ali Khan 2004; Mery 2004; Pieters 1995), RGD peptide matrix, Solcoseryl® (a tissue stimulating agent) (La Grenade 2003));

      c) steroids (such as cortisone - a potent anti-inflammatory) (Rice 1953);

      d) miscellaneous (such as topical opioids) (Ballas 2002).

2. Non-pharmaceutical interventions

   i) **Non-pharmaceutical surgical interventions:**

      a) reconstructive surgery (free flap transfer) (Spence 1985);

      b) cell therapy (e.g. human skin equivalent (Gordon 2003); allogeneic keratinocytes (Amini-Adle 2007));

   ii) **Non-pharmaceutical and non-surgical:**

      a) laser therapy (e.g. InGaP (670 nm) laser) (Lucena 2007);

      b) miscellaneous (e.g. hyperbaric oxygen) (Espinosa 1992).

These interventions might occur alone or in combination. Recently, for example, a combination of treatment approaches (antibacterial agent, zinc oxide, bandages and debridement) have been proposed for improving the care of people with SCLUs (Schleucher 2007).

**Why it is important to do this review**

Even with rigorous conventional care, leg ulcers tend to be indolent and intractable, healing slowly over months or years (Ballas 2002; Minniti 2010). In the USA, the average duration of a SCLU has been reported to exceed three years (Wethers 1994) with recurrence rates ranging from 25% to 52% (Koshy 1989). The decreased quality of life, increased disability, absence from work and high utilization of health care resources can severely affect the lives of people with SCLUs (Cumming 2008; Halabi-Tawil 2008).

In this review, we assessed the clinical effectiveness and safety of interventions for treating leg ulcers in people with SCD.
**OBJECTIVES**

To determine whether any clinical interventions (used either alone or in combination) are effective and safe when treating leg ulcers in people with SCD.

**METHODS**

Criteria for considering studies for this review

**Types of studies**

Randomised controlled trials (RCTs).

**Types of participants**

People with all types of SCD who have been diagnosed with a leg ulcer and treated in a hospital or community setting, or both.

**Types of interventions**

Single or combination treatment regimen (with each treatment classified as pharmaceutical or non-pharmaceutical as detailed below) compared to either conventional care or another treatment regimen for leg ulcers in people with SCD.

1. **Pharmaceutical interventions**

   **Systemic interventions**
   - vascular drugs (such as pentoxifylline (blood viscosity-reducing agent), isoxsuprine hydrochloride (β-adrenergic receptor stimulant), xanthinol nicotinate (vasodilator));
   - antioxidant agents (such as L-carnitine);
   - recombinant agents and related (such as recombinant human erythropoietin (erythropoiesis-stimulating agent which increases the haemoglobin levels), antithrombin III concentrate (potent coagulation inhibitor));
   - growth factors: such as bosentan (a receptor endothelin receptor blocker);
   - pharmacologic stimulation of HbF synthesis agents (such as arginine butyrate);
   - oral zinc sulphate.

   **Topical interventions**
   - antibiotics and antiseptics (such as topical antibiotic, collagen dressing, natural honey);
   - growth factors and related (such as topical granulocyte-macrophage colony-stimulating factor (GM-CSF), RGD peptide matrix, Solcoseryl® (a tissue stimulating agent));
   - steroids (such as cortisone (a potent anti-inflammatory));
   - dressing;
   - debriding agents;
   - compression;
   - miscellaneous (such as topical opioids).

2. **Non-pharmaceutical interventions**

   - reconstructive surgery (free flap transfer);
   - cell therapy (e.g. human skin equivalent; allogeneic keratinocytes);
   - laser therapy (e.g. InGaP (670 nm) laser);
   - miscellaneous (e.g. hyperbaric oxygen).

**Types of outcome measures**

**Primary outcomes**

1. Incidence of complete closure (defined as 100% epithelization or skin closure without drainage)
2. Time to ulcer closure
3. Change in ulcer size (surface area or volume)

**Secondary outcomes**

1. Ulcer-free survival following treatment for SCLUs (free from leg ulcer recurrence)
2. Quality of life measures (based on any item from a validated scale, e.g. SF-36, EuroQoL, WHOQOL-BREF (Asnani 2009))
3. Incidence of amputation
4. Adverse events
   i) any adverse event defined as “any untoward medical occurrence that may present during treatment with a pharmaceutical product but which does not necessarily have a causal relationship with this treatment” (Nebeker 2004).
   ii) adverse drug reactions defined as “a response to a drug which is noxious and uninitiated and which occurs at doses normally used in man for prophylaxis, diagnosis, or therapy of disease, or for the modification of physiologic functions” (Nebeker 2004).

**Search methods for identification of studies**

We searched for trials, irrespective of publication status (trials may be unpublished or published as an article, an abstract, or a letter),
language or country. No limit was applied with respect to the period of follow-up.

**Electronic searches**

We identified relevant trials from the Cystic Fibrosis and Genetic Disorders (CFGD) Group's Haemoglobinopathies Trials Register using the terms: (sickle cell OR (haemoglobinopathies AND general)) AND leg ulcers.

Date of the last search of the Cochrane Cystic Fibrosis and Genetic Disorders Group Haemoglobinopathies Trials Register: 25 May 2012.

The CFGD Group's 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 quarterly 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.

We also searched the specialised register of the Cochrane Wounds Group. Date of the last search of the specialised register of the Cochrane Wounds Group: 10 May 2012.

We also searched the following resources using the keywords: (sickle cell OR (haemoglobinopathies AND general)) AND leg ulcers (to August 2012).

- Latin American and Caribbean Health Sciences Information System (LILACS) (from 1982);
- African Index Medicus (AIM);
- ISI Web of Knowledge (from 1985);
- Clinical Trials Search Portal of the World Health Organization;
- Food and Drug Administration;
- European Medicines Agency;
- Medicines and Healthcare Products Regulatory Agency;
- Scirus;
- CenterWatch;
- Sistema de Información Esencial en Terapéutica y Salud;
- Evidence in Health and Social Care;
- DailyMed.

**Searching other resources**

We also checked the reference lists of all the trials identified by the above methods. We contacted a number of key researchers (by email), asking whether they knew of studies assessing treatments for SCLUs in people with SCD.

**Data collection and analysis**

Data collection and analysis procedures are described below, and follow documented Cochrane Collaboration methodologies (Higgins 2011a).

**Selection of studies**

Two authors independently selected studies for inclusion, and extracted data. A third author was always included when two authors disagreed, with all disagreements resolved by group discussion.

**Data extraction and management**

We extracted the following groups of data:

- demographics (age, sex, country);
- characteristic of the ulcer (anatomic site, size, number of ulcers, presence of infection, how long the patient has had the ulcer);
- sickle cell genotype (SS, SC);
- phenotypic expression (e.g. total haemoglobin, fetal haemoglobin).

One review author, Jennifer M Knight-Madden, interviewed Dr Graham R. Serjeant, an author on four of the included RCTs (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997) to clarify aspects of these trials that were unclear from the published trial reports.

Although potential interventions were broadly classed as pharmaceutical and non-pharmaceutical, we identified only pharmaceutical interventions and present results under two major groupings (systemic and topical). Given the clinical diversity of the included interventions, we do not regard it as appropriate to combine data for meta-analysis at this time.

For future updates, if we are able to find RCTs reporting these data, we plan to report our endpoints at between three and six months, and greater than six months; work from the Jamaican sickle cell cohort suggest that most leg ulcers take a minimum of between three and six months to heal (Serjeant 2005).

**Assessment of risk of bias in included studies**

We followed the domain-based evaluation for risk of bias in included RCTs (Higgins 2011a). All review authors independently assessed the risk of bias of the trials according to the *Cochrane Handbook for Systemic Reviews of Interventions* (Higgins 2011a). The disagreements were resolved through discussion.

We assessed the following domains as low, unclear, or high risk of bias:

1. generation of allocation sequence;
2. allocation concealment;
3. blinding (of participants, personnel and outcome assessors);
4. incomplete outcome data;
5. selective reporting;

**Interventions for treating leg ulcers in people with sickle cell disease (Review)**

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6. Free of other bias.

1. Generation of allocation sequence (checking for possible selection bias)
We described, for each included trial, whether the method used to generate the allocation sequence was reported in sufficient detail to allow an assessment of whether it produced comparable groups. We assessed the method as having one of the following risks of bias:
- low risk (any truly random process, e.g. random number table; computer random number generator);
- high risk (any non-random process, e.g. odd or even date of birth; hospital or clinic record number);
- unclear risk (if the trial was described as randomised, but the method used for the allocation sequence generation was not described).

2. Allocation concealment (checking for possible selection bias)
We described, for each included trial, whether the method used to conceal the allocation sequence was reported in sufficient detail and determined whether group allocation could have been foreseen in advance of, or during recruitment, or changed after assignment. We assessed the methods as having one of the following risks of bias:
- low risk (e.g. telephone or central randomisation; consecutively numbered sealed opaque envelopes);
- high risk (open random allocation; unsealed or non-opaque envelopes, alternation; date of birth);
- unclear risk (if the trial was described as randomised, but the method used to conceal the allocation was not described).

3. Blinding or masking (checking for possible performance bias)
We described, for each included trial, the methods, if any, to blind study participants and personnel from knowledge of which intervention a participant received. We judged trials at low risk of bias if they were blinded, or if we judged that the lack of blinding could not have affected the results. We aimed to assess blinding separately for different outcomes or classes of outcomes. We assessed the methods as having one of the following risks of bias:
- low, high or unclear risk for participants;
- low, high or unclear risk for personnel;
- low, high or unclear risk for outcome assessors.

4. Incomplete outcome data (checking for possible attrition bias through withdrawals, dropouts, protocol deviations)
We assessed the methods as having one of the following risks of bias:
- low risk (the numbers and reasons for dropouts and withdrawals in all intervention groups were described or if it was specified that there were no dropouts or withdrawals);
- high risk (the number or reasons for dropouts and withdrawals were not described).
- unclear risk (the report gave the impression that there had been no dropouts or withdrawals, but this was not specifically stated).

We further examined the percentages of overall dropouts in each trial and per randomisation arm and aimed to evaluate whether an intention-to-treat analysis had been performed or could be performed from the published information.

5. Selective reporting bias (reporting bias due to selective outcome reporting)
We described, for each included trial, how we investigated the possibility of selective outcome reporting bias and what we found. We assessed the methods as having one of the following risks of bias:
- low risk (any one of the following: the study protocol is available and all of the study’s pre-specified (primary and secondary) outcomes that are of interest in the review have been reported in the pre-specified way or the study protocol is not available but it is clear that the published reports include all expected outcomes, including those that were pre-specified (convincing text of this nature may be uncommon));
- high risk (any one of the following: not all of the study’s pre-specified primary outcomes have been reported; one or more primary outcomes is reported using measurements, analysis methods or subsets of the data (e.g. sub scales) that were not pre-specified; one or more reported primary outcomes were not pre-specified (unless clear justification for their reporting is provided, such as an unexpected adverse effect); one or more outcomes of interest in the review are reported incompletely so that they cannot be entered in a meta-analysis; the study report fails to include results for a key outcome that would be expected to have been reported for such a study);
- unclear risk (insufficient information to permit judgement of ‘low risk’ or ‘high risk’).

6. Free of other bias (bias due to problems not covered elsewhere in the table)
We described, for each included study, any important concerns we have about other possible sources of bias and made assessments as follows:
- low risk of bias (the trial appears to be free of other components that could put it at risk of bias);
- high risk of bias (there are other factors in the trial that could put it at risk of bias);
• unclear risk of bias (the trial may or may not be free of other components that could put it at risk of bias); e.g. ascertainment bias, bias in the presentation of data, design bias (Porta 2008). Please refer to the appendices for further details (Appendix 2).

7. Overall risk of bias

We made explicit judgements about whether studies are at high risk of bias, according to the criteria given in the Cochrane Handbook for Systematic Reviews of Interventions (Higgins 2011a). In reference to the six domains above, we assessed the likely magnitude and direction of any bias and whether we considered it was likely to impact on the review findings. Trials that achieved a 'low risk' assessment for adequate generation of allocation sequence, allocation concealment, blinding, handling of incomplete outcome data, and no selective outcome reporting, and that were without other risks of bias, were considered to be at an overall low risk of bias, while trials that were assessed as either 'high risk' or 'unclear risk' on the majority of domains were considered to be at a high risk of bias. We aimed to explore the impact of the level of bias through undertaking sensitivity analyses - see Sensitivity analysis.

Measures of treatment effect

For binary outcome measures (incidence of complete closure, adverse events), we calculated the risk ratio (RR) with 95% confidence intervals (CIs) for each outcome. For continuous outcomes (change in ulcer size measured using surface area or volume), we calculated the mean difference with 95% CIs. For the future update, if the search strategy finds new RCTs, we will plan to assess of treatment effect as follows:

• for binary outcome measures (incidence of complete closure, incidence of amputation, adverse events, and adverse drug reaction), we plan to calculate the risk ratio (RR) with 95% confidence intervals (CIs) for each outcome.
• for continuous outcomes (quality of life), we will use the standardized mean difference with 95% CIs, since different scales may be used to measure quality of life. Change in ulcer size (surface area or volume) will be measured using a pooled estimate of treatment effect by calculating the mean difference with 95% CIs. If statistical information is missing (such as standard deviations), we will try to extract them from other relevant information in the paper, such as P values and CIs.
• for time-to-event outcome (time to ulcer closure and ulcer-free survival following treatment for SCLU), we plan to calculate the hazard ratios (HR) with 95% CIs for each outcome.

Unit of analysis issues

This review has treated the participant as the unit of analysis. This is important as the meta-analytic techniques used assume independence between measurements, and more than one treated ulcer per participant would not be statistically independent. A result of ignoring this unit-of-analysis issue could be overly optimistic confidence intervals. We found six RCTs and of these, four used the ulcer as their unit of analysis (La Grenade 1993; Baum 1987; McMahon 2010; Serjeant 1977). For future updates, and if the review authors receive extra information from studies, we plan to incorporate multiple ulcers per participant into our quantitative analyses using ‘approximate analyses of cluster-randomised trials of meta-analysis’, according to the Cochrane Handbook for Systematic Reviews of Interventions (Higgins 2011b). Due to problems with ensuring a successful treatment ‘washout’ period, we do not consider cross-over trials an appropriate study design for assessing wound healing, and we have not included cross-over trials in this systematic review.

Dealing with missing data

For all included studies, we recorded the levels of participant attrition. Attrition rates were not stated in four of the six trial reports, and for these trials we approached the lead author for additional information on levels of missing data (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997). This has allowed an intention-to-treat analysis for all included studies. For future updates, if necessary, we will seek full reports from authors where studies have been published in abstract form, presented at meetings or reported to the co-authors. Where information is missing or unclear, we will contact 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

For future updates, when more trials are included, we plan to test for heterogeneity between studies using a standard chi-squared test and I² statistic (Higgins 2003). The chi-squared test is a statistical test for heterogeneity, whereas I² assesses the impact of heterogeneity on the meta-analysis. We will use the following I² ranges to interpret 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.

In the presence of substantial heterogeneity we plan to explore this heterogeneity by pre-specified subgroup analysis. We will also evaluate the extent of heterogeneity by visual inspection of the forest plot (Higgins 2011).

Assessment of reporting biases

Comprehensive searches were done by two 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. For future updates, if we are able to include 10 RCTs or more in a single meta-analysis, we will assess whether the review is subject to possible publication bias by using a funnel plot to graphically illustrate variability between trials. If asymmetry is detected, we will explore causes other than publication bias.

Data synthesis
No meta-analysis was undertaken in this review. For future updates, any meta-analyses will be performed separately for each intervention group (pharmaceutical systemic, pharmaceutical topical, non-pharmaceutical surgical, non-pharmaceutical non-surgical). If the eligible trials are sufficiently homogenous, we will summarize their findings using a fixed-effect model. However, if we find statistical heterogeneity using the $I^2$ statistic ($I^2 > 50\%$) we will use a random-effects model (Higgins 2003; Higgins 2011). Clinical variability (variability in types of participants, interventions, or outcomes) may prevent us from pooling trials.

Summary of findings tables
We used the GRADE proposals to assess the quality of the body of evidence associated with the following outcomes: complete closure; change in ulcer size; and safety (total adverse events and related study treatment adverse events) (Guyatt 2011). We constructed summary of findings tables (SoF) using the GRADE-Pro software (Summary of findings for the main comparison; Summary of findings 2; Summary of findings 3; Summary of findings 4) (GRADEPro 2008). The GRADE approach appraises the quality of a body of evidence based on the extent to which one can be confident that an estimate of effect or association reflects the item being assessed. The quality of a body of evidence considers within-study risk of bias (methodological quality), the directness of the evidence, heterogeneity of the data, precision of effect estimates and risk of publication bias (Baldesh 2011; Guyatt 2011a; Guyatt 2011b; Guyatt 2011c; Guyatt 2011d; Guyatt 2011e; Guyatt 2011f; Guyatt 2011g; Guyatt 2011h; Guyatt 2012).

We would have used (and will apply these for future updates, if possible) the principles of the GRADE system to assess the quality of the body of evidence associated with others outcomes of this review which were not assessed by the included RCTs in this review: time to ulcer closure; ulcer-free survival following for SCLUs (free from leg ulcer recurrence); quality of life; and incidence of amputation.

Subgroup analysis and investigation of heterogeneity
For future updates, if we find clinical heterogeneity, we plan to conduct subgroup analyses as follows:
1. SS type versus SC type (Appendix 1);
2. follow-up duration.
These subgroup analyses will be only conducted for primary outcomes.

Sensitivity analysis
No sensitivity analyses were undertaken in this review. If sufficient data had been available, we would have used the following procedures (and will apply these for future updates, if possible). We would have compared RCTs with high versus low methodological quality (studies classified as having a 'low risk of bias' versus those identified as having a 'high risk of bias') and RCTs that performed intention-to-treat versus per-protocol analyses (Higgins 2011). For future updates, we will also evaluate the risk of attrition bias, as estimated by the percentage of participants lost. Trials with a total attrition of more than 30% or where differences between the groups exceed 10%, or both, would be excluded from meta-analysis but included in the review (Higgins 2011a).

RESULTS

Description of studies
See: Characteristics of included studies; Characteristics of excluded studies.

Results of the search
We identified 88 references using our search strategy (see Figure 2 for search result details). Six RCTs met our inclusion criteria (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994). These RCTs were published between 1977 and 2010, four in Jamaica (Baum 1987; La Grenade 1993; Serjeant 1993; Serjeant 1997; Serjeant 1997; Wethers 1994) and two in The United States of America (McMahon 2010; Wethers 1994). There were 250 ulcers studied among 198 participants (trial sample sizes ranged from 15 to 55 participants). A detailed description of the included trials is provided in the Characteristics of included studies table.
Figure 2. Flowchart of last search of the Group’s Trials Register: 25 May 2012.

Included studies

The six included RCTs report the following baseline characteristics (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994).

Clinical characteristics

1. Genotype of SCD

Two RCTs only included patients with HbSS (Baum 1987; La Grenade 1993). Three RCTs included patients with HbSS, HbSC, HbSS-O (Arab), HbSβ thalassaemia, HbSC Harlem, HbS (Serjeant 1977; Serjeant 1997; Wethers 1994). One trial reported 92.3% (24 out of 26) of the participants with HbSS (McMahon 2010).

2. Leg ulceration duration, leg ulcer diameter on entry to the trial and number of ulcer per patient

Two RCTs mentioned leg ulceration duration at baseline (Baum 1987; Wethers 1994). Five RCTs mentioned leg ulcer diameter on trial entry (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1997; Wethers 1994). One trial excluded patients with large leg ulcers, but did not define the limits of inclusion (Baum 1987). Two RCTs reported baseline ulcer size using median area (Baum 1987; La Grenade 1993). Two RCTs reported baseline ulcer size using mean area (McMahon 2010; Wethers 1994). Three RCTs reported the number of ulcers per patient (Baum 1987; McMahon 2010; Serjeant 1977). See Characteristics of included studies for details.

Intervention characteristics

1. Type of intervention

Each trial investigated a different intervention to treat SCLUs. Baum assessed topical antibiotics (Baum 1987), La Grenade assessed Solcoseryl® and DuoDerm as two arms of a three-arm
trial (La Grenade 1993), McMahon assessed arginine butyrate (McMahon 2010), Serjeant assessed isoxsuprine (Serjeant 1977) and propionyl-L-carnitine (Serjeant 1997), and Wether assessed RGD peptide matrix (Wethers 1994).

2. Administration route of intervention

Two RCTs used an oral administration (Serjeant 1977; Serjeant 1997). Three RCTs used topical interventions (Baum 1987; La Grenade 1993; Wethers 1994). One RCT used an intravenous intervention (McMahon 2010).

Outcome characteristics

Three RCTs explicitly stated their primary outcomes (Baum 1987; La Grenade 1993; Wethers 1994). The primary outcomes were pain (Baum 1987), healing rate (La Grenade 1993) and changes in per cent ulcer closure (Wethers 1994). According to Dr Graham Serjeant, interviewed by one author of this review (JKM), ‘ulcer area-change’ was the primary outcome assessed in four trials that he was involved in (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997). No RCTs explicitly described their secondary outcomes. These outcome measurements or definitions, as they were shown in each RCT, are listed within an appendix (Appendix 3).

Methodology characteristics

1. Units of randomisation and of analysis

Five RCTs declared participants as the unit of randomisation (Baum 1987; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994). One RCT chose leg ulcers as the unit of randomisation (La Grenade 1993). However, from the published papers, it was clear that two RCTs used participants as the unit of analysis (Serjeant 1997; Wethers 1994) and that four reported ulcers as the unit of analysis (La Grenade 1993; McMahon 2010; Serjeant 1977; Baum 1987). See the appendices for a summary of the unit of randomisation and the unit of analysis (Appendix 4). For those trials with ulcer as the unit of analysis, data have been narratively reported. We have contacted trial authors requesting further information that would allow us to accommodate the increased correlation that comes from multiple ulcers ‘clustered’ within an individual into our meta-analysis. If received, we would use this information in a future update of the review.

2. Number of comparison groups

Five RCTs were conducted comparing two groups (Baum 1987; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994). One RCT was conducted comparing three groups (La Grenade 1993).

3. A priori sample size estimation

One RCT was conducted with an a priori sample size estimation (Serjeant 1997). Five RCTs did not report information on sample size calculation a priori (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Wethers 1994). Dr Graham Serjeant (04 February 2011) has confirmed that four of the trials were not conducted using sample size a priori (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997).

4. Follow-up period

The follow-up period ranged between two and six months: eight weeks (Baum 1987); ten weeks (Wethers 1994); twelve weeks (La Grenade 1993; McMahon 2010; Serjeant 1997); and six months (Serjeant 1977).

5. Description of the inclusion and exclusion criteria

One RCT did not describe the inclusion criteria (Serjeant 1977) and three RCTs did not describe the exclusion criteria (La Grenade 1993; Serjeant 1977; Serjeant 1997).

6. Multiple trials reports

The search strategy identified three included RCTs with multiple preliminary reports (Baum 1987; McMahon 2010; Wethers 1994).

Excluded studies

Eight studies were excluded (Afifi 1979; Cacciola 1990b; Lucena 2007; Neves 2010; Okany 2004; Paggiaro 2010; Serjeant 1970; Sawyer 1979). Overall, the majority of these studies were non-RCT or case reports. See the Characteristics of excluded studies table for a detailed account of the reasons for exclusion.

Risk of bias in included studies

None of the included RCTs were graded as having an overall low risk of bias. The risk of bias of all domains listed above for all included trials are summarised in Figure 3 and Figure 4.
Figure 3. Risk of bias graph: review authors’ judgements about each risk of bias domain presented as percentages across all included studies.
Figure 4. Risk of bias summary: review authors’ judgements about each risk of bias domain for each included study

<table>
<thead>
<tr>
<th>Study</th>
<th>Random sequence generation (selection bias)</th>
<th>Allocation concealment (selection bias)</th>
<th>Blinding of participants and personnel (performance bias)</th>
<th>Blinding of outcome assessment (detection bias)</th>
<th>Incomplete outcome data (attrition bias)</th>
<th>Selective reporting (reporting bias)</th>
<th>Other bias</th>
</tr>
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<tbody>
<tr>
<td>Baum 1987</td>
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<td>McMahon 2010</td>
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<td>Serjeant 1977</td>
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<td>Serjeant 1997</td>
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<tr>
<td>Wethers 1994</td>
<td>?</td>
<td>+</td>
<td>+</td>
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<td>+</td>
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</table>
Allocation
One RCT used an adequate method for sequence generation (graded as low risk) (McMahon 2010). We assessed five RCTs as having an unclear risk of bias in relation to the method used for generating the allocation sequence (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997; Wethers 1994). No information was reported on allocation concealment by any of the included trials (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994). See Characteristics of included studies for details.

Blinding
We assessed participant and personnel blinding separately. Three RCTs were described within the published papers as blinded trials (Serjeant 1977; Serjeant 1997; Wethers 1994). We judged one trial to have an adequate blinding process (low risk) (Wethers 1994). No information was reported about this domain in the remaining five trials (unclear risk) (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997). See Characteristics of included studies for details.

Incomplete outcome data
One RCT was assessed as having adequately reported outcome data (low risk) (Wethers 1994). Three RCTs were assessed as having an unclear risk in this domain (Baum 1987; McMahon 2010; Serjeant 1997), and two as having a high risk of bias in this domain (La Grenade 1993; Serjeant 1977). See Characteristics of included studies for details.

Selective reporting
Two RCTs show adequate reporting (graded as low risk) (McMahon 2010; Wethers 1994). The remaining four RCTs were assessed as having a high risk of bias for this domain (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997). See Characteristics of included studies for details.

Other potential sources of bias
All of the included RCTs show some type of bias (ascertainment bias, bias in the presentation of data, design bias). See Characteristics of included studies for details. See Appendix 2 for bias definitions.

Effects of interventions
See: Summary of findings for the main comparison Isoxsuprine compared to placebo for leg ulcer in people with sickle cell disease; Summary of findings 2 Arginine butyrate plus standard local care compared to standard local care for sickle cell in people with sickle cell disease; Summary of findings 3 L-carnitine compared to placebo for leg ulcer in people with sickle cell disease; Summary of findings 4 RGD peptide matrix compared to placebo for leg ulcer in people with sickle cell disease
Six RCTs were conducted for treating leg ulcers in people with SCD (Baum 1987; La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994). The results are based on five RCTs (La Grenade 1993; McMahon 2010; Serjeant 1977; Serjeant 1997; Wethers 1994) as Baum did not report on any of the outcomes pre-defined in our protocol (Baum 1987). See Summary of findings for the main comparison; Summary of findings 2; Summary of findings 3; Summary of findings 4. Results were based on 168 participants with 210 ulcers. All RCTs assessed pharmaceutical interventions. Due to the differences in the modes of action of the pharmaceutical interventions used in the trials, we regarded it as inappropriate to combine the data available. Please note, as referred to above, for those trials where ulcers were the unit of analysis (instead of the participant), data have been narratively reported and not included in the meta-analysis (La Grenade 1993; McMahon 2010; Serjeant 1977).

Primary outcomes

I. Pharmaceutical interventions

Systemic interventions
Three trials investigated systemic interventions (McMahon 2010; Serjeant 1977; Serjeant 1997).

Primary outcomes

1. Incidence of complete closure (defined as 100% epithelization or skin closure without drainage)
   - Vascular drugs
   - Isoxsuprine
   One trial (involving 30 patients as the unit of randomisation and 46 ulcers as the unit of analysis) compared isoxsuprine with placebo (Serjeant 1977). This trial reported on: healed ulcers
Pharmacologic stimulation of HbF synthesis agents

Arginine butyrate
One trial (involving 23 patients as the unit of randomisation and 62 ulcers as the unit of analysis) assessed arginine butyrate plus standard local care versus standard local care. This trial reported a complete closure incidence of 30% (11 out of 37) in the arginine butyrate group compared with 8% (2 out of 25) in the control group at 12 weeks of trial (P = 0.056) (McMahon 2010).

Secondary outcomes

1. Ulcer-free survival following treatment for SCLU (free from leg ulcer recurrence)
   This outcome was not reported on in any of the RCTs.

2. Quality of life measures (based on any item from a validated scale, e.g. SF-36, EuroQoL, WHOQOL-BREF)
   This outcome was not reported on in any of the RCTs.

3. Incidence of amputation
   This outcome was not reported on in any of the RCTs.

4. Adverse events
   - Pharmacologic stimulation of HbF synthesis agents

Arginine butyrate (McMahon 2010).
One trial reported that no serious adverse events were reported to be directly related to the study drug (McMahon 2010). Prior to arginine butyrate treatment on this study, port-a-cath infections did occur in two participants who had ports for their standard sickle cell care.

Topical pharmaceutical interventions

Three trials investigated topical interventions (Baum 1987; La Grenade 1993; Wethers 1994). The results are based on two RCTs (La Grenade 1993; Wethers 1994)

Primary outcomes

1. Incidence of complete closure (defined as 100% epithelization or skin closure without drainage)
   - Growth factors and related

One trial of 55 participants and 14 events reported this outcome over the course of the trial with a non-statistically significant increase in complete closure noted in the RGD peptide matrix group compared with the placebo group, RR 0.40, 95% CI 0.15 to 1.04 (P = 0.06) (Wethers 1994) (Analysis 2.1).
3. Change in ulcer size

- Growth factors and related

One trial reported this outcome over the course of the trial with a statistically significant change in ulcer size noted in the RGD peptide matrix group compared with the placebo group, MD 6.60 cm² (95% CI 5.51 to 7.69) (P <0.001) (Wethers 1994) (Analysis 2.2).

A further trial of 32 participants and 49 ulcers reported this outcome over the course of the trial with a non-statistically significant change in ulcer size noted in the Solcoseryl® and hydrocolloid dressing compared with those non-receiving Solcoseryl® and hydrocolloid dressing (La Grenade 1993). It was reported that "The mean difference in reduction of ulcer area between Solcoseryl® and controls was 7.1 cm² (95% IC -0.7 to 14.9), and between hydrocolloid dressing and controls was 4.2 cm² (95% CI -3.6 to 12.0). The mean difference in relative ulcer size between Solcoseryl® and controls was 28.3 cm² (95% CI -1.9 to 58.5), and between hydrocolloid dressing and controls was -1.3 cm² (95% CI -31.5 to 28.9)."

Secondary outcomes

1. Ulcer-free survival following treatment for SCLU (free from leg ulcer recurrence)

This outcome was not reported on in any of the RCTs.

2. Quality of life measures (based on any item from a validated scale, e.g. SF-36, EuroQoL, WHOQOL-BREF)

This outcome was not reported on in any of the RCTs.

3. Incidence of amputation

This outcome was not reported on in any of the RCTs.

4. Adverse events

- Growth factors and related

One trial of 55 participants and 33 events reported the total occurrence of adverse events over the course of the trial with a non-statistically significant increase of this outcome noted in the RGD peptide matrix group compared with the placebo group, RR 0.76 (95% CI 0.50 to 1.17) (P = 0.21) (Wethers 1994) (Analysis 2.3). Wethers (for 33 participants and 5 events) also reported the treatment-related adverse event over the course of the trial, which showed a non-statistically significant increase of this outcome in the RGD peptide matrix group compared with the placebo group, RR 1.41 (95% CI 0.27 to 7.38) (P = 0.68) (Wethers 1994) (Analysis 2.3).

One trial assessing Solcoseryl® and hydrocolloid dressing reported that "Solcoseryl® was well tolerated as was Eusol except for a burning sensation reported by several patients (La Grenade 1993). Hydrocolloid dressing was not well tolerated, four patients expressing dislike because of the large volume of exudate which was smelly and required daily changes of dressing in the early part of the study, and two of these defaulted".
arginine butyrate plus standard local care compared to standard local care for sickle cell in people with sickle cell disease

**Patient or population:** sickle cell in people with sickle cell disease  
**Settings:**  
**Intervention:** arginine butyrate plus standard local care  
**Comparison:** standard local care

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Illustrative comparative risks* (95% CI)</th>
<th>Relative effect (95% CI)</th>
<th>No of Participants (studies)</th>
<th>Quality of the evidence (GRADE)</th>
<th>Comments</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>Assumed risk</td>
<td>Corresponding risk</td>
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<td>standard local care</td>
<td>arginine butyrate plus</td>
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<td></td>
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<td>standard local care</td>
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<tr>
<td><strong>Complete healing</strong></td>
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<tr>
<td>Follow-up: 12 weeks</td>
<td>See comment</td>
<td>See comment</td>
<td>Not estimable</td>
<td>23 participants (62 ulcers)</td>
<td>⬤⬤⬤⬤ very low&lt;sup&gt;1,2,3&lt;/sup&gt; This trial shows inconsistency between units of randomisation (23 participants) and unit of analysis (62 ulcers)</td>
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<td></td>
<td>(1 study; McMahon 2010)</td>
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<tr>
<td><strong>Change in ulcer size</strong></td>
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<td>Not estimable</td>
<td>(1 study; McMahon 2010)</td>
<td>⬤⬤⬤⬤ very low&lt;sup&gt;1,2,3&lt;/sup&gt; This trial shows inconsistency between units of randomisation (23 participants) and unit of analysis (62 ulcers)</td>
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<tr>
<td>Follow-up: 12 weeks</td>
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</table>

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

**GRADE Working Group grades of evidence**  
**High quality:** Further research is very unlikely to change our confidence in the estimate of effect.  
**Moderate quality:** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.  
**Low quality:** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.  
**Very low quality:** We are very uncertain about the estimate.
Unclear allocation concealment

Underpowered for this outcome.

Few participants (23 participants; 62 ulcers) and events (N = 13 complete closure).

CI: confidence interval
### L-carnitine compared to placebo for leg ulcer in people with sickle cell disease

**Patient or population:** leg ulcer in people with sickle cell disease  
**Settings:**  
**Intervention:** L-carnitine  
**Comparison:** placebo  

<table>
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<th>Outcomes</th>
<th>Treated comparison</th>
<th>Placebo comparison</th>
<th>Relative effect (95% CI)</th>
<th>No of Participants (studies)</th>
<th>Quality of the evidence (GRADE)</th>
<th>Comments</th>
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<tbody>
<tr>
<td>Change in ulcer size</td>
<td>See comment</td>
<td>See comment</td>
<td>Not estimable</td>
<td>15 (1 study; Serjeant 1997)</td>
<td>⬤⃝⃝⃝ very low</td>
<td>Mean difference: -3.90 (95% CI -13.44 to 5.64).</td>
</tr>
</tbody>
</table>

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).*

**GRADE Working Group grades of evidence**  
**High quality:** Further research is very unlikely to change our confidence in the estimate of effect.  
**Moderate quality:** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.  
**Low quality:** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.  
**Very low quality:** We are very uncertain about the estimate.

---

1 Random sequence generation: unclear.  
2 Underpowered for this outcome.  
3 Few participants (N = 15). Blinding levels were not described.  
CI: confidence interval.
### RGD peptide matrix compared to placebo for leg ulcer in people with sickle cell disease

**Patient or population:** patients with leg ulcer in people with sickle cell disease  
**Settings:**  
**Intervention:** RGD peptide matrix  
**Comparison:** placebo

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Illustrative comparative risks* (95% CI)</th>
<th>Relative effect (95% CI)</th>
<th>No of Participants (studies)</th>
<th>Quality of the evidence (GRADE)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Complete closure</strong></td>
<td>placebo, See comment</td>
<td>Not estimable</td>
<td>55 (1 study; Wethers 1994)</td>
<td>very low</td>
<td>Risk ratio: 0.40 (95% CI 0.15 to 1.04).</td>
</tr>
<tr>
<td>Follow-up: 10 weeks</td>
<td>RGD peptide matrix, See comment</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Change in size ulcers healed</strong></td>
<td>placebo, See comment</td>
<td>Not estimable</td>
<td>55 (1 study; Wethers 1994)</td>
<td>very low</td>
<td>Mean difference: 6.60 (95% CI 5.51 to 7.69).</td>
</tr>
<tr>
<td>Follow-up: 10 weeks</td>
<td>RGD peptide matrix, See comment</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Total adverse events</strong></td>
<td>placebo, See comment</td>
<td>Not estimable</td>
<td>55 (1 study; Wethers 1994)</td>
<td>very low</td>
<td>Risk ratio: 0.76 (95% CI 0.50 to 1.17).</td>
</tr>
<tr>
<td>Follow-up: 10 weeks</td>
<td>RGD peptide matrix, See comment</td>
<td></td>
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</tr>
<tr>
<td><strong>Related study treatment adverse events</strong></td>
<td>placebo, See comment</td>
<td>Not estimable</td>
<td>33 (1 study; Wethers 1994)</td>
<td>very low</td>
<td>Risk ratio: 1.41 (95% CI 0.27 to 7.38).</td>
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<tr>
<td>Follow-up: 10 weeks</td>
<td>RGD peptide matrix, See comment</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*The basis for the assumed risk (e.g. the median control group risk across studies) is provided in footnotes. The corresponding risk (and its 95% CI) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).

GRADE Working Group grades of evidence  
**High quality:** Further research is very unlikely to change our confidence in the estimate of effect.  
**Moderate quality:** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.  
**Low quality:** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.  
**Very low quality:** We are very uncertain about the estimate.
Sequence generation and allocation concealment: unclear
Underpowered to address this outcome
Few participants (N = 55) and events (N = 14)
CI: confidence interval
DISCUSSION

Summary of main results

This review of interventions for treating leg ulcers in people with SCD included six studies (198 participants with 250 ulcers) which were grouped as systemic pharmaceutical interventions (L-carnitine (Serjeant 1997), arginine butyrate (McMahon 2010), isosuprine hydrochloride (Serjeant 1977)) and topical pharmaceutical interventions (Solcoseryl® cream (La Grenade 1993), RGD peptide dressing (Wethers 1994), and topical antibiotics (Baum 1987). One trial was included but not reported any pre-specified outcome in this Cochrane review (Baum 1987). Of these, RGD peptide dressing was effective in reducing the size of treated ulcers compared with placebo (Wethers 1994). The evidence for the use of interventions to treat people with sickle cell disease is not strong. Although there is some encouraging evidence for the use of RGD peptide matrix to reduce ulcer size, given the inadequacies with the associated trial reports, we currently feel that further confirmatory work is needed. Trials had a high risk of bias and failed to show beneficial effects. Please refer to the following tables for details (Summary of findings for the main comparison; Summary of findings 2; Summary of findings 3; Summary of findings 4).

Overall completeness and applicability of evidence

This review provides inconclusive evidence on the assessed interventions due to both heterogeneity between trials, and inadequate information provided by trial reports (Hopewell 2010). During this review, we have identified the following issues, which we feel are particularly relevant to consider as further work is planned. Generally, heterogeneity between trials prevented the pooling of results, with the main areas of variation between trials being differences in outcome definition, and inconsistency between unit of randomisation and unit of analysis (Appendix 4). In this regard, it has been recently suggested that trials adopt an agreed set of core outcomes for each medical condition (Clarke 2007). This approach could help to reduce the impact of outcome reporting bias (Kirkham 2010). In particular, a lack of reported inclusion and exclusion criteria and a lack of reported outcome definitions hampered the ability to compare trials (Appendix 3).

Quality of the evidence

The main source of bias in the included studies was the lack of detail in describing the generation of the randomisation sequences or the allocation concealment (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997; Wethers 1994). We interviewed an author who was on four of the six included studies, who provided information on outstanding details. Trials also lacked detail about their blinding processes. The review authors’ assessment of the risk of bias of the included studies has been described previously and a summary can be found in Figure 3 and Figure 4. The studies were classified as having a high risk of bias. Uncertainty remains about possible harms from the interventions, due to a lack of information presented on safety data.

Potential biases in the review process

In the process of performing a systematic review, there is a group of biases called significance-chasing biases (Ioannidis 2010). These include publication bias, selective outcome reporting bias, selective analysis reporting bias, and fabrication bias (Ioannidis 2010). Publication bias represents a major threat to the validity of systematic reviews, particularly in reviews such as this one that include small trials. However, we believe that this Cochrane review has a low risk of publication bias due to the thorough trial search process. We contacted the main author of four included RCTs. Selective outcome reporting bias operates through suppression of information on specific outcomes and has similarities to study publication bias, in that ‘negative’ results remain unpublished (Ioannidis 2010). This Cochrane review found that four out of the six included RCTs have high risk of selective outcome reporting (Baum 1987; La Grenade 1993; Serjeant 1977; Serjeant 1997).

AUTHORS’ CONCLUSIONS

Implications for practice

This Cochrane review of interventions for treating leg ulcers in people with sickle cell disease provides some evidence of effectiveness for treatment with RGD peptide (Wethers 1994). However, this trial had a generally high risk of bias due to inadequacies in the trial reports. We recommend that readers interpret the trial results with caution. There was no reported information on the safety of these interventions.

Implications for research

This systematic review has highlighted a need for well-designed, high-quality randomised trials to assess the benefits and harms of interventions to improve the healing of the leg ulcer in patients with SCD. The RCTs should include outcomes such as incidence of complete closure, ulcer-free survival following treatment for SCLU, time to ulcer closure, change in ulcer size (surface area or volume), quality of life measures and adverse events. Future trials should be conducted by independent researchers and reported according to the Consolidated Standards of Reporting Trials (CONSORT) guidelines (Ioannidis 2004; Moher 2010) and using the Foundation of Patient-Centered Outcomes Research recommendations (Anonymous 2012; Gabriel 2012).
ACKNOWLEDGEMENTS

We want to express our gratitude to the following people, whose help made it possible to improve the quality of this review: Tracey Remington, Dr Ian Hambleton, Dr Graham Serjeant, Dr E. Vichinsky, Professor C. Johnson, Dame Professor Sally Davies, Dr Oswald Castro, Dr R.F. Machado, Dr A. Almeida, Professor M. Steinberg and Dr Samir Ballas.

We also want express our gratitude the Cochrane Wounds Group Wounds Group.

REFERENCES

References to studies included in this review

Baum 1987  {published data only}


La Grenade 1993  {published data only}

McMahon 2010  {published data only}


Serjeant 1977  {published data only}

Serjeant 1997  {published data only}

Wethers 1994  {published data only}

References to studies excluded from this review

Afifi 1979  {published data only}

Cacciola 1990b  {published data only}

Lucena 2007  {published data only}
Lucena A, Souza K, Anselmo C, Nascimento A, Araújo A. InGaP (670nm) laser use in the leg ulcer healing in patients with sickle cell anemia [O uso do laser de ingap (670nm) na cicatrização de úlceras de perna em pacientes com anemia falciforme].  Anais da Faculdade de Medicina da Universidade
Okany 2004 [published data only]

Paggiaro 2010 [published data only]

Sawyer 1979 [published data only]

Serjeant 1970 [published data only]

Additional references

Akinsheye 2010

Akinyanju 1979

al-Momen 1991

Alexander 2004

Alikhan 2004

Alleyne 1977

Amini-Adle 2007

Ankra-Badu 1992

Anonymous 2012

Aslan 2007

Asnani 2009

Ballas 2002

Ballas 2010

Balshem 2011

Bardakdjian-Michau 2009

Bender 2009

Bergin 2006
Cacciola 1989

Cacciola 1990a

Chalchal 2001

Clarke 2002

Clarke 2007

Creary 2008

Cumming 2008

Daudt 2007

Durosimni 1991

Eckman 1996

Edwards 2002

Espinoso 1992

Frost 1990

Gabriel 2012

Gordon 2003

GRADEPro 2008
Brozek J, Oxman A, Schünemann H. GRADEpro. 3.2 for Windows 2008.

Gulbis 2006

Guyatt 2011

Guyatt 2011a

Guyatt 2011b

Guyatt 2011c

Guyatt 2011d
Interventions for treating leg ulcers in people with sickle cell disease (Review)

Copyright © 2012 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.
Interactions for treating leg ulcers in people with sickle cell disease (Review)

Copyright © 2012 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.
Vermeulen 2004

Wasiak 2008

Weatherall 2006

Wilkinson 1998

Wood 2008

* Indicates the major publication for the study
### Characteristics of included studies  
**[ordered by study ID]**

**Baum 1987**

| **Methods** | Desing trial: parallel (2 groups).  
Study phase: not described.  
Baseline observation period: no (Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011).  
Follow-up period: 8 weeks.  
Randomisation unit: patient.  
Unit of analysis: ulcers.  
Intention to treat: unclear. |
|---|---|
| **Participants** | Total: 30.  
Antibiotic therapy: 16.  
Placebo: 14.  
Withdrawals: 7% (2/30) (1 from each group).  
Reported leg ulcers at the start of the trial: 40 (data supplied by Dr Graham Serjeant, Jamaica, February 4, 2011)  
Age: median (range). Antibiotic therapy group: 29 years (14 - 44 years); control group: 25 years (17 - 49 years)  
Number of ulcers: 40 (antibiotic therapy: 20; placebo: 20).  
Duration (median):  
Antibiotic therapy group: 3.6 years (range: 0.3 to 15 years).  
Control group: 3.5 years (range: 0.3 to 12 years).  
Mean baseline ulcer area (cm$^2$) ± SD. Antibiotic therapy group: 20.0 (14.9); placebo: 20.8 (19.0)  
Inclusion criteria:  
1. Hb SS;  
2. leg ulceration of at least 3 months duration;  
3. bacterial swabs revealed at least one of the skin pathogen;  
4. to attend at 2-weekly intervals;  
5. to agree to the trial protocol;  
6. multiple ulcers: yes, all used (data supplied by Dr Graham Serjeant, Jamaica, February 4, 2011).  
7. new ulcers discussed: not information (Data supplied by Dr Graham Serjeant, Jamaica, February 4, 2011)  
Exclusion criteria:  
1. large leg ulcers;  
2. chronic renal failure. |
| **Interventions** | Antimicrobial therapy (aerosol preparation, twice daily):  
neomycin, bacitracin and polymyxin B.  
Control (aerosol dispenser, twice daily).  
Placebo: sterile normal saline with 1:1000 red food colouring  
Co-intervention: disinfectant (Eusol). |
Rest with elevation of the affected leg(s). Systematic antibiotics: not allowed. Oral zinc sulphate was allowed.

**Outcomes**

Primary: pain. Likely secondary outcomes (these end points were not described at ‘methods section’): healing rate; bacteriological investigation. According with Dr Graham Serjeant, interviewed by one author of this review (JKM), 'ulcer area-change' was the primary outcome (February 4, 2011)

**Notes**

1. This trial was described as “randomised controlled crossover trial” at Proceedings of the Commonwealth Caribbean medical Research Council (31st Scientific Meeting, 1986, April 16-19, Port Spain, Trinidad and Tobago.
2. Country: University Hospital of the West Indies, Kingston, Jamaica.
3. Study time: 10 June to 9 September 1993.
4. A priori sample size estimation: not stated.
5. 3M Company Incorporated donated Rikospray®, an aerosol preparation of neomycin, bacitracin, and polymyxin B

**Risk of bias**

<table>
<thead>
<tr>
<th>Bias</th>
<th>Authors’ judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random sequence generation (selection bias)</td>
<td>Unclear risk</td>
<td>Quote: “patients were assigned, using size 4 block randomization” (page 847). Comments: there is an imbalance in two characteristics baseline: zinc therapy was only received by control patients and marked pain (14 placebo patients vs 8 patients treatment) Insufficient information about the sequence generation process to permit judgement of ‘low risk’ or ‘high risk’</td>
</tr>
<tr>
<td>Allocation concealment (selection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of ‘low risk’ or ‘high risk’</td>
</tr>
<tr>
<td>Blinding of participants and personnel (performance bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of ‘low risk’ or ‘high risk’</td>
</tr>
<tr>
<td>Blinding of outcome assessment (detection bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of ‘low risk’ or ‘high risk’</td>
</tr>
<tr>
<td>Incomplete outcome data (attrition bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient reporting of attrition and exclusions to permit judgement of ‘low risk’ or ‘high risk’</td>
</tr>
</tbody>
</table>
Selective reporting (reporting bias) | High risk | One or more clinically relevant and reasonably expected outcomes were not reported on; data on these outcomes were likely to have been recorded (i.e. safety). According with Dr Graham Serjeant, interviewed by one author of this review (JKM), ‘ulcer area-change’ was the primary outcome (February 4, 2011)

Other bias | High risk | Ascertainment bias and bias in the presentation of data (Appendix 2). Table of characteristics of patients at entry study only shows 28 patients. There is inconsistency relating to the study design and the healing rate end point. The inconsistency between ‘Randomisation unit: patient’ and ‘Unit of analysis: ulcers’ could generate the design and confusion bias (Appendix 2).

La Grenade 1993


Participants | Sample size: 32 participants. Randomised leg ulcers: 49 (Solcoseryl® 14; hydrocolloid dressing 14; placebo 21). Loss patients reported: 14 (Solcoseryl® 4; hydrocolloid dressing: 8; placebo: 2). (Dr. Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011). 16 (Solcoseryl®: 3); hydrocolloid dressing: 8); (Placebo: 5) Loss (%): 50%. Age: Mean (± SD): Solcoseryl®: 27.0 years (8.2); hydrocolloid dressing: 35.5 years (9.8); placebo: 30.7(8.2) Sex (males): Solcoseryl®: 93%; hydrocolloid dressing: 50%; placebo: 71%. Inclusion criteria: Patients with Hb SS. Age (years): ≥ 15 Leg ulcer duration: ≥ 16 months. Ulcer size: ≥ 3 cm. Patients attending for a 3-month baseline observation period. Exclusion criteria: not described.
Interventions

1. Solcoseryl® (de-proteinized extract of calf blood) supplied in tubes as jelly or ointment, twice daily after cleaning with Eusol, covered with a gauze dressing and supported by an Elastoweb bandage

2. Hydrocolloid dressing (hydroactive dressing) supplied as sheets of occlusive dressing and as granules, and covered with gauze and supported by an Elastoweb bandage. The dressing was replaced at weekly intervals or more often if indicated

Control: Twice-daily cleaning with a mild antiseptic agent (Eusol), wet dressing and a firm supportive elastic bandage

Outcomes

Healing ulcer.

According with Dr Graham Serjeant, interviewed by one author of this review (JKM), 'ulcer area-change' was the primary outcome (February 4, 2011)

Notes

Centre and Country: Sickle-cell Clinic of the University Hospital of the West Indies, Kingston, Jamaica.

Solcoseryl® was supplied by Solco Basle Ltd.

Hydrocolloid dressing by Convatec.

A priori sample size estimation: no described.

Risk of bias

<table>
<thead>
<tr>
<th>Bias</th>
<th>Authors' judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random sequence generation (selection bias)</td>
<td>Unclear risk</td>
<td>Quote: &quot;Unilateral ulcers were randomized into one of three treatment schedules in blocks of three&quot; and &quot;In bilateral ulcers, one leg was randomized to one of the two treatment agents and the other leg to control therapy&quot; (page 121) Insufficient information about the sequence generation process to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Allocation concealment (selection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of participants and personnel (performance bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of outcome assessment (detection bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Incomplete outcome data (attrition bias) All outcomes</td>
<td>High risk</td>
<td>Dropouts: 50% (16/32) (Placebo 24% (5/21); Solcoseryl® 21.4% (3/14); hydrocolloid dressing 57.1% (8/14). Reasons: none described.</td>
</tr>
</tbody>
</table>
La Grenade 1993  (Continued)

<table>
<thead>
<tr>
<th>Bias</th>
<th>High risk</th>
<th>Imbalance between hydrocolloid dressing group and the other groups &gt; 10%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Selective reporting (reporting bias)</td>
<td>High risk</td>
<td>1 or more clinically relevant and reasonably expected outcomes were not reported on; data on these outcomes were likely to have been recorded (i.e. safety)</td>
</tr>
<tr>
<td>Other bias</td>
<td>High risk</td>
<td>Ascertainment and design bias and bias in the presentation of data (Appendix 2). Comments: there is imbalance in sex and age.</td>
</tr>
</tbody>
</table>

McMahon 2010

<table>
<thead>
<tr>
<th></th>
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</thead>
<tbody>
<tr>
<td>Participants</td>
<td>Sample size: 26 participants Randomised leg ulcers: 62 (arginine butyrate plus standard local care: 37; standard local care: 25) Age: median (range): arginine butyrate plus standard local care: 36.6 (21-60); standard local care: 34.7 (20-57) Sex (female): arginine butyrate plus standard local care: 57%; standard local care: 58%. Mean baseline ulcer area (cm²) ± SE: arginine butyrate plus standard local care: 25.7 cm² (8.1); standard local care: 50.6 cm² (13.9). Total number ulcers: arginine butyrate plus standard local care: 37; standard local care: 25. Number of ulcers per patient: arginine butyrate plus standard local care: 3.5; standard local care: 2.1 Inclusion criteria: Age: &gt;18 years. Sickle cell anaemia or sickle beta thalassaemia. Presence of one or more leg ulcers which had been refractory to healing with National Institutes of Health (NIH)-defined standard local care, consisting of twice-daily cleaning and wet-to-dry dressing changes, for at least 6 months. Patients were also eligible to participate if an ulcer had recurred and not healed with at least 3 months of standard care Exclusion criteria: Renal or hepatic compromise. Current chronic transfusion therapy.</td>
</tr>
</tbody>
</table>
**Interventions**

| Intervention* | arginine butyrate at a total daily dose of 500 mg/kg/dose given 5 d/week plus standard local care. | Arginine butyrate formulation: 5% butyric acid (50 g/l) and 7.5% L-arginine (75 g/l) by intravenous infusion through a port-a-cath or peripheral pass-port, generally over 6-8 h/d. | The administration rate: 85 mg/kg/h and those patients infused with rates > 60 mg/kg/h were pre-medicated with acetaminophen or ibuprofen and an anti-emetic to prevent headache or nausea. Patients were encouraged and assisted to ambulate frequently during the 6-h infusion | Control arm: standard local care alone. | Co-intervention: topical antibiotics and surgical debridement. Only 1 subject received concomitant hydroxycarbamide (in the control arm) |

**Outcomes**

| Wound healing: partial healing (decrease in ulcer area by at least 25% of the baseline ulcer area) and complete healing (complete closure of the ulcer (to an area of 0 cm²)). |

**Notes**

* If healing was objectively documented during the first 12-week treatment cycle, as determined by a decrease in measured ulcer area by at least 25% of the baseline area, arginine butyrate could be continued for two additional courses of 8-week cycles, although the responses to the extended treatment were not analysed as study endpoints. Control arm participants could cross-over to the treatment arm if their ulcer did not heal after 12 weeks of closely monitored and supervised standard local care. Their remaining ulcers were then assessed on the treatment arm for 12 weeks. |

A priori sample size estimation: no. |

Country: USA. |

Founders: the Food and Drug Administration, Office of Orphan Product Development (Grant FD-R-000176) and the General Clinical Research Center at Boston University (Grant M01 RR00533) |

**Risk of bias**

<table>
<thead>
<tr>
<th>Bias</th>
<th>Authors' judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random sequence generation (selection bias)</td>
<td>Low risk</td>
<td>Quote &quot;table of random numbers prepared by a blinded statistician.&quot; (page 517)</td>
</tr>
<tr>
<td>Allocation concealment (selection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of participants and personnel (performance bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of outcome assessment (detection bias) All outcomes</td>
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</table>
**McMahon 2010 (Continued)**

<table>
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<tr>
<th>Incomplete outcome data (attrition bias)</th>
<th>Unclear risk</th>
<th>Insufficient reporting of attrition/exclusions to permit judgement of ‘low risk’ or ‘high risk’</th>
</tr>
</thead>
<tbody>
<tr>
<td>All outcomes</td>
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</table>

<table>
<thead>
<tr>
<th>Selective reporting (reporting bias)</th>
<th>Low risk</th>
<th>-</th>
</tr>
</thead>
<tbody>
<tr>
<td>Other bias</td>
<td>High risk</td>
<td>Bias in the presentation of data (Appendix 2). The inconsistency between ‘randomisation unit: patient’ and ‘unit of analysis: ulcers’ could generate the design and confusion bias (Appendix 2).</td>
</tr>
</tbody>
</table>

**Serjeant 1977**

<table>
<thead>
<tr>
<th>Methods</th>
<th>Design trial: parallel (two groups).</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study phase: III.</td>
<td>Study phase: III.</td>
</tr>
<tr>
<td>Baseline observation period: no (Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011).</td>
<td>Baseline observation period: no (Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011).</td>
</tr>
<tr>
<td>Follow-up period: 6 months.</td>
<td>Follow-up period: 6 months.</td>
</tr>
<tr>
<td>Randomisation unit: patients.</td>
<td>Randomisation unit: patients.</td>
</tr>
<tr>
<td>Unit of analysis: ulcers.</td>
<td>Unit of analysis: ulcers.</td>
</tr>
<tr>
<td>Intention-to-treat analysis: no (information was supplied by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011)</td>
<td>Intention-to-treat analysis: no (information was supplied by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Participants</th>
<th>Enrolled: not described.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Randomised: 40 (unclear). This number was confirmed by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011.</td>
<td>Randomised: 40 (unclear). This number was confirmed by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011.</td>
</tr>
<tr>
<td>1. Reported patients: 40</td>
<td>1. Reported patients: 40</td>
</tr>
<tr>
<td>(This number was supplied by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011).</td>
<td>(This number was supplied by Dr Graham Serjeant, interviewed by one author of this review (JKM), on February 4, 2011).</td>
</tr>
<tr>
<td>3. Reported leg ulcers at the start trial: 63</td>
<td>3. Reported leg ulcers at the start trial: 63</td>
</tr>
<tr>
<td>• 3.1. 1 ulcer reported by 23 patients: 23 ulcers.</td>
<td>• 3.1. 1 ulcer reported by 23 patients: 23 ulcers.</td>
</tr>
<tr>
<td>• 3.2. 2 ulcers reported by 12 patients: 24 ulcers.</td>
<td>• 3.2. 2 ulcers reported by 12 patients: 24 ulcers.</td>
</tr>
<tr>
<td>• 3.3. 3 ulcers reported by 4 patients: 12 ulcers.</td>
<td>• 3.3. 3 ulcers reported by 4 patients: 12 ulcers.</td>
</tr>
<tr>
<td>• 3.4. 4 ulcers reported by 1 patient: 4 ulcers.</td>
<td>• 3.4. 4 ulcers reported by 1 patient: 4 ulcers.</td>
</tr>
<tr>
<td>• Total: 63 ulcers.</td>
<td>• Total: 63 ulcers.</td>
</tr>
<tr>
<td>6. Reported leg ulcers completing trial: 46 (75% 30/40 patients).</td>
<td>6. Reported leg ulcers completing trial: 46 (75% 30/40 patients).</td>
</tr>
<tr>
<td>7. Ulcers developed during trial which completed trial: 8 (# patients: no reported)</td>
<td>7. Ulcers developed during trial which completed trial: 8 (# patients: no reported)</td>
</tr>
</tbody>
</table>

| Isoxuprine group: 3.6 years (range: 0.3 to 15). |
| Placebo group: 3.5 years (range: 0.3 to 12). |
Allocated group: not mentioned.
Information about haemoglobin disorder type:
Homozygous sickle cell disease "Diagnosed by haemoglobin electrophoresis on cellulose acetate and agar gel, and quantification of $A_2$ levels": 38.
Sickle cell-haemoglobin C disease: 1.
Sickle cell-haemoglobin O (Arab) disease: 1.

Age (years): 17 to 67.
Sex (% male): 52.5 (21/40).
Inclusion criteria: information was not supplied.
Exclusion criteria: information was not supplied.

Interventions
Experimental group:
Isoxuprine hydrochloride: 40 mg, per oral (bd).
Control group:
Placebo: 40 mg, per oral (bd).
Co-interventions:
Local ulcer therapy: dressings plus mild antiseptic agent (not named): bd

Outcomes
Outcomes were not describe as primary or secondary. They were reported into 'results' section:
Variation ulcer size; healed.
According with Dr Graham Serjeant, interviewed by one author of this review (JKM), 'ulcer area-change' was the primary outcome (February 4, 2011)

Notes
Centre and Country: Medical Research Council Laboratories, University of West India, Jamaica
A priori sample size estimation: no.
Support: Mead Johnson and Company.
Founder role: it supplied isoxsuprine hydrochloride and financial assistance

Risk of bias

<table>
<thead>
<tr>
<th>Bias</th>
<th>Authors' judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random sequence generation (selection bias)</td>
<td>Unclear risk</td>
<td>Quote &quot;patients were allotted serial numbers in a code...&quot; (page 164)</td>
</tr>
<tr>
<td>Allocation concealment (selection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information about the sequence generation process to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of participants and personnel (performance bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>All outcomes</td>
<td></td>
<td>Comment: Dr Searjent referred this paper</td>
</tr>
<tr>
<td>Blinding of outcome assessment (detection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>-----------------------------------------------</td>
<td>--------------</td>
<td>---------------------------------------------------------------------</td>
</tr>
<tr>
<td>All outcomes</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

| Incomplete outcome data (attrition bias)      | High risk    | The number or reasons for dropouts and withdrawals were not described. Comments: this RCT loss 25% (10/40) of the patients and 27% (17/63) of the leg ulcers |
| All outcomes                                  |              |                                                                     |

| Selective reporting (reporting bias)          | High risk    | One or more clinically relevant and reasonably expected outcomes were not reported on; data on these outcomes were likely to have been recorded |
|                                              |              |                                                                     |

| Other bias                                    | High risk    | Quote “The code was kept by the producing company”. Comments. 1. Characteristic baseline table: not shown. 2. It is unclear data on randomisation unit reported by the authors. Dr Graham Serjeant pointed out unit of randomisation were the patients, and unit of analysis were the ulcers (Interviewed by one author of this review (JKM), on February 4, 2011). 3. Ulcers developed during trail were included which were analysed. 4. There is inconsistency regarding numbers of the total ulcers. 5. Ascertaiment bias and bias in the presentation of data (Appendix 2). 6. The inconsistency between 'Randomisation unit: patient' and 'Unit of analysis: ulcers' could generate the design and confusion bias (Appendix 2). |
|                                              |              |                                                                     |
### Methods

Design trial: parallel (2 groups).
Study phase: no described.
Baseline observation period: 12 weeks.
Follow-up period: 12 weeks.
Randomisation unit: patients.
Unit of analysis: patients.
Intention-to-treat analysis: declared. It was unclear, according with Dr Graham Serjeant, interviewed by one author of this review (JKM) on February 4, 2011.

### Participants

1. Enrolled: not described.
2. Randomised: 15 (experimental group: 7; placebo: 8).
3. Excluded: 2 (experimental group: 1; placebo: 1).
4. Active participants: 13 (87%) (experimental group: 6; placebo: 7).
5. Reason for exclusion (2/15 = 13%)
   Quote: “One subject in the placebo group default after 27 days, and 1 participant in the PLC group, admitted to the hospital with acute chest pain after 46 days, failed to take the treatment for 54 days, and then resumed, but compliance was erratic” (Page 492)
6. Reported leg ulcers at the start trial: 34 (Data supplied by Dr Graham Serjeant, Jamaica, February 4, 2011)
7. Sex (male): 12 (80%).
8. Age (years): (arithmetic range):
   Total Group: 17 to 40.
   Experimental group: not described / Placebo group: not described
9. Clinical characteristics of the leg ulcers: Leg ulcers of at least six months’ duration and at least 3 cm diameter at the onset of treatment.
10. Inclusion criteria:
   Patients with sickle cell disease.
   Age (years): 17 to 40.
   Leg ulcer duration: ≥ 6 months.
   Leg ulcer diameter: ≥ 3 cm at the onset of treatment.
11. Exclusion criteria: not described.

### Interventions

Baseline treatment (12-weeks): all patients
2. Photographic images: at 4-week intervals.
3. Dressing with 0.01% potassium permanganate: twice daily.
4. Debridament of dirty ulcers with crushed unripe papaya.
5. Oral zinc sulphate: 200 mg per oral, 3-times-daily, after meals
   Experimental group:
   Propionyl-L-carnitine (PLC): 2 gr/ per oral/ twice-daily.
   Control group:
   Placebo: lactose and microcrystalline cellulose/per oral/twice-daily
   Co-interventions:
   As for baseline period.

### Outcomes

Healing rate.
Quote: “... was standardized to a 12-weeks period and expressed as (1) absolute change in area (change in ulcer area over treatment period divided by number of weeks on treatment ) x 12; and (2) percentage change in area (absolute change in area + area at
Serjeant 1997  (Continued)

According with Dr Graham Serjeant, interviewed by one author of this review (JKM), 'ulcer area-change' was the primary outcome (February 4, 2011)

| Notes | RCT start date: not described. Centre and Country: Conducted at Medical Research Council Laboratories, University of West India, Jamaica A priori sample size estimation: 14 patients (7 patients by group)  
- P-value: 0.05.  
- Healing rate: 15% over three months.  
- Power: 80%.  
- Percentage healing rate: 74%.  
- Trial number: not described.  
- Expected loss: not described. Support: Sigma-Tau Pharmaceuticals, Gaitherborg, Md (Partial grant) Founder role: not described. |

<table>
<thead>
<tr>
<th>Risk of bias</th>
<th>Authors' judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random sequence generation (selection bias)</td>
<td>Unclear risk</td>
<td>Quote &quot;...were randomly assigned to a treatment group, seven in the PLC group and eight in the placebo group&quot; (pages 491-2) Quote &quot;randomisation within small blocks (2-4)&quot; This information was supplied by Dr Serjeant on February 4, 2011 Insufficient information about the sequence generation process to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Allocation concealment (selection bias)</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Blinding of participants and personnel (performance bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk' Comment: Dr Serjeant referred this paper as blinded (Interviewed by one author of this review (JKM), on February 4, 2011)</td>
</tr>
<tr>
<td>Blinding of outcome assessment (detection bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient information to permit judgement of 'low risk' or 'high risk'</td>
</tr>
<tr>
<td>Incomplete outcome data (attrition bias) All outcomes</td>
<td>Unclear risk</td>
<td>Insufficient reporting of attrition/exclusions to permit judgement of 'low risk' or 'high risk'</td>
</tr>
</tbody>
</table>
### Serjeant 1997

<table>
<thead>
<tr>
<th>Selective reporting (reporting bias)</th>
<th>High risk</th>
<th>if one or more clinically relevant and reasonably expected outcomes were not reported on; data on these outcomes were likely to have been recorded</th>
</tr>
</thead>
</table>

| Other bias | High risk | Design bias ([Appendix 2](#)). Comment: there is imbalance: ulcer size before treatment ($cm^2$) ($\pm$SD): 45.4(46.9) [4.9 to 146.28 range] in placebo group vs 19.7 (4.7) [13.7 to 27.9] in experimental group |

### Wethers 1994

#### Methods
- Design trial: parallel (2 groups).
- Multicenter study: yes
- Study phase: no described.
- Follow-up period: 10 weeks.
- Baseline observation period: none declared.
- Randomisation unit: patients.
- Unit of analysis: patients.
- Intention-to-treat analysis: yes.

#### Participants
- Hb genotype: HbSS (50), HbSP thalassaemia (2), HbSC(1), HbSC Harlem (1), HbSD (1)
- Completed study: 48 (87%). RGD peptide matrix (27/32 = 84%), placebo (21/23 = 91%)
- Loss: 7 (RGD peptide matrix: 5/32:16%) / placebo (2/23:9%)
- Imbalance between both groups: 7%

Age: median($\pm$ SE) and range: RGD peptide matrix: 32.9(1.8) years, range 18 - 65 years; placebo: 36.6(2.3) years, range 26 - 65 years

Baseline ulcer duration (months) median($\pm$ SE) and range: RGD peptide matrix: 52.0 months (12.6), range 1 - 312; placebo: 45.9 months (12.4), range 1 - 192.

Baseline ulcer area ($cm^2$) median($\pm$SE) and range: RGD peptide matrix: 14.8(3.3), range 0.4 -91.3; placebo: 11.0(2.4), range 0.9-36.7

Gender (male: female): RGD peptide matrix: 21:11; placebo: 12:11

Inclusion criteria:
- 1. isolated full-thickness lower leg or ankle ulcers that did not involve bone or tendon and had persisted at least 1 month;
- 2. age: $\geq$18 years old;
- 3. informed written consent.

Exclusion criteria:
- 1. medical conditions that might retard healing (immune system, diseases, uncontrolled diabetes, bleeding disorders, neurological disorders, or cancer requiring chemotherapy or radiation treatment);
- 2. receiving medications that might adversely affect healing (systemic corticosteroids
or antineoplastic agents);  
3. history of chronic transfusion therapy within the 3 months preceding study 
commencement.

Interventions

RGD peptide matrix (Argidene gel: formerly Telio-Derm gel, Telios Pharmaceuticals, 
Inc, San Diego, CA).  
Applications and dressing changes: once per week by 10 weeks  
Placebo: saline solution.  
Applications and dressing changes: once per week by 10 weeks  
Co-intervention: debridement and cleaning at each visit.

Outcomes

Primary: changes in per cent ulcer closure.  
Secondary: none declared.

Notes

Country: conducted at USA.  
A priori sample size estimation: no.  
Support: Telios Pharmaceuticals, Inc.  
Founder role: none declared.

Risk of bias

<table>
<thead>
<tr>
<th>Bias</th>
<th>Authors' judgement</th>
<th>Support for judgement</th>
</tr>
</thead>
</table>
| Random sequence generation (selection bias) | Unclear risk       | Quote “sequentially assigned to treatment groups based on a unique randomization number list” (page 1776)  
Comments: imbalance exists on 'baseline ulcer area (cm^2 ).  
No Information reported about how “unique randomization number list” was generated |
| Allocation concealment (selection bias)   | Unclear risk       | Insufficient information to permit judgement of 'low risk' or 'high risk'              |
| Blinding of participants and personnel (performance bias) All outcomes | Low risk | Patient and outcome assessor were blinded. |
| Blinding of outcome assessment (detection bias) All outcomes | Low risk | Patient and outcome assessor were blinded. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Dropouts < 20% |
| Selective reporting (reporting bias)      | Low risk           | -                                                                                     |
| Other bias                                | High risk          | Design bias (Appendix 2).                                                             |
Characteristics of excluded studies  [ordered by study ID]

<table>
<thead>
<tr>
<th>Study</th>
<th>Reason for exclusion</th>
</tr>
</thead>
<tbody>
<tr>
<td>Afifi 1979</td>
<td>Author did not report data from the patients suffering sickle cell thalassaemia</td>
</tr>
<tr>
<td>Cacciola 1990b</td>
<td>Not a randomised clinical trial.</td>
</tr>
<tr>
<td>Lucena 2007</td>
<td>Not a randomised clinical trial.</td>
</tr>
<tr>
<td>Neves 2010</td>
<td>Case report.</td>
</tr>
<tr>
<td>Okany 2004</td>
<td>Not a randomised clinical trial.</td>
</tr>
<tr>
<td>Paggiaro 2010</td>
<td>Case study.</td>
</tr>
<tr>
<td>Sawyer 1979</td>
<td>Case report including three types of ulcers.</td>
</tr>
<tr>
<td>Serjeant 1970</td>
<td>Not a randomised clinical trial.</td>
</tr>
</tbody>
</table>
## DATA AND ANALYSES

### Comparison 1. L-carnitine versus placebo

<table>
<thead>
<tr>
<th>Outcome or subgroup title</th>
<th>No. of studies</th>
<th>No. of participants</th>
<th>Statistical method</th>
<th>Effect size</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Change in ulcer size (surface area or volume)</td>
<td>1</td>
<td></td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>Totals not selected</td>
</tr>
<tr>
<td>1.1 All randomised patients</td>
<td>1</td>
<td></td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>0.0 [0.0, 0.0]</td>
</tr>
</tbody>
</table>

### Comparison 2. RGD peptide matrix versus placebo

<table>
<thead>
<tr>
<th>Outcome or subgroup title</th>
<th>No. of studies</th>
<th>No. of participants</th>
<th>Statistical method</th>
<th>Effect size</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Incidence of complete closure</td>
<td>1</td>
<td></td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>Totals not selected</td>
</tr>
<tr>
<td>2 Change in size of ulcers healed</td>
<td>1</td>
<td></td>
<td>Mean Difference (IV, Fixed, 95% CI)</td>
<td>Totals not selected</td>
</tr>
<tr>
<td>3 Adverse events</td>
<td>1</td>
<td></td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>Totals not selected</td>
</tr>
<tr>
<td>3.1 Total</td>
<td>1</td>
<td></td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>0.0 [0.0, 0.0]</td>
</tr>
<tr>
<td>3.2 Related study treatment</td>
<td>1</td>
<td></td>
<td>Risk Ratio (M-H, Fixed, 95% CI)</td>
<td>0.0 [0.0, 0.0]</td>
</tr>
</tbody>
</table>

### Analysis 1.1. Comparison 1 L-carnitine versus placebo, Outcome 1 Change in ulcer size (surface area or volume).

Review: Interventions for treating leg ulcers in people with sickle cell disease

Comparison: 1 L-carnitine versus placebo

Outcome: 1 Change in ulcer size (surface area or volume)

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>L-carnitine</th>
<th>Placebo</th>
<th>Mean Difference</th>
<th>Mean Difference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N Mean(SD)</td>
<td>N Mean(SD)</td>
<td>IV,Fixed,95% CI</td>
<td>IV,Fixed,95% CI</td>
</tr>
<tr>
<td>1 All randomised patients</td>
<td>7</td>
<td>-3.2 (5.2)</td>
<td>8</td>
<td>0.7 (12.6)</td>
</tr>
</tbody>
</table>

-50 -25 0 25 50
Favours L-carnitine Favours placebo
Analysis 2.1. Comparison 2 RGD peptide matrix versus placebo, Outcome 1 Incidence of complete closure.

Review: Interventions for treating leg ulcers in people with sickle cell disease

Comparison: 2 RGD peptide matrix versus placebo

Outcome: 1 Incidence of complete closure

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>RGD peptide matrix</th>
<th>Placebo</th>
<th>Risk Ratio</th>
<th>Risk Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n/N</td>
<td>n/N</td>
<td>M-H,Fixed,95% CI</td>
<td>M-H,Fixed,95% CI</td>
</tr>
<tr>
<td>Wethers 1994</td>
<td>5/32</td>
<td>9/23</td>
<td>0.40 [0.15, 1.04]</td>
<td></td>
</tr>
</tbody>
</table>

Analysis 2.2. Comparison 2 RGD peptide matrix versus placebo, Outcome 2 Change in size of ulcers healed.

Review: Interventions for treating leg ulcers in people with sickle cell disease

Comparison: 2 RGD peptide matrix versus placebo

Outcome: 2 Change in size of ulcers healed

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>RGD peptide matrix</th>
<th>Placebo</th>
<th>Mean Difference</th>
<th>Mean Difference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N</td>
<td>Mean(SD)</td>
<td>IV,Fixed,95% CI</td>
<td>IV,Fixed,95% CI</td>
</tr>
<tr>
<td>Wethers 1994</td>
<td>32</td>
<td>10.2 (2.8)</td>
<td></td>
<td>6.60 [5.51, 7.69]</td>
</tr>
</tbody>
</table>
Analysis 2.3. Comparison 2 RGD peptide matrix versus placebo, Outcome 3 Adverse events.

Review: Interventions for treating leg ulcers in people with sickle cell disease

Comparison: 2 RGD peptide matrix versus placebo

Outcome: 3 Adverse events

<table>
<thead>
<tr>
<th>Study or subgroup</th>
<th>RGD peptide matrix</th>
<th>Placebo</th>
<th>Risk Ratio</th>
<th>Risk Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n/N</td>
<td>n/N</td>
<td>M-H, Fixed, 95% CI</td>
<td>M-H, Fixed, 95% CI</td>
</tr>
<tr>
<td>1 Total</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wethers 1994</td>
<td>17/32</td>
<td>16/23</td>
<td>0.76 [0.50, 1.17]</td>
<td></td>
</tr>
<tr>
<td>2 Related study treatment</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wethers 1994</td>
<td>3/17</td>
<td>2/16</td>
<td>1.41 [0.27, 7.38]</td>
<td></td>
</tr>
</tbody>
</table>

**APPENDICES**

Appendix 1. Glossary of medical terms.

<table>
<thead>
<tr>
<th>Terms</th>
<th>Definition</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hb SS</td>
<td>An abnormal hemoglobin resulting from the substitution of valine for glutamic acid at position 6 of the beta chain of the globin moiety. The heterozygous state results in sickle cell trait, the homozygous in sickle cell anemia</td>
<td>National Library of Medicine; Mesh home page.</td>
</tr>
<tr>
<td>Hb SC</td>
<td>One of the sickle cell disorders characterized by the presence of both hemoglobin S and hemoglobin C. It is similar to, but less severe than sickle cell anemia</td>
<td>National Library of Medicine; Mesh home page.</td>
</tr>
<tr>
<td>L-carnitine</td>
<td>A derivative of the amino acid lysine, required for the transport of fatty acids into mitochondria for oxidation</td>
<td>Bender 2009</td>
</tr>
<tr>
<td>Solcoseryl®</td>
<td>Solcoseryl® is a tissue metabolism activator, chemically and biologically standardized and deproteinized, non antigenic and pyrogenic hemodil-</td>
<td><a href="http://www.drugspro.org/">http://www.drugspro.org/</a></td>
</tr>
</tbody>
</table>
**DuoDerm**

DuoDerm (Granuflex®, Convatec, Uxbridge, UK, marketed as DuoDerm® in the USA) A hydrocolloid dressing containing colloids and elastomeric and adhesive components.

**Solcoseryl®** contains a large amount of natural low molecular substances - glycolipides, nucleosides, nucleotides, amino acids, oligopeptides, irreplaceable micro elements, electrolytes.

**RGD peptide matrix**

Arginine-glycine-aspartic acid (RGD). RGD peptide matrix is designed to act as a temporary, topical synthetic extracellular matrix that substitutes for the damaged natural matrix and provides support for cell ingrowth into the ulcer site. This synthetic matrix consists of an RGD-containing peptide complexed with sodium hyaluronate in a sterile, non preserved viscous gel. The peptide matrix contains the arginine-glycine-aspartic acid amino acid sequence, by which cells in vivo become attached to extracellular matrix macromolecules via surface integrin receptors. The matrix is a sterile non-preserved clear viscous gel, formulated in phosphate-buffered saline and dispensed from a single-use syringe container. The functional ingredient of RGD peptide matrix is a complex formed by the combination of a synthetic 18 amino acid peptide and sodium hyaluronate. It also contains added unconjugated sodium hyaluronate as a viscosity-increasing agent, and therefore does not require preparation from patient samples.

**Isoxsuprine**


**Oral zinc sulphate**

A salt of zinc used as a supplement for treating zinc deficiency. [Bender 2009](http://www.cancer.gov/drugdictionary)

**Arginine butyrate**

The butyric acid salt of the amino acid arginine. [http://www.cancer.gov/drugdictionary](http://www.cancer.gov/drugdictionary)

**Haemoglobinopathy**

Any of a group of inherited diseases, such as thalassaemia and sickle-cell disease, in which there is an abnormality in the production of haemoglobin. [Martin 2011](http://www.cancer.gov/drugdictionary)
Priapism | Persistent and usually painful erection of the penis that requires urgent decompression; occasionally, however, it can be painless, in which case its treatment is less urgent | Martin 2011

Pulmonary hypertension | Condition in which there is raised blood pressure within the blood vessels supplying the lungs (the pulmonary artery blood pressure is normally much lower than the pressure within the aorta and its branches) | Martin 2011

Nitric oxide | An important member of the group of gaseous mediators, which - together with amine mediators (e.g. adrenaline, noradrenaline, histamine, acetylcholine) and lipid mediators (e.g. prostaglandins) - produce many physiological responses | Martin 2011

Endothelium | The single layer of cells that lines the heart, blood vessels, and lymphatic vessels | Martin 2011

Antithrombin III | Coagulation inhibitory protein | Santen 2004

**Appendix 2. Bias definition**

<table>
<thead>
<tr>
<th>Bias (Porta 2008)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bias in the presentation of data</td>
<td>Error due to irregularities produced by digit preference, incomplete data, poor techniques of measurement, technically poor laboratory procedures, or intentional attempts to mislead.</td>
</tr>
<tr>
<td>Selection bias</td>
<td>Distortions that result from procedures used to select subjects and from factors that influence participation in the study.</td>
</tr>
<tr>
<td>Reporting bias</td>
<td>Selective revealing or suppression of information.</td>
</tr>
<tr>
<td>Design bias</td>
<td>The difference between a true value and that obtained as a result of faulty design of a study.</td>
</tr>
</tbody>
</table>
## Appendix 3. Outcome definitions as described by the RCTs’ author

<table>
<thead>
<tr>
<th>Trials</th>
<th>Assessment of ulcer healing</th>
<th>Change in ulcer size (surface area or volume)</th>
<th>Complete closure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baum 1987</td>
<td>&quot;Ulcer was measured in two dimensions and photographs taken for determination of ulcer&quot; as described by Serjeant 1970.</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>La Grenade 1993</td>
<td>It was analysed from the measured long and short dimensions of the ulcer, arbitrarily assuming and elliptical shape.</td>
<td>It was expressed as in real changes and percentage changes in area</td>
<td>-</td>
</tr>
<tr>
<td>McMahon 2010</td>
<td>&quot;During the weekly clinical visits, the ulcer was traced on acetate film and photographed. All ulcer areas were then calculated by computerized planimetry, using the IMAGEJ software (NIH, Bethesda, MD, USA)&quot;</td>
<td>NA</td>
<td>Complete closure of the ulcer (to an area of 0 cm²).</td>
</tr>
<tr>
<td>Serjeant 1977</td>
<td>&quot;The ulcer periphery was defined on the photograph, cut out, weighed, and the area of the ulcer calculated from the weight/area ratio of the photographic paper&quot;</td>
<td>Default: &quot;defined as less than 4 months satisfactorily attendance at the trial&quot;. No change: &quot;variation of less than 1 cm² in ulcer area during the 6-month period&quot;. Healed ulcers &quot;expressed as a proportion of those ulcers likely to heal on the basis of small size (8 cm²).&quot;</td>
<td>-</td>
</tr>
</tbody>
</table>
| Serjeant 1997   | “Ulcer size was measured by area (measured in square centimeters), computing an assumed ellipse from long and short axis measurements. When more than one ulcer was present, the average area calculated for that patient was used in the analysis” | Period: 12 weeks.  
(1) absolute change in area (change in ulcer area over treatment period divided by number of weeks on treatment) x 12 and (2) percentage change in area (absolute change in area + area at start of treatment)” | -                |
| Wethers 1994    | “Changes in percent ulcer closure occurring between study commencement and endpoint” | -                                             | -                |
## Appendix 4. Unit of randomisation versus unit of analysis

<table>
<thead>
<tr>
<th>Study</th>
<th>Unit of randomisation</th>
<th>Unit of analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baum 1987</td>
<td>participants</td>
<td>ulcer</td>
</tr>
<tr>
<td>La Grenade 1993</td>
<td>ulcer</td>
<td>ulcer</td>
</tr>
<tr>
<td>McMahon 2010</td>
<td>participants</td>
<td>ulcer</td>
</tr>
<tr>
<td>Serjeant 1977</td>
<td>participants</td>
<td>ulcer</td>
</tr>
<tr>
<td>Serjeant 1997</td>
<td>participants</td>
<td>participants</td>
</tr>
<tr>
<td>Wethers 1994</td>
<td>participants</td>
<td>participants</td>
</tr>
</tbody>
</table>

## Appendix 5. LILACS search strategy

**Key words**

- sickle cell disease
- leg ulcer

```
((Pt ENSAIO CONTROLADO ALEATORIO OR Pt ENSAYO CLINICO CONTROLADO OR Mh ENSAYOS CONTROLADOS ALEATORIOS OR Mh DISTRIBUCIÓN ALEATORIA OR Mh METODO DOBLE CIEGO OR Mh METODO SIMPLE-CIEGO OR Pt ESTUDIO MULTICÉNTRICO) or ((tw ensaio or tw ensayo or tw trial) and (tw azar or tw acaso or tw placebo or tw control$ or tw aleat$ or tw random$ or (tw duplo and tw cego) or (tw doble and tw ciego) or (tw double and tw blind)) and tw clinic$)) AND NOT ((Ct ANIMALES OR Mh ANIMALES OR Ct CONEJOS OR Ct RATÓN OR Mh Ratas OR MH Primates OR MH Perros OR MH Conejos OR MH Porcinos) AND NOT (Ct HUMANO AND Ct ANIMALES)) [Palabras] and sickle [Palabras del resumen] and ulcer [Palabras] (Strategy results:10)
```

## Appendix 6. African Index Medicus (AIM) search strategy

**Key words**

- sickle cell disease
- leg ulcer
- wound

Search result: 0 references.
Appendix 7. ISI Web of Knowledge

Key words

sickle
leg ulcer
Search result: 47 references.

HISTORY

Protocol first published: Issue 3, 2010
Review first published: Issue 11, 2012

CONTRIBUTIONS OF AUTHORS

Arturo Martí-Carvajal conceived and drafted the review with comments from Jennifer Knight-Madden and María José Martínez-Zapata.

Arturo Martí-Carvajal acts as guarantor for the review.

DECLARATIONS OF INTEREST

In 2004 Arturo Martí-Carvajal was employed by Eli Lilly to run a four-hour workshop on 'How to critically appraise clinical trials on osteoporosis and how to teach this'. This activity was not related to his work with The Cochrane Collaboration or any Cochrane Review.

In 2007 Arturo Martí-Carvajal was employed by Merck to run a four-hour workshop 'How to critically appraise clinical trials and how to teach this'. This activity was not related to his work with The Cochrane Collaboration or any Cochrane Review.

Jennifer Knight-Madden and María José Martínez-Zapata: none known.

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