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[Intervention Review]

Interventions for treating phosphorus burns

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ABSTRACT

Background

Phosphorus burns are rarely encountered in usual clinical practice and occur mostly in military and industrial settings. However, these burns can be fatal, even with minimal burn area, and are often associated with prolonged hospitalisation.

Objectives

To summarise the evidence of effects (beneficial and harmful) of all interventions for treating people with phosphorus burns.

Search methods

In October 2013 for this first update we searched the Cochrane Wounds Group Specialised Register; the Cochrane Central Register of Controlled Trials (CENTRAL) (The Cochrane Library); Ovid OLDMEDLINE; Ovid MEDLINE; Ovid MEDLINE (In-Process & Other Non-Indexed Citations); Ovid EMBASE; EBSCO CINAHL and Conference Proceedings Citation Index - Science (CPCI-S). We did not apply any methodological filters or restrictions on the basis of study design, language, date of publication or publication status.

Selection criteria

Any comparisons of different ways of managing phosphorus burns including, but not restricted, to randomised trials.

Data collection and analysis

We found two non-randomised comparative studies, both comparing patients treated with and without copper sulphate.

Main results

These two comparative studies provide no evidence to support the use of copper sulphate in managing phosphorus burns. Indeed the small amount of available evidence suggests that it may be harmful.

Authors' conclusions

First aid for phosphorus burns involves the common sense measures of acting promptly to remove the patient's clothes, irrigating the wound(s) with water or saline continuously, and removing phosphorus particles. There is no evidence that using copper sulphate to assist visualisation of phosphorus particles for removal is associated with better outcome, and some evidence that systemic absorption of copper sulphate may be harmful. We have so far been unable to identify any other comparisons relevant to informing other aspects of the care of patients with phosphorus burns. Future versions of this review will take account of information in articles published in languages other than English, which may contain additional evidence based on treatment comparisons.

PLAIN LANGUAGE SUMMARY

Interventions for treating phosphorus burns

Phosphorus is a chemical element sometimes used in a military or industrial context. Phosphorus burns resulting from military or industrial injuries are chemical burns that can be fatal. Although rare, these burns are serious, often very deep and painful, and can be associated with lengthy periods of time in hospital for patients.

The usual procedure for dealing with phosphorus burns is to remove any affected clothing and wash the wounds with water or saline solution. In addition, copper sulphate can be used to make the particles of phosphorus more visible and easier to remove, however, copper sulphate is poisonous and can in itself be fatal if absorbed into the body. This review found two retrospective studies (88 patients) that compared burns treated with or without copper sulphate. The review found no evidence that using copper sulphate improves the outcome of the burn, indeed, based upon the limited available evidence, the review authors suggest that copper sulphate should not be used in the treatment of phosphorus burns.

No other studies were identified that could be used to assess other treatments for this type of burn.

BACKGROUND

Description of the condition

Clinical reports of phosphorus burns have been appearing for more than half a century ([Rabinowitch 1943](#)). These burns are sustained by people injured by bombs or other weapons containing phosphorus, or during the manufacture of munitions or fireworks. For example, in a major study of 276 burned patients treated in US military units over 51 years, white phosphorus was the cause in half of the cases ([Barillo 2004](#)). Phosphorus is a component of insecticides and fertilisers, and burns from these sources have also been reported.

White phosphorus is used as a smoke-producing flare and is a waxy, yellow transparent combustible solid ([Al Barqouni 2010](#)). In the presence of oxygen, white phosphorus ignites spontaneously with a yellow flame and dense smoke and remains ignited until either deprived of oxygen or burned out ([Eldad 1991](#)).

Description of the intervention

Phosphorus burns result from industrial and military injuries therefore they are thankfully rarely encountered in usual clinical practice. These chemical burns are sometimes fatal, however, and often associated with significant morbidity and prolonged hospitalisation ([Davis 2002](#)).

White phosphorus burns are extremely painful ([Chou 2001](#)). They affect areas of exposed skin and typically appear as yellowish, necrotic, full-thickness burns resulting from both chemical and thermal effects. The high lipid solubility (the ability of the phosphorus to penetrate the fatty tissues easily) of phosphorus can result in deep extension of the burn injury into the fatty subcutaneous tissues, and this can substantially delay wound healing. Phosphorus can be lethal if inhaled. It may also be absorbed systemically and this can result in multiple organ dysfunction due to its haemolytic effects on erythrocytes and toxic effects on organs, including the kidneys, liver and heart ([Davis 2002](#); [Eldad 1991](#); [Souba 2007](#)).

Most of the available options for treating phosphorus burns depend on the usual practice of clinician rather than the evidence-based practice; these include:

Initial management

A patient's clothes must be removed immediately, as they may ignite or re-ignite. During transportation of the burned person, the burn wound should be covered with a saline or water-soaked dressing until the patient reaches the hospital or other place of treatment ([Al Barqouni 2010](#)). On arrival, continuous irrigation of the burn site with copious amounts of saline or water is used to minimise any complications of phosphorus burns and pain control provided as needed ([Eldad 1991](#); [Kaufman 1988](#)).

Identification and removal of phosphorus particles

Large, easily identifiable particles of phosphorus must be removed. Ultraviolet light or a solution of copper sulphate is sometimes used to facilitate the identification and removal of smaller or embedded particles. However, the use of copper sulphate may also have adverse effects, such as intravascular haemolysis and renal failure ([Davis 2002](#)).

Excision of the necrotic tissue

Excision involves removal of all dead, non-viable tissues.

Systemic support of the burned patient

In critically ill patients with phosphorus burns, practice frequently involves appropriate fluid replacement, with close monitoring of electrolyte concentrations (mainly serum calcium and phosphorus) and electrocardiograms (ECGs) to identify and reduce predictable complications such as hypocalcaemia, hyperphosphataemia and cardiac arrhythmia ([Eldad 1955](#)).

Long-term management

After completion of the initial treatment period, patients are discharged and followed up in an outpatient setting where further wound care (including pain management), physiotherapy, rehabilitation and psychiatric care should be given. In some patients, skin grafting and specific measures to prevent scar formation and contractures is required ([Spanholtz 2009](#)).

Why it is important to do this review

Despite the limited treatment options available in the management process for phosphorus burns, there is no clarity regarding the most effective treatment approaches, either for their immediate or the ongoing management. No systematic review has been undertaken to summarise evidence of the effects of alternative approaches for managing phosphorus burns. The circumstances in which phosphorus burns occur influence the types of studies likely to be possible. Phosphorus burns from causes other than munitions are likely to be isolated single case reports of industrial or other accidents. Although many more patients suffer burns from bombs and other weapons used during wars, these circumstances may not be conducive to carefully controlled comparisons of alternative management strategies. This reality has prompted some controlled experiments to be done in animals and we are currently preparing a completely separate systematic review of these animal studies.

These methodological challenges cannot alter the reality that people have sustained phosphorus burns over a period of more than 60 years, and that it is unfortunately the case that phosphorus burns will continue to occur accidentally and deliberately in the future. The responsibility of this review is therefore to provide information that will assist in the management of future burns, however they occur, using the results of research, while drawing attention to the important questions that remain inadequately addressed.

OBJECTIVES

To summarise the evidence for the effects (beneficial and harmful) of all interventions for treating people with phosphorus burns.

METHODS

Criteria for considering studies for this review

Types of studies

We considered any comparisons of different ways of managing phosphorus burns, including randomised trials where possible.

Types of participants

Studies involving people of any age and gender with phosphorus burns. We considered studies involving people with any percentage of total body surface area burned (%TBSA). The identification of phosphorus as the causative burn agent depended on the history of phosphorus exposure, and we accepted study authors' definitions of phosphorus burns. We have excluded studies involving non-human participants.

Types of interventions

Studies of all types of interventions, either topical or systemic, for the treatment of phosphorus burns. The review distinguished between emergency and post-emergency interventions. Eligible interventions included (but were not limited to) wound irrigation with saline or water, ultraviolet light, copper sulphate, surgical debridement (removal of dead tissues) and systemic support including fluid and electrolyte replacement and antibiotic therapy.

Types of outcome measures

Primary outcomes

1. Death.
2. Time to complete wound healing/proportion of burns completely healed in a specified period of time.

Secondary outcomes

1. Change in wound surface area over time/proportion of wounds partly healed in a specified time period.
2. Complications, for example, intensive care unit admission, wound infection.
3. Pain.
4. Patient satisfaction (cosmetic appearance and/or function).
5. Quality of life.
6. Length of hospital stay.

Search methods for identification of studies

The search methods used for the original version of this review can be found in [Appendix 1](#).

Electronic searches

In October 2013 for the first update of this review we searched the following databases:

- Cochrane Wounds Group Specialised Register (searched 11 October 2013);
- The Cochrane Central Register of Controlled Trials (CENTRAL) (*The Cochrane Library* 2013, Issue 9);
- Ovid OLDMEDLINE (1946 to 1965);
- Ovid MEDLINE (1950 to October Week 1 2013);
- Ovid MEDLINE (In-Process & Other Non-Indexed Citations October 01, 2013);
- Ovid EMBASE (1974 to 2013 Week 40);
- EBSCO CINAHL (1982 to 10 October 2013)
- Conference Proceedings Citation Index - Science (CPCI-S) (1990 to 11 October 2013)

We searched the Cochrane Central Register of Controlled Trials (CENTRAL) using the following search strategy:

- #1 MeSH descriptor Phosphorus explode all trees
- #2 MeSH descriptor Burns, Chemical explode all trees
- #3 (#1 AND #2)
- #4 phosphorus NEAR/5 burn*:ti,ab,kw
- #5 "white phosphorus":ti,ab,kw
- #6 "red phosphorus":ti,ab,kw
- #7 "yellow phosphorus":ti,ab,kw
- #8 (#3 OR #4 OR #5 OR #6 OR #7)

The search strategies used in Ovid MEDLINE, Ovid EMBASE and EBSCO CINAHL can be found in [Appendix 2](#), [Appendix 3](#) and [Appendix 4](#) respectively. We did not apply any methodological filters or restrictions on the basis of study design, language, date of publication or publication status.

Searching other resources

We handsearched the *Journal of the Royal Army Medical Corps* from 1939 to 1948. We also searched the World Health Organization (WHO) International Clinical Trials Registry Platform Search Portal (www.who.int/trialsearch). We checked the reference lists of all included studies to identify reports that had not been found using the methods outlined above.

Data collection and analysis

Selection of studies

We selected studies in two stages. First, based on the title and abstracts of reports, two review authors (LB, KE) independently selected those that were judged potentially relevant. To decide on eligibility we obtained the full texts of articles written in English that potentially matched our inclusion criteria, and which required further scrutiny. Secondly, two review authors (LB, KE) independently assessed the full texts of potentially eligible reports against the pre-determined eligibility criteria. A third review author (NS) arbitrated any disagreement.

Data extraction and management

Two review authors (LB, NS) independently extracted data from the included studies using a pre-determined data extraction sheet. The information extracted included study population (age, gender, setting); information on the location, severity and extent of the burn; the nature of the interventions; analysis; outcomes; and the characteristics of the study (source of funding, country, setting).

Assessment of risk of bias in included studies

We intended that two review authors would independently assess each included study using the Cochrane Collaboration tool for assessing risk of bias ([Higgins 2011a](#)). This tool assesses the risk of bias in six specific domains: sequence generation, allocation concealment, blinding, incomplete outcome data, selective outcome reporting and other issues (likely to be if the timing of outcome assessments are similar). Each criterion was to be judged using: low risk of bias, high risk of bias or at unclear risk of bias (see [Appendix 5](#) for details of criteria on which the judgement was to be based). We intended to complete a 'Risk of bias' table for each eligible study. We planned to discuss any disagreement amongst all review authors to achieve a consensus.

Measures of treatment effect

We planned to report the quantitative data in individual trials for outcomes listed in the inclusion criteria using risk ratios (RR)

with corresponding 95% confidence intervals (CIs) for dichotomous outcomes, mean differences (MD) with 95% CIs for continuous outcomes, and hazard ratios (HR) and 95% CI for time to event outcomes (e.g. time to healing).

Unit of analysis issues

The unit of randomisation is the individual patient. If RCTs with a cluster-randomised design had been identified, to avoid unit of analysis errors in cluster-randomised trials we planned to re-analyse these studies by calculating the effective sample sizes where possible, according to the methods outlined in [Higgins 2011b](#) and, if necessary, we would have incorporated an estimate of the intra-cluster coefficient (ICC) using external estimates obtained from similar studies.

Dealing with missing data

Where data were missing from published reports, we intended to contact authors for further information. If these data were not available we would have made explicit the assumptions of any methods used to cope with missing data, performed sensitivity analyses to assess how sensitive results are to reasonable changes in the assumptions that are made, and addressed the potential impact of missing data on the findings of the review in the Discussion.

Assessment of heterogeneity

We planned to assess the studies for clinical heterogeneity by checking the inclusion criteria, exclusion criteria, differences in the intervention, differences in the control and differences in the definition used for outcomes. We planned to assess heterogeneity in the effect estimates by visually examining the forest plots, by the Cochrane Q statistic and the I^2 statistic values. The I^2 statistic examines the percentage of total variation across studies due to heterogeneity rather than due to chance. Values of I^2 over 75% indicate a high level of heterogeneity ([Higgins 2003](#)). Statistically significant values would have been interpreted in accordance with recommendations outlined by [Deeks 2009](#).

Assessment of reporting biases

We planned to assess publication bias by preparing a funnel plot and examining it either visually or quantitatively by the rank correlation test and/or the graphical test with or without heterogeneity.

Data synthesis

We planned to pool the results of clinically homogenous groups of studies using the fixed-effect model for meta-analysis. If there had been evidence of some clinical and statistical heterogeneity (I^2 over 50%), we would have used a random-effects model. We would not have pooled studies if statistical heterogeneity (I^2 over 75%) was high. We presented results in a narrative format by type of intervention.

Subgroup analysis and investigation of heterogeneity

We planned to consider subgroup analyses according to the setting (i.e. evaluations of those with burns because of being in a conflict zone versus those exposed to phosphorus as an occupational hazard).

Sensitivity analysis

We planned to conduct a sensitivity analysis to investigate how conclusions might be affected if studies at high risk of bias (trials that did not report adequate allocation concealment or do not have blinded outcome assessment) are excluded from the analyses.

RESULTS

Description of studies

See: [Characteristics of included studies](#); [Characteristics of excluded studies](#).

Results of the search

The search for the update produced 58 citations for screening. As a result a total 431 reports were screened for this review and we identified 19 of these as potentially relevant. Two authors (LB and KE) obtained the full texts of these 19 records for detailed examination. We identified no further studies by searching the reference lists of selected studies or by handsearching. Of these 19 reports, we excluded 16 (see [Excluded studies](#) below) and one paper is awaiting further assessment after translation. This left only two reports in English in which different management strategies were compared and which were included in the review.

Included studies

We identified no RCTs or quasi-randomised RCTs (the eligible designs pre-specified in our protocol). At this point we made a collective decision to deviate from our protocol to include any comparative studies in an effort to find any evidence, however weak. The two included studies were both retrospectively assembled case series reporting comparisons of different ways of managing phosphorus burns. The first involved 77 of 96 patients treated at the US Army Institute of Surgical Research between 1950 and 1986 ([Curreri 1970](#)), and the second 11 patients treated at the US Army Surgical Research Unit in 1965 and 1966 ([Summerlin 1967](#)). In both these reports, comparisons were made between patients in whom copper sulphate had been used and patients in whom it had not been used before, during and after debridement.

Excluded studies

Fourteen studies did not meet the inclusion criteria for the review and we excluded them for various reasons, the most common being that no treatment comparison had been reported. In summary there were three retrospective surveys ([Barillo 2004](#); [Chou 2001](#); [Mozingo 1988](#)) and eleven case reports with no comparison group ([Al Barqouni 2010](#); [Broekhuizen 1982](#); [Carras 1993](#); [Conner 2007](#); [Davis 2002](#); [Fang 1987](#); [Frank 2008](#); [Karunadasa 2010](#); [Konjoyan 1983](#); [Loveall 2007](#); [Saracoglu 2013](#); [Song 1985](#); [Weinberger 1978](#)) see [Table 1](#) and [Table 2](#) for additional information.

Risk of bias in included studies

Both included studies were at high risk of bias. Firstly both were at high risk of selection bias since the comparison groups were not prospectively assembled at random; neither of the included reports described how patients had been selected for treatment with copper sulphate. [Curreri 1970](#) stratified analyses by the %TBSA burned, but not in terms of other factors, such as co-morbidities, or type of treatment. Furthermore there is attrition bias: in [Curreri](#)

1970, only 77 out of 96 patients were included in the study report; the remaining 19 patients were not accounted for.

Effects of interventions

In the report of Curreri 1970, copper sulphate was used in 40 patients with phosphorus burns and not used in 37 patients. In the report by Summerlin 1967, copper sulphate was used in eight patients and not used in three patients.

Outcomes

Neither study reported the primary outcome of wound healing.

Curreri 1970 stratified their analysis by the percentage of total body surface area (%TBSA) burned. The average duration of hospitalisation among patients with 0% to 20% TBSA burned was 101 days after treatment with copper sulphate and 97 days if copper sulphate had not been used. Average duration of hospitalisation among patients with 2% to 40% TBSA burned was 135 days after use of copper sulphate and 112 days among patients in whom copper sulphate had not been used. Neither of these differences between the two groups were statistically significant. No differences were observed in the time to eschar (a dry dark scab) separation or in residual contractures. The primary outcome of death was reported only in relation to the total number of patients who had burns resulting from chemical agents, this includes phosphorus, 5.6% (6/111). It was not reported how many of these deaths were in patients with phosphorus burns.

In Summerlin 1967, haematuria, haemoglobinuria, mild to severe hypocalcaemia, high level of copper in urine, oliguria and finally renal failure 20 to 72 hours after injury developed in three of the eight patients treated with copper sulphate, and one of them required haemodialysis. It was stated that all patients recovered, therefore there were no deaths reported. None of the three patients who did not receive copper sulphate experienced any of these complications.

DISCUSSION

It is a challenge to review evidence relevant to the management of phosphorus burns, as most of the cases occur in acute conflict circumstances, in which doing carefully designed research will be extremely difficult or impossible. We did not expect to find randomised trials. We found only reports in which different management strategies had been compared after assembling case series retrospectively. These provide little assurance that selection and measurement biases had been adequately controlled. It is possible that more and better evidence is contained in the reports awaiting translation from languages other than English.

However, the restricted evidence we have identified so far provides no support for the use of copper sulphate in the management of phosphorus burns, and some evidence of the serious complications that may follow its use. On the basis of possible harmful adverse events of copper sulphate, we suggest the use of alternative approaches, such as ultraviolet light (Wood's lamp), which are likely to be a safer option for identifying and removing small phosphorus particles.

We will update this review when reports published in languages other than English have been translated and assessed. We have also embarked on an analysis of potentially relevant animal research.

AUTHORS' CONCLUSIONS

Implications for practice

The treatment of people with acute phosphorus burns is based on clinical experience, custom and practice rather than research evidence (which is lacking). Removal of patients' clothes, continuous irrigation of their wounds with cold solutions and removal of phosphorus particles are the most important elements of the management of phosphorus burns. Beyond these common sense first aid measures, the research we have reviewed provides little guidance on subsequent management. However, the evidence we had analysed so far suggests that copper sulphate should not be used for visualisation of phosphorus particles as what evidence there is suggests a possible association with adverse outcomes. Ultraviolet light can be used to assist the visualisation of phosphorus particles during the process of debridement as a safer alternative.

Implications for research

The conduct of high-quality randomised controlled trials to address the uncertainties around the management of people with phosphorus burns, is highly desirable, but would be extremely difficult given the context in which such burns occur.

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CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Curreri 1970

Methods	Retrospectively assembled case series based on the clinical records of all 111 patients admitted to the US Army Institute of Surgical Research between 1950 and 1986
Participants	77 out of a total of 96 patients with phosphorus burns
Interventions	Without copper sulphate: 37 patients: wounds irrigated with copious amounts of water or saline, debridement with removal of phosphorus particles at least once, and application of wet dressing with thick gauze pads, and irrigation with water or dilute solution of sodium bicarbonate With copper sulphate: 40 patients treated as above, plus application of copper sulphate before, during and after debridement
Outcomes	Average length of hospitalisation, the time required for eschar separation and incidence of residual contractures
Notes	USA

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Not randomised
Allocation concealment (selection bias)	High risk	No allocation concealment

Summerlin 1967

Methods	Retrospective survey of the clinical records of patients admitted to the US Army Surgical Research Unit between 1965 and 1966
Participants	11 patients with phosphorus burns
Interventions	Without copper sulphate: 3 patients with phosphorus burns treated without copper sulphate With copper sulphate: 8 patients with phosphorus burns treated with copper sulphate
Outcomes	Without copper sulphate: no complications had been experienced With copper sulphate: haematuria, haemoglobinuria and mild to severe hypocalcaemia, high level of copper in urine, oliguria and finally renal failure 20 to 72 hours after injury developed in 3 patients out of 8. One required haemodialysis. Eventually all recovered.
Notes	USA

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Not randomised
Allocation concealment (selection bias)	High risk	No allocation concealment

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Al Barqouni 2010	Methodologically inappropriate, case report (Table 2)
Barillo 2004	Methodologically inappropriate, non-comparative retrospective study (Table 1)
Broekhuizen 1982	Methodologically inappropriate, case report (Table 2)
Carras 1993	Methodologically inappropriate, case report (Table 2)
Chou 2001	Methodologically inappropriate, non-comparative retrospective study (Table 1)
Conner 2007	Methodologically inappropriate, case report (Table 2)
Davis 2002	Methodologically inappropriate, case report (Table 2)
Fang 1987	Methodologically inappropriate, case series, and deal with red phosphorus rather than white.
Frank 2008	Methodologically inappropriate, case report (Table 2)
Karunadasa 2010	Methodologically inappropriate, case report (Table 2)
Konjoyan 1983	Methodologically inappropriate, case report (Table 2)
Loveall 2007	Methodologically inappropriate, case report (Table 2)

Interventions for treating phosphorus burns (Review)

Study	Reason for exclusion
Mozingo 1988	Methodologically inappropriate, non-comparative retrospective study (Table 1)
Saracoglu 2013	Methodologically inappropriate, case report (Table 2)
Song 1985	Methodologically inappropriate, case report (Table 2)
Weinberger 1978	Methodologically inappropriate, case report (Table 2)

Characteristics of studies awaiting assessment [ordered by study ID]

Bonelli 1971

Methods	
Participants	
Interventions	
Outcomes	
Notes	Awaiting translation - study language: Italian

ADDITIONAL TABLES

Table 1. Retrospective surveys with no treatment comparison

Study	Type	Participants	Interventions	Outcome	Other
Chou 2001 Taiwan	Retrospective survey of hospital clinical records between 1984 and 1998	7 out of 326 chemical burn patients resulted from white phosphorus, and 2 of these were presented	Irrigation and dressing Copious normal saline irrigation dressing with saline-soaked pads Identification and removal of phosphorus particles Identification of phosphorus particles Use of copper sulphate or ultraviolet light 1% copper sulphate solution for neutralisation Systemic support Monitoring for electrolytes and cardiac function Long-term management No details provided Others - none	1 of the 7 died from inhalation injury Of the 7 patients, 5 required skin grafting when wounds had not healed within 14 days of presentation. Hypocalcaemia was commonly encountered.	Mentioned the use of cold solution in irrigation

Table 1. Retrospective surveys with no treatment comparison (Continued)

Mozingo 1988 USA	Retrospec- tive survey of clinical records in US Army In- stitute of Surgical Re- search be- tween 1969 and 1985, mostly in Vietnam	49 out of 87 chem- ically burned patients were burned with white phospho- rus	<p>Irrigation and dressing</p> <p>Removal of clothes and copious water irrigation done as soon as possible</p> <p>Identification and removal of phosphorus particles</p> <p>Removal of particles from the skin surface</p> <p>Use of copper sulphate or ultraviolet light</p> <p>Copper sulphate solution (1% or less) or Woods lamp (ultraviolet light) used</p> <p>Systemic support</p> <p>No details provided</p> <p>Long-term management</p> <p>No details provided</p> <p>Others</p> <p>The report does not allow disaggregation of the management of the 49 patients with white phosphorus burns</p>	Cases of white phosphorus burns remained in hospital longer	Noticed the importance of not liquifying the phosphorus particles
Barillo 2004 USA	Retrospec- tive survey of clinical records in US Army In- stitute of Surgical Re- search be- tween 1950 and 2000	146 out of 276 chemical- ly burned patients were burned with white phospho- rus	<p>Irrigation and dressing</p> <p>Wounds thoroughly irrigated and then covered with saline or saline-soaked pads</p> <p>Identification and removal of phosphorus particles</p> <p>Visible particles of white phosphorus removed and placed in cold water to prevent re-ignition</p> <p>Immediate surgical debridement necessary and repeated until all phosphorus particles have been removed.</p> <p>Debrided wounds examined at least twice daily for new particles or smoking areas, which would indicate the need for re-operation</p> <p>Use of copper sulphate or ultraviolet light</p> <p>No details provided</p> <p>Systemic support</p> <p>No details provided</p> <p>Long-term management</p> <p>No details provided</p> <p>Others - none</p>	Mortality increased from 5.4% between 1950 and 1968 to 13.8% between 1969 and 1985. Mortality from 1986 to 2000 was 0%.	Hospital length of stay decreased from a mean of 90 days in the first 19 years of the study to a mean of 15 days in the most recent 15-year period. The chemical responsible for injury was white phosphorus in 146 cases.

Table 2. Case reports, with no treatment comparisons

Study	Type	Participants	Interventions	Outcome	Other
Frank 2008 Germany	4 cases	German civilians picking up phosphorus (sometimes mistaken for amber) from North German sea beach	Irrigation and dressing Submerging affected skin in cold water. Saline used for irrigation. Identification and removal of phosphorus particles Phosphorus particles removed, debridement of necrotic tissue (sometimes under general anaesthesia) Use of copper sulphate or ultraviolet light No details provided Systemic support No details provided Long-term management Defects in skin covered by allogenic grafts in one case. Vacuum-assisted closure in one case. Others - none	All survived. First case discharged home after 4 weeks; 2nd after 12 days; 3rd after 20 days; 4th after 11 days	Refers to the use of white phosphorus in World War I, and to the military jargon 'willie pete' Emphasised the use of cold rather than warm water for irrigation
Karunadasa 2010 Sri Lanka	2 cases	2 soldiers injured by an exploding rocket-propelled grenade	Irrigation and dressing Clothes were not removed, with resulting deep burns. Saline used for irrigation. Identification and removal of phosphorus particles Serial excisions and debridement Use of copper sulphate or ultraviolet light No details provided Systemic support Intravenous antibiotics and blood transfusion Long-term management Grafting Others - none	Both survived. Referred to aggressive rehabilitation after 13 weeks. No contracture observed at 12 months.	
Davis 2002 USA	1 case	50-year old male worker at munitions manufactory suffered 36% total body surface area burns	Irrigation and dressing Clothes were removed. Water used for irrigation. Identification and removal of phosphorus particles Debridement and removing of phosphorus particles by metal forceps Use of copper sulphate or ultraviolet light No details provided Systemic support		

Table 2. Case reports, with no treatment comparisons (Continued)

Table 2: Case reports, with no treatment comparisons (continued)					
No details provided					
Long-term management					
2 skin grafts were placed					
Others - none					
Konjoyan 1983	1 case	21 year old male soldier injured after explosion of a defective mortar in an armed personnel carrier. Another soldier was killed by the blast.	Irrigation and dressing Wounds irrigated and dressed with silver sulphadiazine	Survived. After 23 days, transferred to another government health facility to be closer to his home.	
USA			Identification and removal of phosphorus particles Debridement of wounds on the left knee, with removal of all visible phosphorus particles		
			Use of copper sulphate or ultraviolet light No details provided		
			Systemic support Resuscitation with intravenous fluids, analgesics, oxygen and blood transfusion		
			Long-term management Closure of wounds that were not healed.		
			Others - none		
Song 1985	1 case	41-year old male patient 7% of surface area, 2nd degree burn on lower extremities	Irrigation and dressing Wound rinsed and cleaned with water, covered with wet gauze pad.	The patient gets worse at night, developing nausea and vomiting, haematuria, jaundice and hepatomegaly.	Conclusion to use the silver nitrate solution instead, as it can make phosphorus particles non-flammable for 6 months or more, and also use of wet compress of 3% to 5% sodium bicarbonate
China			Identification and removal of phosphorus particles No details provided		
			Use of copper sulphate or ultraviolet light 0.4% copper sulphate applied immediately and with wet gauze pads	On the 11th day the patient died of acute renal failure as result of copper ion absorption.	
			Systemic support No details provided		
			Long-term management No details provided		
			Others - none		
Conner 2007	1 case	A 19-year-old man presented burns after exposure to an incendiary agent	Irrigation and dressing Burns irrigated with copious saline		
USA			Identification and removal of phosphorus particles Remaining phosphorus particles removed. Surgical debridement if still present.		
			Use of copper sulphate or ultraviolet light		

Table 2. Case reports, with no treatment comparisons (Continued)

			Examination under ultraviolet light if required	
			Systemic support	
			No details provided	
			Long-term management	
			No details provided	
			Others - none	
Loveall 2007	1 case	Patient with hand actively burning	Irrigation and dressing	
USA			Hand rinsed in warm tap water for an hour, then wrapped with gauze, and soaked in sterile water. Hand later submerged in vegetable oil.	
			Identification and removal of phosphorus particles	
			Surgical excision	
			Use of copper sulphate or ultraviolet light	
			Examination under ultraviolet light	
			Systemic support	
			Inpatient intensive care unit management	
			Long-term management	
			No details provided	
			Others - none	
Al Bar- qouni 2010	1 case	18-year old male patient with 30% surface area burned	Irrigation and dressing	8 days af- ter admis- sion, the pa- tient was dis- charged rel- atively well, and after 16 months of follow-up no permanent complications
Palestine			Irrigation with dilute sodium bicarbonate, wet dress- ing applied	
			Identification and removal of phosphorus particles	
			Debridement and removal of phosphorus particles	
			Use of copper sulphate or ultraviolet light	
			No details provided	
			Systemic support	
			Monitoring of electrolytes and cardiac function	
			Long-term management	
			No details provided	
			Others - none	
Carras 1993	1 case	33-year old male with 18% surface area burned, 10% of which were deep burns	Irrigation and dressing	
France			Irrigation with 1.4% sodium bicarbonate	

Table 2. Case reports, with no treatment comparisons (Continued)

			Identification and removal of phosphorus particles	
			Debridement and removal of phosphorus particles	
			Use of copper sulphate or ultraviolet light	
			0.5% copper sulphate	
			Systemic support	
			Inpatient intensive care unit management	
			Long-term management	
			Transferred to rehabilitation unit after 30 days	
			Others - none	
Saracoglu 2013	1 case	6-year old boy accidentally injured by a fire-work, resulting in a full-thickness, 1.5 cm in depth burns in his neck.	Irrigation and dressing Irrigation with sterile distilled water, wet dressing applied 2-3 Identification and removal of phosphorus particles Debridement and removal of necrotic skin and phosphorus particles daily Use of copper sulphate or ultraviolet light ultraviolet light Systemic support Antibiotic therapy, fluid resuscitation with dexamethasone, intermittent bronchodilator and oxygen were administered. monitoring of blood pressure, electrocardiography, peripheral oxygen saturation and temperature Long-term management No details provided Others - high protein diet	The patient gets worse during the first 3 days, developing inspiratory stridor and tachypnoea. Transferred to burn and wound care center. after 9 days survived without any systemic complication, except total left facial nerve paralysis, so that transferred for facial and reconstructive surgery.
Weinberger 1978	3 cases	A case series describing the treatment of three patients with phosphorous burns from a single incident of a phosphorous grenade explosion.	No specific intervention was described, the paper dealt with the overall management of the three patients.	Translated paper
Broekhuizen 1982	1 case	A case report on a munitions worker wounded in the face by	The patient was immediately treated with water, copper sulphate bandages and surgery.	The paper also reported on a BEN HUR

Table 2. Case reports, with no treatment comparisons *(Continued)*

an exploding yellow phosphorus
high explosive
grenade.

solution, designed by Israeli doctors, which is copper sulphate (1%) with Hydroxyethyl cellulose (2%) and sodium bicarbonate (5%). This solution should prevent the uptake of copper sulphate in the bloodstream, and hence lessen the effect of copper sulphate on kidney and liver damage. The authors advocate quick surgical intervention to remove all phosphorous materials from the body.

Translated paper

APPENDICES

Appendix 1. Search strategy - for the original version

For this third update we searched the following electronic databases:

- Cochrane Wounds Group Specialised Register (searched 30 September 2011);
- The Cochrane Central Register of Controlled Trials (CENTRAL) (The Cochrane Library 2011, Issue 3);
- Ovid MEDLINE (1947 to September Week 3 2011);
- Ovid MEDLINE (In-Process & Other Non-Indexed Citations September 29, 2011);
- Ovid EMBASE (1980 to 2011 Week 38);
- EBSCO CINAHL (1982 to 23 September 2011)
- Conference Proceedings Citation Index - Science (CPCI-S) (1990 to 30 September 2011)

Interventions for treating phosphorus burns (Review)

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Appendix 2. Ovid MEDLINE search strategy

1 exp Phosphorus/
2 exp Burns, Chemical/
3 1 and 2
4 (phosphorus adj5 burn*).tw.
5 white phosphorus.tw.
6 red phosphorus.tw.
7 yellow phosphorus.tw.
8 3 or 4 or 5 or 6 or 7

Appendix 3. Ovid EMBASE search strategy

1 exp phosphorus/
2 exp chemical burn/
3 1 and 2
4 (phosphorus adj burn*).tw.
5 white phosphorus.tw.
6 red phosphorus.tw.
7 yellow phosphorus.tw.
8 3 or 4 or 5 or 6 or 7

Appendix 4. EBSCO CINAHL search strategy

S8 S3 or S4 or S5 or S6 or S7
S7 T1 yellow phosphorus or AB yellow phosphorus
S6 T1 red phosphorus or AB red phosphorus
S5 T1 white phosphorus or AB white phosphorus
S4 T1 phosphorus N5 burn* or AB phosphorus N5 burn*
S3 S1 and S2
S2 (MH "Burns, Chemical")
S1 (MH "Phosphorus")

Appendix 5. 'Risk of bias' assessment criteria for RCTs

1. Was the allocation sequence randomly generated?

Low risk of bias

The investigators describe a random component in the sequence generation process such as: referring to a random number table; using a computer random number generator; coin tossing; shuffling cards or envelopes; throwing dice; drawing of lots.

High risk of bias

The investigators describe a non-random component in the sequence generation process. Usually, the description would involve some systematic, non-random approach, for example: sequence generated by odd or even date of birth; sequence generated by some rule based on date (or day) of admission; sequence generated by some rule based on hospital or clinic record number.

Unclear

Insufficient information about the sequence generation process to permit judgement of low or high risk of bias.

2. Was the treatment allocation adequately concealed?

Low risk of bias

Participants and investigators enrolling participants could not foresee assignment because one of the following, or an equivalent method, was used to conceal allocation: central allocation (including telephone, web-based and pharmacy-controlled randomisation); sequentially numbered drug containers of identical appearance; sequentially numbered, opaque, sealed envelopes.

High risk of bias

Participants or investigators enrolling participants could possibly foresee assignments and thus introduce selection bias, such as allocation based on: using an open random allocation schedule (e.g. a list of random numbers); assignment envelopes were used without appropriate safeguards (e.g. if envelopes were unsealed or nonopaque or not sequentially numbered); alternation or rotation; date of birth; case record number; any other explicitly unconcealed procedure.

Unclear

Insufficient information to permit judgement of low or high risk of bias. This is usually the case if the method of concealment is not described or not described in sufficient detail to allow a definite judgement, for example if the use of assignment envelopes is described, but it remains unclear whether envelopes were sequentially numbered, opaque and sealed.

3. Blinding - was knowledge of the allocated interventions adequately prevented during the study?**Low risk of bias**

Any one of the following.

- No blinding, but the review authors judge that the outcome and the outcome measurement are not likely to be influenced by lack of blinding.
- Blinding of participants and key study personnel ensured, and unlikely that the blinding could have been broken.
- Either participants or some key study personnel were not blinded, but outcome assessment was blinded and the non-blinding of others unlikely to introduce bias.

High risk of bias

Any one of the following.

- No blinding or incomplete blinding, and the outcome or outcome measurement is likely to be influenced by lack of blinding.
- Blinding of key study participants and personnel attempted, but likely that the blinding could have been broken.
- Either participants or some key study personnel were not blinded, and the non-blinding of others likely to introduce bias.

Unclear

Any one of the following.

- Insufficient information to permit judgement of low or high risk of bias.
- The study did not address this outcome.

4. Were incomplete outcome data adequately addressed?**Low risk of bias**

Any one of the following.

- No missing outcome data.
- Reasons for missing outcome data unlikely to be related to true outcome (for survival data, censoring unlikely to be introducing bias).
- Missing outcome data balanced in numbers across intervention groups, with similar reasons for missing data across groups.
- For dichotomous outcome data, the proportion of missing outcomes compared with observed event risk not enough to have a clinically relevant impact on the intervention effect estimate.
- For continuous outcome data, plausible effect size (difference in means or standardised difference in means) among missing outcomes not enough to have a clinically relevant impact on observed effect size.
- Missing data have been imputed using appropriate methods.

High risk of bias

Any one of the following.

- Reason for missing outcome data likely to be related to true outcome, with either imbalance in numbers or reasons for missing data across intervention groups.
- For dichotomous outcome data, the proportion of missing outcomes compared with observed event risk enough to induce clinically relevant bias in intervention effect estimate.
- For continuous outcome data, plausible effect size (difference in means or standardised difference in means) among missing outcomes enough to induce clinically relevant bias in observed effect size.
- 'As-treated' analysis done with substantial departure of the intervention received from that assigned at randomisation.
- Potentially inappropriate application of simple imputation.

Unclear

Any one of the following.

- Insufficient reporting of attrition/exclusions to permit judgement of low or high risk of bias (e.g. number randomised not stated, no reasons for missing data provided).
- The study did not address this outcome.

5. Are reports of the study free of suggestion of selective outcome reporting?

Low risk of bias

Any 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.
- 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 of bias

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. subscales) 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

Insufficient information to permit judgement of low or high risk of bias. It is likely that the majority of studies will fall into this category.

6. Other sources of potential bias

Low risk of bias

The study appears to be free of other sources of bias.

High risk of bias

There is at least one important risk of bias. For example, the study:

- had a potential source of bias related to the specific study design used; or
- had extreme baseline imbalance; or
- has been claimed to have been fraudulent; or
- had some other problem.

Unclear

There may be a risk of bias, but there is either:

- insufficient information to assess whether an important risk of bias exists; or
- insufficient rationale or evidence that an identified problem will introduce bias.

WHAT'S NEW

Date	Event	Description
29 May 2014	New search has been performed	First update, new search, three additional studies have been added to the Excluded studies table in the update.
29 May 2014	New citation required but conclusions have not changed	No new studies included, conclusions remain unchanged.

CONTRIBUTIONS OF AUTHORS

Loai Barqouni conceived, designed and co-ordinated the review, extracted and checked quality of extracted data, undertook quality assessment, analysed and interpreted data, performed and checked statistical analysis, completed first draft of the review, performed part of writing or editing the review, made an intellectual contribution, approved final review prior to submission, advised on the review, secured funding, performed previous work that was the foundation of the current review, wrote to study authors/experts/companies, and is guarantor of the review.

Khamis Elessi conceived the review, extracted data, checked quality of data extraction, undertook quality assessment, analysed data, checked quality assessment, performed part of data analysis or interpretation, performed and checked quality of statistical analysis, advised on the review, and secured funding.

Nafiz Abu Shaaban conceived the review, analysed or interpreted data, checked quality assessment, performed part of data analysis or interpretation, advised on the review, secured funding and performed previous work which was the foundation for the current review.

Contributions of editorial base:

Nicky Cullum: edited the review, advised on methodology, interpretation and review content. Approved the final review prior to submission.

Sally Bell-Syer: co-ordinated the editorial process. Advised on methodology, interpretation and content. Edited and copy edited the review and the updated review.

Ruth Foxlee: designed the search strategy, ran the searches and edited the search methods section.

DECLARATIONS OF INTEREST

No conflict of interest.

SOURCES OF SUPPORT

Internal sources

- The Department of Health Sciences, University of York, UK.
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External sources

- The Al-Quds Foundation for Medical Schools in Palestine, UK.

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

The eligibility criteria for type of studies were relaxed to include any comparative studies given the absence of any prospective, controlled trials. The absence of randomised or quasi-randomised evidence can be attributed to the circumstances in which phosphorus burns occur which make the possibility of conducting a randomised controlled trial (RCT) unlikely. However, despite the rarity of such RCTs, phosphorus burns will continue to occur and such a systematic review will provide the practitioner with the best available evidence in treating phosphorus burns.

INDEX TERMS

Medical Subject Headings (MeSH)

*Phosphorus; Antidotes [adverse effects] [*therapeutic use]; Burns, Chemical [*drug therapy]; Copper Sulfate [adverse effects] [*therapeutic use]; Retrospective Studies

MeSH check words

Humans