Antiseptic Agents for Chronic Wounds: A Systematic Review

Koko Barrigah-Benissan 1,*, Jérôme Ory 1,*, Albert Sotto 2, Florian Salipante 3, Jean-Philippe Lavigne 1,* and Paul Loubet 2

Abstract: In many parts of the world, antiseptic agents remain non-indicated in chronic wound care. In the current context of bacterial resistance to antibiotics and the development of new-generation antiseptic agents, wound antisepsis represents an asset for the prevention of wound infection. We aimed to evaluate four common antiseptic agents in chronic wound care complete healing. The review protocol was based on the Cochrane Handbook for Systematic Reviews of Intervention and devised in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analyses (PRISMA) statement guidelines. Five databases and three clinical trials registries were searched from inception to 30 June 2021 without language restrictions. We included randomised trials evaluating the efficacy of antiseptic agents in chronic wound care in adults. Interventions considered were those using antiseptics for cleansing or within a dressing. Risk of bias was assessed using the bias excel tool provided by the Bristol Academy. Evidence quality was assessed using Grading of Recommendation Assessment, Development and Evaluation (GRADE) criteria. Of 838 studies, 6 were finally included, with a total of 725 patients. The included studies assessed iodine (cadexomer or povidone iodine) (n = 3), polyhexamidine (n = 2), and octenidine (n = 1). Limited evidence suggested a better wound healing completion with iodine compared to saline (two randomised controlled trials (RCT), 195 patients, pooled RR 1.85 (95%CI (1.27 to 2.69)), moderate-quality evidence). There was not enough evidence to suggest a difference in wound healing using octenidine or polyhexamidine. None of the antiseptic agents influenced adverse event occurrence compared to saline.

Keywords: antiseptic agents; efficiency; iodine; systematic review; wound healing; wound infection

1. Introduction
Chronic wounds are wounds failing to proceed through the normal phases of healing in an orderly and timely manner. The definition of time without complete or partial healing differs across countries, ranging from 4 weeks to 3 months [1]. The Wound Healing Society defines four types of chronic wounds: diabetic foot ulcers (DFU), vascular ulcers (containing venous and arterial ulcers), and pressure ulcers (PU) [2].

Infection is a common complication of chronic wounds. Historically, the research on wound infection control and improvement was focused on reducing the “pathogen burden”. However, quantitative consideration of microbial load is insufficient for assessing wound improvement or wound risk of infection [3,4]. Microorganisms in a chronic wound are co-aggregated together within a protective extracellular matrix, constituting a biofilm. This biofilm conformation induces a dramatically increased tolerance to host immune defences and a greater resistance to antimicrobials [5]. Biofilm delays wound healing by...
inducing an ineffective host inflammatory response and damaging host tissues [6]. For this reason, a management of the biofilm is more relevant for the treatment of chronic wound [7]. In 2011, Dissemond et al. classified wounds into four categories, depending on wound bed clinical and microbiological situation: (1) contaminated or colonised wounds without risk of infection; (2) colonised wounds at risk (WAR) or critically colonised wounds; (3) wounds with local infection; and (4) systemic infections and infected wounds. The authors suggested the use of antiseptic agents for wounds from the second category alongside other treatments [8].

Chronic wound care commences with wound bed preparation via (i) wound cleansing to create a wet or moist environment, favourable to healing. (ii) Wound debridement via removal of devitalised, contaminated tissue from within or adjacent to a wound, until surrounding healthy tissue is exposed [9,10]. Debridement can be mechanical (sharp debridement, surgical), enzymatical, or bio-surgical (e.g., maggot therapy) [11–15]. Negative pressure wound therapy has also been used for bacterial decontamination and wound bed preparation [16]. (iii) Application of a suitable dressing, according to the type of wound. (iv) Antibiotic treatment, exclusively for infected wounds. Other therapies are beneficial to specific wounds: compression therapy is required for venous leg ulcers (VLU) [17]; arterial revascularisation, offloading foot ulcers, and diabetes control are essential in DFU [18,19]; and skin assessment and care, offloading and pressure redistribution, dressings [19,20], and structured educational program are useful for all types of chronic wounds.

In some countries, tap water or saline remain the only recommended agents for wound cleansing. Antisepsis is a common, yet controversial, wound cleansing method. Some studies consider debridement alone insufficient to reduce the biofilm that delays wound healing and suggest antiseptics to delay biofilm reformation and reduce the risk of infection [7,20]. Antiseptic agents may complement the debridement process and control infection.

The primary mode of action of antiseptics can be pharmacological, metabolic, and/or immunological [21]. For the purpose of this review, antiseptic agents are defined as medication that can prevent the growth or destroy microorganisms in or on a living tissue. Following this definition of medication, antiseptic agents must pass through a drug authorisation procedure with a medicine agency [22]. The main antiseptic agents used in chronic wound care are halogenated compounds, alcohol-based agents, biguanides (e.g., polyhexanide also called polyhexamethylenebiguanide or PHMB, chlorhexidine), and quaternary ammoniums (e.g., octenidine). Halogenated compounds include subfamilies such as the iodine/iodophor agents (e.g., povidone iodine, cadexomer iodine) and chlorous agents (hypochlorite, hypochlorous acid) [21,22]. Alternative therapeutics (e.g., honey, silver), while being antimicrobial agents are not antiseptic agents as they did not go through an authorisation procedure for this purpose [23]. They therefore are not part of the antiseptic agents’ classification. International guidelines recommend against the routine use of topical antiseptics to manage infected chronic wounds [24–27].

The emergence and diffusion of multidrug resistant bacteria and a better formulation of antiseptics with less side effects has renewed consideration of antisepsis. No resistance or adaptation has yet been observed in antiseptic agents with unspecific effects such as iodine agents, polyhexanide, octenidine, or oxidising agents (e.g., hypochlorous acid) [21]. In countries where antiseptic agents are a part of chronic wound care management protocol, there is no consensus on the best antiseptic agents for chronic wound care. Our study aimed to assess the evidence of four common antiseptic agents on chronic wound healing.

2. Results
2.1. Characteristics of the Included Studies

We identified 838 studies including 61 registered clinical trials through database searching and 31 through manual searching. Of these studies, 149 duplicates were removed, and a further 613 studies were excluded by title or abstract. Three studies could not be retrieved and they were withdrawn from the database. We fully screened 73 studies, leaving 6 studies eligible for inclusion (Figure 1) [28–33]. A total of 17 studies were excluded after
unpublished studies were included. We fully screened 73 studies, leaving 6 studies eligible for inclusion (Figure 1).

The included studies are listed in Table S1. No studies assessing chlorous agents met our eligibility criteria. Saline was the comparator in all studies. None of the studies compared two or more antiseptic agents. Three studies evaluated the efficacy of iodine (cadexomer iodine or polyvidone iodine) [28,31,32], two studies evaluated polyhexanide [30,33], and one last study evaluated octenidine [29]. Five studies had two comparative groups of patients, and one had three comparative groups [32]. All studies were multicentre, taking place in 55 centres, both inpatient and outpatient settings across South Korea, India, France, Hungary, the UK, the USA, Germany, Canada, and Italy. Different types of chronic wounds were studied: one including DFU only [28]; one including VLU only [29]; two including PU only [30,31]; one including DFU, VLU, and PU [32]; and the last including chronic wounds without precision of the type of the wound [33]. The minimum duration used to define chronic wounds ranged from 4 weeks [28,29] to 3 months [31], although others did not state the definition, but gave the mean duration of the wounds [32], or merely described wounds as “chronic” [33], or only gave the duration of the comorbidity [30]. Study duration ranged from 4 weeks [30,33] to 24 weeks [31].

The seven domains of risk of bias were assessed following the Cochrane recommendation of 2011 [34] and are presented in Table S2. The most common source of bias was outcome assessment blinding (see Table S2). The Risk of Bias tool (2019) released by the

![Figure 1. Study flow diagram.](image-url)
Cochrane collaboration was also used for the risk of bias assessment and presented in Table S3. The overall risk of bias among included studies was rated with some concerns for five studies [28–30,32,33] and high for one [31] (Table S3). A plot of the percentage of risk of bias assessments per domain is presented in Figure S1, as recommended by the Cochrane collaboration tool for risk of bias assessment in randomised trials (2019 version).

Three studies reported the primary outcome (wound healing) as the proportion of patients with complete wound healing at 4 and 8 weeks [29], at 12 weeks [32], and as healing time from wound size [29]. All studies assessed wound healing rate as wound size reduction by planimetry measurement. Three studies assessed pain as a secondary outcome with various pain scales (e.g., verbal rating scales, visual analogue scales) [30,31,33]. Two studies evaluated bacterial bioburden reduction [31,33]. Four studies reported adverse events (AEs) [28,29,31,32]. No AEs were noted in the last two studies [30,33]. Table 1 reports the summary of findings for the primary outcome and Table S4 the different outcomes reported among the studies.

2.2. Iodine vs. Saline (3 Randomised Controlled Trials (RCT), 260 Patients)

Three studies, with a total of 260 patients, compared iodine to saline [28,31,32]. In 1984, Holloway et al. studied the efficacy of cadexomer iodine in 3 months venous stasis ulcers over 24 weeks [31]. The study was rated at high risk of bias. In 2018, Raju et al. also studied the efficacy of cadexomer iodine in the treatment of various 1 month chronic ulcers over 12 weeks [32]. In 2019, Gwak et al. studied the efficacy of an 8 weeks treatment of povidone iodine in patients with DFU with a mean duration of 7 weeks [28]. The risk of bias was unclear for these studies.

Two studies assessed the primary outcome as the proportion of patients with complete wound healing (Table S1) [28,32]. Gwak et al. observed no significant difference in the two groups at 8 weeks (44.4% vs. 44.1%; \( p = 0.978 \)) [28], whereas Raju et al. found a significant complete wound healing for patients treated with two different formulations of cadexomer iodine compared to saline at 12 weeks (61.9% vs. 20%; \( p < 0.001 \)) [32]. The pooled data showed that iodine has a higher percentage of patients achieving complete wound healing (RR 1.85 (95%CI (1.27 to 2.69), \( n = 2 \) studies, moderate quality evidence) (Table 2). In the last study, the endpoint was the time to complete healing (Table S1) [31]. The authors found no significant difference between the two groups (31.0 days \( \pm \) 14.1 for povidone iodine vs. 33.3 \( \pm \) 12.6 for saline; \( p = 6.54 \)).
Table 1. Description of included studies.

| Study                  | Country                  | Date                          | Setting               | Number of Participants | Diagnosis                                                                 | Main Outcome (Scale Details)                          | Duration of the Study | Intervention                  | Comparator                  |
|------------------------|--------------------------|-------------------------------|-----------------------|------------------------|-----------------------------------------------------------------------------|-------------------------------------------------------|------------------------|-----------------------------|-----------------------------|
| Gwak et al., 2020 [28]  | Korea                    | March 2016–September 2017     | Multicentre 10 sites  | 71                     | DFU ≥ 1 cm² post debridement and no clinical signs of infection             | Proportion of patients with complete wound healing    | 8 weeks                | PVP-I 44.4% (16/36)          | Saline 44.1% (15/35)        |
| Raju et al., 2019 [32]  | India                    | March 2016–March 2017         | Multicentre 15 sites  | 124                    | Chronic ulcers (single VLU, DFU, PU with adequate arterial blood supply)   | Proportion of patients with complete wound healing    | 12 weeks               | Cadexomer iodine ointment: 65.9% (27/41) | Cadexomer iodine powder: 58.1% (25/43) | Saline 20% (8/40) |
| Bellingeri et al., 2016 [30] | Italy                   | June 2010–December 2013       | Multicentre 6 sites   | 289                    | PU less than 80 cm²                                                        | Wound improvement measured by BWAT scale tool         | 4 weeks                | PHMB                        | Saline                      |
| Vanscheidt et al., 2011 [29] | Germany, France, Hungary, UK | November 2007–December 2009   | Multicentre 15 sites  | 126                    | Chronic venous ulcer locally infected                                       | Time to complete wound healing                       | 12 weeks               | Octenidine 92 days 30.6% (15/49) | Saline 87 days 32.0% (16/50) |
| Sibbald et al., 2011 [33] | Canada                   | February 2008–April 2009      | Multicentre 2 sites   | 40                     | Chronic wounds > 1 cm²                                                     | Healing rate                                         | 4 weeks                | PHMB 35% reduction in wound surface | Saline 28% reduction in wound surface |
| Holloway et al., 1989 [31] | USA                      | NG                            | Multicentre 3 sites   | 75                     | At least a venous stasis ulcer present for a minimum of 3 months           | Healing rate                                         | 24 weeks               | Cadexomer iodine 0.95 cm² per week | Saline 0.41 cm² per week |

BWAT, Bates Jensen Wound Assessment Tool; DFU, diabetic foot ulcer; NG, not given; PHMB, polyhexamethylenebiguanide; PU, pressure ulcer; PVP-I, povidone-iodine; VLU, venous leg ulcer.
**Table 2.** Summary of findings table with comparison between iodine and saline solution in chronic wound care.

| Outcomes                                                                 | Anticipated Absolute Effects * (95% CI) | Relative Effect (95% CI) | № of Participants (Studies) | Certainty of the Evidence (GRADE) |
|--------------------------------------------------------------------------|-----------------------------------------|--------------------------|-----------------------------|-----------------------------------|
| Patient or Population: Chronic Wound Care                                | Intervention: Iodine                    |                          |                             |                                   |
| Comparison: Normal Saline                                                |                                         |                          |                             |                                   |
| Proportion of patients with complete wound healing assessed with: visual assessment follow up: range 8 weeks to 12 weeks | Risk with Normal Saline: 307 per 1000 (390 to 824) | RR 1.8478 (1.2706 to 2.6874) | 195 (2 RCTs) | ⊕⊕⊕⊕ MODERATE ⬤ |
| Risk with Iodine: 567 per 1000 (980 to 824)                               |                                         |                          |                             |                                   |
| Adverse events assessed with: report follow-up: range 8 weeks to 24 weeks | Risk with Normal Saline: 115 per 1000 (89 to 308) | RR 1.440 (0.774 to 2.676) | 270 (3 RCTs) | ⊕⊕⊕ MODERATE ⬤ |
| Risk with Iodine: 166 per 1000 (89 to 308)                                |                                         |                          |                             |                                   |
| Ulcer healing rate (healing rate) assessed with: planimetry follow-up: range 8 weeks to 24 weeks | Raju et al. [32] found a significantly ($p < 0.001$) higher percentage of reduction for both formulations of iodine (94.3% and 90.4%) compared to saline (67.8%). Holloway et al. [31] found a rate reported to baseline of 0.04 ± 0.01 cm²/week/cm² for cadexomer iodine and 0.03 ± 0.01 cm²/week/cm² for saline. There was no significant difference ($p = 0.079$). Gwak et al. [28] presented the healing rate with three different visual displays showing the percentage change rate for the length, the width, and the area. They found no difference. | | 270 (3 RCTs) | ⊕⊕⊕⊕ VERY LOW b,c,d |
| Pain evaluation (Pain) assessed with: mean rate of change follow up: mean 24 weeks | The mean rate of change in pain scores were −2.44 ± 0.4 for cadexomer iodine and −2.47 ± 0.3 with saline with a $p = 0.96$. | | (1 RCT) | ⊕⊕⊕ LOW d,e |

**GRADE Working Group grades of evidence**

- **⊕⊕⊕⊕ High certainty:** We are very confident that the true effect lies close to that of the estimate of the effect.
- **⊕⊕⊕ Moderate certainty:** We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.
- **⊕⊕ Low certainty:** Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect.
- **⊕ Very low certainty:** We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect.

* The risk in the intervention group (and 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and 95% CI). CI: confidence interval; RR: risk ratio; RCT: randomised controlled trial. (a) Missing outcome data, around 30% of patient losses during follow-up. (b) One study with a high risk of bias due to methodology risk and both studies at risk to outcome reporting. (c) Different results in the two studies. (d) Selective outcome report. (e) Study with an overall high risk of bias.
Among the three studies, two reported the healing rate (Table 2) [31,32]. Raju et al. presented the healing rate as reduction percentage in ulcer size from baseline to 12 weeks. They observed a significant reduction on wounds treated with both formulations of iodine (94.3% and 90.4%) compared to saline (67.8%) ($p < 0.001$) [32]. Holloway et al. defined the healing rate as the ulcer size reduction per week per baseline size. They noted no significant difference between patients treated with cadexomer iodine versus saline ($0.04 \text{ cm}^2/\text{week/cm}^2 \pm 0.01$ vs. $0.03 \pm 0.01$, respectively; $p = 0.079$) [31]. Finally, Gwak et al. reported three indirect markers of healing rate: the percentage rate of change of wound length, width, and area [28]. They found no difference between the two treatments. The heterogeneity of the data prevented pooling. The quality of the evidence was very low.

Pain was only evaluated in one study [31]. No statistical difference was observed between the two treatments (cadexomer iodine versus saline) ($p = 0.96$) (Table 2). The quality of the evidence was low.

Finally, no studies evaluated the bacterial bioburden.

All three studies reported AEs [28,31,32]. The pooled data showed no significant difference between AE incidence in the iodine group compared to the saline group (RR 1.44 (95%CI (0.77 to 2.68), $n = 3$ studies, moderate quality evidence) (Table 2).

### 2.3. Polyhexanide Compared to Saline (2 RCT, 334 Patients)

Two studies, with a total of 334 patients, compared polyhexanide to saline [30,33], both with unclear risk of bias. In 2011, Sibbald et al. studied the efficacy of polyhexanide solution in patients with chronic wounds over 4 weeks [33]. In 2016, Bellingeri et al. studied the efficacy of polyhexanide solution in patients with PUs or mixed aetiologies of chronic ulcers, over 4 weeks [30]. Neither study reported wound healing, and thus the primary outcome could not be evaluated.

The healing rate is described in Table 3 [30,33]. Bellingeri et al. evaluated healing rate as wound improvement on the 13 item BWAT scale (Bates Jensen Wound Assessment Tool) [30]. They observed a significantly improved healing rate in the polyhexanide group ($p = 0.072$), and a significantly better score at 4 weeks compared to the first week in the experimental group ($p = 0.025$). Sibbald et al. calculated the percentage decrease of the wound surface area by planimetry measurement and comparison [33]. They noted no significant difference between the median reduction of the wound surface in the polyhexanide group versus the saline group (35% vs. 28%; $p = 0.85$). Due to the heterogeneity of the measurement tools, we could not pool the data. The quality of the evidence was low.
Table 3. Summary of findings table with comparison between polyhexanide and saline solution in chronic wound care.

### Comparison Two: Polyhexanide Compared to Saline for Chronic Wound Care

**Patient or Population:** Chronic Wound Care  
**Intervention:** Polyhexanide  
**Comparison:** Saline

| Outcomes | Anticipated Absolute Effects * (95% CI) | Relative Effect (95% CI) | № of Participants (Studies) | Certainty of the Evidence (GRADE) |
|----------|---------------------------------------|--------------------------|-----------------------------|----------------------------------|
| Wound healing follow-up: mean 4 weeks | Risk with Saline: Not measured  
Risk with Polyhexanide: Not measured | RR 0.2024 (0.0098 to 4.1813) | 334 (2 RCTs) | ⊕⊕⊕⊕ LOW a,b |
| Adverse events assessed with: report follow-up: mean 4 days | 12 per 1000 (0 to 50)  
2 per 1000 | | | |
| Healing rate assessed with: planimetry follow-up: median 4 weeks | Bellingeri et al. [30] found a significantly better progression of wounds in the polyhexanide group (p = 0.0025) using the BWAT wound assessment scale.  
Sibbald et al. [33] found no significant difference (p = 0.85) between the two groups by comparing wound surface reduction (35% vs. 28%). | 334 (2 RCTs) | ⊕⊕⊕⊕ LOW a,b |
| Pain assessment assessed with: Pain scales follow-up: mean 4 weeks | Bellingeri et al. [30] found similar pain scores with no significant difference in the two groups (average score = 3 with minimal or no change during follow up).  
Sibbald et al. [33] reported significant pain reduction in the polyhexanide group compared to the saline control group (73.1% vs. 38.1%; p = 0.02). | (2 RCTs) | ⊕⊕⊕⊕ LOW a,b |

**GRADE Working Group grades of evidence**

⊕⊕⊕⊕ *High certainty:* We are very confident that the true effect lies close to that of the estimate of the effect.

⊕⊕⊕ *Moderate certainty:* We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.

⊕⊕ *Low certainty:* Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect.

⊕ *Very low certainty:* We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of the effect.

* The risk in the intervention group (and 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and 95% CI). CI: confidence interval; RR: risk ratio; RCT: randomised controlled trial. (a) Two studies with different results. (b) Very few participants in one study.

Concerning pain, Bellingeri et al. observed no significant difference between groups [30], whereas Sibbald et al. reported a significant pain reduction in the polyhexanide group compared to the saline group (73.1% vs. 38.1%, p = 0.02) [33]. We graded the quality of the evidence low.

Finally, Sibbald et al. assessed the reduction of the bacterial bioburden [33]. They noted polymicrobial microorganisms in 5.3% of the polyhexanide group wounds versus 33% in the control group (p = 0.016).

Bellingeri et al. reported no AEs in either group [30], but Sibbald et al. recorded two AEs as infection in the saline control group [33]. The pooled data showed a significant difference between the incidence of AEs in the polyhexanide group compared to the saline group (RR 0.2 (95%CI (0.01 to 4.18), n = 2 studies, low-quality evidence) (Table 3).
2.4. Octenidine Compared to Saline (1 RCT, 126 Patients)

One study, with 126 patients, compared octenidine to saline [29]. Vanscheidt et al. assessed the efficacy of octenidine in patients with locally infected chronic VLU that was at least 1 month old and who had had no previous or concomitant drug therapy for 12 weeks. Their study was categorised at unclear risk of bias.

Wound healing was assessed as time to complete wound closure and the proportion of patients with complete wound closure (Table S1). The time to complete wound healing was not significantly different between the groups (92 days for octenidine vs. 87 days for saline; $p = 0.952$) (Table 4). Accordingly, the proportion of patients with complete wound healing was similar (30.6% vs. 32%; $p = 0.882$). Interestingly, Vanscheidt et al. reported a significant proportion of healing for patients with ulcers larger than 6 cm$^2$ and older than 6 months in the octenidine group versus the saline group (33.3% vs. 0%, respectively $p=0.022$) [29]. We graded the quality of the evidence as high.

Table 4. Summary of findings table with comparison between octenidine and saline solution in chronic wound care.

| Outcomes                          | Anticipated Absolute Effects * (95% CI) | Relative Effect (95% CI) | No of Participants (Studies) | Certainty of the Evidence (GRADE) |
|-----------------------------------|----------------------------------------|--------------------------|------------------------------|----------------------------------|
| Wound healing assessed with:     |                                        |                          |                              |                                  |
| Proportion of patients with      | 242 per 1000                           | RR 1.0313                | 126                          | ⭐⭐⭐⭐ HIGH                       |
| complete wound healing follow-up:| 250 per 1000                           | (0.5595 to 1.9009)      |                              |                                  |
| mean 12 weeks                    | (136 to 461)                           |                          |                              |                                  |
| Adverse events assessed with:    | 317 per 1000                           | RR 0.5614                | 120                          | ⭐⭐⭐⭐ HIGH                       |
| AE report follow-up:             | 178 per 1000                           | (0.2844 to 1.1081)      |                              |                                  |
| mean 12 weeks                    | (90 to 351)                            |                          |                              |                                  |
| Healing rate assessed with:      |                                        | No difference in the healing rate of the patients in the octenidine group, compared to patients in the saline group (37.9% vs. 40.3%; $p = 0.769$) [29]. | 120                          | ⭐⭐⭐⭐ HIGH                       |
| planimetry follow-up:            |                                        |                          |                              |                                  |
| mean 12 weeks                    |                                        |                          |                              |                                  |
| Pain assessment—not measured      | Not reported                           |                          |                              |                                  |

GRADING Working Group grades of evidence

⭐⭐⭐⭐ High certainty: We are very confident that the true effect lies close to that of the estimate of the effect.
⭐⭐⭐ Moderate certainty: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.
⭐⭐ Low certainty: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect.
⭐ Very low certainty: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of the effect.

* The risk in the intervention group (and 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and 95% CI). CI: confidence interval; RR: risk ratio; RCT: randomised controlled trial.
No difference in the healing rate was noted in the octenidine group compared to the saline group (37.9% vs. 40.3%, $p = 0.769$). The octenidine group achieved better results from 5 weeks. We graded the quality of the evidence high.

The study did not assess pain or bacterial bioburden.

AEs were reported in 10 patients in the octenidine group and 19 patients in the saline group, without significant difference between the groups [29]. We graded the quality of the evidence high.

3. Discussion

We reviewed the RCT evidence for the use of antiseptic agents in chronic wound care in adult patients. Although saline is the main recommended product used in chronic wound cleansing, numerous clinical studies described the benefit of antiseptic agents in this situation [7,17,21,35–37]. A limited number of studies are available on the efficacy of antiseptic agents on chronic wound healing. More limited studies are available on the efficacy of antiseptics on pain. The trials are small, clinically heterogeneous, without clearly defined outcomes, and at high or unclear risk of bias. Of the 838 RCT identified, only 6 studies were included, representing a total of 725 patients.

Our review established a better wound healing with iodine compared to saline (2 RCT, 195 patients, RR 1.85 (1.27 to 2.69)), although the quality of the evidence was moderate. In contrast, no statistical efficacy of octenidine on healing rate (compared to saline) was seen with a high-quality evidence grade (1 RCT, 126 patients RR 1.03 (0.56 to 1.90)). Interestingly, none of the antiseptic agents influenced AE occurrence compared to saline. Notably, over half of the clinical trials have never been published. Most studies had unclear risk of bias, as previously described [38,39].

Of the 838 studies, most of them did not evaluate clinical signs of infection, and mainly focused on bacterial load reduction, a measure long deemed unsuitable [2–4]. Furthermore, the six included studies ignored the effect of biofilm in delaying the healing process. Of the two studies assessing microbiological impact on infection, none assessed biofilm reduction [30,31,33]. For future research, we suggest the use of dynamic models mimicking the wound environment instead of the traditional quantitative microbial load in in vitro studies [40]. This includes non-static models and the consideration of multispecies biofilm reduction over a clinically relevant time (>1 month).

Although most guidelines recommend against the use of antiseptic agents [2,4,25,27], a recent consensus suggested using hypochlorite and polyhexanide in chronic wound care [21]. We found no study demonstrating a significant effect of hypochlorite on the healing of chronic wounds. We could not assess the efficacy of polyhexanide due to the heterogeneity of outcomes between studies. However, this consensus included other types of non-healing wounds such as post-surgical or burn wounds and made no distinction between WAR score categories (colonised and infected wounds). Finally, it also included non-randomised trials, which provide lower evidence than RCTs and different systematic bias are encountered [41]. The main limitation of this guideline is the extrapolation of recommendations from various studied wounds to specific chronic wound care. Another key problem is the attribution of effect to antiseptic agents when antibiotics were systematically used in case of infected wounds.

Following the diverse interpretations of study results in recommendations, future investigations in primary research must focus on value to patients and healthcare professionals, particularly treatment choice. The design of future trials should be driven by high-priority questions. Moreover, good practice guidelines must be followed at each step (e.g., design, implementation, reporting). Assessment of complete wound healing instead of wound healing rate would be more relevant, and time to complete wound healing should be reported as the main endpoint. Future research should be controlled at least against saline, and preferably with another or multiple other antiseptic agents. Another fruitful area of research would be the impact of antiseptic agents according to wound size. Two of our included studies on two different antiseptic agents reported increased healing rate for
wounds larger than 6 cm² versus smaller wounds [29,31]. Further good quality evidence studies may aid decision making about the use of topical antiseptics in the management of chronic wounds.

The main limitation of this review was the great heterogeneity in study designs, methodologies, and outcomes. Overall, the six studies included in this review were too heterogeneous for pooling. We could not compare the four antiseptic agents with one another. Sensitivity analysis could not be performed due to the low number of results nor could subgroup analysis by age or type of chronic wound.

Despite its limitations, this review assesses the quality of the available data on the four selected antiseptic agents on chronic wound healing, using a well-known and robust methodology. It focuses the topic on chronic wound, reducing the bias of specific cares required for the other types of wounds.

The relative effects of topical antiseptic treatment on chronic wounds are unclear. There is insufficient evidence to determine the superiority of one antiseptic agent over the others. We could not assess the effect of hypochlorous agents on chronic wound healing. Moderate evidence suggests an improvement of wound healing with iodine compared to saline. Currently, there is not enough evidence to recommend one antiseptic agent over another in this clinical situation.

4. Materials and Methods
4.1. Selection Criteria and Search Strategy

The study is registered at PROSPERO (CRD42020213494).

The review protocol was based on the Cochrane Handbook for Systematic Reviews of Intervention (version 6.2) [42] and devised in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analyses (PRISMA) statement guidelines [43]. A meta-analysis was initially planned. Due to heterogeneity in included studies, a systematic review with a summary of effect estimates was performed.

Published and unpublished RCT were eligible. Two reviewers (K.B.B., J.O.) independently screened titles and abstracts to determine eligibility and assessed the full text of retained articles. Disagreements were solved by discussion or arbitration by three independent reviewers (A.S., P.L., J.P.L.). Exclusion reasons were recorded. This study followed this PICO strategy:

1. **Population**: Our population was adult patients (≥18 years) with chronic wounds as previously defined [2]. We included studies from primary, secondary, and tertiary clinical settings. We included different types of chronic wounds (leg ulcer, DFU, PU, eschar). We excluded studies containing patients with wounds requiring specific care (acute wounds, burn wounds, systemic infected wound, postsurgical wounds).
2. **Interventions** of interest were those including antiseptics as cleansing method or within a dressing with at least weekly application.
3. **Comparative** regimens included saline solution or another antiseptic. We anticipated that interventions would consist of povidone-iodine, hypochlorite or hypochlorous acid, iodine, polyhexanide, and octenidine in the form of creams, ointments, powders, sprays, or impregnated into dressings. We included intervention schedules applying concurrent therapies (e.g., negative pressure wound therapy) if the therapy was common across study arms. We excluded (i) interventions where the antiseptic agent was not the only systematic difference between treatment groups; (ii) physical and biological therapies with antimicrobial properties, such as heat or larval therapy; (iii) studies evaluating topical antiseptics in prevention of chronic wounds or those using antiseptics as preparation for surgical treatment of ulcers; (iv) studies evaluating non-recommended antiseptics in chronic wound care and those evaluating antiseptic agents alongside antibiotic agents.
4. **Outcome**: The primary outcome was wound healing, evaluated as the proportion of patients with complete healing during follow-up and/or the time to complete wound healing (analysed using survival, time-to-event approaches). An adjustment
for relevant covariates such as baseline wound area or duration were ideally used to evaluate the outcome. The secondary outcomes were healing rate (described as changes or rate of change in wound size, with adjustment for baseline size); mean pain scores; bacterial bioburden reduction; and AEs, including infection.

We searched PubMed (NLM database), MEDLINE (OvidSP), Web of Science (Thomson Reuters), Google Scholar, and Cochrane library databases, as well as 3 clinical trial registries (ClinicalTrials.gov (www.clinicaltrials.gov, accessed on 17 January 2022)), EU Clinical Trials Register (https://www.clinicaltrialsregister.eu, accessed on 17 January 2022), and World Health Organisation (WHO) International Clinical Trial Registry Platform (https://apps.who.int/trialsearch/, accessed on 17 January 2022) up to 30 June 2021 without restrictions for language, study status, date of publication, or country, using a MeSH terms string chain (Figure 1). Furthermore, we searched the reference lists of reviewed studies for relevant studies. The search strategy for all databases is presented in Appendix A.1.

4.2. Data Collection and Analysis

One reviewer (K.B.B.) performed data extraction and quality assessment for the included studies, validated by a second author (J.O.). Disagreements were resolved by arbitration by three independent review authors (A.S., P.L., J.P.L.). We contacted study authors for additional data if necessary. We performed data extraction using a standardised sheet, as recommended by the Cochrane Collaboration’s handbook for systematic review (trial authors, year of publication, patient population characteristics, duration of follow-up, trial design, measured outcomes, including assessment methods, objectives, results, country where trials were performed, number of participants randomly assigned to each treatment group, clinical setting, detail of interventions in each group, details of comparators in each group, source of funding, number of withdrawals, outcomes). Data are presented in Table S2.

4.3. Risk of Bias and Certainty of the Evidence

Two reviewers (K.B.B., J.O.) independently assessed the risk of bias of eligible studies; any disagreements were resolved by arbitration by three independent review authors (A.S., P.L., J.P.L.). Risk of bias was assessed using the bias excel tool (RoB 2 checklist, 2019) [44] as recommended by the Cochrane Handbook for Systematic Reviews of Intervention (version 6.2) [42]. The overall bias risk was rated as low, moderate, high, or unclear (also some concerns). The 7 domains of bias were also assessed for each trial, following the recommendation of the Cochrane collaboration, 2011 [34].

The overall quality of evidence of included studies was assessed using Grading of Recommendations Assessment, Development and Evaluation (GRADE) as recommended by the Cochrane Handbook for Systematic Reviews of Intervention (version 6.2) [44]. Summary tables for each antiseptic were produced with GRADEPro software considering five outcomes: complete wound healing (healing rate or proportion of patients with complete wound healing), rate of change in wound size, pain assessment, bacterial bioburden reduction, and AEs. We calculated the risk ratio (RR) for dichotomous outcomes (wound healing, AEs, infection) with 95% confidence interval (CI).

4.4. Role of the Funding Source

There was no funding source for this study.

5. Conclusions

Iodine compounds showed a better effect on chronic wound healing compared to saline. Octenidine and polyhexanide did not show any difference in this healing compared to saline. Currently, there is not enough evidence to recommend one antiseptic over another in this clinical situation. Future clinical trials assessing antiseptic agents in chronic wounds management should pay attention to include several antiseptic agents in their trial for comparison, respecting the double-blind trial and with a well-defined study population.
They should establish the main efficiency criteria as complete wound healing and not only wound size reduction. A sub-group analysis based on the size of the wound would be relevant to the matter. Finally, following good practice guidelines is mandatory in every step of trials in order to avoid the numerous biases found in the assessed studies.

**Supplementary Materials:** The following supporting information can be downloaded at [https://www.mdpi.com/article/10.3390/antibiotics11030350/s1](https://www.mdpi.com/article/10.3390/antibiotics11030350/s1), Figure S1: Plot of the percentage of risk of bias assessments at each level of risk of bias per domain. Table S1: Summary of the included studies. Table S2: Risk of bias assessment among included studies. Table S3: Review authors’ judgements about each risk of bias item for each included study. Table S4: Different outcomes reported among the studies.

**Author Contributions:** Conceptualisation, A.S., J.-P. L. and P.L.; methodology, K.B.-B., F.S. and P.L.; software, F.S.; validation, J.O., A.S., J.-P. L. and P.L.; formal analysis, K.B.-B. and J.O.; investigation, K.B.-B. and J.O.; writing—original draft preparation, K.B.-B., J.O. and P.L.; writing—review and editing, A.S., F.S. and J.-P. L.; supervision, A.S. and J.-P. L.; project administration, P.L.; funding acquisition, A.S. and J.-P. L. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research received no external funding.

**Acknowledgments:** We thank Sarah Kabani for editing the article. We thank the Nîmes University hospital for its structural, human, and financial support through the award obtained by our team during the internal call for tenders “Thématiques phares”. The authors belong to the FHU InCh (Federation Hospitalo Universitaires Infections Chroniques, Aviesan.

**Conflicts of Interest:** The authors declare no conflict of interest.

**Appendix A**

**Appendix A.1 Medline and Cochrane Library Search**

Our search strategy basis was to focus on the four main antiseptic agents recommended or used in practice: chlorous compounds, octenidine, iodophor compounds, and polyhexanide. Four different searches were conducted then combined:

- (((((chronic wound) OR diabetic foot) OR leg ulcer) OR pressure ulcer) OR eschar) AND (((iodine) OR povidone iodine OR betadine) OR cadexomer iodine).
- (((((chronic wound) OR diabetic foot) OR leg ulcer) OR pressure ulcer) OR eschar) AND (((hypochlorite) OR hypochlorous) OR Dakin) OR Javel).
- (((((chronic wound) OR diabetic foot) OR leg ulcer) OR pressure ulcer) OR eschar) AND (((phMB) OR polyhexanide) OR Polyhexamethylene biguanide) OR Polyhexamethylene).
- (((((chronic wound) OR diabetic foot) OR leg ulcer) OR pressure ulcer) OR eschar) AND (Octenidine).

**References**

1. Järbrink, K.; Ni, G.; Sönnergren, H.; Schmidtchen, A.; Pang, C.; Bajpai, R.; Car, J. Prevalence and incidence of chronic wounds and related complications: A protocol for a systematic review. *Syst. Rev.* 2016, 5, 152. [CrossRef] [PubMed]
2. Kirsner, R. The Wound Healing Society chronic wound ulcer healing guidelines update of the 2006 guidelines—Blending old with new. *Wound Repair Regen.* 2016, 24, 110–111. [CrossRef] [PubMed]
3. Lipsky, B.A.; Berendt, A.R.; Deery, H.G.; Embil, J.M.; Joseph, W.S.; Karchmer, A.W.; LeFrock, J.L.; Lew, D.P.; Mader, J.T.; Norden, C.; et al. Infectious Diseases Society of America. Diagnosis and treatment of diabetic foot infections. *Clin. Infect. Dis.* 2004, 39, 885–910. [CrossRef] [PubMed]
4. Société de Pathologie Infectieuse de Langue Française. Management of diabetic foot infections. Long text. Société de Pathologie Infectieuse de Langue Française. *Med. Mal. Infect.* 2007, 37, 26–50.
5. Bjarnsholt, T. The role of bacterial biofilms in chronic infections. *APMIS* 2013, 121(s136), 1–58. [CrossRef]
6. Pouget, C.; Dunyach-Remy, C.; Pantel, A.; Schuldiner, S.; Sotto, A.; Lavigne, J.P. Biofilms in diabetic foot ulcers: Significance & clinical relevance. *Microorganisms* 2020, 8, 1580.
7. Schultz, G.; Bjarnsholt, T.; James, G.A.; Leaper, D.J.; McBain, A.J.; Malone, M.; Stoodley, P.; Swanson, T.; Tachi, M.; Wolcott, R.D. Global Wound Biofilm Expert Panel. Consensus guidelines for the identification and treatment of biofilms in chronic nonhealing wounds. *Wound Repair Regen.* 2017, 25, 744–757. [CrossRef]
8. Dissemond, J.; Assadian, O.; Gerber, V.; Kingsley, A.; Kramer, A.; Leaper, D.J.; Mosti, G.; Piatkowski de Grzymala, A.; Riepe, G.; Risse, A.; et al. Classification of wounds at risk and their antimicrobial treatment with polyhexamidine: A practice-oriented expert recommendation. *Skin Pharmacol. Physiol.* 2011, 24, 245–255. [CrossRef]

9. Bakker, K.; Apelqvist, J.; Lipsky, B.A.; Van Netten, J.J. International Working Group on the Diabetic Foot. The 2015 IWGDF guidance documents on prevention and management of foot problems in diabetes: Development of an evidence-based global consensus. *Diabetes Metab. Res. Rev.* 2016, 32, 2–6. [CrossRef]

10. Wolcott, R.D.; Kennedy, J.P.; Dowd, S.E. Regular debridement is the main tool for maintaining a healthy wound bed in most chronic wounds. *J. Wound Care.* 2009, 18, 54–56. [CrossRef]

11. Sibbald, R.G.; Elliott, J.A.; Persaud-Jaimangal, R.; Goodman, L.; Armstrong, D.G.; Harley, C.; Coelho, S.; Xi, N.; Evans, R.; Mayer, D.O.; et al. Wound Bed Preparation 2021. *Adv. Skin Wound Care.* 2021, 34, 183–195. [CrossRef]

12. Sun, X.; Chen, J.; Zhang, J.; Wang, W.; Sun, J.; Wang, A. Maggot debridement therapy promotes diabetic foot wound healing by up-regulating endothelial cell activity. *J. Diabetes Complicat.* 2016, 30, 318–322. [CrossRef] [PubMed]

13. Ayello, E.A.; Cuddigan, J.E. Debridement: Controlling the necrotic/cellular burden. *Adv. Skin Wound Care* 2004, 17, 66–75. [CrossRef] [PubMed]

14. Kravitz, S.R.; McGuire, J.; Zinsker, K. Management of skin ulcers: Understanding the mechanism and selection of enzymatic debridging agents. *Adv. Skin Wound Care* 2008, 21, 72–74. [CrossRef] [PubMed]

15. De Francesco, F.; De Francesco, M.; Riccio, M. Hyaluronic Acid/Colagenase Ointment in the Treatment of Chronic Hard-to-Help Wounds: An Observational and Retrospective Study. *J. Clin. Med.* 2022, 11, 537. [CrossRef] [PubMed]

16. Agarwal, P.; Kukrele, R.; Sharma, D. Vacuum assisted closure (VAC)/negative pressure wound therapy (NPWT) for difficult wounds: A review. *J. Clin. Orthop. Trauma.* 2019, 10, 845–848. [CrossRef]

17. O’Meara, S.; Cullum, N.; Nelson, E.A.; Dumville, J.C. Compression for venous leg ulcers. *Cochrane Database Syst. Rev.* 2012, 11, CD000265. [CrossRef] [PubMed]

18. Vouillarmet, J.; Bourron, O.; Gaudric, J.; Lermusiaux, P.; Millon, A.; Hartemann, A. Lower-extremity arterial revascularization: Is there any evidence for diabetic foot ulcer-healing? *Diabetes Metab.* 2016, 42, 4–15. [CrossRef]

19. Schaper, N.C.; van Netten, J.J.; Apelqvist, J.; Bus, S.A.; Hinchliffe, R.J.; Lipsky, B.A. IWGDF Editorial Board. Practical guidelines on the prevention and management of diabetic foot diseases (IWGDF 2019 update). *Diabetes Metab. Res. Rev.* 2020, 36, e3266. [CrossRef]

20. Bowers, S.; Franco, E. Chronic wounds: Evaluation and management. *Am. Fam. Physician.* 2020, 101, 159–166.

21. Kramer, A.; Dissemond, J.; Kim, S.; Willy, C.; Mayer, D.; Papke, R.; Tuchmann, F.; Assadian, O. Consensus on wound antisepsis: Update 2018. *Skin Pharmacol. Physiol.* 2018, 31, 28–58. [CrossRef] [PubMed]

22. Babalska, Z.L.; Korbecka-Paczkowska, M.; Karpinski, T.M. Wound Antiseptics and European Guidelines for Antiseptic Application in Wound Treatment. *Pharmaceuticals* 2021, 14, 1253. [CrossRef] [PubMed]

23. Pouget, C.; Dunyach-Remy, C.; Pantel, A.; Boutet- Dubois, A.; Schuldiner, S.; Sotto, A.; Lavigne, J.P.; Loubet, P. Alternative approaches for the management of diabetic foot ulcers. *Front. Microbiol.* 2021, 12, 747618. [CrossRef] [PubMed]

24. Rayman, G.; Vas, P.; Dhatariya, K.; Driver, V.; Hartemann, A.; Londahl, M.; Piagessi, A.; Apelqvist, J.; Attinger, C.; Game, F. International Working Group on the Diabetic Foot (IWGDF). Guidelines on use of interventions to enhance healing of diabetic foot ulcers in diabetes (IWGDF 2019 update). *Diabetes Metab. Res. Rev.* 2020, 36, e3283. [CrossRef]

25. Norman, G.; Dumville, J.C.; Moore, Z.E.H.; Tanner, J.; Christie, J.; Goto, S. Antibiotics and antiseptics for pressure ulcers. *Cochrane Database Syst. Rev.* 2016, 4, CD001186. [CrossRef]

26. Dogra, S.; Sarangal, R. Summary of recommendations for leg ulcers. *Indian Dermatol. Online J.* 2014, 5, 400–407. [CrossRef]

27. Lipsky, B.A.; Senneville, E.; Abbas, Z.G.; Aragón-Sánchez, J.; Diggle, M.; Embil, J.M.; Kono, S.; Lavery, L.A.; Malone, M.; van Asten, S.A.; et al. International Working Group on the Diabetic Foot (IWGDF). Guidelines on the diagnostic and treatment of foot infection in persons with diabetes (IWGDF 2019 update). *Diabetes Metab. Res. Rev.* 2020, 36, e3280. [CrossRef]

28. Gwak, H.C.; Han, S.H.; Lee, J.; Park, S.; Sung, K.S.; Kim, H.J.; Chun, D.; Lee, K.; Ahn, J.H.; Kwak, K.; et al. Efficacy of a povidone-iodine foam dressing (Betafoam) on diabetic foot ulcer. *Int. Wound J.* 2020, 17, 91–99. [CrossRef]

29. Vanscheidt, W.; Harding, K.; Teot, L.; Siebert, J. Effectiveness and tissue compatibility of a 12-week treatment of chronic venous leg ulcers with an octenidine based antiseptic—A randomized, double-blind controlled study. *Int. Wound J.* 2012, 9, 316–323. [CrossRef]

30. Bellingeri, A.; Falciani, F.; Traspedini, P.; Moscatelli, A.; Russo, A.; Tino, G.; Chiari, P.; Peghetti, A. Effect of a wound cleansing solution on wound bed preparation and inflammation in chronic wounds: A single-blind RCT. *J. Wound Care* 2016, 25, 160–168. [CrossRef]

31. Holloway, G.A.; Johansen, K.H.; Barnes, R.W.; Pierce, G.E. Multicenter trial of cadexomer iodine to treat venous stasis ulcer. *West J. Med.* 1989, 151, 35–38. [CrossRef]

32. Raju, R.; Kethavath, S.N.; Sangavaranapu, S.M.; Kanjarla, P. Efficacy of cadexomer iodine in the treatment of chronic ulcers: A randomized, multicenter, controlled trial. *Wounds* 2019, 31, 85–90. [PubMed]

33. Sibbald, R.G.; Coutts, P.; Woo, K.Y. Reduction of bacterial burden and pain in chronic wounds using a new polyhexamethylene biguanide antimicrobial foam dressing-clinical trial results. *Adv. Skin Wound Care* 2011, 24, 78–84. [CrossRef] [PubMed]
34. Higgins, J.P.T.; Altman, D.G.; Gøtzsche, P.C.; Jüni, P.; Moher, D.; Oxman, A.D.; Savovic, J.; Schulz, K.F.; Weeks, L.; Sterne, J.A.C. Cochrane Bias Methods Group; Cochrane Statistical Methods Group. The Cochrane Collaboration’s tool for assessing risk of bias in randomised trials. *BMJ* 2011, 343, d5928. [CrossRef]
35. O’Meara, S.M.; Cullum, N.A.; Majid, M.; Sheldon, T.A. Systematic review of antimicrobial agents used for chronic wounds. *Br. J. Surg.* 2001, 88, 4–21. [CrossRef]
36. Hämmerle, G.; Strohal, R. Efficacy and cost-effectiveness of octenidine wound gel in the treatment of chronic venous leg ulcers in comparison to modern wound dressings. *Int. Wound J.* 2016, 13, 182–188. [CrossRef]
37. To, E.; Dyck, R.; Gerber, S.; Kadavil, S.; Woo, K.Y. The effectiveness of topical polyhexamethylene biguanide (PHMB) agents for the treatment of chronic wounds: A systematic review. *Surg. Technol. Int.* 2016, 29, 45–51.
38. Dwan, K.; Altman, D.G.; Arnaiz, J.A.; Bloom, J.; Chan, A.W.; Cronin, E.; Decullier, E.; Easterbrook, P.J.; Von Elm, E.; Gamble, C.; et al. Systematic review of the empirical evidence of study publication bias and outcome reporting bias. *PLoS One* 2008, 3, e3081. [CrossRef]
39. Chan, A.W.; Hróbjartsson, A.; Haahr, M.T.; Gøtzsche, P.C.; Altman, D.G. Empirical evidence for selective reporting of outcomes in randomized trials: Comparison of protocols to published articles. *JAMA* 2004, 291, 2457–2465. [CrossRef]
40. Bahamondez-Canas, T.F.; Heersema, L.A.; Smyth, H.D.C. Current status of in vitro models and assays for susceptibility testing for wound biofilm infections. *Biomedicines* 2019, 7, 34. [CrossRef]
41. Schmidt, W.P. Randomised and non-randomised studies to estimate the effect of community-level public health interventions: Definitions and methodological considerations. *Emerg. Themes Epidemiol.* 2017, 14, 9. [CrossRef] [PubMed]
42. Higgings, J.P.T.; Thomas, J.; Chandler, J.; Cumpston, M.; Li, T.; Page, W.J.; Welch, V.A. Cochrane Handbook for Systematic Reviews of Interventions Version 6.2 (Updated February 2021). Cochrane. 2021. Available online: www.training.cochrane.org/handbook (accessed on 17 January 2022).
43. Moher, D.; Liberati, A.; Tetzlaff, J.; Altman, D.G. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *Int. J. Surg.* 2010, 8, 336–341. [CrossRef] [PubMed]
44. Higgins, J.P.T.; Savović, J.; Page, M.J.; Elbers, R.G.; Sterne, J.A.C. Chapter 8: Assessing risk of bias in a randomized trial. In *Cochrane Handbook for Systematic Reviews of Interventions v6.0*; Higgins, J.P.T., Higgins, J.P.T., Thomas, J., Chandler, J., Cumpston, M., Li, T., Page, M.J., Welch, V.A., Eds.; Cochrane: London, UK, 2019; Available online: www.training.cochrane.org/handbook/current/chapter-08 (accessed on 17 January 2022).