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BESS Patient Care Pathway: Frozen Shoulder  
Partial distal biceps tendon tears  
Scapular Dyskinesia and grip strength

# British Elbow and Shoulder Society patient care pathway: Frozen shoulder

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## Abstract

**Background:** Current guidelines from the British Elbow and Shoulder Society (BESS) were published in 2015 for managing frozen shoulders in the primary and secondary care setting. Updated guidelines have been developed using the Grading of Recommendations, Assessment, Development, and Evaluations (GRADE) methodology.

**Methods:** A multi-disciplinary BESS Working Group defined key management questions based on agreed outcome measures and time points. A literature search, conducted up to March 2023 following PRISMA guidelines, identified randomised controlled trials, systematic reviews, and meta-analyses. Quality assessments were performed using the GRADE Decision Framework, considering bias, imprecision, indirectness, and inconsistency. Data were extracted for meta-analysis. In the absence of high-quality trials, narrative reviews were created.

**Results:** Consensus opinions produced statements based on the quality and volume of evidence and the magnitude of desirable and undesirable effects. These statements form a comprehensive framework for managing frozen shoulder.

**Discussion:** This updated guideline provides evidence-based guidance for managing frozen shoulder and identifies key areas for future research.

## Keywords

Frozen shoulder, adhesive capsulitis, guideline, physiotherapy, hydrodistension, hydrodilatation, manipulation under anaesthesia, steroid, arthroscopic capsular release

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## Introduction

Frozen shoulder, also known as adhesive capsulitis, is a painful debilitating condition with insidious onset, typically leading to stiffness and disability in the shoulder, which lasts over 3 months.<sup>1</sup> The cumulative incidence of frozen shoulder is 2.4/1000 per year,<sup>2</sup> with the prevalence estimated to be between 2% and 5%.<sup>3</sup> It typically affects women between the ages of 45 and 60, with a peak age of 56.<sup>4–7</sup> Hormonal changes in peri-menopausal women are thought to contribute to the increased prevalence, although a causal link has not been identified.<sup>8</sup> The non-dominant arm has an increased risk of being affected, although 6%–17% of patients develop contralateral frozen shoulder within 5 years.<sup>6</sup> The strongest association of frozen shoulder is with diabetes mellitus, where an incidence of 10.8% was found in patients with diabetes compared to 2.3% in patients without diabetes.<sup>9</sup> Other co-morbidities such as hypothyroidism, Dupuytren's disease, Parkinson's disease, osteoporosis, stroke, hyperlipidaemia

and patients having undergone cardiac or neurosurgery have also been associated.<sup>1,5</sup> Three overlapping stages of a frozen shoulder have been described as painful, freezing and thawing.<sup>10</sup> However, owing to considerable overlap between stages, there is highly variable reference to the distinct stages in the literature presented, and its use in guidelines for treatment decision-making is limited.<sup>11</sup>

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Frozen shoulder is a clinical diagnosis based on history and examination, commonly presenting as a slow onset of pain near the deltoid insertion combined with a disproportionately severe loss of passive external rotation.<sup>1</sup> Routine radiographs for suspected frozen shoulder have been shown to have a low pick-up rate for other pathology (osteoarthritis or tumours).<sup>12</sup> However it has been shown that in patients aged >40 presenting with pain and stiffness, an underlying tumour can be misdiagnosed as a frozen shoulder, albeit with a low incidence.<sup>13</sup>

Frozen shoulder is a self-resolving pathology.<sup>11</sup> Therefore, treatment goals are centred around improving pain and function as quickly as possible, to the patient's acceptable level. This underlies the need for shared decision-making in treating frozen shoulders.

Two national guidelines for the treatment of frozen shoulder exist in the United Kingdom.<sup>1,11</sup> Guidelines from the British Elbow and Surgery Society (BESS) were last published in 2015,<sup>1</sup> before more recent multi-centred randomised controlled trials (RCTs) were published.<sup>14</sup> The National Institute for Health and Care Excellence (NICE) guidelines were updated in 2022,<sup>11</sup> but targeted primary care only, without discussing treatment delivered in secondary care. Therefore, the BESS Frozen Shoulder Working Group, therefore, agreed it is appropriate to produce an up-to-date review of treatment options for frozen shoulder using a modern guideline methodology for both primary and secondary care.

## Methods

UK-based shoulder surgeons, surgical trainees, specialist physiotherapists, primary care physicians and consultant radiologists were approached and voluntarily agreed to form the working group. Members were identified either as part of the BESS research committee or had expressed interest in reviewing literature in this field. The Working Group agreed on key questions to frame the management of frozen shoulder in the Population, Intervention, Control and Outcomes (PICO) format (Table 1). Based on these questions, a search performed on MEDLINE, EMBASE, and Cochrane Central Register of Controlled Trials for prospective comparative studies was performed. The literature search used the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA) methodology.<sup>15</sup> No limitations on language or publication dates were placed, although untranslatable text was later excluded. Case-control and non-comparative studies were excluded. Publications were limited to cohort studies, RCTs, systematic reviews and meta-analyses. The search data included all publications up until August 2023. Data was extracted to Covidence, where two authors performed primary abstract screening and secondary full-text screening. If there was a discrepancy, a third author reviewed the paper to assess for inclusion.

**Table 1.** A list of the PICO questions addressed in this guideline.

PICO review questions
1. Does in-person physiotherapy improve symptoms faster than home exercises/single session physiotherapy/natural history in patients with frozen shoulder?
2. Is in-person physiotherapy Following Percutaneous or Surgical Intervention Beneficial for People with Frozen Shoulders?
3. Do low-volume (< 20 mL) glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to natural history or physiotherapy?
4. Do high volume ( $\geq$ 20 mL) glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to natural history or physiotherapy
5. Do low volume ( $\leq$ 20 mL) freehand glenohumeral joint steroid injections expedite improvements in pain and function in FS compared US guided injection?
6. Is low-volume glenohumeral joint steroid injection beneficial compared to a low-volume placebo or local anaesthetic injection for a frozen shoulder?
7. Do high-volume (> 20 mL) glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to low-volume glenohumeral joint steroid injections?
8. Do high-volume (>20 mL) glenohumeral joint steroid injections expedite improvements in pain and function versus high-volume placebo (saline) injections or local anaesthetic-only injections?
9. Is MUA (and steroid injection) more effective than non-operative treatment in patients with frozen shoulders?
10. Is arthroscopic release more effective than non-operative treatment in patients with frozen shoulders?
11. Is arthroscopic release more effective than hydrodilatation in patients with frozen shoulder?

## Data extraction

According to the Grading of Recommendations, Assessment, Development and Evaluations (GRADE) methodology,<sup>16</sup> data relevant to each PICO were extracted to Summary of Findings tables based on pre-determined specific outcome measures. The data extracted included study size, outcome effect size and risk of bias assessment using the RoB-2 tool (Cochrane).<sup>17</sup>

## Statistical analysis

Where multiple studies examined identical outcomes, the mean difference was pooled, and random effect meta-analyses were performed, to minimise heterogeneity. If different outcome scores were used, a standardised mean difference was calculated. SPSS (M Corp. Released 2021. IBM SPSS Statistics for Macintosh, Version 28.0. Armonk, NY: IBM Corp) and Meta-Mar (Ashkan Beheshti, 2018, <https://www.meta-mar.com/>) were used for analyses. A narrative review with cohort studies was performed, where RCTS were absent or had incomplete data to answer a PICO.

## Quality assessment

Each outcome underwent a certainty assessment using the GRADE Evidence to Decision framework<sup>16</sup> to determine the confidence that each effect estimate reflected the actual effect. This assessment was based on (1) the risk of bias, (2) inconsistency, (3) imprecision, and (4) indirectness.

## Formulation of recommendations

Four key factors based on GRADE influenced the direction and strength of the recommendations:

- balance between desirable and undesirable outcomes,
- confidence in the effect of an intervention on essential outcomes,
- values and preferences of an intervention,
- resource allocation.

Recommendations were denoted as either ‘for’ or ‘against’, with the certainty being denoted as ‘strong’ or ‘weak’. Where insufficient evidence was available to guide a direction of recommendation, this was denoted as ‘neutral’. Recommendations were agreed upon by discussion within the multidisciplinary working group, to reach a unanimous consensus on the direction and strength of the recommendation based on the literature analysis.

## Summary of literature review

A total of 116 studies were eligible for this review. The PRISMA flowchart of the literature review conducted is shown in Figure 1.

The group agreed on summary statements based on each PICO (Table 2). Based on this evidence, a flowchart was created (Figure 2). A summary of the findings tables can be found in the Supplemental Material.

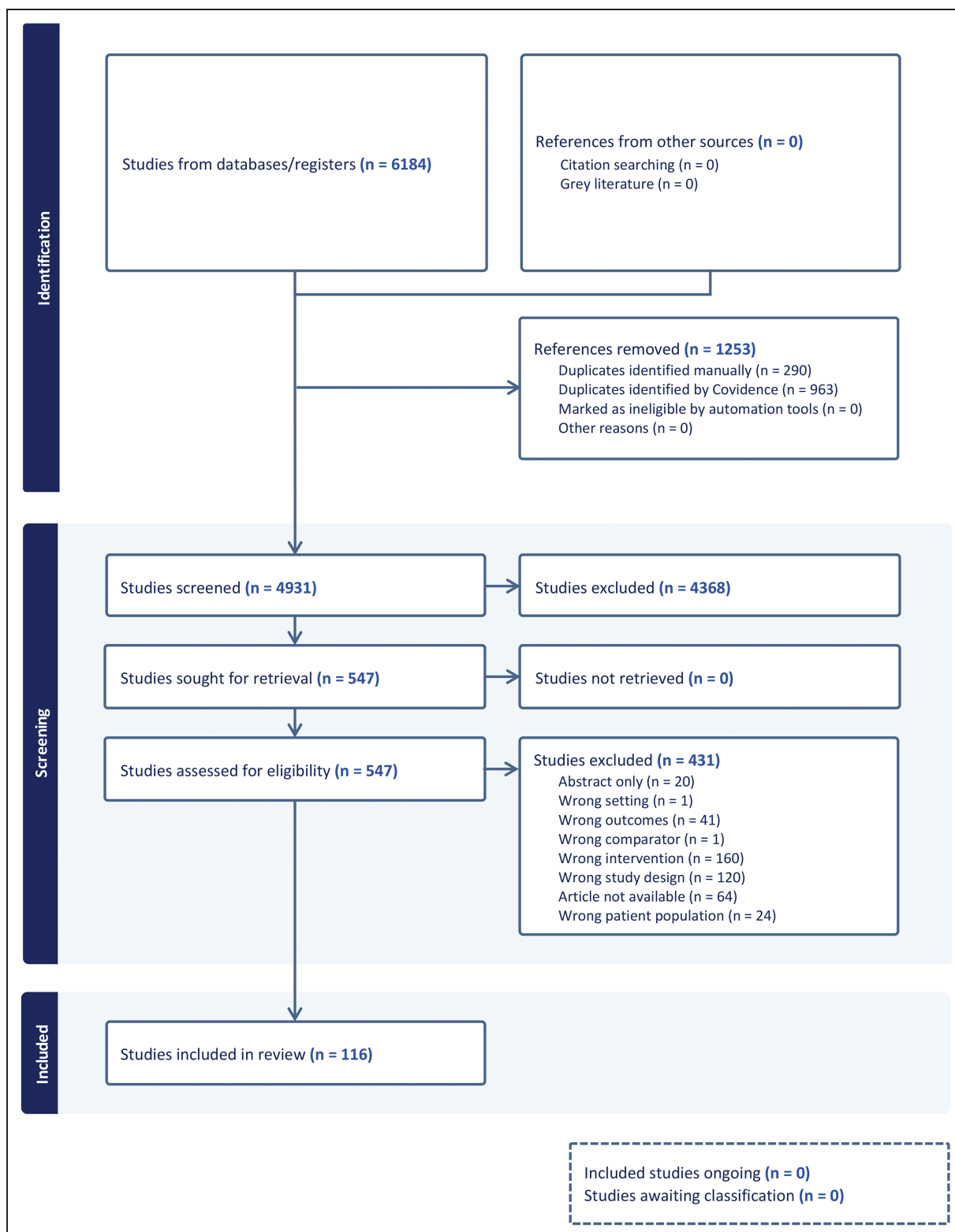
## Evidence-based on PICO questions

PICO 1: Does in-person physiotherapy improve symptoms faster than home exercises/single session physiotherapy/natural history in patients with frozen shoulder?

**Table 2.** Summary statements from the literature review.

Summary statement	Evidence
Glenohumeral steroid injections improve symptoms faster than natural history in the treatment of frozen shoulder.	Strong for
Low-volume (<20 mL) glenohumeral steroid injections can either be delivered using ultrasound guidance or using a free-hand technique by a trained healthcare professional.	Conditional for
Hydrodistension improves short-term pain and range of motion faster than natural history in the treatment of frozen shoulder.	Conditional for
Hydrodistension has better outcomes than low-volume steroid injections in the treatment of a frozen shoulder.	Conditional for
Physiotherapy should be offered following injection therapy.	Conditional for
MUA provides early improved functional outcomes compared to non-operative techniques (steroids and physiotherapy).	Conditional for
Physiotherapy alone improves function faster than natural history in frozen shoulder.	Neutral
In-person physiotherapy should be used over single-session physiotherapy or home-led exercises.	Neutral
Physiotherapy should be offered following surgical treatment.	Neutral
Hydrodistension improves long-term pain and function compared to natural history and physiotherapy.	Neutral
Arthroscopic capsular release provides improved immediate short-term (< 6 weeks) outcomes compared to non-operative techniques.	Neutral
Do high-volume (>20 mL) glenohumeral joint steroid injections expedite improvements in pain and function versus high-volume placebo (saline) injections or local anaesthetic-only injections?	Neutral
MUA provides improved long-term outcomes compared to non-operative techniques.	Conditional against
Arthroscopic capsular release provides improved long-term outcomes compared to non-operative techniques.	Conditional against





**Figure 1.** PRISMA flowchart of literature review performed.

There were no randomised trials comparing physiotherapy to natural history over time. Two studies compared physiotherapy to a placebo glenohumeral joint

injection.<sup>18,19</sup> However, these were excluded from this PICO as the working group determined that a placebo injection was not the same as natural history. A single RCT

compared hospital-based exercise classes to in-person multimodal physiotherapy and home exercises.<sup>20</sup> The study found faster recovery and greater improvement in pain and function with hospital-based exercise classes. However, the study was underpowered.

### Recommendations

It is unknown whether supervised physiotherapy provides any greater benefit than the natural history of a frozen shoulder. Recommendations cannot be made due to the lack of evidence. There is low certainty evidence that hospital-based group exercise classes may lead to a faster recovery and greater improvement in pain and function than in-person multimodal physiotherapy or home exercises.

### Suggestions for future research

There is a need for well-designed studies to:

- Compare physiotherapy to natural history for people with frozen shoulders.
- Compare in-person physiotherapy to home exercise/single-session physiotherapy/self-management for people with frozen shoulders.

PICO 2: Is in-person physiotherapy following percutaneous or surgical intervention beneficial for people with a frozen shoulder?

### Physiotherapy following corticosteroid injection

Three RCTs comparing physiotherapy to no physiotherapy following glenohumeral corticosteroid injection were meta-analysed.<sup>18,19,21</sup> There is low certainty that in-person physiotherapy following corticosteroid injection results in improvement in external rotation, pain and disability at 6 weeks after injection. There is no evidence for a difference at 6 months in pain and disability.

### Physiotherapy following glenohumeral joint distension

Four RCTs related to the role of physiotherapy following glenohumeral joint distension were identified but could not be meta-analysed due to the paucity of data, thus a narrative review was performed.<sup>22–25</sup> Buchbinder et al.<sup>22</sup> compared in-person physiotherapy to a sham intervention, finding no difference in pain, function, or quality of life, but improved shoulder range of movement at 6 weeks, sustained at 6 months. Kwak and Kim<sup>24</sup> compared in-person physiotherapy to home exercise, finding a greater short-term improvement in shoulder range of movement up to 6 weeks from in-person physiotherapy. However, there was

no difference between groups from 12 weeks up to one year. An RCT comparing physiotherapy to no physiotherapy showed improvements in both groups, but this was underpowered and lacked data analysis.<sup>23</sup> Robinson et al.<sup>25</sup> compared in-person physiotherapy to a self-directed home exercise program, finding no difference in primary or secondary outcomes at any time point up to one year. The trial was underpowered, so the conclusion should be interpreted with caution.

### Recommendations

- Current evidence, although low certainty, suggests that in-person physiotherapy should be considered following corticosteroid injection in the treatment of people with frozen shoulders, as it may provide a low-risk short-term benefit.
- The current evidence for in-person physiotherapy after glenohumeral distension to treat a frozen shoulder is limited. Although it may result in greater short-term improvement in the range of movement, it is unlikely to influence the long-term treatment outcome.

### Suggestions for future research

There is a need for well-designed studies to:

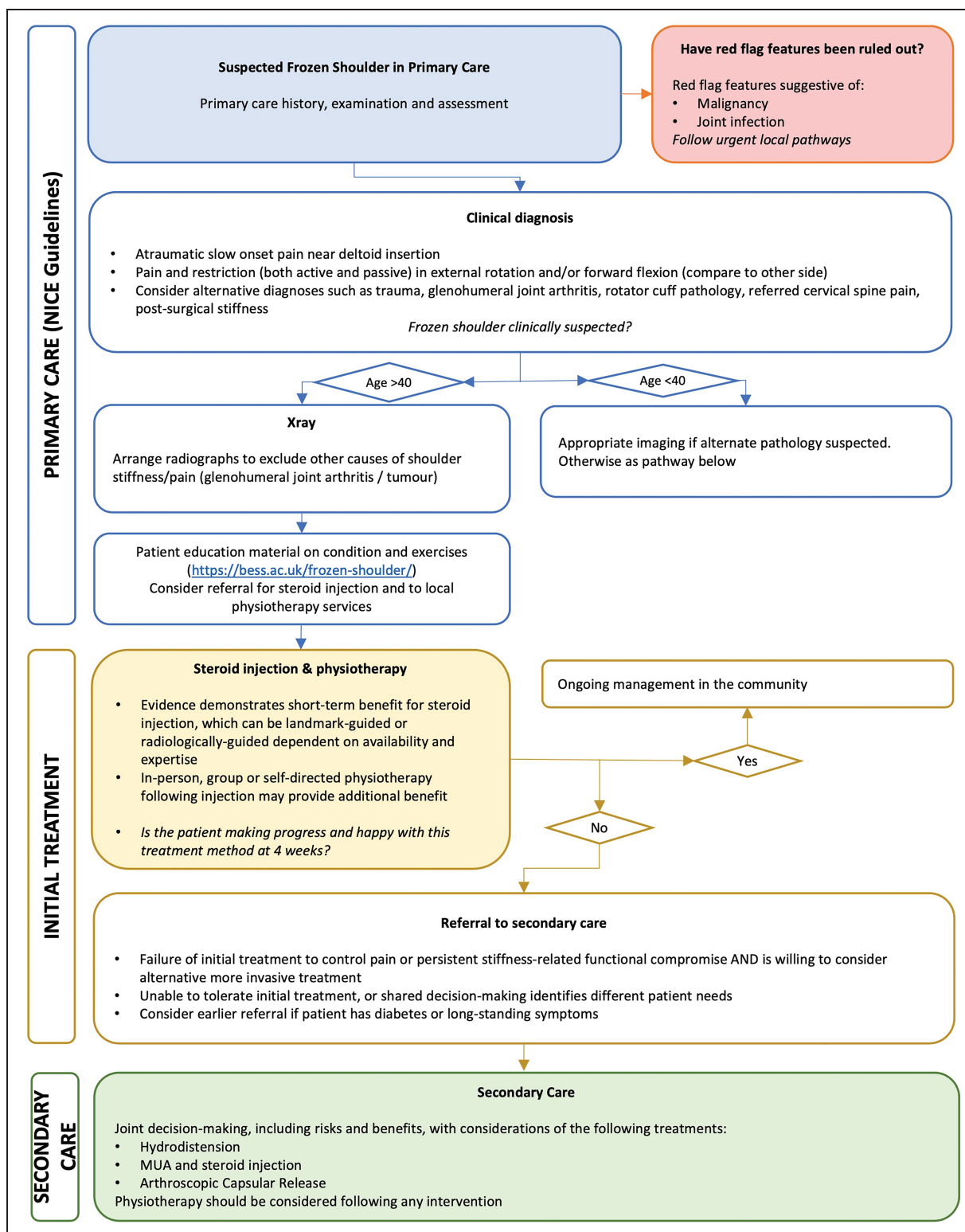
- Determine the effect of physiotherapy following glenohumeral joint injections (low and high volume).

PICO 3: Do low volume (< 20 mL) glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to natural history or physiotherapy

Seven RCTs were identified.<sup>26–33</sup> Although there was heterogeneity in the outcome measures used, the use of VAS scores and passive range of motion assessment demonstrated consistency. Meta-analyses at 6 and 12 months were not possible owing to limited studies assessing these longer-term outcomes.

The effect of steroid injections over physiotherapy alone demonstrated improvement in up to 3 months in pain scores and range of motion (forward flexion, abduction and external rotation). The threshold for this representing a clinically important difference is difficult to assess, but this data demonstrated consistency, directness, and no significant risk of bias. The meta-analysis performed was supported by a sufficient number of papers and participants. The short-term effect of steroids over physiotherapy on PROMs other than pain had limited evidence owing to the heterogeneity of outcomes.

There was no difference between patients who had low-volume steroid injections and those who had physiotherapy alone in pain, range of motion and PROMs in the long term after 3 months. Long-term outcomes were less studied limiting the strength of this evidence.



**Figure 2.** Suggested treated algorithm in from primary to secondary care.

### Recommendations

- Glenohumeral joint steroid injections should be considered as part of the non-operative treatment

pathway for primary frozen shoulder in both primary and secondary care for short-term symptom control.

- The long-term benefit of glenohumeral joint steroid injections after 3 months is not seen, although the strength of this evidence is limited.

### Suggestions for future research

There is a need for well-designed studies to:

- Determine the role of glenohumeral steroid injections in patients with early or late presentation of a frozen shoulder.
- Determine the role of glenohumeral steroid injections and alternative injections in patients with specific risk factors such as diabetes. PICO 4: Do high volume ( $\geq 20$  mL) glenohumeral joint steroid injections (also referred to as 'hydrodilatation' or 'hydrodistention') expedite improvements in pain and function in FS compared to natural history or physiotherapy?

Three randomised controlled studies and one non-randomised prospective comparative study.<sup>23,34–36</sup> Data is limited by a sparsity of RCTs to date investigating hydrodilatation versus physiotherapy or natural progression. The results from this meta-analysis found statistically significant early improvements in abduction following hydrodilatation at up to 3 months compared to conservative management ( $p = 0.0006$ , high certainty). The remaining analyses showed improvement in pain, disability and external rotation following hydrodilatation at up to 3 months (moderate certainty), however, these did not reach statistical significance with data limited by high heterogeneity. Meta-analysis was not possible to assess long-term outcomes.

Elleuch et al.<sup>35</sup> reported significant improvements in pain, function, and range of motion at 1 week with these benefits sustained up to 12 months in their prospective, non-randomised study. Sharma et al.<sup>36</sup> did not demonstrate any statistically significant difference at 12 months in their RCTs.

Transient flushing and after-pain were reported in 14% of patients following hydrodilatation.<sup>36</sup> There were no other adverse events secondary to hydrodilatation reported in any of the included studies.

### Recommendations

- There is evidence to support the use of hydrodilatation over physiotherapy or supportive therapy for frozen shoulder for short to medium-term improvement in pain and abduction range of motion.
- There is insufficient evidence to comment on the long-term outcomes of hydrodilatation versus physiotherapy or supportive therapy.

PICO 5: Is low-volume glenohumeral joint steroid injection beneficial compared to low-volume placebo or local anaesthetic injection for frozen shoulder?

Three studies were included in the synthesis of evidence and meta-analyses.<sup>37–39</sup> Saline injections were used as the control groups in two studies and local anaesthetic in the other study. A fourth RCT was available but provided outcomes as different from the baseline rather than an actual number for the outcome measures and thus was excluded.<sup>19</sup>

Range of motion (external rotation), a PROM and a VAS score were measured for all three RCTs. However, the time intervals differed, except for the 3-month post-interventional point. This time point was used for the meta-analysis across all three studies. The mean difference at 3 months for VAS and PROMS showed no difference between the interventions. The mean difference in external rotation did improve more with steroid injection. With moderate bias in study designs, a moderate level of caution is required in drawing conclusions from this. Consideration should be taken with these findings, as they demonstrate inconsistency when compared to PICO 3 and 6, both of which showed evidence for short-term improvement with steroid injections.

No serious adverse events were reported in two studies,<sup>39,40</sup> and serious adverse events were not mentioned in the third article.<sup>37</sup>

### Recommendations

A meta-analysis found an improvement in external rotation at 3 months with steroid injection compared to placebo or saline, but no difference with VAS or PROMS outcomes. Analysis for other time points was not possible due to the heterogeneity of time points used in the studies.

Whilst meta-analysis of available placebo-controlled trials showed limited differences between steroid and placebo injections at 3 months, this finding should be interpreted cautiously given:

- The small number of studies available for this specific comparison.
- The conflicting evidence from PICO 3 shows the benefit of physiotherapy alone.
- The possibility is that even placebo injections may have therapeutic benefits through capsular distension.

PICO 6: Do low-volume ( $\leq 20$  mL) freehand glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to US-guided low-volume injections?

Four randomised controlled studies were identified.<sup>41–44</sup> Meta-analyses were performed accounting for unstandardised

mean differences. As data was presented differently between papers (two studies presenting absolute changes as opposed to mean score), it was not possible to pool all the data. Lee et al.<sup>41</sup> showed that improvements in pain and ROM after 1 and 4 weeks were more prominent in the US-guided group, but the differences were not statistically significant, except for the changes in extension where the improvements were significantly higher in the US-guided group ( $p=0.01$ ). Raeissadat et al.<sup>43</sup> showed statistically significant improvement in flexion in the US-guided group compared to the landmark-guided group for the first 3 weeks ( $p=0.039$  in the first week,  $p=0.001$  in the second week and  $p=0.025$  in the third week). However, there were no significant differences between the two groups from the fourth week until the end of the study. It was not possible to pool the function data from the two studies as one study<sup>41</sup> presented data as a general function of the shoulder and the second study<sup>43</sup> presented Oxford shoulder score but no difference was identified between the two groups.

Pooled data demonstrate that there is no difference in range of motion, pain, or function between ultrasound-guided or blinded glenohumeral joint injections. Lower-quality evidence suggests that ultrasound-guided injection may result in slightly reduced pain in the first week but certainty is low. The evidence is limited by a lack of high-quality studies and the lack of studies using validated functional outcome measures.

## Recommendations

The available evidence suggests that low-volume steroid injections, however, delivered, are associated with pain relief and improved range of movement (see PICO 3) but there is no difference in clinical outcome if this injection is delivered ultrasound-guided or freehand.

**PICO 7:** Do high-volume ( $\geq 20$  mL) glenohumeral joint steroid injections expedite improvements in pain and function in FS compared to low-volume glenohumeral joint steroid injections?

The authors identified 13 RCTs addressing this subject, embedded within two contemporary systematic reviews with meta-analysis. Saltychev et al. included six studies assessing high-volume injections with steroids to low low-volume injections with steroids.<sup>32,40,45–49</sup> Their inclusion criteria included stiffness of any cause, including osteoarthritis, and the meta-analysis combined all follow-up time-points. For these reasons the data was not used to assess the treatment effect, however, their comprehensive risk of bias review of individual studies informed the data certainty assessment.

Poku et al. included eight RCT studies of high-volume injections with steroids to low-volume injections with steroids.<sup>50–52</sup> Follow-up was quantified as early ( $< 6$  weeks)

and late ( $> 6$  weeks). All studies had some degree of blinding so the risk of bias was low. Some of the studies were underpowered. There was variability with treatment regimes (one vs. three injections), follow-up period and outcome measures. The benefit of high volume was an improvement in pain and disability in the initial outcomes which was not seen at the end of the study period. However, overall, an improvement with external rotation was seen with high volume which was maintained at the final follow-up. Meta-analysis of longer-term follow-up (12 months) was not possible as most studies did not run beyond 6 months.

## Recommendations

The available evidence suggests that high volume ( $\geq 20$  mL) glenohumeral joint steroid injection may provide an improvement in shoulder disability and pain (quantified by patient-reported outcomes), and external rotation when compared to low volume ( $\leq 20$  mL) glenohumeral joint steroid injection in the short term for the treatment of frozen shoulder.

## Suggestions for future research

There is a need for well-designed studies to determine the benefit of hydrodistension in the management of frozen shoulder, versus low-volume steroid injection, in terms of clinical and cost-effectiveness.

**PICO 8:** Do high-volume ( $\geq 20$  mL) glenohumeral joint steroid injections expedite improvements in pain and function versus high-volume placebo (saline) injections or local anaesthetic-only injections?

There were no studies that fulfilled the criteria to assess if the high-volume glenohumeral joint steroid injections expedite improvements in pain and function versus high-volume placebo (saline) injections or local anaesthetic-only injections. A recent systematic review has highlighted the potential value of non-steroid injections in the treatment of frozen shoulder and these may be of particular relevance to patients more susceptible to the side effects of steroid injections (e.g. diabetics).<sup>53</sup>

## Recommendations

Due to a lack of RCTs meeting the criteria and no eligible papers for review, no specific recommendations can be provided based on the available evidence.

## Suggestions for future research

There is a need for a well-designed study to assess the role of steroids, and their alternative, within high-volume glenohumeral injections.



**PICO 9:** Is MUA (and steroid injection) more effective than non-operative treatment in patients with frozen shoulder?

Three randomised controlled studies<sup>14,54,55</sup> and one meta-analysis<sup>56</sup> were identified addressing this subject. The meta-analysis incorporated all three RCTs identified independently and a decision was made to use this meta-analysis of current best evidence.

The authors of the meta-analysis<sup>56</sup> perceived the Jacobs study to incorporate a hydrodilatation as part of the steroid injection and is not part of the specific question for this particular PICO.<sup>54</sup> They also assessed the bias of the papers included with UK FROST having a low risk of bias, except for performance and detection bias.<sup>14</sup> Kivimäki et al.<sup>55</sup> had slightly more concern with performance, detection, attrition, and possible other bias.

No serious adverse events were reported in the Kivimäki paper.<sup>55</sup> UK FROST reported two serious adverse events (1%) in the MUA group, including one septic arthritis (0.5%) and one accident and emergency attendance.<sup>14</sup>

This appears to be a good-quality meta-analysis of the available literature. They have performed a risk of bias assessment on both papers with UK FROST appearing to be the most robust data. Both groups demonstrated improvement in OSS and pain scores, but the mean difference for all presented outcome measures showed no difference between the interventions. It should be noted, that there was a 15% crossover in the non-operative group, of which a third had a subsequent MUA.

A further economic analysis as part of UK FROST, highlighted that despite MUA being more expensive, it was more likely to represent higher cost-effectiveness per quality-adjusted life year than physiotherapy alone.

## Recommendations

We recommend that MUA should be considered as a viable treatment option, particularly when considering its cost-effectiveness. The choice between MUA and conservative treatment should be based on shared decision-making, taking into account patient preference, local waiting times for physiotherapy versus surgical intervention, and the understanding that while final outcomes may be similar, the pathway to achieving these outcomes may differ. Further research is needed to clarify the role of MUA in achieving earlier symptomatic improvement.

**PICO 10:** Is arthroscopic release more effective than non-operative treatment in patients with FS?

Four studies were included in the synthesis of evidence.<sup>14,57–59</sup> Physiotherapy and steroid injection were each used as the control groups in two studies. A fifth RCT was available but had no numerical data available and thus was excluded.<sup>60</sup> Due to missing data and

heterogeneity in outcome use and time intervals, pooled estimates were not produced. A narrative assessment of the treatment effect was therefore undertaken.

No adverse events were reported in either group in either the Smiteman or De Carli paper, but the serious adverse events rate in UK FROST was 4% (ACR) versus 0% (physiotherapy).

Time intervals for the range of motion data differed. The mean difference at 3 and 12 months for forward flexion, abduction and external rotation was higher in arthroscopic capsular release than either steroid or physiotherapy. With moderate bias in study designs, effects from single studies and a lack of confidence interval, a high level of caution is required in drawing conclusions from this.

Within the RCTs available, there was heterogeneity in the outcome measures studied. Of the trials identified, three were felt to be at a moderate risk of bias. UK FROST was the only appropriately powered RCT and was deemed to be of low risk of bias.<sup>14</sup> The results from this study found statistically poorer 3-month outcomes for ACR compared to physiotherapy as quantified by the Oxford Shoulder Score. At 6 months, no difference was observed but at 12 months a statistical difference in favour of ACR was observed. Importantly, neither the 3- or 12-month outcomes exceeded the minimally clinically important difference. Analysis was 'per randomisation', with 15% of the physiotherapy group requiring further treatment. Outcomes based on the chronicity of symptoms were not assessed.

The remaining studies provided limited additional evidence. Although mean difference scores were calculated which demonstrated consistency with improvement in pain scores and PROMs at 12 months, also demonstrated by Zhu et al.,<sup>61</sup> the lack of comprehensive reporting of effect sizes, confidence intervals and moderate risk of bias means that these results should be treated with caution. Meta-analyses, if possible is likely to demonstrate heavy weighting towards UK FROST.

No studies identified the effect of the intervention or control in the short term (< 1 week). The impact of diabetes or refractory frozen shoulder could not be assessed as no studies performed additional subgroup analysis.

## Recommendations

- Shared decision making with current best evidence should be undertaken prior to arthroscopic capsular release, as outcomes do not suggest a superior minimum clinical importance improvement at 12 months, compared to alternative treatment.
- Current research has not addressed confounders such as acute versus chronic frozen shoulder, diabetic status, or the presence of concomitant pathology.

**PICO 11:** Is arthroscopic release more effective than hydrodilatation in patients with FS?

A total of 1 study was identified.<sup>62</sup> There is insufficient evidence at present to determine whether ACR is more effective than HD in the treatment of frozen shoulder. It should be noted that the risk profile between both interventions has not been directly compared in a single study, but the serious adverse events rate for ACR was 4% in UKFROST,<sup>14</sup> whereas studies included in PICO 4 and 7 showed a low serious risk profile for HD.

## Recommendations

Patients should be aware that there is insufficient evidence at present to recommend one treatment over another and the risks and benefits of each.

## Future research

Initially, determine the optimum method of hydrodilatation; volume, location of injection, method of delivery, dose of steroids, role of rupturing capsule, complications, recurrence rates, and costs. Following this, an adequately powered RCT to include the stage of disease, duration of symptoms, and diabetes status. There is a need for well-designed studies to determine the benefit of arthroscopic release over hydrodistension in the management of frozen shoulder in terms of clinical and cost-effectiveness.

## Discussion

The recommendations made within this article are based on the available evidence and through the conduct of the literature review it has become clear that there are notable deficiencies in the research which underpins the treatment of this very common, and often disabling condition.

The role of glenohumeral corticosteroid injections presents an interesting paradox in our findings. Whilst PICO 3 demonstrated a clear benefit of steroid injection over physiotherapy alone in the short term, PICO 5 found limited differences between steroid and placebo injections at 3 months. This apparent contradiction may be explained by several factors. Firstly, the mechanical effect of any injection (steroid or placebo) may itself be therapeutic through capsular distension. Secondly, the placebo-controlled trials were fewer in number and generally smaller than those comparing injection to physiotherapy. Additionally, PICO 6 demonstrated that the method of delivery (ultrasound-guided vs. freehand) did not significantly impact outcomes, suggesting that the act of injection itself may be important. These findings, when taken together, support the use of glenohumeral injection as a therapeutic intervention, whilst highlighting the complexity of determining the relative contributions of mechanical distension versus pharmacological effect.

Discussions about optimal management extend further than the work presented here, with questions raised about

the optimal contents of any injectate used in the treatment of frozen shoulder and this question has been addressed in a recent systematic review.<sup>53</sup>

The recommendations in these guidelines are based on a systematic review of the available evidence, utilising the GRADE methodology to assess the quality and strength of evidence. While we acknowledge significant practical challenges in healthcare delivery, including variable waiting times for different treatment modalities, these guidelines intentionally focus on clinical effectiveness rather than service delivery constraints. This approach ensures the recommendations remain applicable across different healthcare settings and timeframes. The pragmatic implementation of these guidelines, including considerations of local resource availability and access times, must be determined by local care teams. Only UK FROST specifically incorporated service delivery considerations into its methodology, finding that despite being more expensive, MUA was more likely to be cost-effective per quality-adjusted life year than physiotherapy alone. However, the interpretation of timing-dependent outcomes from this and other studies could support different treatment approaches depending on local service delivery capabilities and costs per treatment in different settings. These guidelines are to present the evidence for clinical effectiveness, allowing healthcare providers to develop local protocols that optimise patient care within their specific service delivery constraints. Future research would benefit from more explicit consideration of how treatment timing and access affect outcomes,<sup>63</sup> as this could help inform both clinical recommendations and service delivery planning.

The work presented here aims to deliver an evidence-based algorithm treatment that is applicable 'today' whilst highlighting the strength and fragility of some of the data on which these recommendations are based on, and importantly highlighting key areas for future research.

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## Category of paper

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## Disclaimer

The information set out in this publication is intended for use as a guide of a general nature and may not be relevant to a particular patient's circumstances. This publication is not exhaustive of the subject matter and when implementing any recommendations contained in this publication, please exercise your own independent judgement or seek appropriate professional advice relevant to your own particular circumstances. These recommendations cannot guarantee discharge of the duty of care owed to patients. The text is directed to health professionals possessing appropriate qualifications and skills in ascertaining and discharging their professional (including legal) duties. It is not to be regarded as clinical advice and is no substitute for a full examination and consideration of medical history in reaching a diagnosis and treatment.

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