

Radiofrequency vs Steroid Injections for Spinal Facet and Sacroiliac Joint Pain: A Systematic Review and Meta-Analysis

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Purpose: Pain management for spinal facet joint (SFJ) and sacroiliac joint (SIJ) pain is challenging, often requiring interventions like radiofrequency ablation (RFA) or corticosteroid injections (CI). This study aims to assess and compare the effectiveness of CI and RFA in treating SFJ and SIJ pain. We combine these treatments due to their shared pathophysiology, similar therapeutic interventions, and the necessity for an integrated approach to spinal pain management.

Patients and methods: Literature search from PubMed, Scopus, CENTRAL and Google Scholar for published studies upto 31st December 2023, and reporting data of patients who were treated using CI or RFA for SFJ and SIJ pain. Pooled standardized mean difference (SMD) with a 95% Confidence Interval (CI) was calculated.

Results: Our meta-analysis incorporated thirteen studies. Overall, patients, treated with CI had a higher pain intensity score compared to patients treated with RFA (SMD=0.92; 95% CI: 0.19 to 1.65) at 3 months, and at 6 months (SMD=1.53; 95% CI: 0.66 to 2.40) after the treatment. No significant association was reported at 12 months (SMD=1.47; 95% CI: -0.03 to 2.97). Subgroup analysis based on joint types revealed increased pain intensity scores in patients who were treated with CI for SIJ (SMD=1.25; 95% CI: 0.39 to 2.11) and SFJ (SMD=1.33; 95% CI: 0.09 to 2.57) pain. A negative but not significant effect was detected in patients, treated with CI for cervical joint pain (SMD=-0.40; 95% CI: -0.90 to 0.10). Patients treated with CI exhibited higher functional disability score compared to patients treated with RFA at 3 months (SMD=1.28; 95% CI: 0.20 to 2.35) post-treatment.

Conclusion: This study suggests that RFA may offer superior pain relief with longer duration compared to steroid injections for spinal facet and sacroiliac joint pain. Decision regarding specific interventions should be individualized and consider patient preferences, clinical context, and potential risks.

Keywords: radiofrequency ablation, steroid injections, spinal facet pain, sacroiliac joint pain, systematic review, meta-analysis, pain management

Introduction

Pain management for spinal facet joint (SFJ) and sacroiliac joint (SIJ) pain is challenging, and interventions, aimed to alleviate discomfort and improve patients' quality of life are crucial.^{1,2}

Over 25% of individuals with chronic lower back pain are reported to have SFJ, or more specifically, lumbar facet joint pain that is predominantly linked to degenerative osteoarthritis.^{3,4} SIJ pain accounts for up to 20% of low back pain cases, and poses its own set of challenges.⁵ Valid clinical and diagnostic tests for SIJ pain remain elusive, leading to diverse diagnostic approaches.⁶

Both SFJ and SIJ pain result from similar underlying mechanisms, including joint degeneration, inflammation, and nerve irritation, suggesting that therapies targeting one condition may also be effective for the other. An integrated

approach is essential for effectively managing spinal pain, given the interconnected nature of spinal structures and the common occurrence of multi-faceted pain syndromes. Although the SIJ and facet joints differ in size and function—the SIJ being larger and primarily responsible for load transfer between the spine and lower limbs, while facet joints facilitate spinal movement and stability—there may be overlapping pain mechanisms and treatment responses. Investigating these conditions concurrently allows for the evaluation of shared diagnostic tools and therapeutic approaches, potentially leading to more efficient and effective management strategies. Additionally, comparing these distinct but related pain sources can highlight similarities and differences in their pathophysiology, enhancing the precision of interventions tailored to each condition.

Recently, radiofrequency ablation (RFA) and corticosteroid injections (CI) have emerged as prominent therapeutic modalities in pain management of such patients.^{7–9} The primary interventions for both conditions, RFA and CI, work by reducing inflammation and disrupting pain signals, supporting a combined therapeutic strategy. Injection with corticosteroids have shown promise in managing SFJ-related lower back pain.¹⁰ Additionally, the application of RFA in patients with SFJ pain has demonstrated effectiveness in reducing pain scores and enhancing overall quality of life.^{11,12} Current treatment options for patients with SIJ pain include pharmacotherapy, chiropractic manipulation, CI injections, and surgical fixation.¹³ Recently, RFA has gained popularity for SIJ pain management, with studies reporting promising results.^{14,15}

However, due to variability in methodologies, including differences in techniques and selection criteria, the comparative effectiveness of RFA and CI in pain management of patients with SFJ and SIJ problem is still unclear. Given the significant impact of chronic spinal pain on quality of life, physical function, and healthcare resources, identifying the optimal treatment modality is essential. Insurance coverage for these procedures often varies, with some insurers being more inclined to cover steroid injections over RFA or vice versa. This variability can affect patients' access to the most effective treatments. By providing a systematic comparison of the efficacy of these interventions, our study offers valuable insights that can inform insurance policy decisions. This meta-analysis aims to comparing the efficiency of CI and RFA in managing pain associated with SFJ and SIJ. Our results may provide valuable insights into the optimal approach for pain management of these conditions, guiding clinicians in making informed decisions for the benefit of their patients, reducing reliance on opioids, and enhancing overall management strategies for spinal pain.

Methods

Protocol Registration

We adhered to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA)¹⁶ guidelines. The study is registered with PROSPERO with the identifier no. **CRD42024496067**.

Search Strategy

An electronic search was conducted on PubMed, Scopus, CENTRAL (Cochrane Central Register of Controlled Trials), and Google Scholar databases for English language papers published up to 31st December 2023. The search key terms included variations of “radiofrequency ablation”, such as “RF ablation” and “Radiofrequency”, and terms related to steroid injections, such as “Steroid injections” and “Corticosteroid injections”. The focus extended to spinal facet joint pain, represented by terms like “Spinal facet joint pain” and “Facet joint syndrome”, as well as sacroiliac joint pain, identified by terms like “Sacroiliac joint pain” and “Sacroiliitis”. Additionally, we included terms associated with osteoarthritis, such as “osteoarthritis”, “degenerative joint disease”, and “OA”, and terms related to axial pain, like “axial pain”, “axial back pain”, and “axial spine pain”. Our focus also extended to terms representing spinal facet joint pain, including “spinal facet joint pain” and “facet joint syndrome”, and sacroiliac joint pain, identified by terms like “sacroiliac joint pain” and “sacroiliitis”. Bibliography of retrieved studies and relevant reviews were further screened for the potential inclusion of additional studies.

Inclusion Criteria (PICOS)

Population (P): Patients with spinal facet joint (SFJ) or sacroiliac joint (SIJ)

Intervention (I): Radiofrequency ablation

Comparison (C): Corticosteroid injections
Outcome (O): Pain intensity
Study Design (S): Randomized controlled trials (RCTs).

Exclusion Criteria

1. Non-English or non-human studies
2. Studies lacking adequate data or with short follow-up periods
3. Mixed interventions, inappropriate comparators, or high risk of bias
4. Irrelevant conditions or duplicate data, published beyond a predefined date.

Data Extraction

During the initial phase of our systematic review, databases were searched, and titles and abstracts of identified studies were independently assessed by the two authors. Subsequently, full texts of papers meeting eligibility criteria were obtained for further analysis. The comprehensive data extraction process included key details such as author information, publication year, age distribution within the studied population, sample size, intervention specifics, pain intensity scores, functional disability scores, and any reported adverse events.

Outcomes

The primary outcome, pain intensity, was evaluated by visual analogue scale (VAS) or numerical rating scale (NRS). Pain intensity measurements were categorized into three distinct follow-up periods (3, 6, and 12 months after the procedure). Functional disability was measured using the Oswestry Disability Index (ODI) at 3 months after the treatment.

Quality Assessment

We used Grading of Recommendations, Assessment, Development, and Evaluations (GRADE), and specifically, GRADEpro tool, to assess evidence certainty for the outcomes. This tool considers factors such as study design, risk of bias, inconsistency, indirectness, imprecision, and publication bias.

Publication Bias

Publication bias was assessed by funnel plot analysis¹⁷ and Egger's regression test.¹⁸

Statistical Analysis

All statistical analyses were done by STATA, version 12.0 (Stata Statistical Software, Release 12; StataCorp LP, College Station, TX) and Review Manager (RevMan v5.3 2014; Cochrane, London, England).

Results were presented as standardized mean difference (SMD) with 95% confidence interval (CI) for pain intensity and functional disability. Heterogeneity was assessed by the I^2 statistic. In cases of significant ($I^2 > 50\%$) heterogeneity, random effects model was used. For $I^2 < 50\%$, fixed-effect model was used. Sensitivity analyses were done to validate the pooled observed effect. $P < 0.05$ was statistically significant.

Results

The PRISMA flow diagram of our study (Figure 1) illustrates the systematic process of identifying and selecting relevant studies. Our initial systematic search across various databases yielded a total of 117 studies. After removing duplicate entries, we were left with 46 unique articles. These full texts were then meticulously screened for eligibility based on our inclusion and exclusion criteria. During this screening process, 31 studies were deemed irrelevant as they did not meet our criteria, and 2 studies were excluded because they did not report outcomes of interest. Finally, 13 studies^{19–31} were incorporated in the analysis.

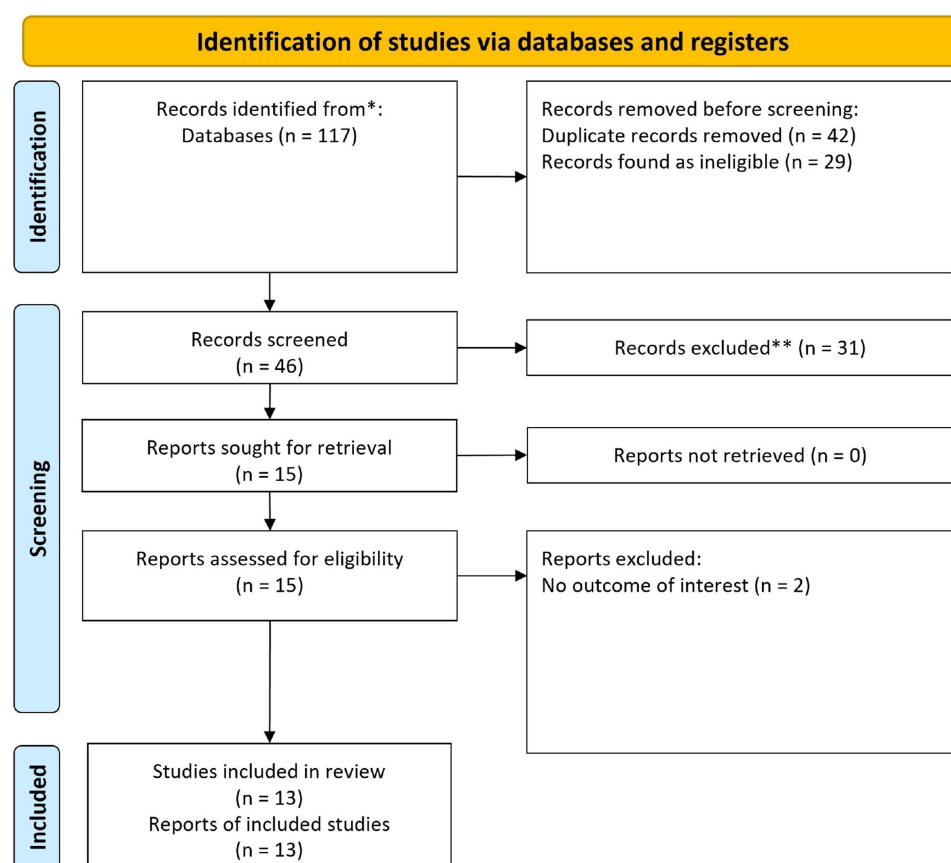


Figure 1 PRISMA 2020 Flow diagram for the selection of studies and specific reasons for exclusion from the present meta-analysis.

Note: *Databases: PubMed, Scopus, CENTRAL and Google Scholar; **31 studies were deemed irrelevant as they did not meet our criteria; irrelevant populations (n=14), different interventions (n=5), lacking a comparator group (n=7), and non-randomized design (n=5).

Characteristics of Studies

As shown in Table 1, eight out of 13 studies focused on patients with lumbar pain, two on cervical pain, and three on sacroiliac joint pain. The publication years of the studies ranged from 2012 to 2023, with sample sizes varying between 11 and 50 participants, and follow-up durations spanning from 6 to 12 months. Six studies were conducted in Asian populations, while seven were carried out in Caucasian populations.

All studies incorporated combined regimen of corticosteroid and local anesthetics for the corticosteroid groups. Methylprednisolone was used in six trials, typically in dosages ranging from 20 mg to 40 mg and often combined with varying amounts of bupivacaine. Betamethasone, used in three trials, had dosages ranging from 3 mg to 12 mg, while dexamethasone, used in two trials, was consistently dosed at 10 mg. Additionally, triamcinolone was used at 10 mg. These steroids were frequently combined with local anesthetics such as bupivacaine or lidocaine to enhance their effectiveness. Eight trials utilized pulsed radiofrequency (PRF), and five trials applied continuous RF (CRF). Adverse events were not reported in seven trials. A study by Lim et al²⁵ reported an increase in lower back pain (LBP) for two patients during the follow-up period, while another study by Do et al¹⁹ reported hyperglycemia in one patient.

Main Findings

Pain Intensity at 3 Months

Nine studies reported data for pain intensity score at 3 months (Figure 2a). CI correlated with higher pain intensity score compared to RFA (SMD=0.92; 95% CI: 0.19 to 1.65), with substantial heterogeneity ($I^2 = 93\%$). Subgroup analysis based on type of joints showed that CI correlated with higher pain intensity score compared to RFA in patients with sacroiliac region (SMD=1.25; 95% CI: 0.39 to 2.11) and lumbar region (SMD=1.33; 95% CI: 0.09 to 2.57) pain.

Table 1 Characteristics of Included Studies Investigating for Radiofrequency vs Steroid Injections for Spinal Facet and Sacroiliac Joint Pain

S. No	Study	Country	Joint Type	Inclusion Criteria	Sample Size		Male/Female		Procedures		Adverse Events		Mean Age \pm SD		Measured Scale	Imaging Modality	Follow up Duration (months)
					CI	RFA	CI	RFA	CI	RFA	CI	RFA	CI	RFA			
1.	Civelek et al, 2012 ²⁰	Turkey	Lumbar	Failure to respond 6 wks after therapy	50	50	11/39	17/33	MBB mixture of MPS40 mg + 2 mL of bupivacaine 0.25–0.50%	MBB CRF 80°C in 120 sec	None	Neuropathy in 2 subjects, Minor burns in few subjects;	56.5 \pm 17.7	51.8 \pm 17	VNS, NASS and EQ-5D	Fluoroscopy	12
2.	Duger et al, 2012 ²¹	Turkey	Lumbar	\geq 6 months history of LBP	40	40	-	-	IA 5 mL mixture of MPS20 mg + 5 mg bupivacaine	PRF at 40°C in 360 sec	-	-	-	-	VAS	-	-
3.	Lakemeier et al, 2013 ²⁴	Germany	Lumbar	\geq 24 months history of non-specific LBP	26	26	16/10	17/9	IA mixture of 1 mL of betamethasone 3mg + 0.5 mL of bupivacaine 0.5%	MBB CRF at 80°C in 90 sec	None	None	56.3 \pm 10.8	57.6 \pm 12.8	RMQ,VAS, ODI	Fluoroscopy	6
4.	Hashemi et al, 2014 ²³	Iran	Lumbar	\geq 6 months history of LBP	40	40	28/12	29/11	MBB using MPS 40 mg+ 0.5 mL bupivacaine 0.5%	MBB PRF at 42°C in 120 sec (2 shots of 45 V/s)	NR	NR	63.85 \pm 11.46	64.32 \pm 13.2	NRS,ODI	Fluoroscopy	12
5.	Martinez et al, 2016 ³²	Spain	Sacroiliac	Intense SIJ > 3 months	20	20	6/14	5/15	2 USG guided IA SIJ blocks with 3 mL 5% levobupivacaine and 12 mg betamethasone sodium, with a 7-day interval	Six 10 cm,10 mm active tip, 22G cannula and 4 electrodes were used to create 2 overlapping bipolar lesions with 4 cannula (90°C,3 min).	None	Moderate Pain	52.6 \pm 14.1	53.2 \pm 13.9	VAS	USG	12
6.	Zhou et al, 2016 ³¹	China	Lumbar	\geq 6 months history of LBP	40	40	23/17	21/19	MBB and IA injection using 5 mL mixture of betamethasone 6 mg + lidocaine 20 mg	MBB CRF at 80°C in 90 sec	None	None	54.6 \pm 7.5	56.5 \pm 8.7	VAS. Schober index	X-ray	6
7.	Shin et al, 2017 ²⁹	Korea	Cervical	Complaint of suboccipital neck pain predominantly for at least 6 months	11	12	3/8	4/8	2% lidocaine, 0.75 mL + Triamcinolone 10 mg, 0.25 mL	PRF 5 Hz and 5 ms pulse width for 360 sec at 55 V	None	None	47.6 \pm 6.6	47.6 \pm 6.8	NRS	Fluoroscopy	6
8.	Do et al, 2017 ¹⁹	Korea	Lumbar	6 months history of LBP	30	30	12/18	12/18	IA 0.5 mL mixture of dexamethasone 10 mg + 0.25 mL of bupivacaine 0.125%	IA PRF at 42°C in 360 sec (5 Hz, a5-ms pulsed width at 55 V)	Hyperglycemia in 1 subject	None	63 \pm 10.9	66.9 \pm 9.6	NRS	Fluoroscopy	6

(Continued)

Table I (Continued).

S. No	Study	Country	Joint Type	Inclusion Criteria	Sample Size		Male/Female		Procedures		Adverse Events		Mean Age \pm SD		Measured Scale	Imaging Modality	Follow up Duration (months)
					CI	RFA	CI	RFA	CI	RFA	CI	RFA	CI	RFA			
9.	Yasar et al, 2018 ³⁰	Turkey	Lumbar	≥ 3 months history of LBP	50	50	34/16	40/10	2.5 mL mixture of MPS40 mg + bupivacaine 0.25%	PRF at 42°C in 120sec (2 shots of 45 V/s)	NR	NR	43.1 \pm 8.3	47.4 \pm 11.1	VAS, ODI	x-ray	12
10.	Dutta et al, 2018 ²²	India	Sacroiliac	LBP ≥ 3 months	15	15	10/5	4/11	3 mL solution containing 2 mL of bupivacaine 0.5% and 1 mL of 40 mg/mL DEPO-MEDROL	PRF parameters were 45 V for 180 s at all levels using the RF generator. Temperatures varied from 38°C to 42°C.	None	None	41.6 (27–60)	43.1 (19–76)	NRS, ODI	Fluoroscopy	6
11.	Salman et al, 2016 ²⁸	Egypt	Sacroiliac	LBP for 6 months	15	15	7/8	5/10	3 mL solution containing 2 mL of Lidocaine 2%- and 1-mL bupivacaine 0.5%	RF probes were sequentially inserted into the cannula and 90s 80 °C lesions were performed	NR	NR	51.8 \pm 13.1	51.9 \pm 13.6	VAS	Fluoroscopy	6
12.	Lim et al, 2017 ²⁵	Korea	Cervical	3-month history of axial cervical pain without radicular symptoms	20	20	10/10	7/13	0.3 mL of contrast into CFJ space, injected 10 mg (0.25 mL) of dexamethasone, mixed with 0.25 mL of 0.125% bupivacaine	5 Hz and a 5-ms pulsed width, for 360 seconds, at 55 V, under the condition that the electrode tip temperature did not exceed 42°C.	Minor in 2 patients	None	52.7 \pm 12.1	52.8 \pm 12.7	NRS	Fluoroscopy	6
13.	McCormick et al, 2023 ²⁷	USA	Lumbar	Unilateral or bilateral axial (non-radicular) LBP for 3 months	12	20	4/8	10/10	0.5 mL of 40 mg/mL Kenalog and 0.5mL of 2% preservative-free lidocaine	C-RFA lesions performed for 165 seconds, with the RFA generator temperature set to 60 °C (intralesional temperature >80 °C)	None	None	62.4 \pm 14.6	62.2 \pm 12.4	NPRS	Fluoroscopy	12

Abbreviations: CI, corticosteroid injection; CRF, continuous radiofrequency; IA, intraarticular; LBP, Low back pain; LFJ, lumbar facet joint; MBB, medial branch block; NRS, numeric rating scale; VAS, visual analog scale, ODI, Oswestry Disability Index; RMQ, Roland, Morris Low Back Pain and Disability Questionnaire; EQ-5D, EuroQol, 5 Dimension; PRF, pulsed radiofrequency; SFJ, spinal facet joint; SIJ, sacroiliac joint; IA, intraarticular; MBB, medial branch block; NR, Not Reported; Sec, seconds; MPS, Methylprednisolone; USG, Ultrasonography; ms, millisecond; v, Volts; wks, weeks; NPRS, numerical pain rating scale.

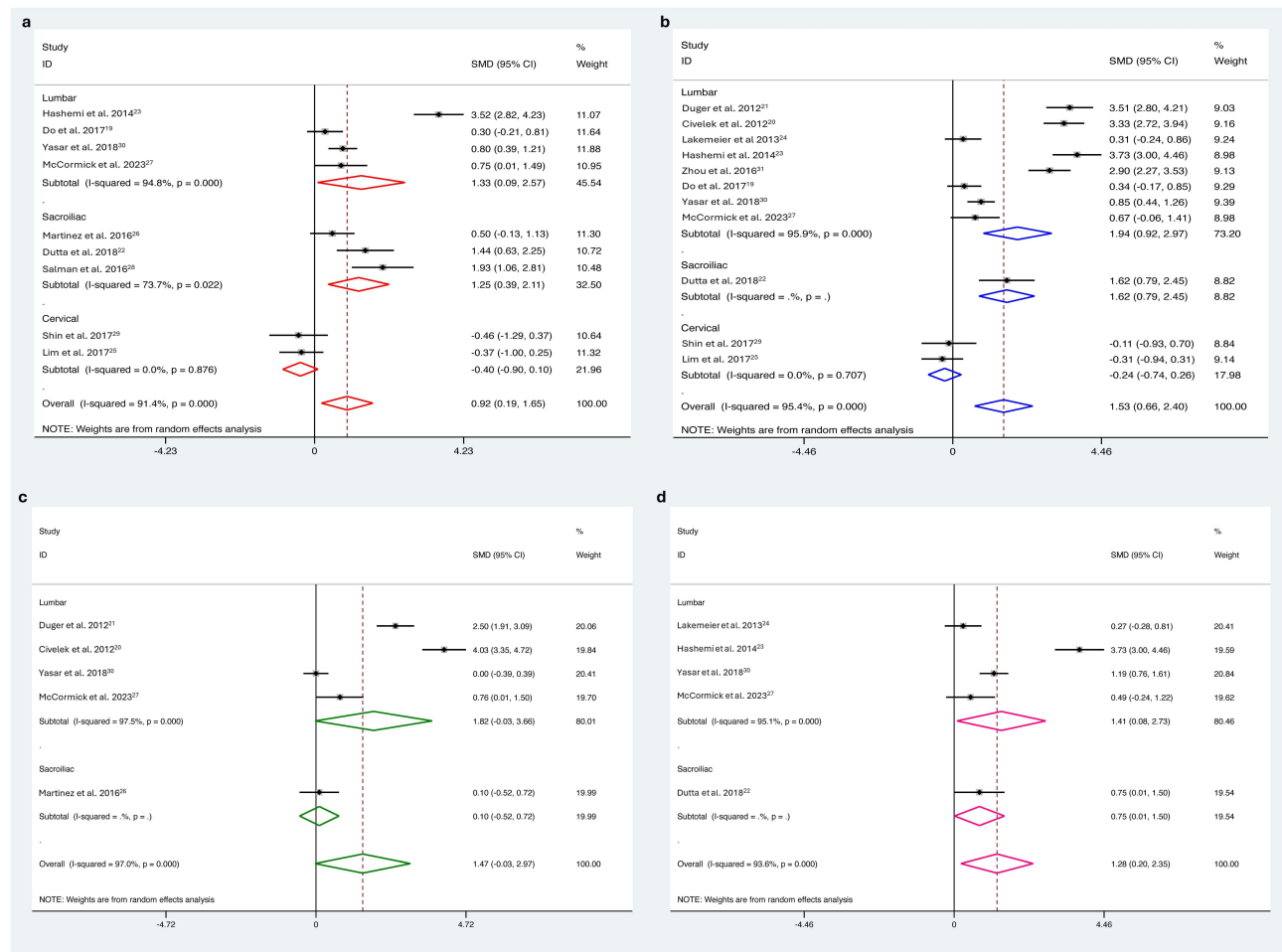


Figure 2 (a-d) Forest Plot for comparing the pain intensity in Radiofrequency vs steroid injections for spinal facet and sacroiliac joint pain across various time points (a) Pain intensity at 3 months, (b) Pain intensity at 6 months, (c) Pain intensity at 12 months and (d) Functional disability at 3 months.

Negative, although not statistically significant effect of CI was reported in patients with cervical region pain (SMD=-0.40; 95% CI: -0.90 to 0.10).

Pain Intensity at 6 Months

Eleven studies reported data of pain intensity score at 6 months (Figure 2b). CI led to higher pain intensity score than RFA (SMD=1.53; 95% CI: 0.66 to 2.40), with substantial heterogeneity ($I^2 = 95.4\%$). Subgroup analysis based on the type of joints showed higher pain intensity score in CI group compared to the RFA group in patients with the sacroiliac region (SMD=1.62; 95% CI: 0.79 to 2.45) and lumbar region (SMD=1.94; 95% CI: 0.92 to 2.97) pain. CI treatment was associated with negative effect in patients with the cervical region pain (SMD=-0.24; 95% CI: -0.74 to 0.26), but the difference was not statistically significant.

Pain Intensity at 12 Months

Five studies reported data of pain intensity at 12 months after the treatment (Figure 2c). Pain scores on the CI and the RFA groups were comparable (SMD=1.47; 95% CI: -0.03 to 2.97), with substantial heterogeneity ($I^2 = 97\%$). Subgroup analysis based on type of joints also showed comparable pain intensity scores in patients with sacroiliac region (SMD=0.10; 95% CI: -0.52 to 0.72) and lumbar region (SMD=1.82; 95% CI: -0.03 to 3.66) pain. No data were available for pain scores of patients with cervical region pain.

Functional Disability at 3 Months

Five studies reported data of functional disability scores at 3 months after the treatment. As shown in Figure 2d, patients in the CI group had a higher functional disability score compared to RFA group (SMD=1.28; 95% CI: 0.20 to 2.35), with substantial heterogeneity ($I^2 = 93.6\%$). Subgroup analysis based on type of joints also showed the CI correlated with significantly higher functional disability than RF ablation in patients with the sacroiliac region (SMD=0.75; 95% CI: 0.01 to 1.50) and lumbar region (SMD=1.41; 95% CI: 0.08 to 2.73) pain. No data were available for patients with the cervical region pain.

Sensitivity Analysis

Sensitivity analysis demonstrated that the pooled SMD estimates were not significantly influenced by the removal of any individual study (Figure 3). These findings indicate the robustness of our meta-analysis results, reassuring that they are not driven by any single study, enhancing confidence in the validity of our findings.

Publication Bias

Funnel plot analysis revealed no apparent signs of asymmetry, with studies evenly distributed around the estimated effect. This suggests the absence of significant publication bias that could impact the results. Funnel plots for pain intensity at 3 months ($p = 0.55$), 6 months ($p = 0.35$), and 12 months ($p = 0.30$) and functional disability at 3 months ($p = 0.69$) are shown in Figure 4a–d.

Quality Assessment

Figure 5 illustrates the assessment of bias across various domains. The evaluation considered the method of randomization, adherence to interventions, management of missing data, precision in measurement, and reporting acceptability in a total of nine studies. Table 2 provides a summary of findings with each outcome based on GRADEpro, enhancing the understanding of the reliability of findings from the studies.

Discussion

Our findings emphasize the potential superiority of RFA in providing sustained pain relief for patients with chronic LBP. Chronic conditions such as LBP often necessitate long-term investment in conservative treatments, which are frequently

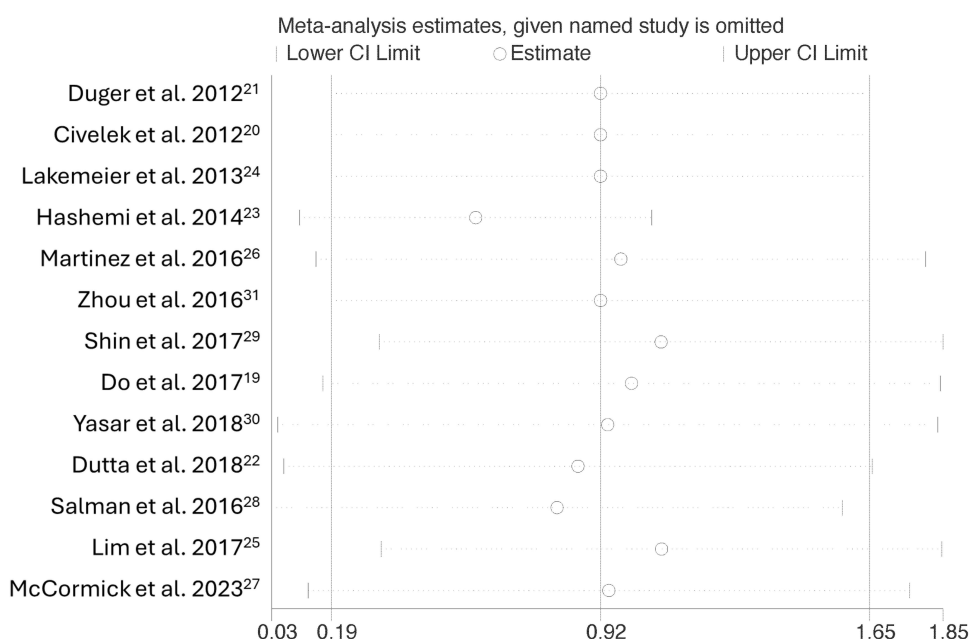


Figure 3 Sensitivity plot for comparing the pain intensity in radiofrequency vs steroid injections for spinal facet and sacroiliac joint pain.

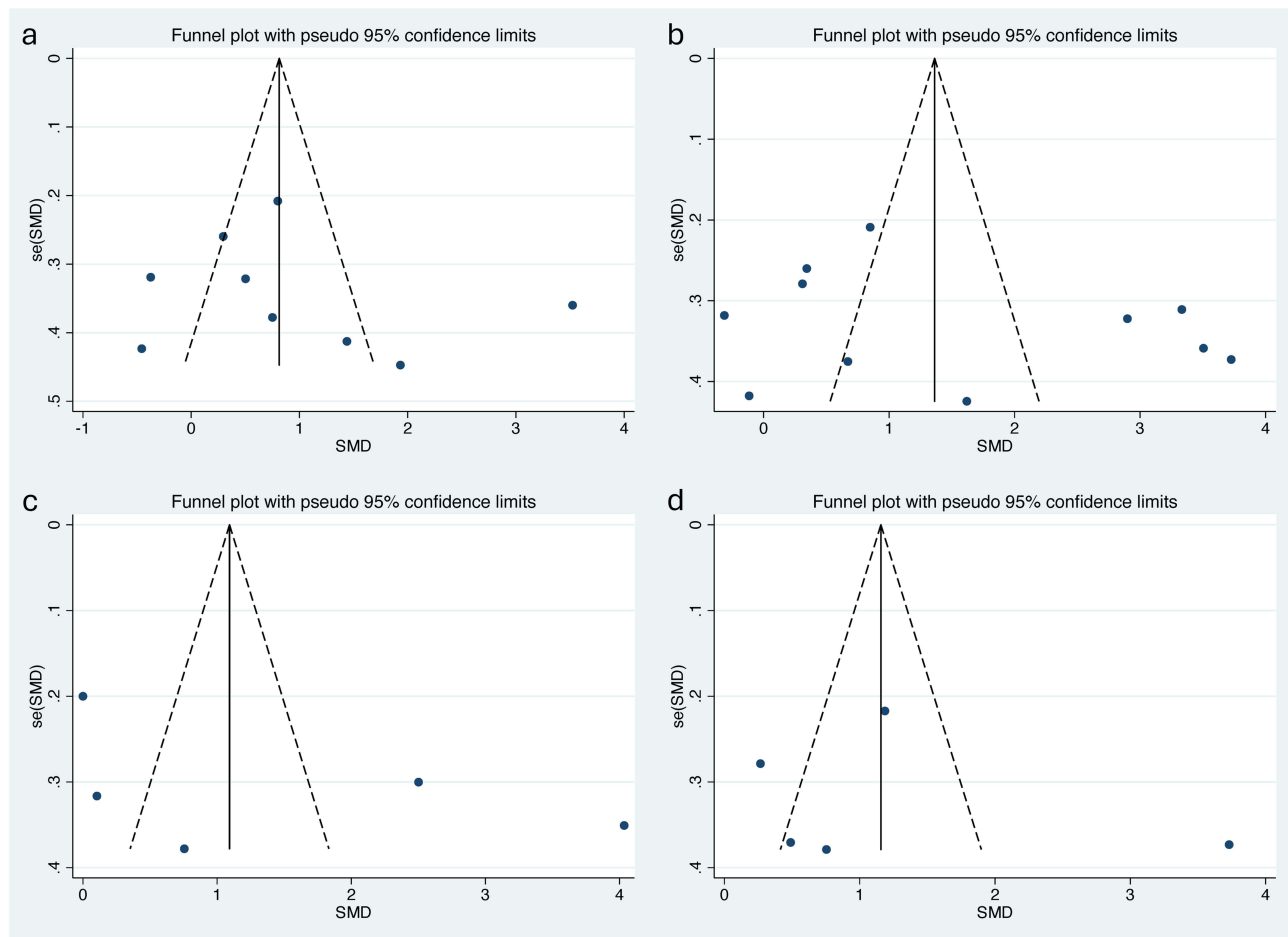


Figure 4 (a-d) Funnel Plot of for comparing the pain intensity in radiofrequency vs steroid injections for spinal facet and sacroiliac joint pain at various time point.

limited in their efficacy. In this context, radiofrequency thermocoagulation presents itself as both an effective and relatively cost-effective alternative to traditional surgery.¹² Conventional radiofrequency thermocoagulation induces localized injury with well-defined boundaries by applying radiofrequency current to heat adjacent tissue.³³ Studies have shown that temperatures up to 80°C can lead to the dissolution and necrosis of unmyelinated nerve C fibers, blocking neuralgia-conducting nerves for extended periods. In contrast, pulsed radiofrequency does not induce neuro-destructive processes like conventional radiofrequency,³⁴ a distinction particularly relevant for sensitive nerves in regions such as the head, neck, dorsal root ganglia, and trigeminal ganglia.

Intraarticular (IA) injections, while potentially offering greater accuracy due to varied facet joint innervation, pose challenges due to their difficulty and increased pain compared to medial branch block (MBB). The choice between IA and MBB approaches may necessitate a trade-off between accuracy and patient comfort.³⁵ The included studies employed diverse techniques for RFA, lacked standardized approaches, and reported varied adverse events. Different pain assessment scales and follow-up intervals were utilized, with no consensus on a superior technique or standardization. Stricter criteria for MBBs have been associated with improved outcomes in lumbar RFA, while more relaxed standards have shown reduced efficacy. The absence of a standardized approach or superior technique indicates a lack of conclusive evidence supporting one method over another, highlighting the need for further research and standardization in RFA procedures to enhance clarity for clinicians and improve patient outcomes.

The variations in steroid dosages and methodologies across the included studies present significant challenges in drawing firm conclusions about their efficacy and safety. Different types and dosages of steroids were combined with various local anesthetics in differing concentrations, making direct comparisons difficult. The methodologies employed

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Civelek et al. 2012 ²⁰	+	−	+	+	−	+	+
Do et al. 2017 ¹⁹	+	+	?	+	?	+	+
Duger et al. 2012 ²¹	+	−	?	?	−	?	?
Dutta et al. 2018 ²²	+	+	+	?	+	+	+
Hashemi et al. 2018 ²³	?	?	?	?	+	+	+
Lakemeier et al. 2013 ²⁴	+	+	+	+	+	+	+
Lim et al. 2017 ²⁵	?	?	+	?	+	+	?
Martinez et al. 2016 ²⁶	+	?	?	?	?	+	?
McCormick et al. 2023 ²⁷	−	−	−	−	+	+	?
Salman et al. 2016 ²⁸	+	+	+	+	+	+	+
Shin et al. 2017 ²⁹	+	+	+	+	+	+	+
Yasar et al. 2018 ³⁰	+	+	+	+	+	+	?
Zhou et al. 2016 ³¹	?	+	+	+	+	+	?

Figure 5 Risk of Bias Summary.

Table 2 Summary of Evidences Using GRADEpro

Certainty assessment							№ of Patients		Effect		Certainty	Importance
№ of Studies	Study Design	Risk of Bias	Inconsistency	Indirectness	Imprecision	Other Considerations	[intervention]	[comparison]	Relative (95% CI)	Absolute (95% CI)		
Pain Intensity at 3 months (follow-up: median 12 months; Scale from: 0 to 10)												
9	Randomised trials	Not serious	Not serious	Not serious	Not serious	None	339	348	-	SMD 0.92 SD higher (0.19 higher to 1.65 higher)	⊕⊕⊕⊕High	
Pain Intensity at 6 months (follow-up: mean 12 months; Scale from: 0 to 10)												
11	Randomised trials	Not serious	Not serious	Not serious	Not serious	None	369	378	-	SMD 1.53 SD higher (0.66 higher to 2.4 higher)	⊕⊕⊕⊕High	
Pain Intensity at 12 Months (follow-up: mean 12 months; Scale from: 0 to 10)												
5	Randomised trials	Not serious	Not serious	Not serious	Not serious	None	192	200	-	SMD 1.47 SD higher (0.03 higher to 2.97 higher)	⊕⊕⊕⊕High	

in these studies also differed, impacting the ability to assess long-term outcomes. Additionally, the inclusion criteria and patient populations were not uniform, which could influence the generalizability of the results. The lack of standardized measures for outcomes and the reporting of adverse events further complicates the analysis, making it challenging to perform a meta-analysis or systematic review that provides definitive conclusions.

Corticosteroids can exert their anti-inflammatory effects to alleviate synovial inflammation and attenuate the excitability of nociceptive nerve fibers, thus providing pain relief. However, potential adverse effects, such as inhibition of the pituitary-adrenal axis, hyperadrenocorticism, osteoporosis, avascular necrosis, and steroid myopathy, pose safety concerns.^{36,37} In contrast, RFA therapy eliminates the need for corticosteroid administration, thereby avoiding these side effects.

It is essential to acknowledge limitations for a comprehensive interpretation of our findings. Small sample sizes observed in most of the included studies may compromise the generalizability and statistical robustness of the results. Larger sample sizes are necessary to improve reliability and accommodate potential variability in patient responses. Additionally, there is substantial heterogeneity among the studies due to variations in patient populations, procedural techniques, and outcome measures. The diversity in procedures and agents used across trials contributes to this heterogeneity, particularly evident within the comparison between intervention groups involving IA and MBB approaches. Moreover, the choice between pulsed and continuous RFA methods may impact outcomes, potentially influencing the depth of analysis. These limitations should be considered when interpreting the study findings, and further research is needed to address these challenges effectively.

Conclusion

This study suggests that RFA may offer superior pain relief and a longer duration of efficacy compared to steroid injections for spinal facet and sacroiliac joint pain. However, the choice between these interventions should be individualized, considering patient preferences, clinical context, and potential risks. Further research, particularly large-scale randomized controlled trials, is needed to strengthen the evidence and refine clinical recommendations.

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Disclosure

The author(s) report no conflicts of interest in this work.

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