

# Dorsal and Volar Approaches for Proximal Interphalangeal Joint Replacement: Comparing Outcomes Through Systematic Review and Meta-Analysis

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Gagandeep Mahi<sup>1</sup> , Eric Ansong<sup>2</sup>, Eugene Gan<sup>3</sup>, Sasan Dehbozorgi<sup>3</sup>, and Han Hong Chong<sup>3</sup> 

## Abstract

**Background:** The 2 primary surgical approaches for proximal interphalangeal joint (PIPJ) arthroplasty, dorsal or volar, have been extensively described in the literature. However, the ongoing debate regarding which approach offers superior results or is associated with fewer complications persists. This systematic review aims to compare the outcomes of PIPJ arthroplasty between the dorsal and volar approaches. **Methods:** A comprehensive search of multiple databases was conducted, and studies meeting predetermined criteria were included. Data extraction, assessment of bias risk, and statistical analysis were performed to compare treatment modalities. Outcome measures included range of motion (ROM), patient-reported outcome measures (PROMs), revisions, and reported complications. **Results:** Among 368 screened articles, 5 studies involving 302 patients (310 implants) were eligible for final review. No significant differences were observed between the 2 approaches regarding postoperative ROM (mean difference [MD] 2.24; 95% confidence interval [CI] –3.83, 8.32;  $P = .47$ ) and PROMs (standardized mean difference [SMD] 0.18; 95% CI –0.12, 0.48;  $P = .25$ ). Complication rates, including revision/fusion, persistent pain, stiffness, infection, and dislocation, did not significantly differ between the approaches. Notably, dorsal approach was associated with higher risk of swan-neck deformity (9 out of 82 implants), while no such cases were reported in the volar approach (0 out of 101 implants). **Conclusion:** Despite limitations and heterogeneity in the literature, both dorsal and volar approaches for PIPJ arthroplasty appear to yield equivalent outcomes for patients.

**Level of Evidence:** II, therapeutic.

**Keywords:** digits, arthroplasty, arthritis, osteoarthritis, outcomes, surgery

## Introduction

Arthritis affecting the proximal interphalangeal joint (PIPJ) is a prevalent condition, with reported population-based studies estimating its incidence at around 18% to 20%, while symptomatic cases range from 0.7% to 6%.<sup>1–3</sup> Typically, patients can manage their symptoms through activity modification and medication such as analgesics or anti-inflammatories. In cases of exacerbation, intra-articular steroid injections may provide relief. However, as symptoms progress, surgical intervention becomes necessary.

Traditionally, arthrodesis of the PIPJ has been the standard surgical approach, particularly for achieving stability and strength in pinch and grip, notably in the index and middle fingers. However, this procedure can be disabling for the ring and little fingers, which require flexion for

better function.<sup>4</sup> Joint replacement has emerged as an alternative, aiming to alleviate pain while preserving or enhancing mobility.<sup>5</sup> Over the past 4 decades, various PIPJ replacement implants have been developed, ranging from flexible silastic hinges to constrained hinges and recently, Pyrocarbon anatomical surface replacements.<sup>5–7</sup>

<sup>1</sup>Kettering General Hospital NHS Foundation Trust, Kettering, UK

<sup>2</sup>Brighton and Sussex NHS Foundation Trust, Brighton, UK

<sup>3</sup>Cardiff and Vale University Health Board, Cardiff, UK

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### Corresponding Author:

Han Hong Chong, University Hospital Llandough, Cardiff and Vale University Health Board, Penlan Road, Llandough, Penarth CF64 2XX, UK.  
Email: chonghh90@doctors.org.uk

The 2 primary surgical approaches for PIPJ replacement, dorsal and volar, have been extensively described, along with variations to facilitate exposure and protect soft tissue stabilizers. The dorsal approach provides wide exposure and access to the joint, with potential for rebalancing the extensor mechanism. However, it carries a risk of joint contracture due to extensor tendon adhesion and possible extension lag. In contrast, the volar approach maintains the integrity of the extensor mechanism and allows for early postoperative rehabilitation.<sup>8,9</sup> While the lateral approach has been mentioned in the literature, its outcomes remain uncertain due to limited reported data.<sup>10,11</sup>

Despite the growing popularity of PIPJ replacement, the evidence base remains limited, primarily consisting of retrospective case series with small sample sizes. A meta-analysis by Adams et al<sup>12</sup> in 2012 highlighted the lack of established effectiveness for PIPJ replacement due to study design and outcome reporting heterogeneity. Subsequently, Yamamoto et al in 2017 conducted a systematic review on surgical approaches and implants in PIPJ replacement but focused solely on descriptive analysis without direct comparative analysis.<sup>9</sup>

Given the gaps in current literature, the purpose of this systematic review was to evaluate the available literature on the 2 common surgical approaches for PIPJ replacement. We aim to perform statistical analyses to compare the patients' outcomes and complications rates between the 2 approaches in PIPJ replacement.

## Methods

This systematic review was performed in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 2020 statement (Supplemental Appendix 1).<sup>13</sup>

### Protocol Setting

We systematically included all comparative studies directly contrasting dorsal and volar approaches for PIPJ replacement in our search. This encompassed randomized controlled trials (RCTs), as well as prospective and retrospective observational studies and series. Only studies available in English or with a translation were considered eligible for inclusion. Our criteria dictated that participants must be over 18 years old, diagnosed with primary PIPJ arthritis, and have undergone PIPJ replacement. We excluded studies focusing on individuals with PIPJ arthritis secondary to trauma or cancer, as well as those investigating revision PIPJ replacement.

The primary outcome of interest revolved around patient-reported outcome measures (PROMs), such as the Disabilities of the Arm, Shoulder, and Hand score (DASH/*QuickDASH*), Michigan Hand Outcomes Questionnaire (MHQ), and Patient-Rated Wrist/Hand Evaluation (PRWHE). Additionally, we

analyzed secondary outcomes, including data comparing range of motion (ROM), revision rates, and complications.

### Identification of Studies

The search strategy of our literature search is detailed in Supplemental Appendix 2. We conducted searches in Medline, Embase, and the Cochrane Central Register of Controlled Trials (CENTRAL) from their inception to August 2023. Additionally, we reviewed the reference lists of identified studies and previous systematic reviews to identify any relevant studies for potential inclusion.

### Selection of Studies

The screening and selection of studies were performed by 3 authors (GM, EA, and EG), and any disagreements were resolved by the fourth author (SD). Initially, articles obtained from the searches underwent screening based on a review of their titles and abstracts, followed by a thorough assessment of the full texts.

### Risk of Bias Assessment

Three authors (GM, EA, and EG) independently evaluated each study for risk of bias using the Cochrane Risk of Bias in Non-randomized Studies—of Interventions (ROBINS-I) tool.<sup>14</sup> Any discrepancies were resolved through discussion with fourth authors (SD) to achieve consensus. The findings were visually depicted through traffic light plots and summary plots using the robvis online tool.<sup>15</sup>

### Data Extraction

Data were extracted from the included studies and collated into a standardized proforma. The extracted data, comparing the 2 approaches, encompassed study characteristics, patient demographics, implant types, PROMs, ROM, implant revisions, complications, and duration of follow up.

### Data Synthesis and Statistical Analysis

We utilized Cochrane Review Manager (RevMan) v5.4 for conducting meta-analysis calculations and generating forest plots. In cases where data extracted from the included studies were incomplete, efforts were made to reach out to the original corresponding authors for any missing information.

For the meta-analysis of PROMs, we employed the standardized mean difference (SMD) and 95% confidence intervals (CIs) to enable the pooling of various reported outcomes. Mean difference (MD) was employed for analyzing ROM, while relative risk (RR) was used to analyze categorical data for revision and complications. In each analysis, a

random-effect model was applied. Statistical significance was defined as  $P < .05$ .

To evaluate study heterogeneity, we utilized the  $I^2$  statistic. An  $I^2$  value of zero indicated perfect homogeneity among the data from the included studies, while an  $I^2$  closer to 100% indicated significant heterogeneity among the studies.

## Results

Following the initial literature search and removal of duplicate results, we identified 202 individual studies. After applying our eligibility criteria, conducting title and abstract screening, and completing the final full-text review, 5 studies were deemed eligible for inclusion in this study.<sup>16-20</sup> The process is illustrated in the PRISMA flowchart, depicted in Figure 1.

Table 1 delineates the particulars of each study. In total, 302 patients (310 implants) were included, with sample sizes varying from 32 to 100 patients, and follow-up durations ranging from 1 month to 5 years.

### Risk of Bias

The authors' judgment assessing studies' quality and risk of bias is demonstrated in Supplemental Appendix 3. In summary, 4 studies were classified as having a low risk of bias, while the remaining one was categorized as moderate risk.

### PROMs

Two studies, Van Nuffel 2014 and Bodmer 2019, utilized PROMs as their outcome measures, specifically the DASH score and MHQ, respectively.<sup>18,20</sup> Postoperatively, no significant differences between the 2 approaches were observed (SMD 0.18; 95% CI -0.12, 0.48;  $P = .25$ ;  $I^2$  2%), as illustrated in Figure 2.

### ROM

ROM was assessed as an outcome measure in 5 studies.<sup>16-20</sup> Postoperatively, no significant differences between the 2 approaches were found (MD 2.24; 95% CI -3.83, 8.32;  $P = .47$ ;  $I^2$  50%) (Figure 3).

### Revision and Complication

Four studies provided data on the revision of implants to either new implants or arthrodesis.<sup>16-18,20</sup> There was no difference in relative risk between the 2 approaches (RR 0.77; 95% CI 0.30, 1.96;  $P = .58$ ;  $I^2$  0%).

We analyzed 5 common complications reported by the included studies, namely persistent pain, stiffness, infection, dislocation, and swan-neck deformity. There were no statistically significant differences in relative risk between

both approaches regarding persistent pain, stiffness, infection, and dislocation postoperatively. Dorsal approach was shown to have higher risk of developing swan-neck deformity (9 out of 82 implants) postoperatively compared to volar approach (0 out of 101 implants) (RR 0.12; 95% CI 0.02, 0.88;  $P = .04$ ;  $I^2$  0%) (Figure 4).

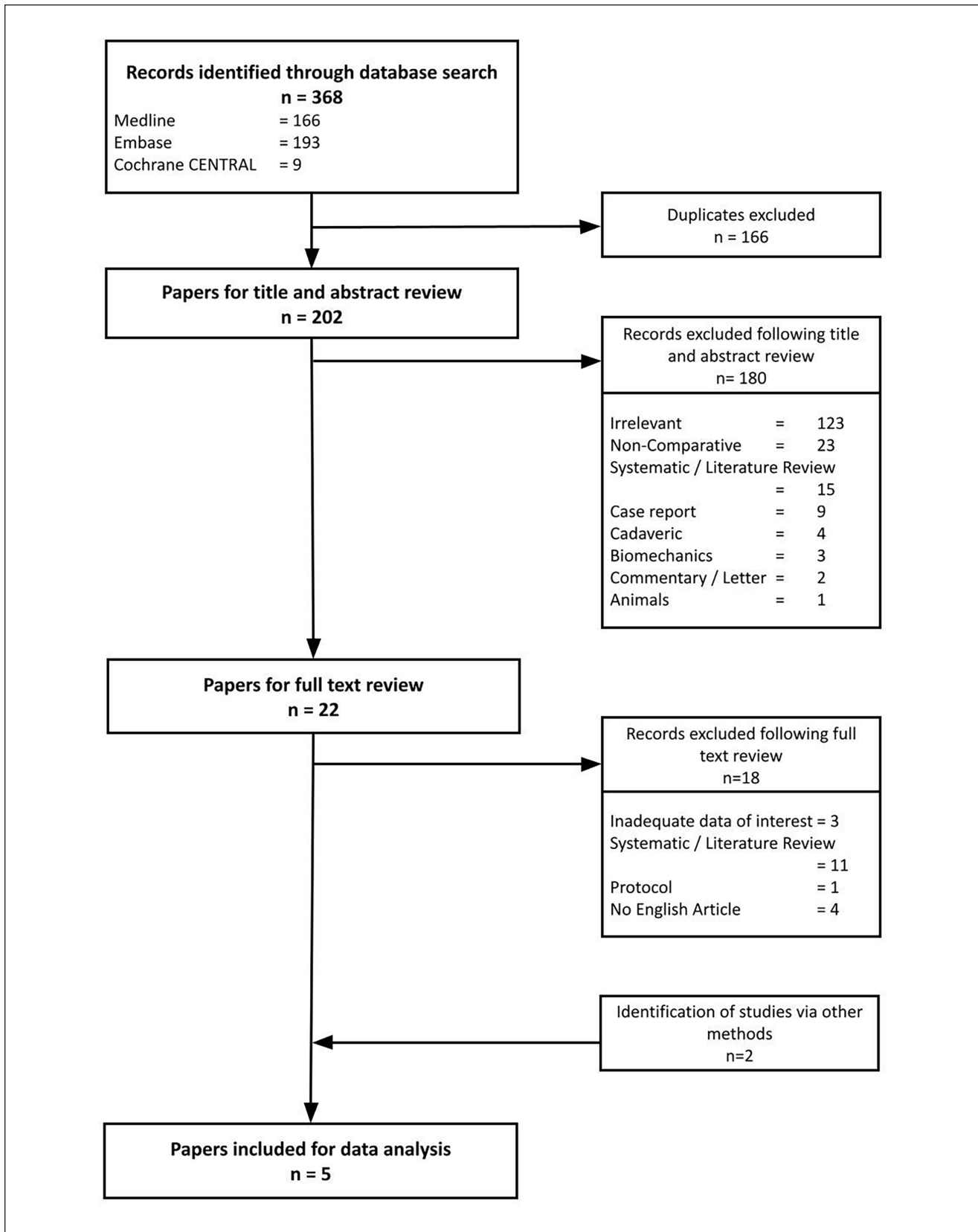
## Discussion

Our study aimed to assess the outcomes of PIPJ replacement, specifically comparing the dorsal and volar approaches. Our analysis revealed no significant differences between the 2 approaches in terms of PROMs, ROM, revision rates, and common postoperative complications. However, it is noteworthy that the dorsal approach exhibited a slightly higher risk of developing swan-neck deformity compared to the volar approach.

The dorsal approach remains the predominant technique for PIPJ replacement, primarily due to its technical simplicity and favorable access to the joint. Furthermore, the majority of current implant instrumentation systems are designed to facilitate this approach.<sup>8</sup> However, recent publications have highlighted the promising outcomes associated with the volar approach.<sup>21-23</sup> Advocates of the volar approach suggest that it preserves the extensor mechanism, enabling early postoperative rehabilitation to prevent complications such as adhesions, extensor lag, and joint contracture.<sup>8,9</sup> Despite these theoretical advantages, our meta-analysis did not detect a statistically significant difference in postoperative range of motion between the 2 approaches. This may be attributed to advancements in surgical techniques, implant designs, and tailored rehabilitation strategies for each approach.

Yamamoto et al<sup>9</sup> conducted a systematic review examining various implants and approaches for PIPJ replacement. Their analysis of 40 studies suggested that silicone implants with the volar approach yielded the best ROM outcomes with fewer complications compared to other implant designs and surgical approaches. Although their review provided valuable insights, limitations such as limited data and inability to perform meta-analysis for each subcategory combination of implant and approach were acknowledged. In contrast, our study focused specifically on comparing the dorsal and volar approaches, allowing for a more detailed and focused analysis. Our findings complement Yamamoto et al's work by providing additional evidence regarding the comparative effectiveness and safety profiles of these 2 approaches.

The findings of this systematic review provide essential insights for surgeons performing PIPJ replacement. Both dorsal and volar approaches demonstrate comparable safety and efficacy in terms of surgical complications and functional outcomes, which underscores the importance of a personalized approach in surgical planning. Surgeons



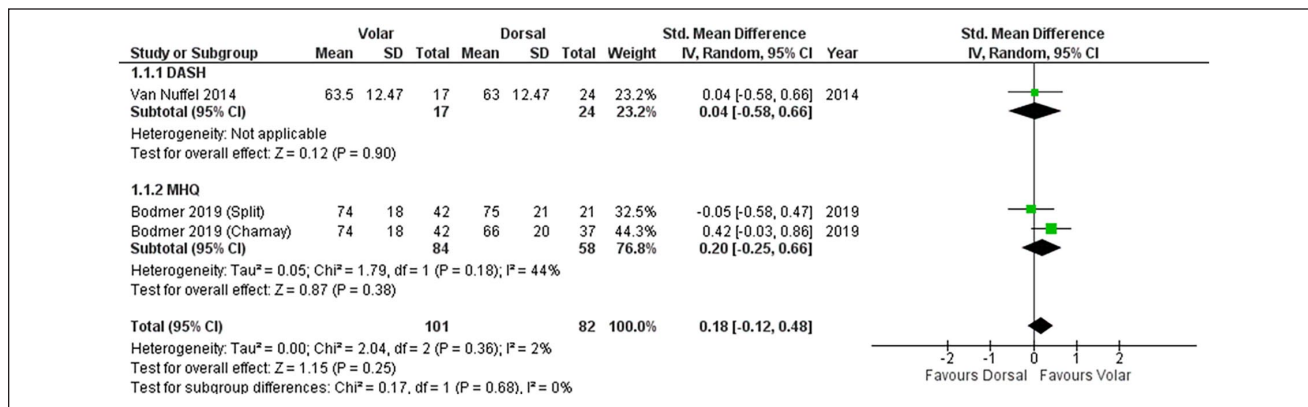
**Figure 1.** Preferred Reporting Items for Systematic Reviews and Meta-Analyses flowchart.

**Table 1.** Characteristics of the Included Studies.

Study / method	Age (years)	Total patients, n=	Type of approach	Total implant, n =	Type of implants	Reported outcome of interest	Follow up
Heren <sup>19</sup> Prospective	65.7 (51-79)	38	Volar	38	Swanson Silicone	ROM	28.2 months (SD 16.1)
	65 (49-75)		Dorsal	21			51.4 months (SD 22.4)
Natera <sup>17</sup> Retrospective	56	66	Volar	8	Avanta Silicone	ROM	29 months
			Dorsal (Lluch modification of Chamay)	14			
Van Nuffel <sup>20</sup> Retrospective	59.5 (39-80)	32	Volar	17	Swanson Silicone	DASH, ROM	Not specified
	60 (45-71)		Dorsal (Chamay)	24	Ascension PyroCarbon		
Bodmer <sup>18</sup> Prospective	66 (SD 11)	100	Volar	42	CapFlex-PIP	MHQ, ROM	2 years
			Dorsal (Chamay)	37	Metal-on-poly		
			Dorsal (Tendon splitting)	21			
Tranchida <sup>16</sup> Retrospective	64.2 (SD 11.8)	66	Volar	45	Silicone, Pyrocarbon, Metal-on-poly	ROM	132 days (30-365)
			Dorsal	43	Silicone, Pyrocarbon		

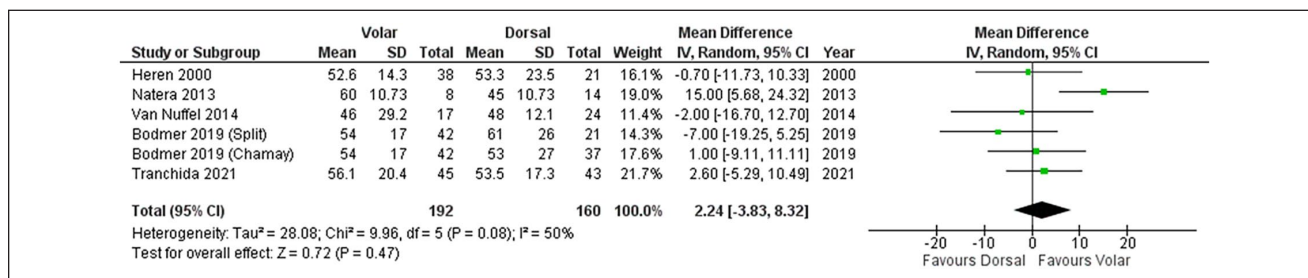
Data are presented as n, mean (range) or mean (SD) unless otherwise indicated.

Abbreviations: DASH, disabilities of arm, shoulder and hand; MHQ, Michigan Hand Questionnaire; ROM, range of motion.



**Figure 2.** Forest plot of the comparison between 2 approaches for PROMs.

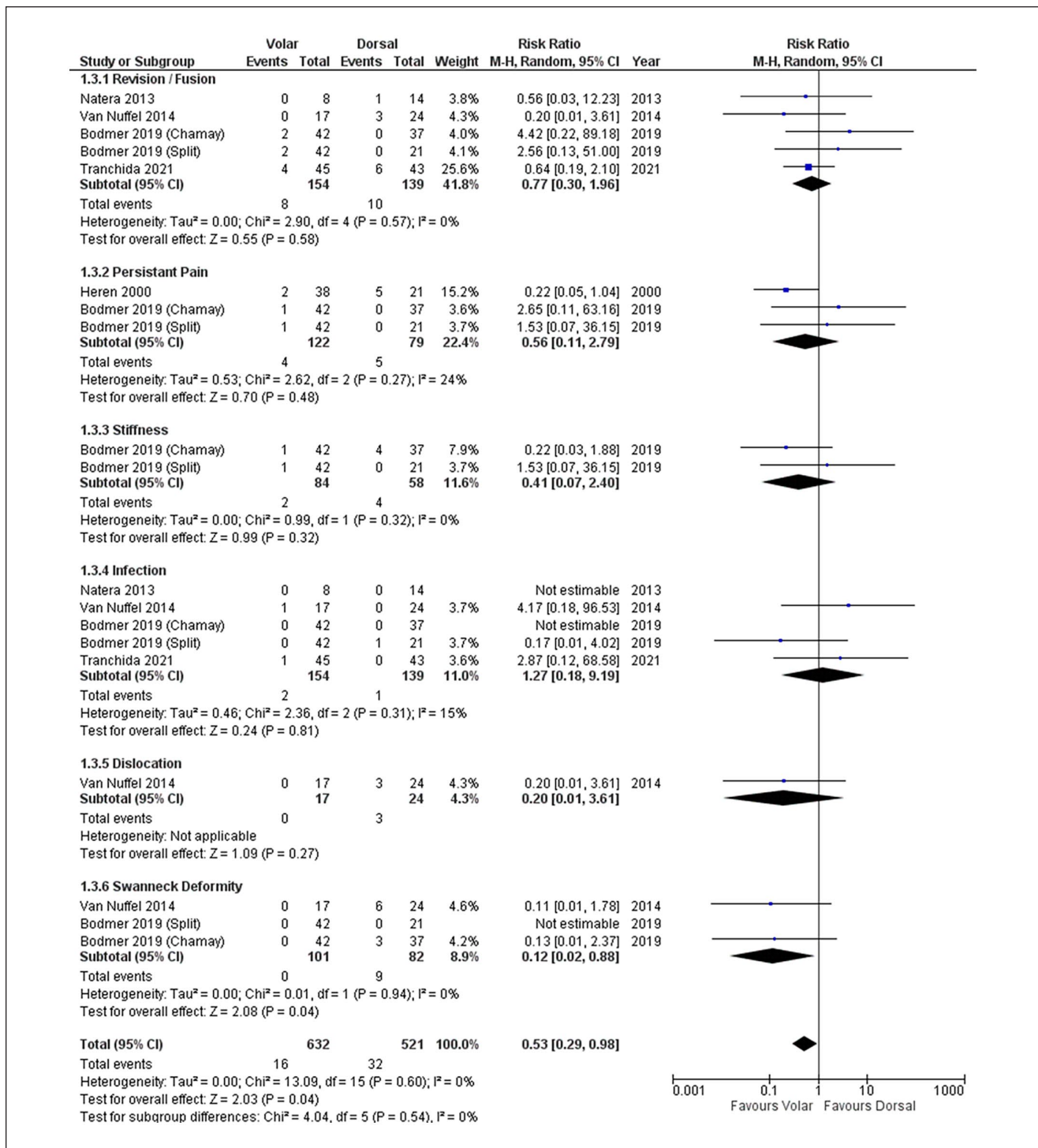
DASH, Disabilities of the Arm, Shoulder, and Hand; MHQ, Michigan Hand Outcomes Questionnaire; PROMs, patient reported outcome measures.



**Figure 3.** Forest plot of the comparison between 2 approaches for ROM.

ROM, range of motion.





**Figure 4.** Forest plot of the comparison between 2 approaches for revision and complications.

can confidently select the approach based on individual patient anatomy, specific details of the joint pathology, and their own surgical experience without compromising the outcomes.

Our study is not without limitations. Firstly, the retrospective nature of the included studies introduces inherent biases that may impact the reliability of our results. Additionally, the heterogeneity among the studies, including variations in

patient populations, preoperative deformities, surgical techniques, surgeon experience, implant usage, and reported outcome measures, may limit the comparability of our findings. Furthermore, the relatively short follow-up durations in some patients may not capture long-term outcomes and complications associated with PIPJ replacement. Lastly, the limited number of studies available for inclusion may restrict the breadth of our analysis and the generalizability of our findings. These limitations underscore the need for further prospective studies with standardized protocols to elucidate the optimal approach for PIPJ replacement.

## Conclusion

Our comparative analysis of dorsal and volar approaches for PIPJ replacement revealed no significant differences in patient's outcome. To validate these findings and overcome the limitations of our study, it is imperative to conduct prospective studies with standardized protocols and extended follow-up periods. Future research should prioritize examining outcomes beyond the immediate postoperative phase and investigating how various implant designs and surgical techniques influence long-term functional outcomes and complication rates. Establishing a national registry database and fostering collaboration among medical centers could facilitate this endeavor.

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## Ethical Approval

Ethical approval is not required for this article.

## Statement of Human and Animal Rights

This article does not contain any studies with human or animal subjects.

## Statement of Informed Consent

Informed consent is not required for this article.

## Declaration of Conflicting Interests

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## ORCID iDs

Gagandeep Mahi  <https://orcid.org/0000-0001-6487-4805>

Han Hong Chong  <https://orcid.org/0000-0003-0102-2167>

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