

POCUS-guided suprapubic bladder aspiration in children: a systematic review and meta-analysis

Punção suprapúbica guiada por POCUS para coleta de urina em crianças: uma revisão sistemática

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Abstract

Objective: Point-of-care ultrasound (POCUS) has been employed to assist with invasive techniques, such as suprapubic aspiration (SPA). We aimed to conduct a meta-analysis to compare the effectiveness of suprapubic bladder aspiration in children, with and without the use of POCUS.

Methods: We conducted a systematic search across the MEDLINE, EMBASE and Cochrane Central Register of Controlled Trials databases for articles published through February 2025. Eligible studies included randomized controlled trials (RCT) comparing SPA with and without POCUS in pediatric populations. Data extraction focused on overall procedural success rates, as well as rates of success at first and second attempts.

Results: We analyzed seven trials collectively evaluating 468 suprapubic bladder aspirations. POCUS-guided SPA showed higher overall success rates (OR = 4.88; 95% CI: 2.36–10.09; $p < 0.0001$; $I^2 = 10\%$). Although no significant difference was found in first-attempt success, POCUS improved success within two attempts (OR = 7.75; 95% CI: 1.54–39.09; $p = 0.01$; $I^2 = 61\%$), although without significantly reducing the mean number of attempts. However, sensitivity analysis showed a benefit after the exclusion of one outlier study.

Conclusions: POCUS improves SPA success in children and may enhance sterile urine collection, although training and equipment access remain challenges.

Keywords: Point-of-care systems; Ultrasonography; Urine specimen collection; Infant; Infant, newborn.

Resumo

Objetivo: A ultrassonografia à beira-leito (POCUS) é uma ferramenta utilizada para auxiliar em técnicas invasivas, como a aspiração suprapúbica (ASP). O objetivo deste estudo foi realizar uma meta-análise para comparar a efetividade da aspiração vesical suprapúbica em crianças com e sem o uso de POCUS. **Métodos:** Foi realizada uma busca sistemática nas bases de dados MEDLINE, EMBASE e Cochrane (CENTRAL) até fevereiro de 2025. Foram incluídos ensaios clínicos randomizados que compararam a ASP com e sem o uso de POCUS em populações pediátricas. A extração de dados concentrou-se nas taxas de sucesso do procedimento em geral e nas primeiras e segundas tentativas.

Resultados: Sete estudos ($n = 468$) foram incluídos na análise. A ASP guiada por POCUS apresentou maiores taxas de sucesso geral (OR = 4,88; IC 95%: 2,36–10,09; $p < 0,0001$; $I^2 = 10\%$). Não houve diferença significativa no sucesso à primeira tentativa, porém o sucesso em até duas tentativas foi maior (OR = 7,75; IC 95%: 1,54–39,09; $p = 0,01$; $I^2 = 61\%$). O POCUS não reduziu significativamente o número médio de tentativas, embora a análise de sensibilidade tenha demonstrado benefício após a exclusão de um estudo discrepante. **Conclusões:** O uso do POCUS aumenta as taxas de sucesso da ASP em crianças e pode favorecer a obtenção de amostras urinárias estéreis, embora o treinamento e o acesso ao equipamento ainda representem desafios.

Unitermos: Sistemas automatizados de assistência junto ao leito; Ultrassonografia; Coleta de urina; Lactente; Recém-nascido.

Running head: POCUS-guided suprapubic bladder aspiration in children

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INTRODUCTION

Urinary tract infections are one of the most common bacterial infections in children and the leading cause of bacterial infections in infants⁽¹⁻³⁾. Ten percent of children experience at least one infection between 1 month and 6 years of age⁽⁴⁾. Although early diagnosis is important given the risk of long-term renal injury, obtaining a sterile urine specimen can be challenging, especially in children who are not toilet trained⁽⁵⁾. Urine collection, which can be performed by using noninvasive methods, such as a bag or clean-catch technique, or invasive methods, such as bladder catheterization or suprapubic aspiration (SPA), is essential for confirming the diagnosis⁽⁶⁾. However, each of those techniques has its limitations, particularly regarding contamination rates, which can lead to false positives⁽⁷⁾. Clinical guidelines generally recommend clean-catch sampling or SPA as preferred methods⁽⁸⁾, with SPA presenting the lowest contamination rates in urine sample collection, as well as exhibiting high sensitivity^(9,10).

Point-of-care ultrasound (POCUS) has become an important procedural tool in clinical practice, providing a noninvasive approach to assess bladder size and anatomical positioning prior to invasive techniques, such as SPA⁽¹¹⁾. The use of POCUS facilitates visualization of the urinary bladder, aiding in localization and ensuring adequate urine sample collection⁽¹²⁾. The use of ultrasound can also increase urine collection by precisely locating a full bladder, providing real-time visualization during the procedure, which may reduce complications and improve the SPA success rate⁽¹³⁾. However, despite the increasing use of POCUS in multiple clinical contexts, there is a lack of consensus among studies evaluating the effectiveness of this tool for SPA, particularly for improving procedural success.

Although some single-center studies suggest that the use of POCUS improves SPA success, robust evidence of its superiority over the blind technique remains limited. Therefore, the aim of this systematic review and meta-analysis was to address this gap by comparing SPA performed with and without POCUS, in children. Specifically, we evaluated differences in success rates (overall and at first or second attempt), as well as the

mean number of attempts required in order to obtain a successful sample, providing a critical analysis that may directly impact clinical practices for urine collection in children. By critically analyzing these outcomes, we hope to generate evidence to improve clinical practice for urine collection in pediatric patients.

MATERIALS AND METHODS

This was a systematic review and meta-analysis, performed in accordance with the Cochrane Collaboration and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines⁽¹⁴⁾. The protocol for this review was registered in the International Prospective Register of Systematic Reviews (Identifier: CRD420250652095)

Search strategy and inclusion criteria

A systematic digital search was conducted across the MEDLINE, EMBASE and Cochrane Central Register of Controlled Trials databases by two authors (S.M. and B.S.R.), working independently, up to February 2025. The search strategy included the terms "Ultrasonography", "Ultrasound", "Point of care ultrasonography", "POCUS", "Suprapubic aspiration", "Bladder aspiration", "Bladder puncture", "Suprapubic puncture", "Infant", "Infants", "Newborn", "Neonate", "Pediatrics", "Pediatric", "Child", and "Children". The complete search strategy can be found in Appendix 1. The same two researchers (B.S.R. and S.M.), working independently, screened titles and abstracts for eligibility, consulting a senior investigator in case of disagreements. Full texts, supplementary materials, online appendices, and reference lists of eligible studies were reviewed to ensure compliance with the inclusion criteria and identify additional relevant studies. Discrepancies were resolved through discussion with other reviewers until consensus was reached.

To be included, articles had to meet all of the following criteria: including data from prospective randomized controlled trials (RCTs); having enrolled pediatric patients (< 24 months of age) who underwent SPA; having compared the SPA procedure with and without the use of POCUS; having been published as a full-text article. Observational studies that did not compare POCUS-guided SPA with traditional SPA were excluded.

Data analysis and quality assessment

A comprehensive data extraction process was conducted by two investigators (E.F.M. and L.B.S.C.), working independently, including population data such as the number of patients, mean age, gender distribution, mean weight, and indication for SPA. Specific information regarding SPA was also collected, covering aspiration criteria when POCUS was used, materials for the procedure, the definition of a successful sample, and the ultrasound device employed. Each of the studies included defined “successful sample” differently, as detailed in Table 1. Definitions included aspiration of urine within up to three attempts, obtaining any amount of urine, requiring a minimum volume of 1 cc, or considering the bladder empty after aspiration. In three studies, the criteria for success were not explicitly stated. These variations were recorded during data extraction and were considered in the qualitative synthesis and sensitivity analyses. The main outcome collected was success rate, with success at first attempt, success up to second attempt and mean number of attempts as secondary outcomes. The risk of bias was assessed by using the Cochrane risk of bias tool for randomized trials, designated the RoB 2 tool⁽¹⁴⁾. Potential bias was evaluated across five domains—randomization process, deviations from intended interventions, missing outcome data, measurement of the outcome, and selection of the reported result—each study thus being classified as having a high, low, or unclear risk of bias.

Statistical analysis

Odds ratios with 95% confidence intervals were combined using a random-effects model to evaluate treatment effects for categorical outcomes. Heterogeneity was assessed through the use of the Cochran Q test and calculation of the I^2 statistic, with significance defined as $p < 0.10$ and $I^2 > 25\%$, respectively. Statistical analyses were performed using the software Review Manager, version 5.4⁽¹⁵⁾ and R-Studio, version R-4.3.3^(16,17). A leave-one-out sensitivity analysis was conducted to determine the impact of individual studies on the overall meta-analysis results.

RESULTS

Study selection and baseline characteristics

The systematic search identified 2,781 studies; after duplicates had been removed, 2,600 remained. Through title and abstract screening, an additional 2,566 studies were removed. The remaining 34 studies underwent full-text evaluation, which resulted in the exclusion of 27 studies. Therefore, seven RCTs were included in the final analysis^(11,12,18–22). The screening and eligibility process is illustrated in Figure 1.

The studies included a collective total of 468 partici-

pants, with 248 allocated to the POCUS groups and 220 allocated to the control groups. The population consisted of neonates and infants, with mean ages ranging from 0.47 to 9.7 months and mean weights ranging from 1.25 to 7.23 kg. Among the seven studies included, age was reported in four and weight was reported in three.

Success rates

The pooled analysis demonstrated significantly higher odds of success for POCUS-guided SPA than for traditional SPA (OR = 4.88; 95% CI: 2.36–10.09; $p < 0.0001$; $I^2 = 10\%$). A forest plot (Figure 2) shows that most studies favored the POCUS approach, with the odds ratio being highest for the study conducted by Mahdipour et al.⁽²¹⁾: 24.13; 95% CI: 2.98, 196.11. The overall success rate for POCUS-guided SPA was 89.9%, compared with 63.6% for the control (traditional SPA), representing a relative difference of approximately 41%.

A secondary analysis revealed that POCUS guidance does not have a significant impact on first-attempt odds of success (OR = 4.83; 95% CI: 0.31–76.21; $p = 0.17$; $I^2 = 86\%$). That analysis also showed that there was high heterogeneity among the studies ($I^2 = 86\%$), suggesting substantial variation in the results (Figure 3), as further discussed below. There was a statistically significant difference in the odds of success within up to two attempts (OR = 7.75; 95% CI: 1.54–39.09; $p = 0.01$; $I^2 = 61\%$), further supporting the superiority of POCUS in this context (Figure 4). Leave-one-out sensitivity analysis was not performed due to the low I^2 indicated.

Mean number of attempts

POCUS guidance did not have a significant impact on reducing the number of attempts, with a mean difference (MD) of -0.77 (95% CI: -1.64 to 0.09 ; $p = 0.08$; $I^2 = 97\%$). When the study conducted by Kiernan et al.⁽¹⁸⁾ was excluded, the leave-one-out sensitivity analysis showed a significant benefit of POCUS guidance (MD = -0.35 ; 95% CI: -2.02 to 0.09 ; $I^2 = 73\%$).

Sensitivity analysis

A sensitivity analysis performed for the overall success rate showed that omitting the studies conducted by Chu et al.⁽²²⁾ and Mahdipour et al.⁽²¹⁾ had a significant influence on heterogeneity, with ORs of 5.78 (95% CI: 3.24–10.30; $I^2 = 0.0\%$) and 4.27 (95% CI: 2.46–7.42; $I^2 = 0.0\%$), respectively (Figure 6). A sensitivity analysis for first-attempt success did not find a specific study responsible for the heterogeneity (Figure 7). A sensitivity analysis for mean number of attempts identified the Kiernan et al. study⁽¹⁸⁾ as the primary source of heterogeneity, and exclusion of that study resulted in the achievement of statistical significance (MD = -0.35 ; 95% CI: -2.02 to

Table 1—Baseline characteristics of studies and patient populations.

Study	RCT Design	Country	Group								Ultrasound device used	Definition of successful sample collection	Reason for aspiration	Criteria for aspiration when POCUS was used	Materials used to perform aspiration	
			POCUS			Control		All subjects								
			(n)	Age (months)	Mean $\pm$ SD	(n)	Age (months)	Mean $\pm$ SD	Age (months)	Male						Weight (kg)
				Mean $\pm$ SD			(%)									
Choi et al. ⁽¹⁸⁾	Single-center	South Korea	46	9.4 $\pm$ 0.7	32	9.7 $\pm$ 0.7	n/a	53.8	n/a	LOGIQ 400	Urine aspirated in up to 3 attempts	Suspected UTI	10 $\times$ 20 mm of urine in the bladder	23 G needle attached to a 10-cc syringe		
Chu et al. ⁽²¹⁾	Single-center	Hong Kong	30	5.2 $\pm$ 3.4	30	4.2 $\pm$ 3.1	4.7 $\pm$ n/a	45.0	n/a	Aloka 620	n/a	Fever and a urine sample testing positive for leukocyte esterase or nitrites, with or without leukocytes (>5 mm ³) OR a positive bacterial culture in a previous bag urine sample	Bladder volume estimated to be greater than 3 mL	21 G needle		
Gochman et al. ⁽¹¹⁾	Single-center	USA	35	1.7 $\pm$ 2.8	31	2.3 $\pm$ 2.9	1.98 $\pm$ n/a	60.6	4.98 $\pm$ n/a	ScanMate II	Any amount of urine aspirated	Judgment of the examining physician	Anteroposterior and transverse maximum diameters $\geq$ 2 cm	22 G spinal needle attached to a 3-mL syringe		
Kiernan et al. ⁽¹⁷⁾	Single-center	USA	28	n/a	25	n/a	n/a	n/a	n/a	Pie Medical Imaging	n/a	Determined by the clinical team	Bladder measured at least 1 $\times$ 1 cm (longitudinal and transverse views)	Sterile 23 G needle attached to a 3-mL syringe		
Lee et al. ⁽¹⁹⁾	Single-center	South Korea	32	n/a	28	n/a	5.28 $\pm$ n/a	70.0	7.23 $\pm$ n/a	Sonoace 8800	n/a	Suspected UTI	Urine present	21 G needle attached to a 10-mL syringe.		
Mahdipour et al. ⁽²⁰⁾	Single-center	Iran	38	n/a	38	n/a	1.17 $\pm$ 0.47	59.6	4.19 $\pm$ 1.25	Ultrasonix	$\geq$ 1 cc of urine aspirated	Suspected UTI	n/a	10-cc syringe with a 24 G needle		
Munir et al. ⁽¹²⁾	Single-center	Australia	39	n/a	36	n/a	n/a	37.3	n/a	n/a	Bladder considered empty	Uncontaminated urine specimen required for microscopy and culture	10 ml of urine in the bladder	23 G needle attached to a 2- or 5-mL syringe		

UTI, urinary tract infection.

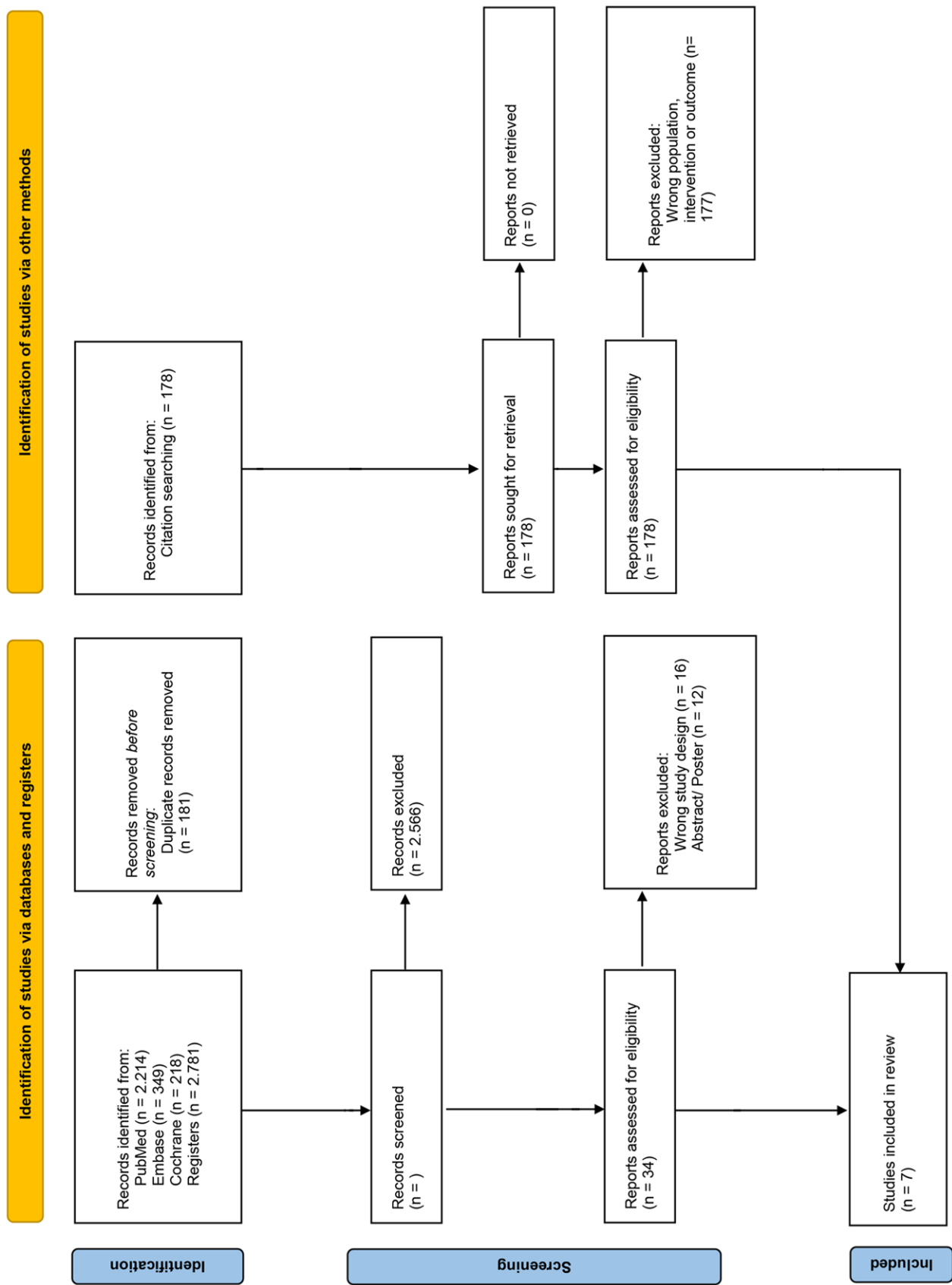


Figure 1. Flow chart of the article selection process, conducted in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses recommendations.

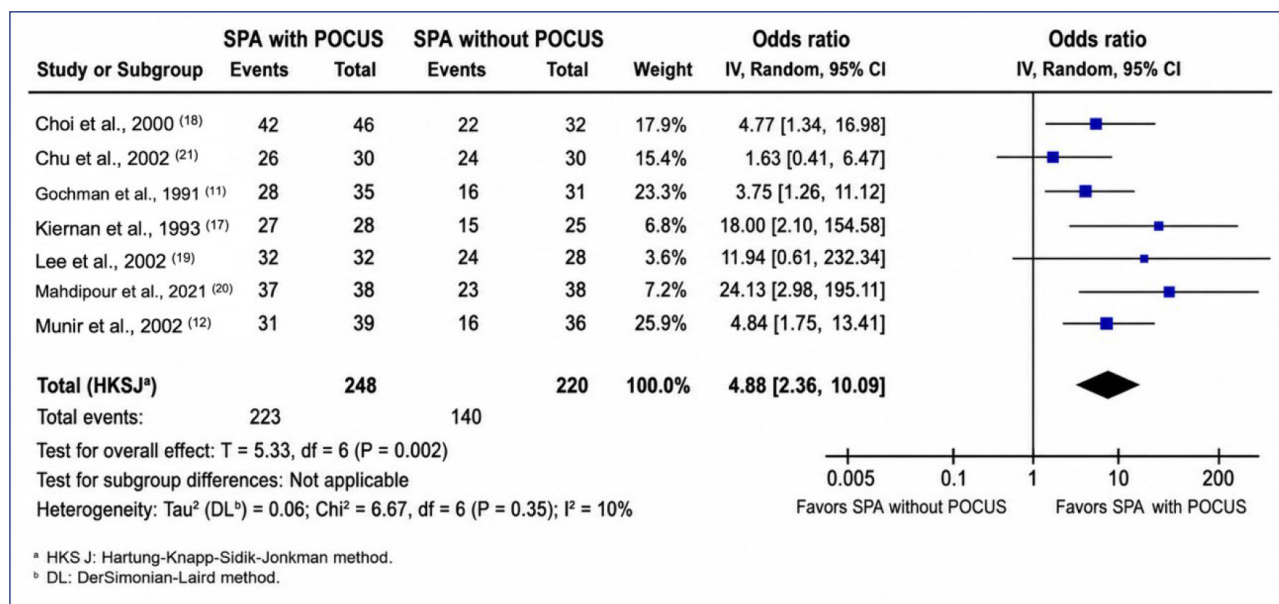


Figure 2. Forest plot showing the overall success rate of SPA, comparing the POCUS group with the control group. HKSJ, Hartung-Knapp-Sidik-Jonkman; REML, restricted maximum likelihood; df, degrees of freedom.

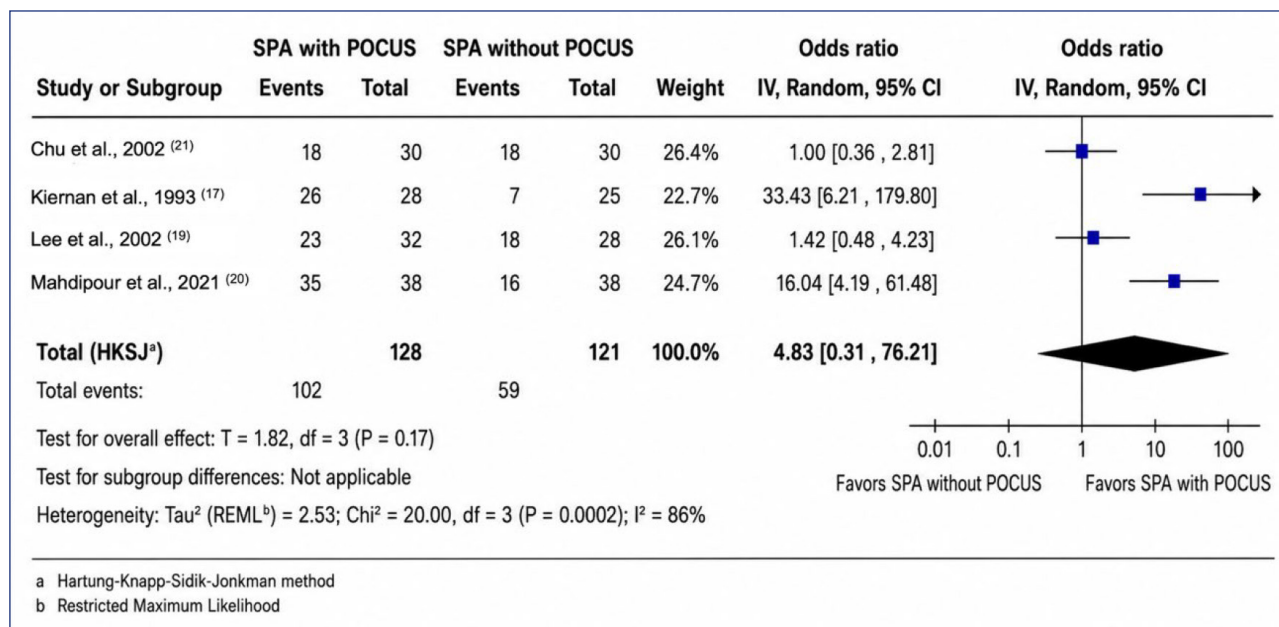


Figure 3. Forest plot of SPA success at first attempt, comparing the POCUS group with the control group. HKSJ, Hartung-Knapp-Sidik-Jonkman; REML, restricted maximum likelihood; df, degrees of freedom.

0.09; I² = 73%). Nonetheless, high heterogeneity persisted across studies (Figure 8).

Risk of bias assessment

The risk of bias across the five domains assessed for RCTs (selection, performance, detection, attrition, and reporting biases), evaluated by using the RoB 2 tool, was low and showed concordance between the two evaluators (E.F.M and L.B.S.C.), as illustrated in Figure 9.

DISCUSSION

In this systematic review and meta-analysis, the pooled results of 468 patients demonstrated a significant improvement in the success rate of SPA due to POCUS guidance in pediatric patients. Overall, considering summarized raw data across studies, the success rate was 89.9% with POCUS and 63.6% without, representing a relative difference of approximately 41%. Success at first

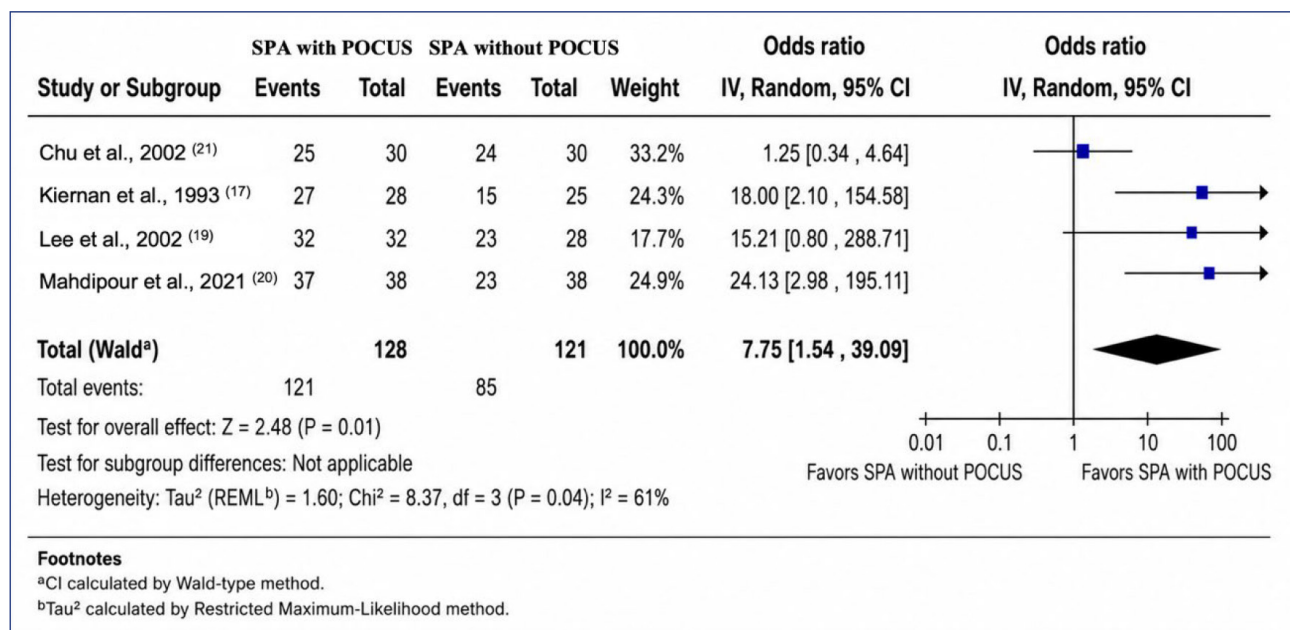


Figure 4. Forest plot of SPA success up to two attempts, comparing the POCUS group with the control group. REML, restricted maximum likelihood; df, degrees of freedom.

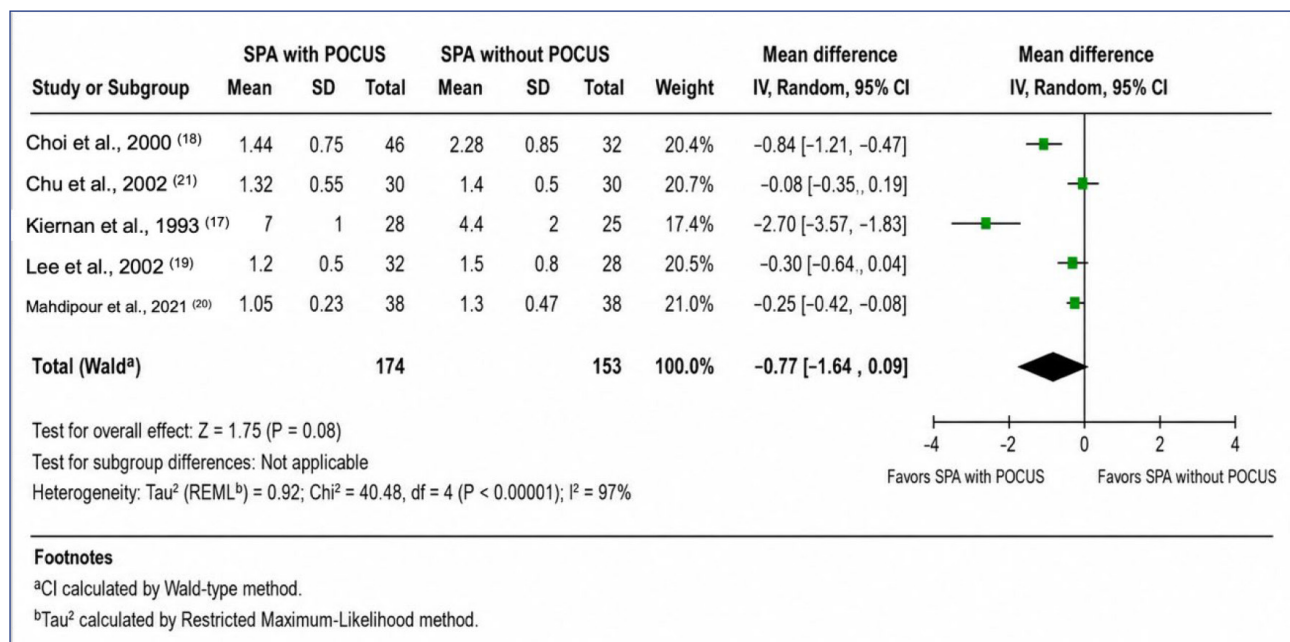


Figure 5. Forest plot of mean number of attempts to collect urine comparing the POCUS group with the control group. REML, restricted maximum likelihood; df, degrees of freedom.

attempt occurred in 102 (79.7%) of the 128 cases in which POCUS was employed, compared with 59 (48.8%) of the 121 cases in which it was not. These findings indicate that using POCUS may improve the efficiency of SPA and reduce patient distress associated with the procedure by reducing the number of attempts needed. In addition to these benefits, POCUS offers practical advantages such

as estimating bladder dimensions, confirming adequate bladder filling, and enabling real time visualization of the needle during the procedure, all of which contribute to a higher rate of procedural success.

Despite the benefits of POCUS in the overall success rate of SPA, some important considerations must be addressed. The first is the variability in operator ex-

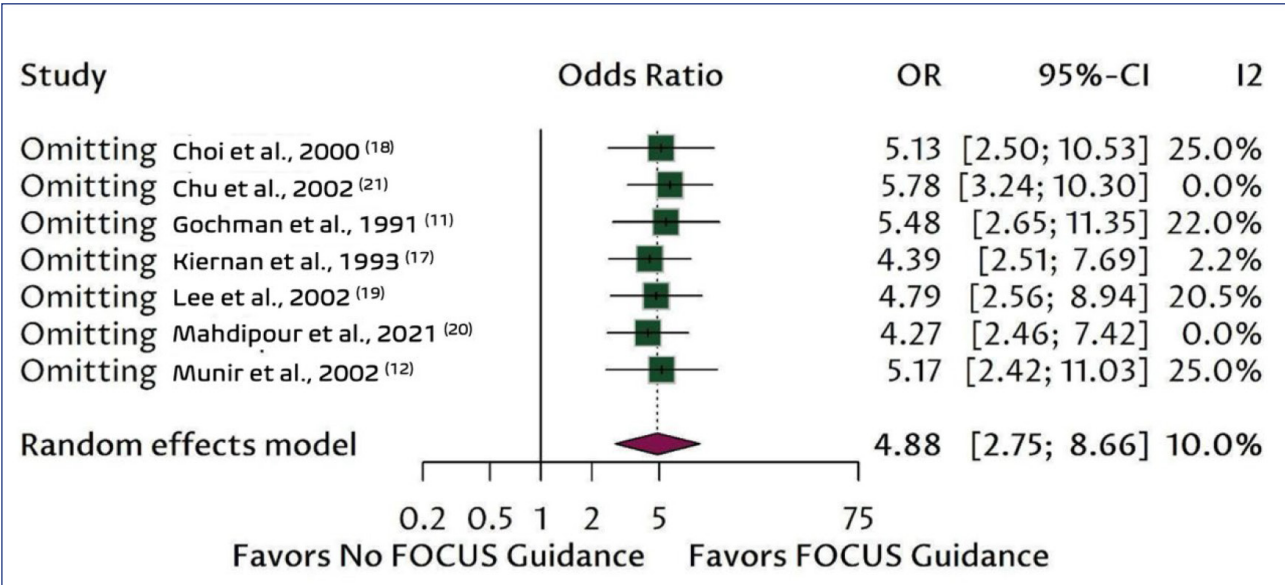


Figure 6. Sensitivity analysis of the overall success rate, examining the stability of the pooled overall success rate for suprapubic bladder aspiration. By sequentially omitting one study at a time, the analysis evaluated the impact of each individual study on the combined estimate, aiming to detect potential outliers or influential studies.

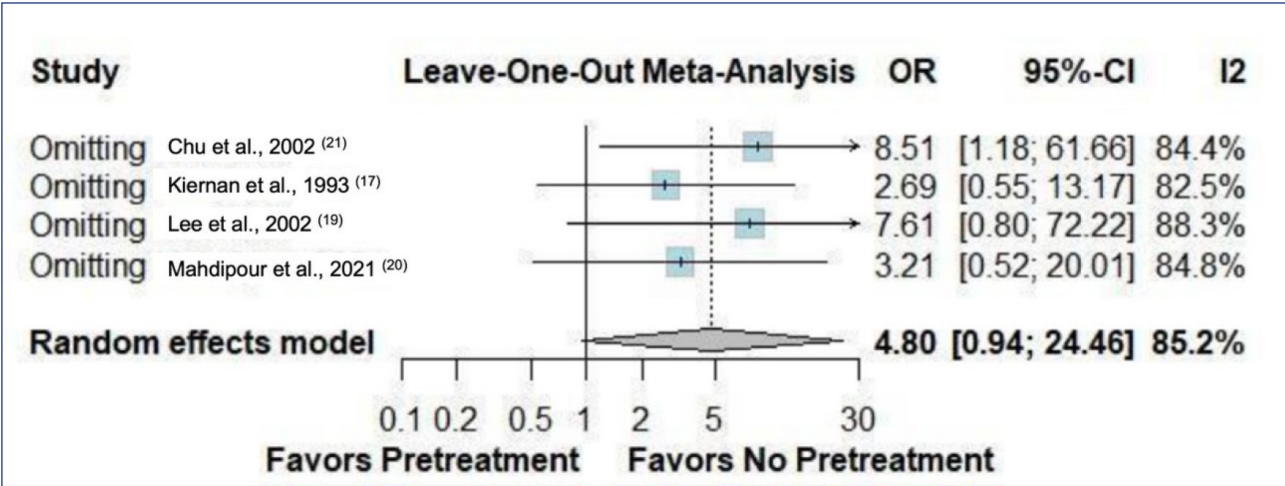


Figure 7. Sensitivity analysis of success at first attempt, evaluating the robustness of the pooled estimate for first-attempt success in suprapubic bladder aspiration. Each iteration involves the exclusion of one study at a time to examine its influence on the overall effect size. The analysis helps identify whether any single study disproportionately affects the pooled outcome.

perience, which may influence success rates and adoption of the method. Studies conducted in Brazil and Saudi Arabia, highlight a significant gap in medical education regarding POCUS training. A study conducted in Saudi Arabia showed that only 21.4% of final-year medical students had received formal POCUS training, revealing a deficiency in ultrasound proficiency among future physicians⁽²³⁾. Similarly, research involving members of the Brazilian Society of Nephrology revealed that the majority (64%) of participants did not have the opportunity to develop ultrasound examination skills during their medical

training⁽²⁴⁾. Despite the well-documented benefits of POCUS, few medical programs offer structured training in this skill, emphasizing the urgent need to integrate ultrasound education into medical curricula to enhance the diagnostic and procedural capabilities of physicians⁽²⁵⁾. In addition, there were notable methodological discrepancies among the trials included. The fact that only one study—that conducted by Mahdipour et al.⁽²¹⁾—explicitly reported which health care professional performed each procedure limited our ability to assess the influence of operator expertise.

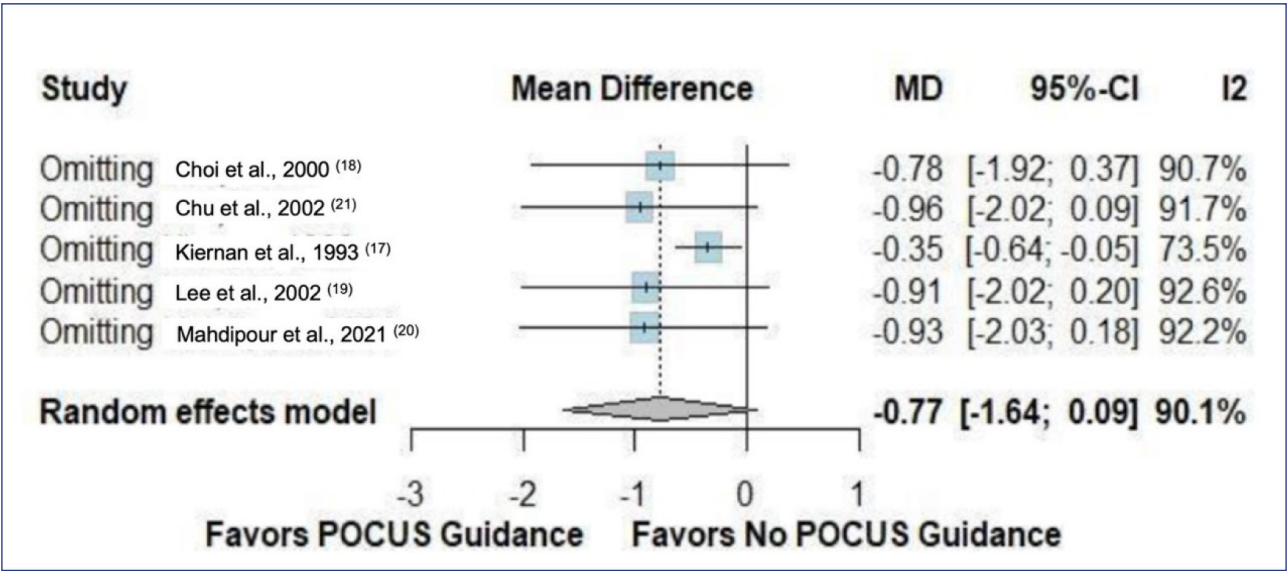


Figure 8. Sensitivity analysis of the mean number of attempts, assessing the robustness of the pooled mean number of attempts across the studies included. Variations in effect estimates were explored by sequentially excluding individual studies.

	Risk of bias					Overall
	D1	D2	D3	D4	D5	
Choi et al. (18)	+	+	+	+	+	+
Chu et al. (21)	+	+	+	+	+	+
Gochman et al. (11)	+	+	+	+	+	+
Kiernan et al. (17)	+	+	+	+	+	+
Lee et al. (19)	+	+	+	+	+	+
Mahdipour et al. (20)	+	+	+	+	+	+
Munir et al. (12)	+	+	+	+	+	+

Judgement
+ Low

Figure 9. Risk of bias summary, providing a visual overview of the risk of bias across the included studies, evaluating domains such as randomization process, deviations from intended interventions, missing outcome data, measurement of the outcome, and selection of the result reported.

Availability is another important factor that may limit the widespread implementation of POCUS in SPA. While ultrasound devices are becoming more accessible, their availability remains inconsistent across different health care settings, particularly in low-resource settings⁽¹³⁾. Institutions with limited funding may lack portable ultrasound machines or may not have enough trained personnel to ensure real-time imaging during SPA procedures. In contrast, traditional (blind) SPA, although less precise, does not require additional equip-

ment. Therefore, cost-effectiveness analyses comparing POCUS-guided SPA with alternative urine collection methods are warranted in order to guide policy decisions and optimize resource allocation.

This study has some limitations that should be considered. The majority of the studies included were published between 1991 and 2002, with the notable exception of that conducted by Mahdipour et al.⁽²¹⁾, which was published in 2021. Given the advances in technology and widespread use of POCUS, including

integration of artificial intelligence, generalizability and contemporary applicability may be limited⁽²⁶⁾. However, these technological innovations may have further improved SPA success rates, resulting in an underestimation of the impact of POCUS in our study. Bladder volume is also an important source of variability across studies, given that it represents an essential determinant of procedural success. Munir et al.⁽¹²⁾ demonstrated that the SPA success rate reaches nearly 90% when bladder volume exceeds 10 mL on ultrasound, highlighting the need for adequate bladder filling before the procedure. However, the remaining studies included in this meta-analysis did not apply a standardized bladder volume threshold, which may have contributed to the heterogeneity observed in diagnostic accuracy. Future methodologically rigorous RCTs should adopt consistent definitions and measurement protocols for bladder volume to more accurately evaluate its influence on the performance and success of ultrasound-guided SPA.

The accuracy of POCUS for SPA in infants depends on the technical capabilities of the ultrasound system. Older-generation machines with low resolution, analog interfaces, and minimal pediatric optimization likely increase reliance on operator skill and contribute to increasing the number of failed or uncertain punctures. In contrast, newer digital systems, introduced in the early 2000s, offer better image quality and usability, improving the effectiveness and ease of the procedure. In comparison, the device used by Gochman et al.⁽¹¹⁾ in their study published in 1991 — the ScanMate II ultrasound system — was limited to a 60° scan angle, and a 10-cm depth range. In contrast, Mahdipour et al.⁽²¹⁾, in their 2021 study, used the Ultrasonix platform, which offered significantly higher resolution and advanced technologies like SonixDAQ, enabling over 10,000 frames per second and photoacoustic imaging. These technological developments appear to improve image clarity and facilitate real-time targeting, as well as potentially reducing operator dependency during procedures such as SPA. Enhanced bladder visualization, including in small infants, could potentially increase procedural accuracy. Such evolving capabilities may, in part, contribute to the heterogeneity observed across studies as ultrasound technology has advanced over time.

Heterogeneity was observed, particularly in first-attempt success rates ($I^2 = 86\%$; $p > 0.001$), potentially due to variations in operator expertise, ultrasound equipment, and reason for aspiration. For example, Gochman et al.⁽¹¹⁾ and Kiernan et al.⁽¹⁸⁾ used the judgement of the

medical team to determine the reason for aspiration, whereas Chu et al.⁽²²⁾ used objective criteria: a urine sample testing positive for leukocyte esterase or nitrites; or the presence of leukocytes ($> 5 \text{ mm}^3$) on microscopic examination. Although the mean number of attempts needed to achieve a successful SPA was lower among the POCUS-guided procedures than among the traditional procedures, statistical significance was not reached. This outcome appears to be influenced by substantial heterogeneity, primarily driven by a single outlier study — that conducted by Kiernan et al.⁽¹⁸⁾ — the results of which were markedly divergent from those of the other studies included in this review. Sensitivity analysis with the leave-one-out method suggests that exclusion of this study would have resulted in a statistically significant difference favoring POCUS, underscoring the considerable impact of the outlier on the overall findings.

Although the present meta-analysis provides valuable insights, it is important to acknowledge that most of the studies included are relatively dated and exhibit considerable methodological heterogeneity, which may limit the generalizability and current applicability of the results. Future research should prioritize the development of well-designed, contemporary RCTs that address these limitations. In particular, key methodological aspects such as operator expertise, ultrasound technology, and bladder volume at the time of assessment should be standardized or systematically evaluated to ensure that the findings are more reliable and reproducible across studies.

5. CONCLUSION

This meta-analysis demonstrates the superiority of POCUS-guided SPA over traditional blind SPA in terms of the overall success rates of the procedure in pediatric patients. Compared with blind SPA, POCUS-guided SPA remains a more successful approach for obtaining a sterile urine sample in children. Despite the heterogeneity observed among the studies included in this review, the overall evidence supports the integration of POCUS into routine pediatric practice.

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