

Techniques and Technologies to Improve Peripheral Intravenous Catheter Outcomes in Pediatric Patients: Systematic Review and Meta-Analysis

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OBJECTIVE: Insertion and function of pediatric peripheral intravenous catheters (PIVCs) present challenges. We systematically reviewed techniques and technologies to improve PIVC outcomes (first-time insertion success, overall insertion success, time to insertion, dwell time, failure, and complications).

DATA SOURCES: Cochrane Central Register of Controlled Trials (CONTROL), Cumulative Index to Nursing and Allied Health (CINAHL), US National Library of Medicine, and Embase.

STUDY SELECTION: English-language pediatric trials published post 2010 reporting PIVC outcomes.

DATA EXTRACTION: Following Cochrane standards, two authors screened, extracted, and critiqued study quality (Grading of Recommendations Assessment, Development and Evaluation approach) data, random effects analysis, results expressed as risk ratios (RR), mean differences (MD) and 95% CIs.

RESULTS: Twenty-one studies (3237 children; 3098 PIVCs) were included. First-time insertion success significantly

increased with ultrasound guidance (compared with landmark insertion; RR, 1.60; 95% CI, 1.02-2.50). Use of ultrasound guidance (compared with landmark insertion) did not improve overall PIVC insertion success (RR, 1.10; 95% CI, 0.94-1.28). There was no evidence of an effect of near-infrared (compared with landmark) on first-time insertion success (RR, 1.21; 95% CI, 0.91-1.59) or number of attempts (MD, -0.65; 95% CI, -1.59 to 0.29); however, it significantly reduced PIVC insertion time (MD, -132.47; 95% CI, -166.68 to -98.26) and increased first-time insertion success in subgroup analysis of patients with difficult intravenous access (RR, 2.72; 95% CI, 1.02-7.24).

LIMITATIONS: Few studies per intervention, small sample sizes, and inconsistent outcome measures precluded definitive conclusions.

CONCLUSIONS: Ultrasound and near-infrared appear to improve pediatric PIVC insertion. High-quality studies examining the full extent of techniques and technologies are needed. Registration: CRD42020175314 *Journal of Hospital Medicine* 2021;16:XXX-XXX. © 2021 Society of Hospital Medicine

Peripheral intravenous catheters (PIVCs) are fundamental to the healthcare practitioners' ability to provide vital intravenous fluids, medications, and blood products, and as a prophylactic measure prior to some procedures, making insertion of these devices the most common in-hospital invasive procedure in pediatrics.^{1,2} Despite the prevalence and ubiquity of PIVCs,¹ successful insertion in pediatrics is problematic,³⁻⁵ and device dysfunction prior to completion of treatment is common.^{3,6} The inability to attain timely PIVC access and maintain postinsertion function has significant short- and long-term sequelae, including pain and anxiety for children and their parents,^{3,7} delays in treatment,³ prolonged hospitalization,⁸ and increased healthcare-associated costs.⁸⁻¹⁰

Approximately 50% of pediatric PIVC insertions are challenging, often requiring upwards of four insertion attempts,

and a similar proportion fail prior to treatment completion.^{3,11} Exactly why PIVC insertion is difficult in children, and the mechanisms of failure, are unknown. It is likely to be multifaceted and related to factors concerning the patient (eg, comorbidities, age, gender, adiposity),^{11,12} provider (eg, insertion practice, care, and maintenance),^{3,13,14} device (eg, size, length, catheter-to-vein ratio),^{15,16} and therapy (eg, vessel irritation).^{11,13,17} Observational studies and randomized controlled trials (RCTs) in hospitalized pediatric patients report that the average PIVC dwell is approximately 48 hours, suggesting multiple PIVCs are required to complete a single admission.^{3,18}

Conventionally, PIVC insertion involved physical assessment through palpation and visualization (landmark approach), and although postinsertion care varies among healthcare facilities, minimal requirements are a dressing over the insertion site and regular flushes to ensure device patency.^{1,3,19} Recently, clinicians have investigated insertion and management practices to improve PIVC outcomes. These can be grouped into techniques—the art of doing (the manner of performance, or the details, of any surgical operation, experiment, or mechanical act) and technologies—the application of scientific knowledge for practical purposes.²⁰ Individual studies have examined the outcomes of new techniques and technologies;

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however, an overall estimation of their clinical significance or effect is unknown.^{11,18} Therefore, the aim of this review was to systematically search published studies, conduct a pooled analysis of findings, and report the success of various techniques and technologies to improve insertion success and reduce overall PIVC failure.

METHODS

Design

The protocol for this systematic review was prospectively registered with PROSPERO (CRD42020165288). This review followed Cochrane Collaboration systematic review methods²¹ and was reported in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement.²²

Inclusion and Exclusion Criteria

Studies were eligible for inclusion if they met predefined criteria: (1) RCT design; (2) included standard-length PIVC; (3) participants aged 0 to 18 years, excluding preterm infants (less than 36 weeks' gestation); (4) required PIVC insertion in an inpatient healthcare setting; and (5) reported PIVC insertion outcomes (described below). Studies were excluded if they were cluster or crossover RCTs, published before 2010, or not written in English.

Interventions

Interventions were PIVC insertion and management techniques, defined as "the manner of performance, or the details, of any surgical operation, experiment, or mechanical act" (eg, needle-tip positioning, vein selection [site of insertion], comfort measures, and flushing regimen), or technologies, defined as "the application of scientific knowledge for practical purpose" (eg, vessel visualization, catheter material, and catheter design), compared with current practice, defined as commonly known, practiced, or accepted (eg, landmark PIVC insertion).²⁰

Primary and Secondary Outcomes

The primary outcome was first-time insertion success (one skin puncture to achieve PIVC insertion; can aspirate and flush PIVC without resistance).²³ Secondary outcomes included: (1) overall PIVC insertion success²³; (2) all-cause PIVC failure (cessation of PIVC function prior to treatment completion)⁶; (3) dwell time¹⁴; (4) PIVC insertion time; (5) insertion attempts²³; (6) individual elements of failure (dislodgement, extravasation, infection, occlusion, pain, phlebitis, and thrombosis)⁶; and (7) patient/parent satisfaction. Some outcomes evaluated were author defined within each study (patient/parent satisfaction, pain score).

Systematic Search

A search of the Cochrane Library and Central Register of Controlled Trials (CENTRAL), Cumulative Index to Nursing and Allied Health (CINAHL), US National Institutes of Health National Library of Medicine (PubMed), and Embase databases between 2010 to 2020 was undertaken on June 23, 2020, and

updated March 4, 2021. Medical Subject Heading (MeSH) terms and relevant keywords and their variants were used in collaboration with a healthcare librarian (Appendix Table 1). Additional studies were identified through hand searches of bibliographies.¹⁹ Studies were included if two authors (TMK and JS) independently agreed they met the inclusion criteria.

Data Extraction

Two authors (TMK/JS) independently abstracted study data using a standardized form managed in Microsoft Excel.

Quality Assessment

Included studies were assessed by two authors (TMK and JS) for quality using the Cochrane risk of bias (RoB2) tool.^{21,24} The overall quality of evidence for each outcome was assessed using the Grading of Recommendations Assessment, Development and Evaluation (GRADE)²⁵ approach. Individual RCTs began at high quality, downgraded by one level for "serious" or two levels for "very serious" study limitations, including high risk of bias, serious inconsistency, publication bias, or indirectness of evidence.

Data Analysis and Synthesis

Where two or more trials with evidence of study homogeneity (trial interventions and population) were identified, meta-analysis using RevMan 5 (version 5.4.1)²⁶ with random effects was conducted. Descriptive statistics summarized study population, interventions, and results. For dichotomous outcomes, we calculated risk ratio (RR) plus 95% CI. For continuous outcomes, we planned to calculate the mean difference (MD) plus 95% CI and the standardized mean difference (SMD) (difference between experimental and control groups across trials) reported as the summary statistic.

Subgroup analyses, where possible, included: difficult intravenous access (DIVA), defined by study authors; age (0-3 years; >3 years up to 18 years); hospital setting during PIVC insertion (awake clinical environment vs awake emergency department vs asleep operating room setting); and by operator (bedside nurse, anesthesiologist).

RESULTS

Search Strategy

Figure 1 describes study selection in accordance with the PRISMA guidelines.²² We identified 1877 records, and 18 articles met the inclusion criteria. An additional 3 studies were identified in the updated search, totaling 21 studies included in the final review.

Study Characteristics

Collectively, 3237 patients and 3098 successful PIVC insertions were reported. In the included studies, 139 patients did not receive a PIVC owing to failed insertion. Ten studies examined techniques (needle-tip positioning,²⁷ vein choice for PIVC insertion,²⁸ flushing regimen,²⁹⁻³¹ nonpharmacological^{32,33} dressing and securement,^{34,35} and pharmacological comfort measures³⁶), and 11 studies examined technologies

(vessel visualization including ultrasound,^{4,37-40} near-infrared [image of vein projected onto the skin],^{37,41-44} transillumination [transmission of light through the skin],⁴⁵ and catheter design⁴⁶). Table 1 outlines characteristics of included studies. Most trials were single center and conducted in an acute inpatient pediatric-specific setting^{4,27-34,36-41,44-46} or dedicated pediatric unit in a large public hospital^{35,43,44}; one study was a multicenter trial.³⁶ All trials described evidence of ethical review board approval and participant consent for trial participation.

Study Quality

The certainty of evidence at the outcome level varied from moderate to very low. Table 2 and Table 3 outline the summary of findings for landmark insertion compared with ultrasound-guided and landmark insertion compared with near-infrared PIVC insertion, respectively. The remaining summary-of-findings comparisons that included more than one study or addressed clinically relevant questions can be found in Appendix Tables 2, 3, 4, 5, 6, 7, and 8. At the individual study level, most domains were assessed as low risk of bias (Appendix Figure 1).

Effectiveness of Interventions

Technology to Improve PIVC Outcomes

Landmark compared with ultrasound-guided PIVC insertion. Five studies compared PIVC insertion success outcomes when traditional landmark technique was used in comparison with ultrasound guidance (Appendix Figure 2). Four studies (592 patients)^{4,37,38,40} assessed the primary outcome of first-time insertion success. Appendix Figure 2.1 demonstrates PIVCs were 1.5 times more likely to be inserted on first attempt when ultrasound guidance was used compared with landmark insertion (RR, 1.60; 95% CI, 1.02-2.50). When examining only studies that included DIVA,^{4,38,40} the effect size increased and CIs tightened (RR, 1.87; 95% CI, 1.56-2.24). No evidence of effect was demonstrated when comparing this outcome in children aged 0 to 3 years (RR, 1.39; 95% CI, 0.88-2.18) or >3 years (RR, 0.72; 95% CI, 0.35-1.51). Two studies^{4,38} demonstrated that first-time insertion success with ultrasound (compared with landmark) was almost twice as likely (RR, 1.87; 95% CI, 1.44-2.42) after induction of anesthesia in contrast to no effect in studies undertaken in the emergency department^{37,40} (RR, 1.32; 95% CI, 0.68-2.56). One study³⁹ (339 patients) reported the secondary outcomes of extravasation/infiltration and phlebitis. Extravasation/infiltration was nearly twice as likely with ultrasound compared with landmark insertion (RR, 1.80; 95% CI, 1.01-3.22); however, there was

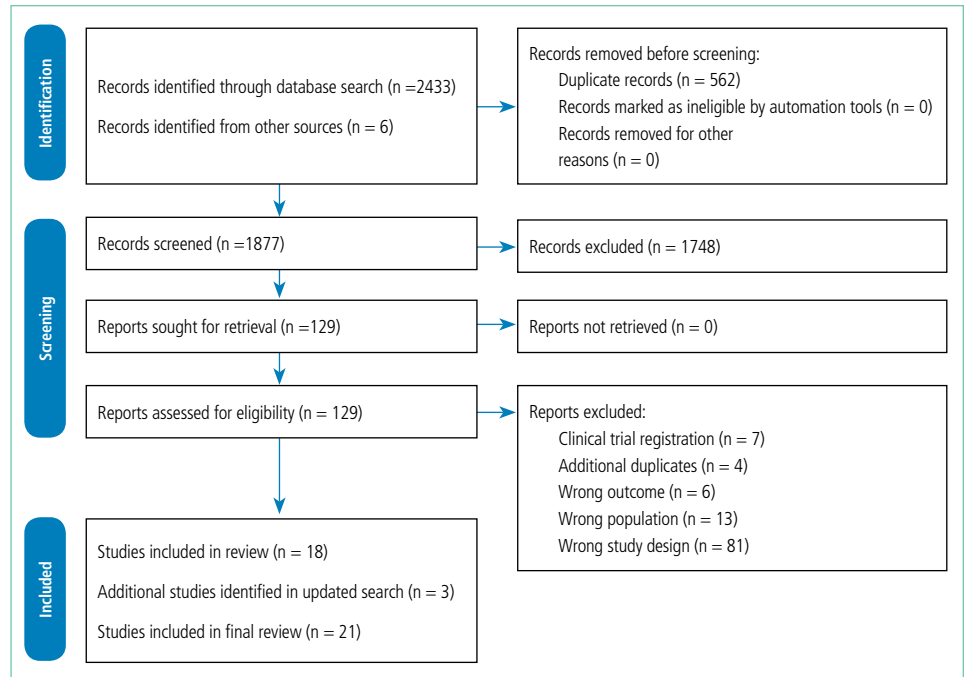


FIG. PRISMA Flow Chart

no evidence of effect related to phlebitis (RR, 0.32; 95% CI, 0.07-1.50).

Four studies^{4,38-40} compared the review's secondary outcome of PIVC insertion success (Appendix Figure 2.2), with no evidence of an effect (RR, 1.10; 95% CI, 0.94-1.28). No improvement in overall insertion success was demonstrated in the following subgroup analyses: patients with DIVA (RR, 1.18; 95% CI, 0.95-1.47), children under 3 years of age (RR, 1.23; 95% CI, 0.90-1.68), and PIVCs inserted by anesthesiologists (RR, 1.25; 95% CI, 0.91-1.72). One study measured this outcome in children aged >3 years (RR, 1.13; 95% CI, 0.99-1.29) with no effect and in the emergency department (RR, 1.09; 95% CI, 1.00-1.20), where ultrasound guidance improved overall PIVC insertion success.

Landmark compared with near-infrared PIVC insertion. First-time insertion success (Appendix Figure 3.1) was reported in five studies^{37,41-44} and 778 patients with no evidence of effect (RR, 1.21; 95% CI, 0.91-1.59). Subgroup analysis by DIVA⁴¹⁻⁴⁴ demonstrated first-time insertion success more than doubled with near-infrared technology compared with landmark (RR, 2.72; 95% CI, 1.02-7.24). Subgroup analysis by age did not demonstrate an effect in children younger than 3 years or children older than 3 years. Subgroup analysis by clinician inserting did not demonstrate an effect. Of the five studies reporting time to insertion,^{37,41-44} two^{41,42} reported median rather than mean, so could not be included in the analysis. Of the remaining three studies,^{37,43,44} near-infrared reduced PIVC time to insertion (Appendix Figure 3.2).

Four studies^{37,42-44} reported the number of attempts required for successful PIVC insertion where no difference was detected; however, subgroup analysis of patients with DIVA^{43,44} and insertion by bedside nurse^{43,44} demonstrated fewer PIVC inser-

TABLE 1. Key Characteristics of Included Studies

Author, country	Method	Participants	Interventions	Outcomes	Notes
A. Technologies					
Avelar et al ³⁹ Brazil	Single-center RCT	302 patients (aged 1 d-18 y) admitted to surgical inpatient unit	Ultrasound-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; catheter dwell time; incidence of infiltration and phlebitis	
Benkhadra et al ³⁸ France	Single-center RCT	40 patients (aged <3 y) with PIVC insertion post-inhalational anesthesia induction	Ultrasound-guided PIVC insertion Landmark PIVC insertion	Overall PIVC insertion success; first-time insertion success; time to cannulation; number of insertion attempts; type of catheter used	
Curtis et al ³⁷ Canada	Single-center RCT	418 patients (up to age 16 y) presenting to the emergency department	Ultrasound-guided PIVC insertion Near-infrared-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; number of insertion attempts; time to successful PIVC insertion	Funding support from the Women's and Children's Health Research Institute, Stollery Children's Hospital Foundation
Demir and Inal ⁴⁴ Turkey	Single-center RCT	129 patients (aged 3-8 y) with peripheral intravenous access assessed as easy (n=50), intermediate (n=35), and difficult (n=44) ⁵¹ admitted to the pediatric unit	Near-infrared-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; number of insertion attempts; time to successful PIVC insertion	
Gümüş and Basbakkal ⁴⁵ Turkey	Single-center RCT	112 patients (aged 1-10 y) presenting to the emergency department	Transilluminator-guided PIVC insertion Landmark-guided PIVC insertion	First-time insertion success; number of insertion attempts; time to successful PIVC insertion	
Hanada et al ⁴ United States	Single-center RCT	102 patients (weight ≥3 kg and aged <4 y); operating room suite	Ultrasound-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; success rate of PIVC insertion within 10 min	
Inal and Demir ⁴³ Turkey	Single-center RCT	54 patients (aged, 0-36 mo) with peripheral intravenous access assessed as easy (n=20), intermediate (n=12), and difficult (n=22) ⁵¹ admitted to the pediatric unit	Near-infrared-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; number of insertion attempts; time to successful PIVC insertion	
Kaddoum et al ⁴² United States	Single-site RCT	146 patients (aged 0-18 y) scheduled for elective surgery; PIVCs inserted after inducing a state of anesthesia with anesthetic gas	Near-infrared-guided PIVC insertion Landmark PIVC insertion	Number of insertion attempts; time to successful PIVC insertion	
Kim et al ⁴¹ Korea	Single-site RCT	111 patients (aged 1 mo-16 y) with peripheral intravenous access assessed as easy (n=36), intermediate (n=28), and difficult (n=47) ⁵¹ in inpatient pediatric unit	Near-infrared-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; time to successful PIVC insertion	
Qin et al ⁴⁶ Australia	Single-center pilot RCT	72 children (aged 1-17 y) undergoing surgery and requiring 48 h of postoperative intravenous therapy	Long PIVC: received an 8-cm (3.1-in) 22-G Leaderflex catheter (Vygon GmbH & Co.) Standard PIVC: 2.5-cm (1-in) 22-G PIVC (Introcan Safety; B. Braun)	Catheter failure due to any complication; individual elements of catheter complication, including infiltration, phlebitis, dislodgement, and occlusion	
Vinograd et al ⁴⁰ United States	Single-center RCT	167 pediatric patients in an urban tertiary pediatric emergency department and requiring PIVC insertion	Ultrasound-guided PIVC insertion Landmark PIVC insertion	First-time insertion success; time to insertion; number of insertion attempts; overall PIVC dwell; PIVC complications; parental satisfaction	

Continued on page E4

tion attempts and a reduction in insertion time, respectively, with the use of near-infrared technology (Appendix Figure 3.3).

Landmark compared with transillumination PIVC insertion. One study⁴⁵ (112 participants) found a positive effect with the use of transillumination and first-time insertion success (RR, 1.29; 95% CI, 1.07-1.54), reduced time to insertion (MD, -9.70; 95% CI, -17.40 to -2.00), and fewer insertion attempts (MD, -0.24; 95% CI, -0.40 to -0.08) compared with landmark insertion.

Long PIVC compared with short PIVC. A single study⁴⁶ demonstrated a 70% reduction in PIVC failure (RR, 0.29; 95% CI, 0.14-0.59) when long PIVCs were compared with standard PIVCs. Specifically, PIVC failure due to infiltration was reduced with the use of a long PIVC (RR, 0.08; 95% CI, 0.01-0.61). There was no difference in insertion success (RR, 1.00; 95% CI, 0.95-1.05) or phlebitis (RR, 1.00; 95% CI, 0.07-15.38).

Technique to Improve PIVC Outcomes

Static ultrasound-guided compared with dynamic needle-tip PIVC insertion. In a single study comparing variation in ultrasound-guided PIVC insertion technique²⁷ (60 patients), dynamic needle-tip positioning improved first-time insertion success (RR, 1.44; 95% CI, 1.04-2.00) and overall PIVC insertion success (RR, 1.42; 95% CI, 1.06-1.91).

Variation in vein choice for successful PIVC insertion. Insertion of PIVC in the cephalic vein of the forearm improved insertion success in a single study²⁸ of 172 patients compared with insertion in the dorsal vein of the hand (RR, 1.39; 95% CI, 1.15-1.69) and great saphenous vein (RR, 1.27; 95% CI, 1.08-1.49).

Variation in PIVC flush. Heparinized saline compared with 0.9% sodium chloride flush²⁹ did not reduce infiltration (RR, 0.31; 95% CI, 0.03-2.84), occlusion (RR, 1.88;

TABLE 1. Key Characteristics of Included Studies (continued)

Author, country	Method	Participants	Interventions	Outcomes	Notes
B. Techniques					
Buyukyilmaz et al ³⁵ Turkey	Single-center RCT	60 pediatric patients (aged 2-24 mo) in pediatric clinical unit within a large public hospital undergoing PIVC insertion into metacarpal vein to infuse penicillin to treat lower respiratory tract infection	1. Primary PIVC dressing 2. Primary PIVC dressing + protective covering (I.V. House)	PIVC dwell time; phlebitis	
Hartling et al ³³ Canada	Single-center RCT	42 patients (aged 3-11 y) presenting to the emergency department	1. No intervention 2. Music as method of distraction	PIVC insertion success; patient distress measured using the OSBD-R ⁵² ; change in self-reported pain; heart rate; parent and healthcare provider satisfaction	
Kleidon et al ³¹ Australia	Single-center pilot 2×2 factorial RCT	55 pediatric patients (aged birth-18 y) who were medical or surgical inpatients	1. Low-frequency flush: every 24 h 2. High-frequency flush: every 6 h 3. High-volume flush: 10 mL sodium chloride 4. Low-volume flush: 3 mL sodium chloride	Feasibility, including parent or caregiver and staff satisfaction; PIVC failure and individual elements of PIVC failure due to infection, occlusion, infiltration, dislodgement, phlebitis; PIVC dwell; safety	Funding through unrestricted industry grant
Kleidon et al ³⁴ Australia	Single-center pilot RCT	330 patients (aged birth-18 y) who were medical or surgical inpatients	1. BPU dressing (TegadermAdvanced 1683 or 1682; 3M) 2. Integrated securement dressing SorbaView SHIELD micro SV226UDT or small SV254UDT (Centurion Medical Products Corporation) 3. Tissue adhesive (Histoacryl; B. Braun Melsungen AG) covered with TegadermAdvanced (3M)	Feasibility including parent or caregiver and staff satisfaction; PIVC failure; individual elements of PIVC failure due to infection, occlusion, infiltration, dislodgement, phlebitis; PIVC dwell; safety	Funding through unrestricted industry grant
Schmitz et al ³⁶ United States	Multisite randomized, double-blind, placebo-controlled trial	504 pediatric patients (aged 3-18 y) who required venous cannulation at the antecubital fossa or back of hand as part of their standard medical care	1. Active lidocaine spray (ie, needle-free powder lidocaine delivery system) 2. Sham placebo group: contained an identical delivery system without lidocaine	PIVC insertion success; analgesic efficacy; safety assessment, including skin damage	Funding for trial and other support, including trial design, interpretation of data, and statistical support, from Anesiva, a late-stage biopharmaceutical company
Schreiber et al ³⁰ Italy	Single-center RCT	198 pediatric patients (aged 1-17 y) who required intravenous access for at least 24 h without continuous infusion	1. Low-frequency flush: PIVC flushed with 0.9% prefilled sodium chloride syringe every 24 h 2. High-frequency flush: PIVC flushed with 0.9% prefilled sodium chloride syringe every 12 h	Catheter patency; extravasation; pain; erythema; swelling	
Takeshita et al ²⁷ Japan	Single-center RCT	60 pediatric patients (aged <2 y) in the pediatric intensive care unit who required insertion of a PIVC	1. Dynamic needle tip 2. Static needle tip	PIVC insertion success; first-time insertion success; PIVC insertion within 10 min; number of PIVC insertion attempts	
Takeshita et al ²⁸ Japan	Single-center RCT	172 pediatric patients (weight <20 kg) scheduled for elective surgery; PIVCs were inserted after inducing a state of anesthesia with anesthetic gas	1. Ultrasound-guided PIVC insertion at dorsum of hand 2. Ultrasound-guided PIVC insertion at cephalic vein of forearm 3. Ultrasound-guided PIVC insertion at great saphenous vein	PIVC insertion success; number of PIVC insertion attempts; time to successful PIVC insertion	
White et al ²⁹ United States	Single-center RCT	62 children (aged 1 mo-17 y) admitted to a large urban hospital and who required insertion of PIVC	Heparinized saline flush/normal saline flush	PIVC patency; redness; swelling; blood in catheter; bruising; leakage; pain; burning; clotting; infiltration	
Wolyniez et al ³² Israel	Single-center pilot RCT	47 children (aged 3-16 y) who required insertion of PIVC or venipuncture in the emergency department, excluding children with developmental disabilities	1. Medical clown distraction 2. Usual departmental procedural distraction	PIVC insertion success; pain score; adult anxiety; number of insertion attempts; time to successful PIVC insertion	

Abbreviations: BPU, bordered polyurethane; OSBD-R, Observational Score of Behavioural Distress-revised; PIVC, peripheral intravenous catheter; RCT, randomized controlled trial.

95% CI, 0.18-19.63) during dwell, or hematoma (RR, 0.94; 95% CI, 0.06-14.33) at insertion.

Two studies^{30,31} (253 participants) compared PIVC flush frequency (daily compared with more frequent flush regimes). There was no reduction in overall PIVC failure, extravasation/infiltration, phlebitis, or occlusion during dwell (Appendix Fig-

ure 4.1-4.4). Additionally, no effect was demonstrated when a single study³¹ investigated volume of flush on extravasation/infiltration, dislodgement, phlebitis, or occlusion.

Variation in dressing and securement. One trial (330 participants)³⁴ demonstrated that integrated securement and dressing (ISD) product reduced PIVC failure (RR, 0.65;

TABLE 2. Summary of Findings: Landmark Compared With Ultrasound-Guided PIVC Insertion

Patient or population: Pediatrics
 Setting: Hospitalized pediatric patients
 Intervention: Landmark insertion
 Comparison: Ultrasound-guided insertion

Outcomes	Anticipated absolute effects ^a (95% CI)		Relative effect (95% CI)	No. of participants (studies)	Certainty of the evidence (GRADE) ^b
	Risk with ultrasound-guided insertion	Risk with landmark insertion			
First-time insertion success	790/1000	498/1000 (316-775)	RR, 0.63 (0.40-0.98)	592 (4 RCTs)	⊗○○○ (Very low) ^{c,d}
Overall insertion success	895/1000	814/1000 (698-957)	RR, 0.91 (0.78-1.07)	691 (4 RCTs)	⊗○○○ (Very low) ^{c,d}
Extravasation/infiltration	129/1000	211/1000 (130-323)	OR, 1.80 (1.01-3.22)	339 (1 RCT)	⊗⊗○○ (Moderate) ^d
Phlebitis/pain	39/1000	13/1000 (3-59)	RR, 0.32 (0.07-1.50)	339 (1 RCT)	⊗⊗○○ (Low) ^e
Time to insertion	The mean time to insertion was 0	MD, 0.9 higher (0.03 higher to 1.77 higher)	–	283 (1 RCT)	⊗⊗○○ (Low) ^{d,f}

^aThe risk in the intervention group (and its 95% CI) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).
^bGRADE Working Group grades of evidence:
 High certainty: we are very confident that the true effect lies close to that of the estimate of the effect.
 Moderate certainty: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.
 Low certainty: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.
 Very low certainty: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.
^cDowngraded 2 levels: inconsistency (heterogeneity >75%).
^dDowngraded 1 level: imprecision (small sample size).
^eDowngraded 2 levels: imprecision (small sample size and wide CI).
^fDowngraded 1 level: indirectness (incomplete outcome measure).
 Abbreviations: GRADE, Grading of Recommendations Assessment, Development and Evaluation; MD, mean difference; OR, odds ratio; PIVC, peripheral intravenous catheter; RCT, randomized controlled trial; RR, relative risk.

TABLE 3. Summary of Findings: Landmark Compared With Near-Infrared PIVC Insertion

Patient or population: Pediatrics
 Setting: Hospitalized pediatric patients
 Intervention: Landmark insertion
 Comparison: Near-infrared insertion

Outcomes	Anticipated absolute effects ^a (95% CI)		Relative effect (95% CI)	No. of participants (studies)	Certainty of the evidence (GRADE) ^b
	Risk with near- infrared insertion	Risk with landmark insertion			
First-time insertion success	687/1000	570/1000 (433-749)	RR, 0.83 (0.63-1.09)	778 (5 RCTs)	⊗○○○ (Very low) ^{c,d,e,f}
Time to insertion	The mean time to insertion was 0	MD, 132.47 lower (166.68 lower to 98.26 lower)	–	183 (2 RCTs)	⊗○○○ (Very low) ^{c,f}
Number of attempts	The mean number of attempts was 0	MD, 0.45 lower (1.6 lower to 0.7 higher)	–	556 (3 RCTs)	⊗○○○ (Very low) ^{c,d,f}

^aThe risk in the intervention group (and its 95% CI) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).
^bGRADE Working Group grades of evidence:
 High certainty: we are very confident that the true effect lies close to that of the estimate of the effect.
 Moderate certainty: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.
 Low certainty: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.
 Very low certainty: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.
^cDowngraded 2 levels: risk of bias (random sequence generation and allocation concealment).
^dDowngraded 1 level: inconsistency (heterogeneity >75%).
^eDowngraded 1 level: indirectness (incomplete outcome definitions).
^fDowngraded 1 level: imprecision (small sample size).
 Abbreviations: GRADE, Grading of Recommendations Assessment, Development and Evaluation; MD, mean difference; RCTs, randomized controlled trials; RR, relative risk.

95% CI, 0.45-0.93) and occlusion (RR, 0.35; 95% CI, 0.13-0.94) compared with bordered polyurethane (BPU). There was no difference in the proportion of PIVC failure between BPU compared with tissue adhesive (TA) (RR, 0.74; 95% CI, 0.52-1.06). When comparing individual elements of PIVC failure, there was no evidence of effect between BPU and ISD in reducing infiltration (RR, 0.74; 95% CI, 0.43-1.27), dislodgement (RR, 0.49; 95% CI, 0.15-1.58), or phlebitis/pain (RR, 0.54; 95% CI, 0.21-1.39); similarly, the use of TA compared with BPU did not reduce failure due to infiltration (RR, 0.78; 95% CI, 0.45-1.33), dislodgement (RR, 0.37; 95% CI, 0.10-1.35), occlusion (RR, 0.91; 95% CI, 0.45-1.84), or phlebitis/pain (RR, 0.42; 95% CI, 0.17-1.05).

A comparison of protective covering³⁵ (60 participants) did not demonstrate a significant improvement in PIVC dwell (RR, 0.83; 95% CI, 0.25-1.41).

Pharmacological and nonpharmacological interventions.

A comparison of nonpharmacological comfort techniques, including music during insertion (one trial, 42 participants), did not improve first-time insertion success between the two groups (RR, 0.74; 95% CI, 0.53-1.03). Similarly, incorporation of a clown³² (47 patients) as method of distraction did not demonstrate an effect on PIVC insertion success (RR, 0.90; 95% CI, 0.77-1.06) or time to PIVC insertion (MD, -0.20; 95% CI, -1.74 to 1.34). In a double-blinded, placebo-controlled RCT³⁶ of pharmacological techniques to reduce PIVC insertion-related pain (504 participants), no evidence of effect was established between the placebo control group and the active analgesia in overall PIVC insertion success (RR, 1.01; 95% CI, 0.97-1.04).

DISCUSSION

Despite their pervasiveness, PIVC insertion in children is problematic and premature device failure is common, yet effective strategies to overcome these challenges have not been systematically reviewed to date. This systematic review (including meta-analysis) examines techniques and technologies to improve PIVC insertion success and reduce overall failure. We demonstrated ultrasound-guided PIVC insertion significantly improved first-time insertion success in general pediatrics.

Analogous to a previous systematic review in adult patients (1660 patients, odds ratio, 2.49; 95% CI, 1.37-4.52; $P = .003$; $P, 69%$),⁴⁷ we confirm ultrasound improves first-time PIVC insertion success, most notably in pediatric patients with difficult intravenous access. However, widespread use of ultrasound-guided PIVC insertion is limited by operator skills, as it requires practice and dexterity, especially for DIVA patients.^{5,47} Healthcare facilities should prioritize teaching and training to support acquisition of this skill to reduce the deleterious effects of multiple insertion attempts, including vessel damage, delayed treatment, pain, and anxiety associated with needles.

Other vessel-visualization technologies (near-infrared and transillumination) did not improve PIVC insertion in generic pediatrics.⁵ However, they significantly improved first-time insertion, time to insertion, and number of insertion attempts in patients with DIVA and should be considered in the absence of ultrasound-proficient clinicians.

Although vessel-visualization technologies provide efficient PIVC insertion, complication-free PIVC dwell is equally important. Few studies examined both insertion outcomes and PIVC postinsertion outcomes (dwell time and complications during treatment). One study reported more postinsertion complications (eg, infiltration) with ultrasound compared with landmark technique.³⁹ Vessel-visualization tools should be used to assess the vein to guide PIVC choice. Pandurangadu et al¹⁵ reported increased PIVC failure when less than 65% of the catheter length resides within the vein; this was consistent with the single RCT⁴⁶ included in this review that demonstrated reduced infiltration with long PIVCs compared with standard-length PIVCs. To reduce this knowledge practice gap, it is critical that clinicians continue to evaluate and publish findings of novel techniques to improve PIVC outcomes.

The review findings have important implications for future research, clinical practice, and policy. Unlike earlier reviews,⁴⁸ vessel-visualization technologies, particularly ultrasound, improved PIVC insertion success; however, during-dwell outcomes were inconsistently reported, and future research should include these. In addition, while there is evidence to support these new technologies, adequate training and resources to ensure a sustained, skilled workforce to optimize PIVC insertion are necessary for successful implementation.

Our study had some limitations, including the methodological quality of included studies (small sample size and significant clinical and statistical heterogeneity). Subgroup analyses were undertaken to reduce the heterogeneity inherent in pediatric populations; however, future studies should stratify for patient (age, DIVA, indication for insertion) and setting (conscious/unconscious, emergent/nonemergent) factors. Incomplete or absent outcome definitions and varied reporting measures (eg, median vs mean) prevented calculation of the pooled incidence of catheter failure and dwell time.

Our review also has notable strengths. Two independent investigators performed a rigorous literature search. Only RCTs were included, ensuring the most robust methods to inform clinically important questions. The primary and secondary outcomes were derived from patient-centered outcomes.

CONCLUSION

This systematic review and meta-analysis describes the pooled incidence of PIVC insertion success and outcomes, including complication and failure in pediatric patients. PIVC insertion with ultrasound should be used to improve insertion success in generic pediatric patients, and any form of vessel-visualization technology (ultrasound, near-infrared, transillumination) should be considered for anticipated difficult insertions.

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