DOI: 10.1111/iwj.13393

ORIGINAL ARTICLE

WILEY

Are high-risk patient and revision arthroplasty effective indications for closed-incisional negative-pressure wound therapy after total hip or knee arthroplasty? A systematic review and meta-analysis

Jun-Ho Kim¹ | Dae-Hee Lee²

¹Department of Orthopedic Surgery, Seoul Medical Center, Seoul, South Korea

²Department of Orthopaedic Surgery, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul, South Korea

Correspondence

Dae-Hee Lee, MD, PhD, Department of Orthopaedic Surgery, Samsung Medical Center, Sungkyunkwan University School of Medicine, 81 Irwon-ro, Gangnam-gu, Seoul 06351, South Korea. Email: eoak22@empal.com

Abstract

To determine the effective indications of closed-incisional negative-pressure wound therapy (ciNPWT) following total hip or knee arthroplasty, this systematic review and meta-analysis was conducted. The systematic search was performed on MEDLINE, Embase, and Cochrane Library, and 11 studies were included. The studies comparing between ciNPWT and conventional dressings were categorised into following subgroups based on patient risk and revision procedures: routine vs high-risk patient; primary vs revision arthroplasty. Pooled estimates were calculated for wound complication and surgical site infection (SSI) rates in the subgroup analyses using Review Manager. In highrisk patients, the overall rates of wound complication (odds ratio [OR] = 0.38; 95% confidence interval [CI] 0.15-0.93; P = .030) and SSI (OR = 0.24; 95% CI = 0.09-0.64; P = .005) were significantly lower in the ciNPWT; however, there were no differences in routine patients. In cases involving revision arthroplasties, the overall rates of wound complication (OR = 0.33; 95% CI = 0.18-0.62; P < .001 and SSI (OR = 0.26; 95% CI = 0.11-0.66; P = .004) were significantly lower in the ciNPWT; however, there were no differences in cases involving primary arthroplasties. In summary, ciNPWT showed a positive effect in decreasing the rates of wound complication and SSI in high-risk patients and in revision arthroplasties.

KEYWORDS

closed-incisional negative-pressure wound therapy, surgical site infection, total hip arthroplasty, total knee arthroplasty, wound complication

1 | INTRODUCTION

Total hip arthroplasty (THA) and total knee arthroplasty (TKA) are the most common and successful operations in modern medicine.^{1,2} However, persistent surgical site complications (SSCs), such as wound complications and surgical site infection (SSI) after THA and TKA, are

major sources for periprosthetic joint infection (PJI) and remain concerns to orthopaedic surgeons.³⁻⁶ Despite the low incidence, deep PJI has a devastating impact on not only the heath burden but also the distress or the economic burden to patients.^{3,7-9}

Given its considerable burden, great efforts to identify preoperative risk factors and to prevent SSCs and PJI in various ways have been made to date. Previous studies have demonstrated that the risk of SSCs and PJI can be significantly different based on the individual patient and the surgical risk factors.^{2,6,10,11} Tan et al² demonstrated that a patient's comorbidities and the revision procedures should be considered as valid risk factors for PJI and the incidence of developing PJI can vary from 0.6% to 20.6% based on the risk factors. Because patients with certain risk factors are frequently associated with SSCs, a variety of dressing materials were applied to prevent SSCs.^{4,12,13} However, the proper indication and the best choice of dressing materials for wound management after THA and TKA still remains unclear.

Closed-incisional negative-pressure wound therapy (ciNPWT) has been recently developed and has shown better efficacy in decreasing SSCs than conventional dressings after THA or TKA.^{3,4,10,14} Although a recent study showed the routine application of ciNPWT to all patients to be a cost-effective intervention to reduce SSCs after primary THA and TKA,¹⁵ the cost of ciNPWT application is substantially increased over that of conventional dressings.

Therefore, we designed a systematic review and metaanalysis to determine the effective indication for ciNPWT in wound management following THA or TKA. We asked the following questions: Does the use of ciNPWT following THA or TKA compared with conventional dressings reduce the incidence of wound complication or SSI in (a) high-risk patients compared with routine patients? and (b) revision arthroplasties compared with primary arthroplasties?

2 | MATERIALS AND METHODS

2.1 | Literature search

The present systematic review followed the recommendation of the Cochrane review methods. Based on the Preferred Reporting Items for Systematic Reviews and Metaanalyses (PRISMA) guidelines,¹⁶ multiple comprehensive literature databases, including PubMed (MEDLINE), Embase, and the Cochrane Library were searched for studies that reported on the outcomes of ciNPWT in wound management following THA or TKA up to September 1st, 2019 using a prior search strategy. There were no restrictions on language or the year of publication. The search terms used in the title, abstract, Medical Subjects Heading, and keywords fields included the following search methodology: ("ciNPT" OR "ciNPWT" OR "closed incisional negative pressure therapy" OR "closed incisional negative wound therapy" OR "negative pressure wound therapy" OR "NPWT" OR "vacuum assisted

Key Messages

- the application of closed-incisional negativepressure wound therapy (ciNPWT) reduced the incidence of wound complication and surgical site infection (SSI) in high-risk patients and in revision procedures after total hip arthroplasty or total knee arthroplasty compared with conventional dressings
- our findings would support the evidence to determine effective indication for ciNPWT application in high-risk patients and in revision arthroplasties
- the wound complication and SSI were significantly less likely to occur in the high-risk patients or in revision arthroplasties using ciNPWT compared with conventional dressings
- there were no significant differences of wound complication and SSI in routine patients or in primary arthroplasties between ciNPWT and conventional dressings

closure" OR "VAC") AND [("TKA" OR "total knee arthroplasty" OR "total knee replacement" OR "arthroplasty, replacement, knee") OR ("THA" OR "total hip arthroplasty" OR "total hip replacement" OR "arthroplasty, replacement, hip")]. Manual searches were also performed for articles that could have been missed by the electronic search.

2.2 | Study selection

Two reviewers independently evaluated titles and abstracts of the identified studies and selected eligible studies for a full review. If the abstract showed insufficient information for a decision, the full text of the article was reviewed. Articles that satisfied the following criteria were selected in this systematic review: (a) patients who underwent THA or TKA using ciNPWT for their surgical incisions; (b) studies that directly compared ciNPWT and conventional dressings in terms of wound complications and SSI; and (c) studies that fully reported the complete numbers of patients or enabled the calculation of the number and proportion of patients regarding wound complications and SSI. Studies not clearly reporting data regarding either wound complication or SSI, indicating vague definition of terms between wound complication and SSI, biomechanical and cadaveric studies, technical notes, letters to the editor, expert opinions, review articles, meta-analyses, scientific conference abstracts, and case reports were excluded. A study of cohorts undergoing ciNPWT for periprosthetic fractures of THA and TKA was also excluded.

2.3 | Data extraction

Two investigators independently extracted data from each article using a predefined data extraction form. Any disagreements between two reviewers were solved by discussion. The extracted outcomes were SSCs including wound complications and SSI. Wound complications included wound discharge, wound dehiscence, hematoma, and seroma. The number of overall wound complications was reported in most included studies, if not, we added the number of specific wound complications. SSIs included both superficial and deep infection. Patient demographic, characteristic, and population data including sample size, mean age, sex, mean body mass index (BMI), and follow-up period were recorded for each included study. If the follow-up periods for wound complications and SSI were different, each follow-up period was separately recorded. Details of wound management such as the specific material and duration of dressing changes were extracted from each included study. Details of study indications were also extracted from pooled studies such as whether routine patients were included or high-risk patients having comorbidities were included, and whether primary and/or revision and THA and/or TKA was performed.

2.4 | Assessment of methodological quality

Two investigators independently assessed the methodological quality of each study using the methodological index for non-randomised studies (MINORS).¹⁷ Using the MINORS checklist, the maximum score is 24 for a comparative study. Furthermore, MINORS has validity to assess the qualities of randomised controlled trials (RCTs) as well as non-randomised studies. Any discrepancies in the scores between the two reviewers were resolved by discussion.

3 | **STATISTICS**

Wound complication and SSI data recorded in the included studies were pooled. The main outcomes of the present study were mean differences in wound complication and SSI between ciNPWT and conventional dressings based on subgroups of patients as follows: routine patients vs high-risk patients; primary arthroplastis vs revision arthroplastis. Thus, subgroup analyses of the studies were performed to determine the effective indication of ciNPWT after THA or TKA. Random-effects metaanalyses were performed to pool the outcomes across the included studies. Binary outcomes, such as the rates of wound complication and SSI were reported as odds ratios (ORs) and 95% confidence intervals (CIs). Heterogeneity was determined by estimating the proportion of betweenstudy inconsistencies because of actual differences between studies, rather than differences because of random error or chance, using the I^2 statistic, where 25% was considered low heterogeneity, 50% was considered moderate heterogeneity, and 75% was considered high heterogeneity. Forest plots were used to show the outcome, pooled estimate of effect, and overall summary effect of each study and constructed using the Review Manager software (RevMan version 5.3; Copenhagen, The Nordic Cochrane Centre, The Cochrane Collaboration). A meta-regression analysis was performed to assess the effects of age, sex, and follow-up period on wound complication and SSI. Analyses were performed using RevMan version 5.3 and Open Meta-Analyst (http:// www.cebm.brown.edu/openmeta). Statistical significance was set at P < .05.

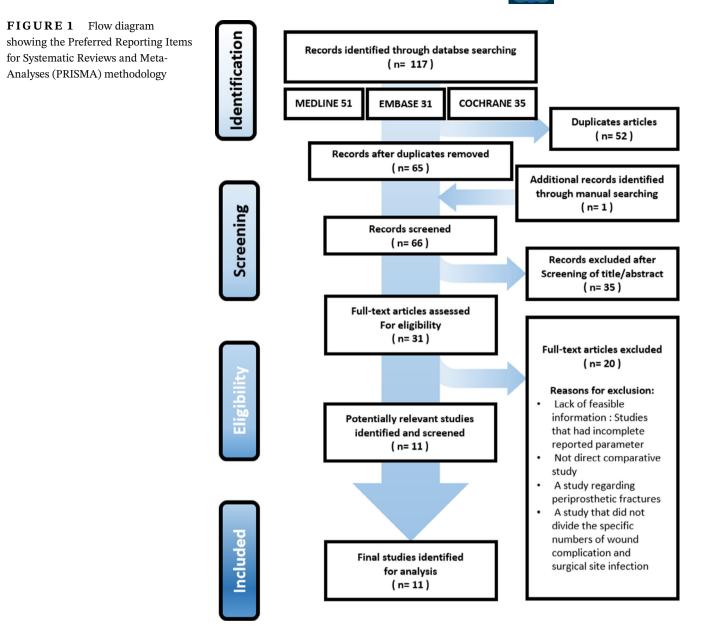
4 | RESULTS

4.1 | Identification of studies

Figure 1 shows the detail of the study identification, inclusion, and exclusion. An electronic search yielded 51 studies in PubMed (MEDLINE), 31 in Embase, and 35 in the Cochrane Library. An additional study was identified through manual searching. After removing 52 duplicate studies, 66 studies remained. After screening the titles and abstracts, and reading the full text, 55 studies were excluded. Thus, 11 studies were finally included in the present study, of which eight RCTs^{3,4,18-23} and three cohort studies^{10,14,24} were eligible for data extraction and meta-analysis.

4.2 | Study characteristics and methodological quality assessment

A total of 1997 cases of THA or TKA were reported including 763 cases with ciNPWT management and 1234 cases with conventional wound management. The details of the study design and patient and population



characteristics including age, percentage of female, mean BMI, follow-up period, and MINORS quality score of each included study are summarised in Table 1. The median MINORS score of the included studies was 20 of 24 (range 16-24). The details of wound management in ciNPWT and conventional wound dressings, the specific study indications for patients at risk, and the type of surgery are described in Table 2. Publication bias was not investigated for as it is not generally necessary when meta-analyses include fewer than 10 studies.²⁵

4.3 | Routine patients vs high-risk patients

In terms of wound complication, there were 6 studies with a total of 548 and 801 routine patients who received

ciNPWT and conventional dressings, respectively. The overall rate of wound complication in routine patients was not significantly different between the ciNPWT and the conventional dressings (OR = 0.52; 95% CI = 0.21-1.33; P = .17). Four studies included a total of 165 and 383 high-risk patients for wound complication who received ciNPWT and conventional dressings, respectively. The overall rate of wound complication in highrisk patients was significantly lower in the ciNPWT group than in the conventional dressings, and the summary OR was 0.38 (95% CI = 0.15-0.93; P = .03) (Figure 2A). In terms of SSI, there were five studies with a total of 539 and 791 routine patients who received ciNPWT and conventional dressings, respectively. The overall rate of SSI in routine patients was not significantly different between the ciNPWT and the conventional dressings (OR = 0.49; 95% CI = 0.22-1.11; P = .09). Five studies

lies ^a
d stuc
include
of the in
scores o
ta and quality sco
a and
t dat
demographic
ails of
The det
1
FABLE 1
ΓA

		0 I	1									
		Study	Sample size (n)	e (n)	Mean age (years)	(years)	Percentage of female	of	BMI (kg/m ²)	1 ²)		MINDRS
Author	Year		ciNPWT	Control	ciNPWT	Control	ciNPWT	Control	ciNPWT	Control	Follow-up	score
						Main findings	ndings					
Cooper and Bas	2016	RCS	30	108	71.7	70.9	NR		31.3	29.6	Wound: 4 weeks SSI: over 34 months	19
	ciNPW	T may decrease	wound com	plications ar	nd SSIs in hig	sh-risk patien	nts with mul-	tiple risk fac	stors for SSI	undergoing	ciNPWT may decrease wound complications and SSIs in high-risk patients with multiple risk factors for SSI undergoing revision TKA or TKA.	
Curley et al	2018	RCS	32	159	63.4	59.5	NR		NR		Retrospective chart review	16
	A lowé althc	lower infection rate was observed for the ciNalthough this difference was not statistically	was observe nce was not	d for the ciN statistically s	JPWT patients significant.	s who had h	igh-risk factc	ors for SSI u	ndergoing pı	rimary TKA	A lower infection rate was observed for the ciNPWT patients who had high-risk factors for SSI undergoing primary TKA as opposed to the dry sterile dressing patients, although this difference was not statistically significant.	dressing patients,
Giannini et al	2018	RCT	50	50	66	66.8	62.0	64.0	27.7	28.2	1 week	20
	The re. high	he results of this study do not support the routine use of ciNPWT high-risk factors for wound complications have been determined.	y do not sur wound com	port the rou olications ha	tine use of ci ve been detei	NPWT follor rmined.	wing revisior	al THA or	TKA. Howev	⁄er, it could l	The results of this study do not support the routine use of ciNPWT following revisional THA or TKA. However, it could be beneficial for selected patients once high-risk factors for wound complications have been determined.	ients once
Gillespie et al	2015	RCT	35	35	63.8	62.5	42.9	51.4	29.9	29.8	6 weeks	21
	A redu after	reduction of 3% in SS after primary THA.	SI incidence	suggests tha	t a definitive	trial require	es approxima	tely 900 pat.	ients per gro	up. Yet, the	A reduction of 3% in SSI incidence suggests that a definitive trial requires approximately 900 patients per group. Yet, there is uncertainty around the benefit of NPWT after primary THA.	benefit of NPWT
Howell et al	2011	RCT	24	36	NR		NR		NR		Wound: 1 week SSI: 12 months	21
	ciNPW	ciNPWT did not appear to prove lower wound	r to prove lo		complication	s in high-risl	k patients fol	lowing prin	nary TKA; hu	owever, it wa	complications in high-risk patients following primary TKA; however, it was associated with blisters.	
Karlakki et al	2016	RCT	102	107	69	69.2	52.0	48.6	30.1	28.4	6 weeks	23
	ciNPW	ciNPWT has a beneficial role in patients undergoing primary THA or TKA to minimise wound complications.	al role in pa	tients under§	soing primar.	y THA or TE	XA to minim.	ise wound c	omplication	S.		
Keeney et al	2019	RCT	185	213	60.6	60.5	60.5	57.3	34.6	36.5	Wound: 12 weeks SSI: 24 months	20
	ciNPW not s grou	NPWT improved wound complication rates compare not significant difference. Patients with a body mass group may be helpful to define the value of iNPWT.	und complic ence. Patien I to define th	ation rates co ts with a bod ne value of ir	ompared wit ly mass index vPWT.	h conventior <>35 kg/m ²	aal dressings showed to b	in patients e more susc	following pri eptible to wc	imary or rev ound compli	ciNPWT improved wound complication rates compared with conventional dressings in patients following primary or revisional THA or TKA; however, SSI rate was not significant difference. Patients with a body mass index >35 kg/m ² showed to be more susceptible to wound complications. Specific study in this high-risk patient group may be helpful to define the value of iNPWT.	er, SSI rate was s high-risk patient
Manoharan	2016	RCT	21	36	66		42.4		29.8		1.5 weeks	17
et al	There prote	There was no benefit in wound healing or cost protection.	n wound he		with NPWT	post-TKA. T.	here was son	ne benefit ir	ı ciNPWT or	ı quality of l	with NPWT post-TKA. There was some benefit in ciNPWT on quality of life factors less wound leakage and better	ge and better
Newman et al	2019	RCT	79	80	65	65	49.4	43.8	33.4	33.4	12 weeks	24
	ciNPW	'T may decrease	the rate of \mathfrak{p}	oostoperative	wound com	plications in	patients wh	o are at an i	ncreased risl	k of such wo	ciNPWT may decrease the rate of postoperative wound complications in patients who are at an increased risk of such wound issues after revision THA or TKA.	A or TKA.
Pachowsky et al	2012	RCT	6	10	66.2	70.5	NR		NR		1.5 weeks	18
	There	was a decreased	developmer	it of postope	rative seromé	as in the wor	ind and imp	roved woun	d healing in	patients wh	There was a decreased development of postoperative seromas in the wound and improved wound healing in patients who used ciNPWT following primary THA	imary THA.

		Study	Sample size (n)	(e (n)	Mean age (years)	(years)	Percentage of female	e of	BMI (kg/m ²)	n ²)		MINORS
Author	Year		ciNPWT	ciNPWT Control	ciNPWT	Control	ciNPWT Control ciNPWT Control	Control	ciNPWT	Control	ciNPWT Control Follow-up	score
Redfern et al	2017	RCS	196	400	66.9	66.8	65.8	54.0	30.5	30.9	Wound: 6 weeks SSI: 2 months	21
	ciNPV	VT for THA and	d TKA in a co	mprehensive	e patient pop	ulation redu	iced overall i	ncidence of	wound comp	olication, bu	ciNPWT for THA and TKA in a comprehensive patient population reduced overall incidence of wound complication, but did not significantly impact the rate of SSI.	pact the rate of SSI.
^a BMI, body mass index; ciNPWT, closed incision negative pressure therapy; MINORS, methodological items for non-randomi ies; RCT, randomised controlled trials; SSI, surgical site infection; THA, total hip arthroplasty; TKA, total knee arthroplasty.	ndex; ciN.	PWT, closed incollection of the second statement of the second se	cision negativ surgical site i	e pressure th infection; TF	rerapy; MINC IA, total hip :	JRS, methoo arthroplasty	dological iten ; TKA, total	ns for non-ra knee arthroj	andomised st plasty.	udies; NR, r	^a BMI, body mass index; ciNPWT, closed incision negative pressure therapy; MINORS, methodological items for non-randomised studies; NR, not reported; RCS, retrospective comparison stud- ies; RCT, randomised controlled trials; SSI, surgical site infection; THA, total hip arthroplasty; TKA, total knee arthroplasty.	ective comparison stud-

included a total of 215 and 433 high-risk patients who received ciNPWT and conventional dressings, respectively. The overall rate of SSI in high-risk patients was significantly lower in the ciNPWT group than in the conventional dressings, and the summary OR was 0.24 (95% CI = 0.09-0.64; P = .005) (Figure 2B).

4.4 | Primary arthroplasty vs revision arthroplasty

In terms of wound complication, there were 8 studies with a total of 561 and 955 patients who received ciNPWT and conventional dressings, respectively, following primary arthroplasty. The overall wound complication rate in primary arthroplasty was not significantly different between the ciNPWT and the conventional dressings (OR = 0.59; 95% CI = 0.25-1.40; P = 0.23). Three studies included a total of 152 and 229 patients for wound complication who received ciNPWT and conventional dressings, respectively, after revision arthroplasty. The overall wound complication rate in revision arthroplasty was significantly lower in the ciNPWT group than in the conventional dressings, and the summary OR was 0.33 (95% CI = 0.18-0.62; P < .001) (Figure 3A). In terms of SSI, there were 7 studies with a total of 552 and 945 patients who received ciNPWT and conventional dressings, respectively, following primary arthroplasty. The overall rate of SSI in primary arthroplasty was not significantly different between the ciNPWT and the conventional dressings (OR = 0.54; 95% CI = 0.26-1.12; P = .10). Four studies included a total of 202 and 279 patients who received ciNPWT and conventional dressings, respectively, following revision arthroplasty. The overall rate of SSI in revision arthroplasty was significantly lower in the ciNPWT group than in the conventional dressings, and the summary OR was 0.26 (95% CI = 0.11-0.66; P = .004) (Figure 3B).

4.5 | Meta-regression analysis

The results of the meta-regression analyses are shown in Table 3. Patient characteristics including age, sex, and follow-up were not significantly associated with the rates of wound complication and SSI.

5 | DISCUSSION

Recent studies, differing in indications of surgical or patient risk factor, have reported outcomes of ciNPWT after THA or TKA. Although most outcomes were shown

1 incision negative-pressure therapy, conventional wound dressings, and details of study indication ^a	
Summary of closed incisio	
TABLE 2	

		ciNPWT		Conventi	Conventional wound dressings	Indication	
Author	Year	Material	Duration (day)	Material	Dressing changes	Surgery indication	Patients indication
Cooper and Bas	2016	Prevena (125 mm Hg, continuous)	9.2	AQUACEL Ag	Leave the dressing for a minimum of first 5 days	Revision THA or TKA	High-risk factors for SSI: morbid obesity, multiple significant medical or social comorbidities, treatment of an infected joint arthroplasty, and wound closure under tension.
Curley et al	2018	Prevena (125 mm Hg, continuous)	Ч	Standard sterile gauze dressing	Depending on the wound leakage	Primary TKA	High-risk factors for SSI: increased body mass index, smoking status, history of infection, and numerous comorbidities.
Giannini et al	2018	PICO (80 mm Hg, continuous)	7	Povidone-iodine gauze dressing	Depending on the wound leakage	Revision THA or TKA	High-risk factors for SSI (at least one risk factor): Age > 65 years, diabetes, smoking, obesity (BMI ≥30 kg/m ²), hypertension, pulmonary disease, and vascular disease.
Gillespie et al	2015	PICO (80 mm Hg, continuous)	Ŋ	Hydrocolloid dressing	Depending on the wound leakage	Primary THA	Routine application to patients following arthroplasty.
Howell et al	2011	VAC (125 mm Hg, continuous)	7	Standard sterile gauze dressing	Leave the dressing on the second postoperative day	Primary TKA	High-risk factors for SSI: Obesity (BMI $>$ 30 kg/m ²) and enoxaparin sodium for deep venous thrombosis prophylaxis.
Karlakki et al	2016	PICO (80 mm Hg, continuous)	7	Mepore or Tegaderm	Mean days of dressing were 4.2	Primary THA or TKA	Routine application to patients following arthroplasty.
Keeney et al	2019	PICO (80 ± 20 mm Hg, continuous)	7	Standard sterile gauze dressing	Subsequent dressing changes every 3 to 5 days	Primary or revision THA or TKA	Routine application to patients following arthroplasty.
Manoharan et al	2016	Prevena (125 mm Hg, continuous)	œ	Standard sterile gauze dressing	Dressing changes on day 1 postoperatively	Primary TKA	Routine application to patients following arthroplasty.
Newman et al	2019	Prevena (125 mm Hg, continuous)	≥2	AQUACEL Ag	Dressing for 7 days	Revision THA or TKA	High-risk factors for SSI: Obesity (BMI >35 kg/m ²), use of anticoagulants other than aspirin, peripheral vascular disease, depression, diabetes mellitus, current smoker, history of a PJI in the limb undergoing revision

		ciNPWT		Conventi	Conventional wound dressings	Indication	
Author	Year	Material	Duration (day)	Material	Dressing changes	Surgery indication	Patients indication
							surgery, on immunomodulators or corticosteroids, current history of cancer or haematological malignancy, inflammatory arthritis, renal failure or dialysis, malnutrition, liver disease, history of organ transplant, or HIV infection.
Pachowsky et al	2012	Prevena (125 mm Hg, continuous)	S	Standard sterile gauze dressing	Standard dressing changes	Primary THA	Routine application to patients following arthroplasty.
Redfern et al	2017	Prevena (125 mm Hg, continuous)	7.1	Standard sterile gauze dressing	Standard dressing changes	Primary THA or TKA	Routine application to patients following arthroplasty.
^a BMI, body ma total hip arthrc	iss index; o	^a BMI, body mass index; ciNPWT, closed-incisional r total hip arthroplasty; TKA, total knee arthroplasty.	negative-press	ure wound therapy; HI	.V, human immunodeficiency virus; F	PJI, periprosthetic joint in	^a BMI, body mass index; ciNPWT, closed-incisional negative-pressure wound therapy; HIV, human immunodeficiency virus; PJI, periprosthetic joint infection; SSI, surgical site infection; THA, total hip arthroplasty; TKA, total knee arthroplasty.

to be effective, the application of ciNPWT to all postoperative wounds would lead to considerable economic burden. Therefore, we performed this systematic review and meta-analysis to assess the effective indications for ciNPWT in wound management following THA or TKA. The most important findings of this study are that wound complication and SSI were significantly less likely to occur in high-risk patients or in revision arthroplasties using ciNPWT compared with conventional dressings. Conversely, there were no significant differences of wound complication and SSI in routine patients or in primary arthroplasties between ciNPWT and conventional dressings.

IWJ

The number of the arthroplasties is expected to increase dramatically with the ageing population of late,²⁶ and the ageing population is likely to have several comorbidities. Patients with comorbidities have been shown to increase the risk of both SSC and PJI^{2,6,27}: thus. accurate risk stratification of patients for SSC and PJI following THA or TKA is essential. Bozic et al²⁸ identified specific patient comorbidities such as rheumatoid disease, obesity, coagulopathy, and preoperative anaemia that were independently associated with an increased risk of PJI following THA. Namba et al²⁷ analysed 56 216 TKAs and demonstrated obesity, diabetes mellitus, male sex, American Society of Anesthesiologists score of \geq 3, and posttraumatic arthritis are patient factors associated with PJI. Furthermore, several studies have similarly shown that comorbidities associated with immune deficiency, such as renal, rheumatologic, and liver disease, are related with SSC and PJI.^{2,29-31} Although many studies have sought to identify risk factors for SSC and PJI, there are relatively little studies that have suggested a method to reduce SSC and PJI in patients having those risk factors, except for controlling or compensating for the comorbidities. Thus, the results of our study indicate that ciNPWT could be a new solution for reducing the wound complication and SSI risk in high-risk patients with comorbidity after THA or TKA.

The prevalence of revision THA and TKA have been increasing with time as primary arthroplasties that have been performed in the past decades require revision and the surgical indications for primary arthroplasties had been broadened recently.^{10,14,26,32} Compared with primary arthroplasties, the revision procedure of THA or TKA requires a longer surgical time, has a longer surgical incision and results in more difficult wound healing because of the previous scar, which frequently causes SSC and consequently increases the risk of PJI.^{2,4,10,18,33} Many studies have identified that the revision procedure is one of the most crucial risk factors for SSC and PJI after THA or TKA, resulting in poorer clinical outcomes, longer hospital

(Continued)

TABLE 2

(A) Wound complication

			-				
	CiNP		Contr			Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% Cl
1.1.1 Routine patients							
Gillespie et al. 2015	24	35	15	35	22.0%	2.91 [1.09, 7.74]	
Karlakki et al. 2016	2	102	9	107	16.0%	0.22 [0.05, 1.03]	• • • • • • • • • • • • • • • • • • •
Keeney et al. 2019	26	185	42	213	26.5%	0.67 [0.39, 1.14]	→
Manoharan et al. 2016	0	21	1	36	6.4%	0.55 [0.02, 14.13]	· · · · ·
Pachowsky et al. 2012	4	9	9	10	9.7%	0.09 [0.01, 1.03]	<
Redfern et al. 2017	3	196	22	400	19.4%	0.27 [0.08, 0.90]	_
Subtotal (95% CI)	0	548	~~		100.0%	0.52 [0.21, 1.33]	
Total events	59	010	98		1001070	0.02 [0.2.1, 1.00]	
Heterogeneity: Tau ² = 0.		16 66		- 0 000	N- IZ - 600	v.	
Test for overall effect: Z=			ui – 5 (r -	- 0.000	0,1 - 003	/0	
restion overall ellect. 2 -	- 1.50 (1 -	0.17)					
1.1.2 High-risk patients	1						1.1
Cooper and Bas. 2016	2	30	29	108	27.0%	0.19 [0.04, 0.87]	
Curley et al. 2018	2	30	10	159	9.0%	0.19 [0.04, 0.87]	← → ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓
	2	24	10	36			11
Howell et al. 2011	2 8	24 79	19		11.8%	3.18 [0.27, 37.20]	
Newman et al. 2019	8		19	80	52.2%	0.36 [0.15, 0.88]	
Subtotal (95% CI)		165		282	100.0%	0.38 [0.15, 0.93]	
Total events	12		59				
Heterogeneity: Tau ² = 0.			f= 3 (P =	0.28);1	*= 21%		i i
Test for overall effect: Z =	= 2.12 (P =	0.03)					
							0.1 0.2 0.5 1 2 5 1
							ciNPT Control
							CINPT Control
) Surgical site infe		т	Contr	ol		Odds Ratio	
) Surgical site infe	ciNP		Contr Events		Weight	Odds Ratio M-H, Random, 95% Cl	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup	ciNP Events				Weight		Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients	ciNP Events	Total	Events	Total		M-H, Random, 95% Cl	Odds Ratio
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015	ciNP Events	Total 35	Events 3	Total 35	11.4%	M-H, Random, 95% Cl 0.31 [0.03, 3.17]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016	ciNP Events 1 1	Total 35 102	Events 3 6	Total 35 107	11.4% 13.2%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019	ciNP Events	<u>35</u> 102 185	Events 3	<u>35</u> 107 213	11.4%	M-H, Random, 95% Cl 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016	ciNP Events 1 1 9	Total 35 102 185 21	Events 3 6 11	35 107 213 36	11.4% 13.2% 50.8%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017	ciNP Events 1 1 9 0	<u>35</u> 102 185	Events 3 6 11 0	35 107 213 36 400	11.4% 13.2%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% CI)	ciNP Events 1 1 9 0 2	35 102 185 21 196	Events 3 6 11 0 14	35 107 213 36 400	11.4% 13.2% 50.8% 24.6%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% Cl) Total events	ciNP Events 1 1 9 0 2 13	35 102 185 21 196 539	Events 3 6 11 0 14 34	35 107 213 36 400 791	11.4% 13.2% 50.8% 24.6% 100.0%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0.	ciNP Events 1 1 9 0 2 13 13; Chi ² = 1	Total 35 102 185 21 196 539 3.61, dt	Events 3 6 11 0 14 34	35 107 213 36 400 791	11.4% 13.2% 50.8% 24.6% 100.0%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% Cl) Total events	ciNP Events 1 1 9 0 2 13 13; Chi ² = 1	Total 35 102 185 21 196 539 3.61, dt	Events 3 6 11 0 14 34	35 107 213 36 400 791	11.4% 13.2% 50.8% 24.6% 100.0%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z =	ciNP Events 1 1 9 0 2 13 13; Chi ² = : = 1.70 (P =	Total 35 102 185 21 196 539 3.61, dt	Events 3 6 11 0 14 34	35 107 213 36 400 791	11.4% 13.2% 50.8% 24.6% 100.0%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients	ciNP <u>Events</u> 1 1 9 0 2 13 13; Chi ² = : 1.70 (P =	Total 35 102 185 21 196 539 3.61, di 0.09)	3 6 11 0 14 f= 3 (P =	Total 35 107 213 36 400 791 0.31);1	11.4% 13.2% 50.8% 24.6% 100.0% F = 17%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016	CiNP <u>Events</u> 1 1 9 0 2 13 13; Chi ² = = 1.70 (P = 1	Total 35 102 185 21 196 539 3.61, dt 0.09) 30	Events 3 6 11 0 14 34 f = 3 (P = 20	Total 35 107 213 36 400 791 0.31);1	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018	ciNP Events 1 1 9 0 2 13 13; Chi ² = 1 = 1.70 (P = 1 0	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32	Events 3 6 11 0 14 14 34 f = 3 (P = 20 9 9 20 9 9	Total 35 107 213 36 400 791 0.31);1 108 159	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7% 12.1%	M-H, Random, 95% CI 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.11]	Odds Ratio M-H, Random, 95% Cl
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z= 1.2.2 High-risk patients Cooper and Bas. 2018 Giannini et al. 2018	ciNP Events 1 1 9 0 2 13; Chi ² = = 1.70 (P = 1 0 0 0	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50	Events 3 6 11 0 14 34 f= 3 (P = 20 9 5	Total 35 107 213 36 400 791 0.31);1 108 159 50	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7% 12.1% 11.7%	<u>M-H, Random, 95% CI</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2016 Redfern et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Howell et al. 2011	ciNP Events 1 1 9 0 2 13 13; Chi ² = 1.70 (P = 1 0 0 0 1	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50 24	Events 3 6 11 0 14 34 f = 3 (P = 20 9 5 1	Total 35 107 213 36 400 791 0.31);1 108 159 50 36	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7% 12.1% 11.7% 12.5%	<u>M-H, Random, 95% Cl</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% CI) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Howell et al. 2011 Newman et al. 2019	ciNP Events 1 1 9 0 2 13; Chi ² = = 1.70 (P = 1 0 0 0	Total 35 102 185 21 196 539 3.61, di 0.09) 3.61, di 0.09) 30 32 50 24 79	Events 3 6 11 0 14 34 f= 3 (P = 20 9 5	Total 35 107 213 36 400 791 0.31);1 108 159 50 36 80	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7% 12.1% 11.7% 12.5% 39.9%	M-H, Random, 95% CI 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56] 0.23 [0.05, 1.14]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Giannini et al. 2018 Howell et al. 2011 Newman et al. 2019 Subtotal (95% Cl)	CiNP Events 1 1 9 0 2 13 13; Chi ² = = 1.70 (P = 1 0 0 1 2	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50 24	Events 3 6 11 0 14 f= 3 (P = 20 9 5 1 8	Total 35 107 213 36 400 791 0.31);1 108 159 50 36 80	11.4% 13.2% 50.8% 24.6% 100.0% ² =17% 23.7% 12.1% 11.7% 12.5%	<u>M-H, Random, 95% Cl</u> 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Giannini et al. 2018 Howell et al. 2011 Newman et al. 2019 Subtotal (95% Cl) Total events	CiNP Events 1 1 9 0 2 13 13; Chi ² = = 1.70 (P = 1 0 0 1 2 4	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50 24 79 215	Events 3 6 11 0 14 14 f = 3 (P = 20 9 5 1 8 43	Total 35 107 213 36 400 791 0.31);1 0.31);1 108 159 50 366 80 433	11.4% 13.2% 50.8% 24.6% 100.0% P=17% 23.7% 12.1% 11.7% 12.5% 39.9% 100.0%	M-H, Random, 95% CI 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56] 0.23 [0.05, 1.14]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Giannini et al. 2018 Howell et al. 2019 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0.	ciNP Events 1 1 9 0 2 13; Chi ² = = 1.70 (P = 1 0 0 0 1 2 4 00; Chi ² =	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50 24 79 215 2.40, dt	Events 3 6 11 0 14 34 f= 3 (P = 20 9 5 1 8 43 f= 4 (P =	Total 35 107 213 36 400 791 0.31);1 0.31);1 108 159 50 366 80 433	11.4% 13.2% 50.8% 24.6% 100.0% P=17% 23.7% 12.1% 11.7% 12.5% 39.9% 100.0%	M-H, Random, 95% CI 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56] 0.23 [0.05, 1.14]	Odds Ratio M-H, Random, 95% CI
Study or Subgroup 1.2.1 Routine patients Gillespie et al. 2015 Karlakki et al. 2016 Keeney et al. 2019 Manoharan et al. 2017 Subtotal (95% Cl) Total events Heterogeneity: Tau ² = 0. Test for overall effect: Z = 1.2.2 High-risk patients Cooper and Bas. 2016 Curley et al. 2018 Giannini et al. 2018 Howell et al. 2011 Newman et al. 2019 Subtotal (95% Cl) Total events	ciNP Events 1 1 9 0 2 13; Chi ² = = 1.70 (P = 1 0 0 0 1 2 4 00; Chi ² =	Total 35 102 185 21 196 539 3.61, dt 0.09) 30 32 50 24 79 215 2.40, dt	Events 3 6 11 0 14 34 f= 3 (P = 20 9 5 1 8 43 f= 4 (P =	Total 35 107 213 36 400 791 0.31);1 0.31);1 108 159 50 366 80 433	11.4% 13.2% 50.8% 24.6% 100.0% P=17% 23.7% 12.1% 11.7% 12.5% 39.9% 100.0%	M-H, Random, 95% CI 0.31 [0.03, 3.17] 0.17 [0.02, 1.41] 0.94 [0.38, 2.32] Not estimable 0.28 [0.06, 1.26] 0.49 [0.22, 1.11] 0.49 [0.22, 1.11] 0.15 [0.02, 1.18] 0.24 [0.01, 4.29] 0.08 [0.00, 1.52] 1.52 [0.09, 25.56] 0.23 [0.05, 1.14]	Odds Ratio M-H, Random, 95% CI

FIGURE 2 Forest plots showing the overall rates of wound complication, A, and surgical site infection (SSI), B, between the ciNPWT and control groups in routine patients and high-risk patients. In routine patients, there were no differences in the rates of wound complication and SSI between the two groups. However, in high-risk patients, the overall rates of wound complication (odds ratio [OR] = 0.38; 95% CI = 0.15-0.93; P = .03) and SSI (OR = 0.24; 95% CI = 0.09-0.64; P = .005) were significantly lower in the ciNPWT group. ciNPWT, closed-incisional negative-pressure wound therapy

stay, and greater economic burden.^{1,2,28,34-36} Although several attempts to decrease the infection risk during the revision procedure have been shown such as using antibioticladen cement, an irrigation solution of antibiotics, and prophylactic antibiotics, the efficacy of those attempts remains unclear.^{27,37,38} Given wound-related complications are a great concern for revision arthroplasty, our

results identified that ciNPWT significantly reduced wound complication and SSI in revision THA or TKA compared with conventional dressings.

0.1 0.2

0.5

ciNP

ż

Control

10

Increasingly, ciNPWT systems have been applied to high-risk wounds in various fields and showed a notable efficacy of reducing SSC.³⁹⁻⁴² Specific to patients following THA or TKA in our study, the use of ciNPWT

(A) Wound complication

	ciNP	т	Contr	ol		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% Cl	M-H, Random, 95% Cl
1.3.1 Primary arthropla	sty						
Curley et al. 2018	0	32	10	159	6.6%	0.22 [0.01, 3.83]	
Gillespie et al. 2015	24	35	15	35	18.9%	2.91 [1.09, 7.74]	
Howell et al. 2011	2	24	1	36	8.2%	3.18 [0.27, 37.20]	+ + +
Karlakki et al. 2016	2	102	9	107	13.7%	0.22 [0.05, 1.03]	• • • • • • • • • • • • • • • • • • •
Keeney et al. 2019	20	142	30	172	22.2%	0.78 [0.42, 1.44]	· · · · · ·
Manoharan et al. 2016	0	21	1	36	5.5%	0.55 [0.02, 14.13]	
Pachowsky et al. 2012	4	9	9	10	8.2%	0.09 [0.01, 1.03]	< <u>·</u>
Redfern et al. 2017	3	196	22	400	16.7%	0.27 [0.08, 0.90]	
Subtotal (95% CI)		561		955	100.0%	0.59 [0.25, 1.40]	
Total events	55		97				i i l
Heterogeneity: Tau ² = 0.1	76; Chi ² =	17.73,	df = 7 (P :	= 0.01)	l ² = 61%		
Test for overall effect: Z =	= 1.19 (P =	0.23)					
	_						i i l
1.3.2 Revision arthropla	sty						
Cooper and Bas. 2016	2	30	29	108	17.6%	0.19 [0.04, 0.87]	
Keeney et al. 2019	6	43	12	41	33.0%	0.39 [0.13, 1.17]	
Newman et al. 2019	8	79	19	80	49.4%	0.36 [0.15, 0.88]	
Subtotal (95% CI)		152		229	100.0%	0.33 [0.18, 0.62]	
Total events	16		60				i i
Heterogeneity: Tau ² = 0.1	00; Chi ² =	0.63, d	f=2(P=	0.73);1	² = 0%		
Test for overall effect: Z =	= 3.43 (P =	0.0000	3)				
							i i
							0.1 0.2 0.5 1 2 5 10

(B) Surgical site infection

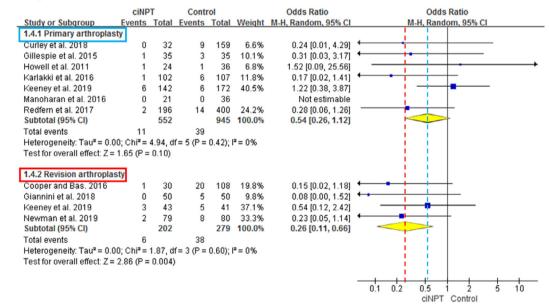


FIGURE 3 Forest plots showing the overall rates of wound complication, A, and surgical site infection (SSI), B, between the ciNPWT and control groups in primary arthroplasty and revision arthroplasty. In primary arthroplasty, there were no differences in the rates of wound complication and SSI between the two groups. However, in revision arthroplasty; the overall rates of wound complication (odd ratio [OR] = 0.33; 95% CI = 0.18-0.62; *P* < .001) and SSI (OR = 0.26; 95% CI = 0.11-0.66; *P* = .004) were significantly lower in the ciNPWT group. ciNPWT, closed-incisional negative-pressure wound therapy

similarly showed significantly lower rates of wound complication and SSI in high-risk patients and in revision arthroplasties than with conventional dressings. Clearly, ciNPWT offers several potential benefits to improve wound healing and prevent SSI of closed surgical incisions. One important explanation may be a reduction of the relative motion on incisional edges by mechanical stabilisation.^{43,44} Another explanation includes reducing dead space, subcutaneous hematoma, and seroma, and improving perfusion and lymphatic flow, all of which contribute to a better environment for wound healing.^{23,44,45} Other explanations may include that ciNPWT keeps surgical wounds sterile in the role of a mechanical barrier and requires fewer dressing changes and a longer duration from the initial application.^{10,46-48}

ciNPT Control

We acknowledge several limitations of this study. The heterogeneity of the demographic data among included studies, including differences in age, sex distribution, as

Variable	Coefficient	SE	P value	95% CI
Wound complication				
Age	-0.254	0.158	.107	-0.564 to 0.055
Percentage of female	-0.112	0.122	.360	-0.352 to 0.128
Follow-up (week)	-0.091	0.265	.731	-0.611 to 0.429
Surgical site infection				
Age	-0.197	0.174	.257	-0.539 to 0.144
Percentage of female	-0.084	0.103	.412	-0.285 to 0.117
Follow-up (month)	0.089	0.095	.349	-0.097 to 0.274

TABLE 3Meta-regressionanalysis for the assessment of theinfluence of age, percentage of female,body mass index, and follow-up periodon the wound complication andsurgical site infection^a

^aBMI, body mass index; CI, confidence interval.

well as differences in follow-up duration, may be potential confounding factors. However, our meta-regression analysis showed that age, sex distribution, BMI, and follow-up were not significantly associated with rates of wound complication and SSI. Although surveillance for the first 12 months after THA and TKA recommended,^{49,50} only three included studies^{3,10,20} had more than 12 months of follow-up. However, woundrelated SSI is likely to occur in the acute setting⁵¹ and a follow-up duration of less than 12 months might be acceptable to evaluate the efficacy of the ciNPWT system on wound management. Second, studies differing in the indication of patient comorbidities might include selection bias. Although a small difference in the specific indication of comorbidities might be a potential selection bias, the details of the indication had a similarity among the included studies of high-risk patients according to Table 2. Third, included studies were fewer in subgroups of high-risk patients and revision arthroplasty than in the subgroups of routine patients and primary arthroplasty, which might include confounding factors. However, the present study has a strength as the first systematic review and meta-analysis regarding this topic because ciNPWT has recently started to be administered to patients undergoing THA or TKA. Finally, we could not perform a costeffectiveness analysis of the ciNPWT system because of a lack of published studies.

In conclusion, the current study showed that the application of ciNPWT reduced the incidence of wound complication and SSI in high-risk patients and in revision procedures after THA or TKA compared with conventional dressings. Our findings suggest that ciNPWT should be considered for high-risk patients and in revision procedures for wound management following THA or TKA.

ACKNOWLEDGEMENTS

We thank to the library team in Samsung Medical Center for gathering the data. The authors did not receive any outside funding or grants in support of their research for or preparation of this work.

CONFLICT OF INTEREST

The authors declare that they have no conflict of interest.

ETHICAL APPROVAL

This article does not contain any studies with human participants or animals performed by any of the authors.

REFERENCES

- 1. Lavernia CJ, Alcerro JC. Quality of life and cost-effectiveness 1 year after total hip arthroplasty. *J Arthroplasty*. 2011;26(5): 705-709.
- Tan TL, Maltenfort MG, Chen AF, et al. Development and evaluation of a preoperative risk calculator for periprosthetic joint infection following Total joint arthroplasty. *J Bone Joint Surg Am.* 2018;100(9):777-785.
- Keeney JA, Cook JL, Clawson SW, Aggarwal A, Stannard JP. Incisional negative pressure wound therapy devices improve shortterm wound complications, but not long-term infection rate following hip and knee arthroplasty. *J Arthroplasty*. 2019;34(4):723-728.
- 4. Newman JM, Siqueira MBP, Klika AK, Molloy RM, Barsoum WK, Higuera CA. Use of closed incisional negative pressure wound therapy after revision total hip and knee arthroplasty in patients at high risk for infection: a prospective, randomized clinical trial. *J Arthroplasty*. 2019;34(3):554-559.e551.
- Weiss AP, Krackow KA. Persistent wound drainage after primary total knee arthroplasty. J Arthroplasty. 1993;8(3):285-289.
- Patel VP, Walsh M, Sehgal B, Preston C, DeWal H, Di Cesare PE. Factors associated with prolonged wound drainage after primary total hip and knee arthroplasty. *J Bone Joint Surg Am.* 2007;89(1):33-38.
- Kurtz SM, Lau E, Schmier J, Ong KL, Zhao K, Parvizi J. Infection burden for hip and knee arthroplasty in the United States. *J Arthroplasty*. 2008;23(7):984-991.
- Kurtz SM, Lau E, Watson H, Schmier JK, Parvizi J. Economic burden of periprosthetic joint infection in the United States. *J Arthroplasty*. 2012;27(8 Suppl):61-65. e61.
- Ong KL, Mowat FS, Chan N, Lau E, Halpern MT, Kurtz SM. Economic burden of revision hip and knee arthroplasty in Medicare enrollees. *Clin Orthop Relat Res.* 2006;446:22-28.

- Cooper HJ, Bas MA. Closed-incision negative-pressure therapy versus antimicrobial dressings after revision hip and knee surgery: a comparative study. *J Arthroplasty*. 2016;31(5):1047-1052.
- 11. Bozic KJ, Lau E, Kurtz S, Ong K, Berry DJ. Patient-related risk factors for postoperative mortality and periprosthetic joint infection in medicare patients undergoing TKA. *Clin Orthop Relat Res.* 2012;470(1):130-137.
- Poultsides LA, Ma Y, Della Valle AG, Chiu YL, Sculco TP, Memtsoudis SG. In-hospital surgical site infections after primary hip and knee arthroplasty—incidence and risk factors. *J Arthroplasty*. 2013;28(3):385-389.
- Cai J, Karam JA, Parvizi J, Smith EB, Sharkey PF. Aquacel surgical dressing reduces the rate of acute PJI following total joint arthroplasty: a case-control study. *J Arthroplasty*. 2014;29(6): 1098-1100.
- Redfern RE, Cameron-Ruetz C, O'Drobinak SK, Chen JT, Beer KJ. Closed incision negative pressure therapy effects on postoperative infection and surgical site complication after total hip and knee arthroplasty. *J Arthroplasty*. 2017;32(11): 3333-3339.
- Nherera LM, Trueman P, Karlakki SL. Cost-effectiveness analysis of single-use negative pressure wound therapy dressings (sNPWT) to reduce surgical site complications (SSC) in routine primary hip and knee replacements. *Wound Repair Regen*. 2017;25(3):474-482.
- Moher D, Shamseer L, Clarke M, et al. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. Syst Rev. 2015;4:1.
- Slim K, Nini E, Forestier D, Kwiatkowski F, Panis Y, Chipponi J. Methodological index for non-randomized studies (minors): development and validation of a new instrument. *ANZ J Surg.* 2003;73(9):712-716.
- Giannini S, Mazzotti A, Luciani D, et al. Postoperative wound management with negative pressure wound therapy in knee and hip surgery: a randomised control trial. *J Wound Care*. 2018;27(8): 520-525. https://doi.org/10.12968/jowc.2018.27.8.520.
- Gillespie B, Rickard C, Thalib L, et al. Use of negative-pressure wound dressings to prevent surgical site complications after primary hip arthroplasty: a pilot RCT. *Surg Innov.* 2015;22(5): 488-495. https://doi.org/10.1177/1553350615573583.
- Howell R, Hadley S, Strauss E, Pelham F. Blister formation with negative pressure dressings after total knee arthroplasty. *Curr Orthop Prac.* 2011;22(2):176-179. https://doi.org/10.1097/ BCO.0b013e31820b3e21.
- 21. Karlakki S, Hamad A, Whittall C, Graham N, Banerjee R, Kuiper J. Incisional negative pressure wound therapy dressings (inpWTd) in routine primary hip and knee arthroplasties: a randomised controlled trial. *Bone J Res.* 2016;5(8):328-337. https://doi.org/10.1302/2046-3758.58.BJR-2016-0022.R1.
- 22. Manoharan V, Grant AL, Harris AC, Hazratwala K, Wilkinson MP, McEwen PJ. Closed incision negative pressure wound therapy vs conventional dry dressings after primary knee arthroplasty: a randomized controlled study. *J Arthroplasty*. 2016; 31(11):2487-2494.
- Pachowsky M, Gusinde J, Klein A, Lehrl S, Schulz-Drost S, Schlechtweg P. Negative pressure wound therapy to prevent seromas and treat surgical incisions after total hip arthroplasty. *International orthopaedics*. 2011;36(4):719-722. https://doi.org/ 10.1007/s00264-00011-01321-00268.

- 24. Curley AJ, Terhune EB, Velott AT, Argintar EH. Outcomes of prophylactic negative pressure wound therapy in knee arthroplasty. *Orthopedics*. 2018;41(6):e837-e840.
- 25. Higgins JPT. Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0. The Cochrane Collaboration; 2011. http://handbook.cochrane.org/
- Kurtz S, Ong K, Lau E, Mowat F, Halpern M. Projections of primary and revision hip and knee arthroplasty in the United States from 2005 to 2030. *J Bone Joint Surg Am.* 2007;89(4): 780-785.
- Namba RS, Inacio MC, Paxton EW. Risk factors associated with deep surgical site infections after primary total knee arthroplasty: an analysis of 56,216 knees. *J Bone Joint Surg Am*. 2013;95(9):775-782.
- Bozic KJ, Lau E, Kurtz S, et al. Patient-related risk factors for periprosthetic joint infection and postoperative mortality following total hip arthroplasty in Medicare patients. *J Bone Joint Surg Am.* 2012;94(9):794-800.
- Bozic KJ, Ong K, Lau E, et al. Estimating risk in Medicare patients with THA: an electronic risk calculator for periprosthetic joint infection and mortality. *Clin Orthop Relat Res.* 2013;471(2):574-583.
- McCleery MA, Leach WJ, Norwood T. Rates of infection and revision in patients with renal disease undergoing total knee replacement in Scotland. *J Bone Joint Surg Br.* 2010;92(11): 1535-1539.
- Jiang SL, Schairer WW, Bozic KJ. Increased rates of periprosthetic joint infection in patients with cirrhosis undergoing total joint arthroplasty. *Clin Orthop Relat Res.* 2014;472(8): 2483-2491.
- 32. Ricciardi BF, Liu AY, Qiu B, Myers TG, Thirukumaran CP. What is the association between hospital volume and complications after revision total joint arthroplasty: a large-database study. *Clin Orthop Relat Res.* 2019;477(5):1221-1231.
- Keswani A, Lovy AJ, Robinson J, Levy R, Chen D, Moucha CS. Risk factors predict increased length of stay and readmission rates in revision joint arthroplasty. *J Arthroplasty*. 2016;31(3): 603-608.
- 34. Goltz DE, Baumgartner BT, Politzer CS, DiLallo M, Bolognesi MP, Seyler TM. The American College of Surgeons National Surgical Quality Improvement Program Surgical Risk Calculator has a role in predicting discharge to postacute care in total joint arthroplasty. *J Arthroplasty*. 2018;33 (1):25-29.
- 35. Schairer WW, Sing DC, Vail TP, Bozic KJ. Causes and frequency of unplanned hospital readmission after total hip arthroplasty. *Clin Orthop Relat Res.* 2014;472(2):464-470.
- 36. Surin VV, Sundholm K, Backman L. Infection after total hip replacement. With special reference to a discharge from the wound. *J Bone Joint Surg Br.* 1983;65(4):412-418.
- Jamsen E, Huhtala H, Puolakka T, Moilanen T. Risk factors for infection after knee arthroplasty. A register-based analysis of 43,149 cases. *J Bone Joint Surg Am.* 2009;91(1):38-47.
- AlBuhairan B, Hind D, Hutchinson A. Antibiotic prophylaxis for wound infections in total joint arthroplasty: a systematic review. *J Bone Joint Surg Br.* 2008;90(7):915-919.
- Hyldig N, Birke-Sorensen H, Kruse M, et al. Meta-analysis of negative-pressure wound therapy for closed surgical incisions. *Br J Surg.* 2016;103(5):477-486.

1322 WILEY IWJ

- 40. Semsarzadeh NN, Tadisina KK, Maddox J, Chopra K, Singh DP. Closed incision negative-pressure therapy is associated with decreased surgical-site infections: a meta-analysis. *Plast Reconstr Surg.* 2015;136(3):592-602.
- 41. Stannard JP, Volgas DA, McGwin G 3rd, et al. Incisional negative pressure wound therapy after high-risk lower extremity fractures. *J Orthop Trauma*. 2012;26(1):37-42.
- Blackham AU, Farrah JP, McCoy TP, Schmidt BS, Shen P. Prevention of surgical site infections in high-risk patients with laparotomy incisions using negative-pressure therapy. *Am J Surg.* 2013;205(6):647-654.
- Wilkes RP, Kilpad DV, Zhao Y, Kazala R, McNulty A. Closed incision management with negative pressure wound therapy (CIM): biomechanics. *Surg Innov*. 2012;19(1):67-75.
- 44. Kilpadi DV, Lessing C, Derrick K. Healed porcine incisions previously treated with a surgical incision management system: mechanical, histomorphometric, and gene expression properties. *Aesthetic Plast Surg.* 2014;38(4):767-778.
- 45. Kilpadi DV, Cunningham MR. Evaluation of closed incision management with negative pressure wound therapy (CIM): hematoma/seroma and involvement of the lymphatic system. *Wound Repair Regen*. 2011;19(5):588-596.
- Liu D, Zhang L, Li T, et al. Negative-pressure wound therapy enhances local inflammatory responses in acute infected softtissue wound. *Cell Biochem Biophys.* 2014;70(1):539-547.
- Morykwas MJ, Argenta LC, Shelton-Brown EI, McGuirt W. Vacuum-assisted closure: a new method for wound control and treatment: animal studies and basic foundation. *Ann Plast Surg.* 1997;38 (6):553-562.

- Blum ML, Esser M, Richardson M, Paul E, Rosenfeldt FL. Negative pressure wound therapy reduces deep infection rate in open tibial fractures. *J Orthop Trauma*. 2012;26(9): 499-505.
- Mangram AJ, Horan TC, Pearson ML, Silver LC, Jarvis WR. Guideline for prevention of surgical site infection, 1999. Hospital Infection Control Practices Advisory Committee. *Infect Control Hosp Epidemiol.* 1999;20(4):250-278.
 Mangram AJ, Horan TC, Pearson ML, Silver LC, Jarvis WR.
- Mangram AJ, Horan TC, Pearson ML, Silver LC, Jarvis WR. Guideline for prevention of surgical site infection, 1999. Centers for Disease Control and Prevention (CDC) Hospital Infection Control Practices Advisory Committee. Am J Infect Control. 1999;27(2):97-132.
- 51. Scalise A, Calamita R, Tartaglione C, et al. Improving wound healing and preventing surgical site complications of closed surgical incisions: a possible role of incisional negative pressure wound therapy. A systematic review of the literature. *Int Wound J.* 2016;13(6):1260-1281.

How to cite this article: Kim J-H, Lee D-H. Are high-risk patient and revision arthroplasty effective indications for closed-incisional negative-pressure wound therapy after total hip or knee arthroplasty? A systematic review and meta-analysis. *Int Wound* J. 2020;17:1310–1322. <u>https://doi.org/10.1111/iwj</u>. 13393