## REVIEW

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## Effects of Intraoperative Saline-Soaked Pharyngeal Packing on Nausea, Vomiting, and Throat Pain After Nasal Surgery: A Systematic Review and Meta-Analysis

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**Background and Objectives:** The purpose of this study was to conduct a meta-analysis of the effects of intraoperative pharyngeal packing on postoperative nausea, vomiting, and sore throat in nasal surgery patients.

**Methods:** Databases were searched from inception to December 2022. Randomized controlled trials comparing saline-soaked pharyngeal packing (packing group) with no packing (control group) during intubation in patients undergoing nasal surgery were included. The primary outcomes of interest were the incidence of postoperative nausea, vomiting, and sore throat at 24 hours.

**Results:** Eleven studies, including a total of 931 patients, were included. There was no significant difference between the two groups in the incidence of postoperative nausea and vomiting and severity scores at 2, 6, and 24 hours postoperatively. The incidence of throat pain was higher in the packing group than in the control group immediately after surgery and at 24 hours postoperatively. However, no significant difference was observed between the two groups in the incidence of sore throat at 6 and 12 hours postoperatively.

**Conclusion:** Intraoperative saline-soaked pharyngeal packing did not significantly decrease postoperative nausea and vomiting. However, the use of pharyngeal packing was associated with a higher incidence of sore throat in the initial recovery period.

Keywords: Pharynx; Nausea; Vomiting; Pharyngitis; Nasal surgical procedures.

## **INTRODUCTION**

Nausea and vomiting are frequent complications of nasal surgery. There are several causes of nausea and vomiting, including patient factors, anesthetics, and opiate analgesics, but ingested blood is also known to be a major cause. Cuffed endotracheal tubes are not thought to be completely effective in protecting against the aspiration of hypopharyngeal blood [1]. Surgery in the nasal cavity and sinuses can cause severe bleeding because these are areas with abundant blood vessels [2]. Therefore, pharyngeal packing is often performed by clinicians during nasal surgery to minimize blood entry into the esophagus and thus reduce postoperative nausea and vomiting [3].

However, pharyngeal packing can also cause complications related to the act itself, such as sore throat after surgery. Sore throat is an adverse effect that occurs due to compression and irritation of the packing on the pharyngeal mucosa [4]. The incidence of throat pain after nasal surgery varies from 14.4% to 50%, and has been reported in up to 60% of patients who receive pharyngeal packing [5]. Therefore, debate continues regarding whether it is beneficial to perform pharyngeal packing during nasal surgery. In fact, some studies have reported that pharyngeal packing had no effect on reducing the incidence of postoperative nausea and vomiting, but rather was associated with an increased incidence of sore throat. Therefore, the purpose of this meta-analysis was to evaluate the efficacy of using pharyngeal packing to prevent postoperative nausea and vomiting compared to patients who did not use

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pharyngeal packing, and to determine whether it also affects the occurrence of throat pain.

## **METHODS**

#### **Selection of studies**

Studies published before December 2022 in PubMed Central, MEDLINE, Scopus, and the Cochrane Central Register of Controlled Trials with the keywords "pharyngeal packing," "postoperative nausea," "vomiting," "throat pain," "sore throat," and "nasal surgery" (including functional endoscopic sinus surgery, septoplasty, septal surgery, septorhinoplasty, turbinectomy, turbinoplasty, nasal polypectomy, and nasal adhesions lysis) were searched. Two reviewers independently evaluated the data extracted from the database, and articles irrelevant to the research topic were excluded based on the title and abstract. If the items selected by the two reviewers were different, the final choice regarding inclusion was made through discussion with a third reviewer. We limited inclusion to randomized controlled trials in which saline-soaked pharyngeal packing was performed on patients during nasal surgery under general anesthesia. Studies were excluded from the analysis if: 1) additional surgery was performed (e.g., middle ear surgery or pharyngeal surgery); 2) patients had a previous history of postoperative nausea and vomiting; 3) patients had underlying diseases such as systemic or malignant diseases; 4) multiple reports were based on the same test data; or 5) data necessary for the analysis were missing or incomplete, making it impossible to extract and calculate appropriate data. The overall article selection process is presented in Fig. 1.

#### Outcome items and risk of bias assessment

The primary outcome was the incidence and severity of nausea and vomiting on the first day after nasal surgery under general anesthesia. The control group comprised patients who did not perform pharyngeal packing. The incidence and severity of throat pain were also assessed as secondary outcomes.

The extracted data were organized using a standardized extraction form and listed as follows: the number of patients with postoperative nausea, vomiting, and sore throat, the incidence (as a percentage), and the p-value for the comparison between the packing and control groups. Quality assessment was conducted using the Cochrane Risk of bias tool.

#### **Statistical analysis**

Meta-analysis was performed using the R Statistical Software (R-4.2.3; R Foundation for Statistical Computing, Vienna, Austria). When the extracted data were continuous, a metaanalysis was performed using the standardized mean difference (SMD). Odds ratios (ORs) were used to analyze incidence.

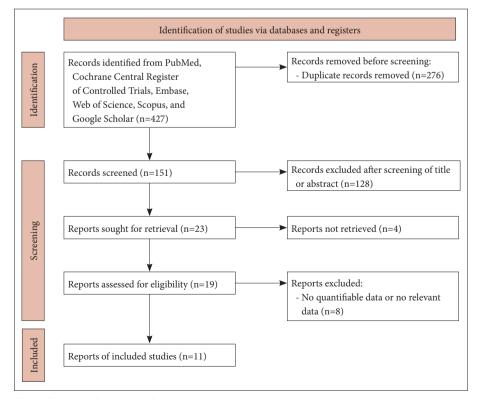


Fig. 1. Diagram of selection of studies.

Table 1. Summary of :	studies and r	Table 1. Summary of studies and risk of bias assessment					
Study (year)	Sample size	Age, yr (mean, range or SD)	Sex (male-to-female ratio)	Study design	Comparison	Outcome measure analyzed	Risk of bias of randomized studies
Basha et al. [1] (2006)	93	34 (16-81)	69 Males and 24 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Unclear risk
Meco et al. [8] (2016)	201	45 (11)	111 Males and 90 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting	Unclear risk
Korkut et al. [2] (2010)	100	30.5 (11.95)	Not declared	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting	Low risk
Rizvi et al. [9] (2015)	40	35 (15-60)	NA	RCT	Pharyngeal packing (saline) vs. nasal packing	Throat pain	Unclear risk
Solmaz et al. [10] (2016)	105	29 (20), 2 not declared	NA	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Unclear risk
Razavi et al. [5] (2015)	89	27.18 (7.08)	89 Males and 35 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Low risk
Fennessy et al. [3] (2011)	32	39 (19–62)	NA	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Low risk
Karbasforushan et al. [4] (2014)	140	20-40	61 Males and 79 females	RCT	Pharyngeal packing (saline) vs. no packing	Throat pain	Unclear risk
Al-Lami et al. [6] (2017)	80	42 (18-72)	57 Males and 23 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Low risk
Temel et al. [11] (2019)	88	30.18 (8.98)	56 Males and 32 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting	Low risk
Borna et al. [7] (2022)	101	35.0 (18-61)	81 Males and 20 females	RCT	Pharyngeal packing (saline) vs. no packing	Postoperative nausea and vomiting, throat pain	Low risk

Table 1. Summary of studies and risk of bias assessment

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A funnel plot and the Egger test were conducted together to assess publication bias. We also adjusted for missing studies using the Duval and Tweedie trim-and-fill method and corrected the overall effect size for publication bias. We also performed a sensitivity analysis to estimate the impact of each individual study on the overall meta-analysis results.

## **RESULTS**

Eleven studies, with 931 participants, were reviewed for eventual inclusion in this meta-analysis [1-11]. The study characteristics and quality assessment are presented in Table 1.

# Effect of pharyngeal packing on postoperative nausea and vomiting

No significant differences in postoperative nausea and vomiting severity scores were found at postoperative 2 hours (SMD=-0.13, 95% CI: -0.39–0.13,  $I^2$ =0%), 6 hours (SMD= 0.08, 95% CI, -0.20–0.35,  $I^2$ =0%), and 24 hours (SMD=-0.07, 95% CI: -0.35–0.21,  $I^2$ =0%) between patients who received pharyngeal packing and patients who did not receive packing (control group) (Fig. 2). Similarly, the incidence of postoperative nausea and vomiting in the packing group were not significantly different from that in the control group immediately after surgery (OR=1.52, 95% CI: 0.78–2.94,  $I^2$ =14%), postoperative 2 hours (OR=0.80, 95% CI: 0.52–1.23,  $I^2$ =9%),

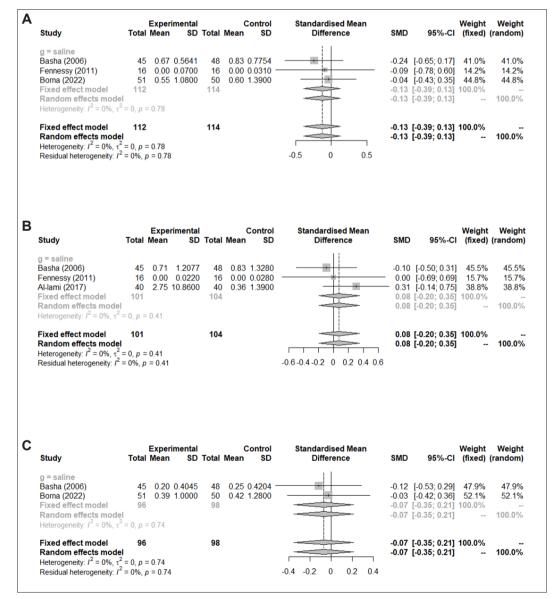


Fig. 2. Severity scores of postoperative nausea and vomiting at 2 hours (A), 6 hours (B), and 24 hours (C) postoperatively [1,3,6,7]. Experimental: pharyngeal packing, control: no packing. SD, standard deviation; SMD, standardized mean difference; CI, confidence interval.

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	Basha (2006)	15	45	7				[1.06; 8.07]	42.8%	39.7%
	Razavi (2015)	3	44					[0.25; 9.90]	13.0%	14.3%
	Meco (2016)	3	48	5			0.60	[0.14; 2.66]	19.8%	20.9%
	Solmaz (2016)	5	35	5	35		1.00	[0.26; 3.81]	24.5%	25.2%
	Fixed effect model		172		178			[0.78; 2.94]		
	Random effects model							[0.71; 3.03]		100.0%
	Heterogeneity: $I^2 = 14\%$ , $\tau^2$		n = 0	32		-		[0.1 1, 0.00]		100.07
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	Fixed effect model Random effects model	I	172		178			[0.78; 2.94] [0.71; 3.03]	100.0%	100.0%
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	Basha (2006)	29	45	32	48	<u> </u>	0.91	[0.39; 2.13]	25.0%	24.79
	Korkut (2010)	29	50	33		<u>+</u>		[0.32; 1.60]	27.8%	27.29
	Razavi (2015)	3	44					[0.20; 5.37]	6.7%	7.29
	Solmaz (2016)	6	35	14				[0.20, 0.97]	14.9%	15.59
		26								
	Temel (2019)	20	44	23				[0.57; 3.06]	25.7%	25.49
	Fixed effect model		218		222	Ĩ		[0.52; 1.23]	100.0%	100.00
	<b>Random effects model</b> Heterogeneity: $I^2 = 9\%$ , $\tau^2$		n = 0	25			0.80	[0.51; 1.25]		100.0%
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	Fixed effect model Random effects model	I	218		222	$\rightarrow$		[0.52; 1.23] [0.51; 1.25]	100.0%	100.0%
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	Study g = saline Basha (2006) Korkut (2010)	Experim Events 20 17	0.35 nental Total 45 50	C Events 27 14	<b>Total</b> 48 50		0.62		(fixed) 30.9% 28.7%	(random 30.99 28.79
.,	Study g = saline Basha (2006) Korkut (2010) Razavi (2015)	Experim Events	0.35 nental Total 45 50 44	C Events 27 14 0	<b>Total</b> 48 50 45		0.62 1.32	[0.27; 1.41] [0.57; 3.10]	(fixed) 30.9% 28.7% 0.0%	(random 30.99 28.79 0.09
	<b>Study</b> g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016)	Experim Events	0.35 nental Total 45 50 44 35	C Events 27 14 0 9	<b>Total</b> 48 50 45 35		0.62 1.32 0.86	[0.27; 1.41] [0.57; 3.10] [0.29; 2.56]	(fixed) 30.9% 28.7% 0.0% 17.3%	(random 30.99 28.79 0.09 17.39
•	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019)	Experim Events	0.35 nental Total 45 50 44 35 44	C Events 27 14 0 9	<b>Total</b> 48 50 45 35 44		0.62 1.32 0.86 - 1.59	[0.27; 1.41] [0.57; 3.10] [0.29; 2.56] [0.61; 4.10]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1%	(random 30.99 28.79 0.09 17.39
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ŝ	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Residual heterogeneity: $I^2$ Study g = saline Basha (2006) Korkut (2010)	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0%, p = Experime Events	0.35 nental Total 45 50 44 218 .44 218 .44 0.44 ental Total 45 50 44 218 .44 0.44	C Events 27 14 0 9 10 9 10 20 5 20 5 20 5 20 5 20 5 20 5 20 5 2	Total 48 50 45 35 44 222 222 222 222	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0R 0.58	[0.27; 1.41] [0.57; 3.10] [0.62; 2.56] [0.61; 4.10] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0%  Weighi (fixed) 60.9% 0.0%	(random 30.99 28.79 0.09 17.39 23.19 100.09 100.09
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	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Residual heterogeneity: $I^2$ Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016)	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0, p = 0 = 0, p = 0 Experime Events	0.35 nental 45 50 44 218 .44 218 .44 0.44 ental Total 45 50 0.44 .44 0.44 .44 0.44	C Events 27 14 0 9 10 10 20 5 20 5 20 5 20 5 20 5 20 5 20	Total   48   50   45   35   44   222   222   222   222   223   224   225   226   227   228   229   220   221   222   222   222   222   235	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0.00 0.58 2.10 5.30	[0.27; 1.41] [0.57; 3.10] [0.62; 2.56] [0.61; 4.10] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60] [0.64; 1.60]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0%  Weight (fixed) 60.9% 0.0% 10.1% 6.4%	(random 30.99 28.79 0.09 17.39 23.19 100.09 100.09 t Weig ) (random (random) 42.5 0.00 0.00 17.10 10.09
	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $J^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $J^2 = 0\%, \tau^2$ Residual heterogeneity: $J^2$ Study g = saline Basha (2006) Korkut (2010) Bazavi (2015) Solmaz (2016) Femel (2019)	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0%, p = 0 Experime Events	0.35 nental Total 45 50 44 218 44 218 44 0.44 ental Total 45 50 44 35 44 218 45 50 44 218 45 50 44 218 44 218 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 45 50 44 50 45 45 45 45 45 45 45 45 45 45	Co Events 27 14 0 9 10 10 20 5 27 14 0 9 10 20 5 20 5 20 5 20 5 20 5 20 5 20 5 2	Total 48 50 45 35 44 222 222 222 222 222 222 222 222 222	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0 0 0 5.30 5.30 3.97	[0.27; 1.41] [0.57; 3.10] [0.57; 3.10] [0.64; 1.60] [0.64; 1.60]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0%  Weight (fixed) 60.9% 0.0% 10.1% 6.4% 22.6%	(random 30.99 28.79 0.00 17.39 23.19 100.09 100.09 t Weig ) (random 42.5 0.00 17.1 0.00 17.1 0.00 17.0 100.09
	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Fixed effect model Heterogeneity: $I^2 = 0\%, \tau^2$ Residual heterogeneity: $I^2$ Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Femel (2019) Fixed effect model	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0, p = 0 = 0, p = 0 Experime Events	0.35 nental 45 50 44 218 .44 218 .44 0.44 ental Total 45 50 0.44 .44 0.44 .44 0.44	C Events 27 14 0 9 10 10 20 5 20 5 20 5 20 5 20 5 20 5 20	Total   48   50   45   35   44   222   222   222   222   223   244   50   44   50   48   50   45   35	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0.68 2.10 5.30 5.30 1.18	[0.27; 1.41] [0.57; 3.10] [0.57; 3.10] [0.64; 4.10] [0.64; 1.60] [0.64; 2.56]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0% 60.9% 0.0% 10.1% 6.4% 22.6% 100.0%	(random 30.99 28.79 0.09 17.33 23.19 100.09 100.09 t Weig ) (random ) (random ) (random ) (random ) 23.31 100.09
	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $J^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $J^2 = 0\%, \tau^2$ Residual heterogeneity: $J^2$ Study g = saline Basha (2006) Korkut (2010) Bazavi (2015) Solmaz (2016) Femel (2019)	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0%, p = Experim Events 8 0 2 7	0.35 hental 45 50 44 218 .44 218 .44 ental Total 45 50 44 218 .44 218 .44 218 .44 218 .44 .44 .44 .44 .44 .44 .44 .4	Co Events 27 14 0 9 10 10 2 2	Total 48 50 45 35 44 222 222 222 222 222 222 222 222 222	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0.68 2.10 5.30 5.30 1.18	[0.27; 1.41] [0.57; 3.10] [0.57; 3.10] [0.64; 1.60] [0.64; 1.60]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0% 60.9% 0.0% 10.1% 6.4% 22.6% 100.0%	(random 30.99 28.79 0.09 17.39 23.19 100.09 100.09 t Weig (random 42.5 0.00 (random 23.19 100.09
	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $J^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $J^2$ Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $J^2 = 42\%, \tau^2$	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0%, p = Experim Events 8 0 2 7	0.35 nental Total 45 50 44 218 .44 218 .44 0.44 ental Total 45 50 44 0.44 ental 70 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 45 50 44 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 47 218 79	Co Events 27 14 0 9 10 10 2 2	Total 48 50 45 35 44 222 222 222 222 222 222 222 222 222	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0 0 0.58 2.10 5.30 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.530 1.5400 1.540 1.540 1.5400 1.5400 1.5400 1.540	[0.27; 1.41] [0.57; 3.10] [0.57; 3.10] [0.64; 1.60] [0.64; 1.60] [0.64	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0%  100.0%  Weight (fixed) 60.9% 0.0% 10.1% 64.4% 22.6% 100.0%	(random 30.99 28.79 0.00 17.39 23.19 100.09 100.09 t Weig ) (random 42.5 0.00 17.1 0.00 17.1 0.00 17.0 100.09
	Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Temel (2019) Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%, \tau^2$ Fixed effect model Random effects model Heterogeneity: $I^2$ Study g = saline Basha (2006) Korkut (2010) Razavi (2015) Solmaz (2016) Femel (2019) Fixed effect model Random effects model	Experim Events 20 17 0 8 14 = 0, p = 0 = 0, p = 0 = 0%, p = Experim Events 8 0 2 7	0.35 hental 45 50 44 218 .44 218 .44 ental Total 45 50 44 218 .44 218 .44 218 .44 218 .44 .44 .44 .44 .44 .44 .44 .4	Co Events 27 14 0 9 10 10 2 2	Total 48 50 45 35 44 222 222 222 222 222 222 222 222 222	Odds Ratio	0.62 1.32 0.86 - 1.59 1.01 1.01 1.01 0.68 2.10 5.30 3.97 1.18 1.63 1.18	[0.27; 1.41] [0.57; 3.10] [0.57; 3.10] [0.64; 4.10] [0.64; 1.60] [0.64; 2.56]	(fixed) 30.9% 28.7% 0.0% 17.3% 23.1% 100.0% 100.0% 100.0% 60.9% 0.0% 10.1% 64.4% 22.6% 100.0%	(random 30.99 28.76 0.09 17.33 23.19 100.09 100.09 t Weig (random (random ) (random ) (random ) (random ) (random ) 12.0 12.0 12.0 12.0 12.0 12.0 12.0 12.0

Fig. 3. Incidence of postoperative nausea and vomiting immediately after surgery (A) and at 2 hours (B), 6 hours (C), and 24 hours (D) postoperatively [1,2,5,8,10,11]. Experimental: pharyngeal packing, control: no packing. OR, odds ratio; CI, confidence interval.

	Study g = saline Karbasforushan (2014) Meco (2016)	70		al D Total	Control Mean SD	Standardised Mean Difference	SMD	95%-CI		Weight (random)
	Karbasforushan (2014) Meco (2016)									
	Meco (2016)								00.404	
		48			1.38 0.5000 0.00 1.5000			[0.20; 0.88]		29.3% 24.7%
	Al-lami (2017)	40						[0.00; 0.89]		21.5%
	Borna (2022)		3.65 1.470					[0.17; 0.97]		24.5%
	Fixed effect model	209		210				[0.20; 0.59]		-
	<b>Random effects model</b> Heterogeneity: $I^2 = 43\%$ , $\tau^2$		305. p = 0.15				0.39	[ 0.13; 0.65]		100.0%
	Fixed effect model	209		210			0.40	[ 0.20; 0.59]	100.0%	
l	<b>Random effects model</b> Heterogeneity: $I^2 = 43\%$ , $\tau^2$	= 0.03	305  n = 0.15				0.39	[ 0.13; 0.65]		100.0%
I	Residual heterogeneity: I <sup>2</sup> :	= 43%,	<i>p</i> = 0.15			-0.5 0 0.5				
В			Experimenta	al	Control	Standardised Mean			Weight	Weigh
:	Study	Total		D Total	Mean SD	Difference	SMD	95%-CI		(random
	g = saline Basha (2006)	15	2.93 1.684	2 48	1.98 1.0604		- 0.62	[0.26; 1.10]	43.0%	36.9%
	Fennessy (2011)		0.00 0.029					[-0.84; 0.55]		27.0%
	Al-lami (2017)		2.10 2.400					[-0.51; 0.36]		36.1%
	Fixed effect model	101		104				[-0.03; 0.52]		
	<b>Random effects model</b> Heterogeneity: $I^2 = 73\%$ , $\tau^2$	= 0.17	727, p = 0.03				0.18	[-0.37; 0.74]		100.0%
	Fixed effect model Random effects model	101		104				[-0.03; 0.52] [-0.37; 0.74]		100.0%
H	Heterogeneity: $I^2 = 73\%$ , $\tau^2$ Residual heterogeneity: $I^2$	= 0.17	727, p = 0.03			-1 -0.5 0 0.5 1				
С			Experimenta		Control	Standardised Mean			Weight	
S	Study	Total	Mean SE	) Total	Mean SD	Difference	SMD	95%-CI	(fixed)	(random
	j = saline Basha (2006)	45	2.69 1.8068	3 48	1.73 1.3788		0.59	[0.18; 1.01]	44.8%	35.9%
	Fennessy (2011)		0.00 0.0180		0.02 0.0420			[-1.47; -0.03]		28.79
	Al-lami (2017)		1.40 1.6000		1.60 2.4000	<u> </u>		[-0.54; 0.34]		35.49
	ixed effect model	101		104				[-0.16; 0.39]	100.0%	-
	Random effects model leterogeneity: $I^2 = 83\%$ , $\tau^2$	= 0.31	50, p < 0.01				-0.04	[-0.74; 0.67]		100.0%
	ixed effect model	101		104		L.	0 11	[-0.16; 0.39]	100.0%	
R	Random effects model			104				[-0.74; 0.67]		100.0%
H R	deterogeneity: $I^2 = 83\%$ , $\tau^2$ Residual heterogeneity: $I^2 =$	= 0.31 83%,	50, p < 0.01 p < 0.01			-1 -0.5 0 0.5 1				
D			-							
\$	Study		Experimenta Mean SI	l D Total	Control Mean SD	Standardised Mean Difference	SMD	95%-CI		Weight (random)
	g = saline									
	Basha (2006)	45			1.12 1.0423			[-0.06; 0.76]		27.8%
Ē	Karbasforushan (2014)	70 51			1.26 0.7000 2.48 1.5500			[-0.11; 0.56] [ 0.06; 0.85]		42.3% 29.9%
Ē	Soma (2022)	<b>D</b> I	3.10 1.510	J 50 168	2.40 1.0000			[0.06; 0.85]		29.9%
E	Borna (2022) Fixed effect model						0.00	[ 0.11, 0.00]	.00.070	
E F F	Borna (2022) Fixed effect model Random effects model Heterogeneity: I <sup>2</sup> = 0%, τ <sup>2</sup> =	166	= 0.68	100			0.33	[ 0.11; 0.55]		100.0%
E F F F	Fixed effect model Random effects model Heterogeneity: $I^2 = 0\%$ , $\tau^2 =$ Fixed effect model	166	= 0.68	168			0.33	[ 0.11; 0.55]		100.0%
E F F F F	<b>Fixed effect model</b> <b>Random effects model</b> Heterogeneity: $I^2 = 0\%$ , $\tau^2 =$	166 0, p = <b>166</b>					0.33	- / -		100.0%  100.0%

Fig. 4. Severity scores of throat pain immediately after surgery (A) and at 2 hours (B), 6 hours (C), and 24 hours (D) postoperatively [1,3,4,6-8]. Experimental: pharyngeal packing, control: no packing. SD, standard deviation; SMD, standardized mean difference; CI, confidence interval.

		Experim			ontrol					Weigh
Stu	dy	Events	Total	Events	Total	Odds Ratio	OR	95%-CI	(fixed)	(random
	saline									
	ha (2006)	33	45	16	48	<u></u>		[2.25; 13.43]		34.5%
Raz	avi (2015)	17	44	12	45	++++	1.73	[0.71; 4.25]	38.0%	34.3%
Rizv	ri (2015)	19	20	13	20		- 10.23	[1.12; 93.34]	6.3%	9.8%
	naz (2016)	7	35	4	35			[0.51; 7.33]		21.5%
		'	144	-	148					
	ed effect model		144		148	Ť		[1.77; 5.35]		-
	ndom effects model					$\sim$	3.14	[1.49; 6.62]		100.0%
Hete	erogeneity: $I^2 = 38\%, \tau^2$	<sup>-</sup> = 0.2132,	p = 0.	19						
Fixe	ed effect model		144		148		3.08	[1.77; 5.35]	100.0%	
	dom effects model				140	- A A A A A A A A A A A A A A A A A A A		[1.49; 6.62]		100.0%
	erogeneity: $I^2 = 38\%$ , $\tau^2$		p = 0.	19				,		
Res	idual heterogeneity: / <sup>2</sup> :	= 38%, p =	= 0.19			0.1 0.51 2 10				
		Experime	ental	Co	ntrol				Weight	Weigh
Stud		Events 1				Odds Ratio	OR	95%-CI		(random
a = 6	aline					13				
<u> </u>		40	45	40	40		0.64	10 10 2 00	10.40/	40.00
	na (2006)	42	45	46	48			[0.10; 3.82]		
	avi (2015)	14	44	8	45	12-		[0.80; 5.83]		
	(2015)	19	20	12	20	1 <del>1</del>	- 12.67	[1.40; 114.42]	8.4%	13.9
Solm	naz (2016)	10	35	10	35		1.00	[0.35; 2.82]	38.0%	33.5
	d effect model		144		148	$\Rightarrow$		[0.85; 3.04]		
	dom effects model				140	1 in the second se		[0.66; 4.38]		
		- 0 4470	0 4	0			1.70	[0.00, 4.30]		100.0
Heter	rogeneity: $I^2 = 46\%, \tau^2$	= 0.4172,	p = 0.1	3						
Fixe	d effect model		144		148		1.60	[0.85; 3.04]	100.0%	
							1 70	[0.66; 4.38]		100.09
Ran	dom effects model				_		1.70	[0.00, 4.00]		
Heter	rogeneity: $I^2 = 46\%, \tau^2$	= 0.4172,	p = 0.1	3	0.01		1	[0.00, 4.00]		
Heter	dom effects model rogeneity: $I^2 = 46\%$ , $\tau^2$ dual heterogeneity: $I^2 =$	= 0.4172, = 46%, p =	p = 0.1 0.13	3	0.01	0.1 1 10 10	1.70 00	[0.00, 4.00]		
Heter	rogeneity: $I^2 = 46\%, \tau^2$	= 0.4172, <i>p</i> = 46%, <i>p</i> = Experim	0.13		⊂ 0.01 ontrol	0.1 1 10 10	1			Weight
Heter Resid	rogeneity: $I^2 = 46\%$ , $\tau^2$ dual heterogeneity: $I^2 =$	= 46%, p =	0.13 nental	C	ontrol	0.1 1 10 10 Odds Ratio	1		Weight	Weight
Heter Resid	rogeneity: $I^2 = 46\%$ , $\tau^2$ dual heterogeneity: $I^2 =$	= 46%, <i>p</i> = Experim	0.13 nental	C	ontrol		00 OR	95%-CI	Weight (fixed) (	Weight
Heter Resid	rogeneity: / <sup>2</sup> = 46%, 5 <sup>2</sup> dual heterogeneity: / <sup>2</sup> = <b>dy</b> saline	Experim Events	0.13 nental Total	Co Events	ontrol Total		00 OR	95%-CI	Weight (fixed) (	Weight random)
Heter Resid Stu g = Bas	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006)	Experim Events	0.13 nental Total 45	Co Events 42	ontrol Total 48		00 OR 3.07 [	<b>95%-Cl</b> 0.59; 16.09]	Weight (fixed) ( 16.1%	Weight random) 19.6%
Heter Resid Stu g = Bas Raz	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006) ravi (2015)	= 46%, <i>p</i> = Experim Events 43 7	0.13 nental Total 45 44	Co Events 42 5	ontrol Total 48 45		00 OR 3.07 [ 1.51	<b>95%-Cl</b> 0.59; 16.09] [0.44; 5.19]	Weight (fixed) ( 16.1% 29.1%	Weight random) 19.6% 28.4%
Heter Resid Stu g = Bas Raz Rizv	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006) tavi (2015) <i>i</i> (2015)	= 46%, <i>p</i> = Experim Events 43 7 18	0.13 nental Total 45 44 20	Co Events 42 5 11	ontrol Total 48 45 20		OR 00 3.07 [ 1.51 - 7.36 ]	<b>95%-Cl</b> 0.59; 16.09] [0.44; 5.19] 1.34; 40.55]	Weight (fixed) ( 16.1% 29.1% 15.2%	Weight random) 19.6% 28.4% 18.8%
Heter Resid Stu g = Bas Raz Rizv Soli	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006) tavi (2015) <i>ii</i> (2015) maz (2016)	= 46%, <i>p</i> = Experim Events 43 7	0.13 nental Total 45 44 20 35	Co Events 42 5	<b>ontrol</b> <b>Total</b> 48 45 20 35		00 00 3.07 [ 1.51 - 7.36 ] 0.87	<b>95%-Cl</b> 0.59; 16.09] [0.44; 5.19] 1.34; 40.55] [0.30; 2.48]	Weight (fixed) ( 16.1% 29.1% 15.2% 39.7%	Weight random) 19.6% 28.4% 18.8% 33.3%
stu g = Bas Raz Rizv Soli Fixo	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006) tavi (2015) <i>ii</i> (2015) <i>maz</i> (2016) ed effect model	Experim Events 43 7 18 9	0.13 nental Total 45 44 20	Co Events 42 5 11	ontrol Total 48 45 20		OR 3.07 [ 1.51 - 7.36 [ 0.87 1.73 ]	<b>95%-Cl</b> 0.59; 16.09] [0.44; 5.19] 1.34; 40.55] [0.30; 2.48] (0.89; 3.36]	Weight (fixed) ( 16.1% 29.1% 15.2% 39.7% 100.0%	Weight random) 19.6% 28.4% 18.8% 33.3%
g = Bas Raz Rizv Solu Fixo Rar	rogeneity: $I^2 = 46\%, \tau^2$ dual heterogeneity: $I^2 =$ <b>dy</b> saline sha (2006) tavi (2015) <i>ii</i> (2015) <i>ia</i> (2015) ed effect model dom effects model	Experim Events 43 7 18 9	0.13 eental Total 45 44 20 35 144	Co Events 42 5 11 10	<b>ontrol</b> <b>Total</b> 48 45 20 35		OR 3.07 [ 1.51 - 7.36 [ 0.87 1.73 ]	<b>95%-Cl</b> 0.59; 16.09] [0.44; 5.19] 1.34; 40.55] [0.30; 2.48]	Weight (fixed) ( 16.1% 29.1% 15.2% 39.7%	Weight random) 19.6% 28.4% 18.8% 33.3%
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Fig. 5. Incidence of throat pain immediately after surgery (A) and at 2 hours (B), 6 hours (C), and 24 hours (D) postoperatively [1,5,9,10]. Experimental: pharyngeal packing, control: no packing. OR, odds ratio; CI, confidence interval.

6 hours (OR=1.01, 95% CI: 0.64–1.60,  $I^2=0\%$ ), and 24 hours (OR=1.18, 95% CI: 0.54–2.56,  $I^2=42\%$ ) (Fig. 3). No statistically significant inter-study heterogeneity ( $I^2<50\%$ ) was found in these outcomes. The Duval and Tweedie trim-and-fill method and the Egger test regarding postoperative nausea and vomiting at all time points were not conducted due to an insufficient number of enrolled studies (<10).

# Effect of pharyngeal packing on postoperative throat pain

Throat pain scores at recovery status immediately after surgery (SMD=0.40, 95% CI: 0.20-0.59, I<sup>2</sup>=43%) and at postoperative 24 hours (SMD=0.33, 95% CI: 0.11-0.55, I<sup>2</sup>=0%) were significantly higher in the packing group than in the control group, except at postoperative 2 hours (SMD=0.18, 95% CI: -0.37-0.74, I<sup>2</sup>=73%) and 6 hours (SMD=-0.04, 95% CI: -0.74-0.67, I<sup>2</sup>=83%) (Fig. 4). Similarly, the incidence of throat pain immediately after surgery (OR=3.08, 95% CI: 1.77-5.35, I<sup>2</sup>=38%) and at 24 hours (OR=2.23, 95% CI: 1.22-4.10,  $I^2=0\%$ ) were statistically significantly higher in patients who had received pharyngeal packing than in the control group (Fig. 5). In contrast, no significant differences were found in the incidence of postoperative nausea and vomiting between the two groups at postoperative 2 hours (OR=1.60, 95% CI: 0.85-3.04, I<sup>2</sup>=46%) and 6 hours (OR=1.73, 95% CI: 0.89 - 3.36,  $I^2 = 39\%$ ).

No statistically significant inter-study heterogeneity was found ( $I^2$ <50%) in these outcomes except for pain scores at postoperative 2 hours and 6 hours. The Duval and Tweedie trim-and-fill method and the Egger test for throat pain at alltime points were not conducted due to an insufficient number of enrolled studies (<10).

## Sensitivity analyses

Sensitivity analyses were performed to assess whether the pooled estimates of the incidence of postoperative nausea, vomiting, and sore throat differed by repeating the meta-analysis while excluding studies one at a time. The results were consistent with those presented above.

## **DISCUSSION**

Nausea and vomiting after surgery are uncomfortable and can be caused by bleeding, dehydration, electrolyte and acidbase imbalances, and lung aspiration. In addition, nausea and vomiting also prolong the time the patient stays in the treatment room, with longer anesthesia, and discharge may be delayed. Postoperative nausea and vomiting can also induce anxiety and reduce patients' satisfaction with nasal surgery. Therefore, routine patient management protocols should include evaluating and controlling symptoms of nausea and vomiting after nasal surgery [4].

We used detailed temporal categories in this meta-analysis to accurately reflect changes over time in the effect of pharyngeal packing on postoperative nausea and vomiting. Sore throat was also evaluated as a side effect.

The meta-analysis confirmed that postoperative nausea and vomiting symptoms showed no significant differences between groups, while sore throat in the recovery room was more frequent in the packing group than in the control group. It is known that pharyngeal packing may cause local trauma and inflammation of the pharyngeal mucosa, which may be associated with pharyngeal plexus injury and tongue swelling [1]. Trauma to the pharynx can irritate the vagus nerve nucleus in the brainstem and cause vomiting [2]. Considering this relationship, pharyngeal packing may in fact cause nausea and vomiting due to sore throat during postoperative recovery.

Although the results of the early (immediate recovery after surgery) and mid-term (2 hours after surgery) showed conflicting patterns, the total number of antiemetic administrations within 24 hours after surgery was statistically similar between the two groups. Ingested blood after surgery can induce very strong vomiting, but actual postoperative nausea and vomiting may be affected by a variety of factors, including individual patient factors, intraoperative management, and postoperative management [8]. Therefore, performing pharyngeal packing during nasal surgery can theoretically reduce blood product intake and reduce postoperative nausea and vomiting. However, other factors, such as damage to the pharyngeal mucosa and postoperative pain control, may also affect the incidence of postoperative nausea and vomiting. The analysis of patterns at multiple time points within 24 hours postoperatively in this study showed that inserting pharyngeal packing did not reduce postoperative nausea and vomiting, but increased throat discomfort in patients after surgery.

Our study had several limitations. The study included various surgical procedures, such as septoplasty, rhinoplasty, and endoscopic sinus surgery. The amount of bleeding and packing time may vary depending on the severity of the condition or differences in individual surgical methods. This issue may explain the high heterogeneity in the values of throat pain. Nonetheless, although several studies (either case series or case-control studies) have presented mixed results regarding the effectiveness of pharyngeal packing, only randomized controlled trials were included in this meta-analysis to improve the validity of the findings.

## CONCLUSION

This study confirmed that intraoperative pharyngeal pack-

ing during nasal surgery may not effectively reduce postoperative nausea and vomiting, but may cause throat pain.

#### **Ethics Statement**

Ethical approval and informed consents does not apply to this article.

#### Availability of Data and Material

The datasets generated or analyzed during the study are available from the corresponding author on reasonable request.

#### **Conflicts of Interest**

Do Hyun Kim and Se Hwan Hwang who are on the editorial board of the *Journal of Rhinology* were not involved in the editorial evaluation or decision to publish this article.

#### **Author Contributions**

Conceptualization: Se Hwan Hwang. Data curation: Do Hyun Kim, Se Hwan Hwang. Formal analysis: Se Hwan Hwang. Methodology: Se Hwan Hwang. Project administration: Do Hyun Kim. Supervision: Se Hwan Hwang. Validation: Do Hyun Kim. Visualization: Se Hwan Hwang. Writing—original draft: Do Hyun Kim. Writing—review & editing: Do Hyun Kim, Se Hwan Hwang.

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