Which heat and moisture exchanger (HME) with bacteria-filtering property should be used?

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Study population: mechanical ventilated ICU patients

Comparison: different HMEs with bacteria-filtering property compared to each other

Outcome: ventilator-associated pneumonia

Methods

Data sources

Publications were retrieved by a search of Medline and the Cochrane Library up to february 2006. Terms included were 'pneumonia' and 'ventilator*' and 'heat and moisture exchanger*'. To identify randomised controlled trials in Medline the following search strategy was used: (humid* OR humidification OR circuit* OR humidity OR humidifier OR humidifiers OR heat and moisture exchanger* OR artificial nose) AND (((ventilator associated pneumonia) OR (VAP AND (pneumonia OR pneum*)) OR ("Respiration, Artificial"[MAJR] AND pneumonia) OR (ventilated AND pneumonia) OR (ventilation AND pneumonia)) AND (((randomized controlled trial[pt] OR controlled clinical trial[pt] OR randomized controlled trials[mh] OR random allocation[mh] OR double-blind method[mh] OR single-blind method[mh] OR clinical trial[pt] OR clinical trials[mh] OR ("clinical trial"[tw]) OR ((singl*[tw] OR doubl*[tw] OR tripl*[tw]) AND (mask*[tw] OR blind*[tw]))) OR ("latin square"[tw]) OR placebos[mh] OR placebo*[tw] OR random*[tw] OR research design[mh:noexp] OR comparative study[mh] OR evaluation studies[mh] OR follow-up studies[mh] OR prospective studies[mh] OR cross-over studies[mh] OR control*[tw] OR prospective*[tw] OR volunteer*[tw]) NOT (animal[mh] NOT human[mh]))))). Additionally, all reference lists of identified trials were examined.

Selection criteria

All randomised and quasi-randomised trials comparing different brands of HMEs and ventilator-associated pneumonia as the outcome measure.

Review methods

Data were extracted by two reviewers independently and compared. Disagreements were resolved by discussion. Data from the original publications were used to calculate the relative risk of ventilator-associated pneumonia. Data for similar outcomes were combined in the analysis where appropriate, using a random-effects model.

Results

Two parallel-group randomised controlled trials were included (1, 2).

Study population, interventions and outcome definitions See Table I

Validity assessment See Table II

Summary estimates of associations between treatment and control group See Figure I

Table I: Study population, interventions and outcome definitions

	Participants	Interventions	Definition of ventilator associated pneumonia (VAP)
Thomachot et al. 1999	Incl: all ICU patients, ventilation ≥ 24 hrs Excl: not reported	Treatment (77 analyzed): Humid-Vent Filter Light (Louis Gibek AB, Upplands-Vasby, Sweden)	VAP was defined as purulent ETS or worsening of PaO ₂ and new infiltrates and a positive quantitative culture from a distal airway sample (BAL $\geq 10^4$ CFU/ml or PSB $\geq 10^3$ CFU/ml)
	Mean number of ventilation days (SD): T: 11.1(6.8); C: 12.3(7.8)	Control (63 analyzed): Clear ThermAl 1841 (Intersurgical, Fontenay-sousbois, France)	
		Note: 1) HMEs with different humidification compounds and different filter membranes; 2) HMEs changed daily	
		End of the study protocol: not reported	

Thomachot et al. 1998	Incl: all ICU patients, ventilation ≥ 24 hrs, no contraindications of HMEFs (bronchopleuro-cutaneous fistula, hemoptysis, significant gas leakage around the tracheal tube cuff	Treatment (66 analyzed): Humid-Vent Filter Light (Gibeck, Upplands Vaesby, Sweden) VAP: T: 21/66	VAP was defined as purulent ETS or worsening of PaO ₂ and new infiltrates and a positive quantitative culture from a distal airway sample (BAL $\geq 10^4$ CFU/ml or PSB $\geq 10^3$ CFU/ml)
	Excl: not reported	Control (70 analyzed): Pall BB 100 (Pall, Newquay, UK) VAP: C: 26/70	
	Mean number of ventilation days (SD): T: 11.7 (11.0); C: 12.2 (12.0)	End of the study protocol: not reported	

Table II: Data on quality assessment

Thomachot et al. 1999	Generation of allocation sequence:	Not reported
	Concealment of allocation:	Unclear
	Blinding attending physician:	No
	Blinding outcome assessors:	No
	Description of dropouts:	No
	Analysis by intention-to-treat:	Unclear
Thomachot et al. 1998	Generation of allocation sequence:	Not reported
	Concealment of allocation:	Unclear
	Blinding attending physician:	No
	Blinding outcome assessors:	No
	Description of dropouts:	No
	Analysis by intention-to-treat:	Unclear

Figure I: Summary estimates of associations between treatment and control group expressed as relative risk (RR) and 95% confidence interval (CI) using a random effects model

Review: VAP - HMEF / different brands

Comparison: 01 HMEF vs HMEF

Outcome: 01 Ventilator-associated pneumonia

Study or sub-category	Treatment n/N	Control n/N		RR (rand 95% (,	Weight %	RR (random) 95% CI
Thomachot '98	21/66	26/70		_	-	51.62	0.86 [0.54, 1.37]
Thomachot '99	24/77	21/63		-	_	48.38	0.94 [0.58, 1.51]
Total (95% CI)	143	133		•		100.00	0.89 [0.64, 1.25]
Total events: 45 (Treatment	t), 47 (Control)			1			
Test for heterogeneity: Chi ²	$I = 0.07$, df = 1 (P = 0.80), $I^2 = 0$	0%					
Test for overall effect: Z = 0	0.66 (P = 0.51)						
			0.1 0.2	0.5 1	2	5 10	
			Favours	treatment I	Favours c	control	

Conclusion

The evidence of two trials indicates that different brands of HMEs have not any effect on the incidence of ventilator-associated pneumonia. The evidence, however, was low because of small sample sizes and insufficient methodological quality.

References

- 1. Thomachot L, Viviand X, Arnaud S, Boisson C, Martin CD. Comparing two heat and moisture exchangers, one hydrophobic and one hygroscopic, on humidifying efficacy and the rate of nosocomial pneumonia. Chest 1998;114:1383-1389.
- 2. Thomachot L, Vialet R, Arnaud S, Barberon B, Michel-Nguyen A, Claude M. Do the components of heat and moisture exchanger filters affect their humidifying efficacy and the incidence of nosocomial pneumonia? Crit Care Med 1999;27(5):923-928.