

Table 2: Characteristics of the two studies

	Study period I (3)	Study period II (6)
Years in which inclusion took place	2004-2006	2009-2013
Participating hospitals (n)	13	16
Patients included (n)	5939	11997
Duration of intervention		
SDD	6 months	12 months
SOD	6 months	12 months
Standard care	6 months	NA
Inclusion criteria to receive SDD/SOD	Mechanical ventilation > 48 hours and/or Length of stay > 72 hours	Mechanical ventilation > 48 hours
Antibiotics used in SDD/SOD		
Tobramycin	Mouthpaste: 2% concentration Suspension: 80mg <sup>a</sup>	<i>idem</i>
Colistin	Mouthpaste: 2% concentration Suspension: 100mg <sup>a</sup>	<i>idem</i>
Amphotericin	Mouthpaste: 2% concentration Suspension: 500mg <sup>a</sup>	<i>idem</i>
Systemic prophylaxis	Cefotaxim 4000mg daily <sup>a</sup>	<i>idem</i>
<b>Methods</b>		
Point prevalence surveys available (n)	18	24
Culture media used for screening in point prevalence surveys	McConkey with - polymyxin B 50iu/ml (5mg/l) - tobramycin 8mg/l - cefotaxim 8mg/l - ciprofloxacin 2mg/l	McConkey with - polymyxin E (colistin) 4mg/l - tobramycin 8mg/l ESBL chromogenic agar VRE chromogenic agar

<sup>a</sup> only during SDD

Abbreviations: SDD: selective digestive tract decontamination; SOD: selective oropharyngeal decontamination; NA: not applicable; ESBL: extended spectrum beta-lactamase; VRE vancomycin resistant enterococcus