

Appendix 1. Study settings and designs of the included studies (n= 26) presented in alphabetical order.

Reference	Ward or pharmacy based preparation or administration	Objective and Study design	Aseptically prepared medicines	Contamination/ results	Method of microbiological analysis
Original research/studies (n=19)					
Austin and Elia 2013	Ward based preparation	To examine whether different aseptic techniques affect the contamination rates of intravenous doses prepared on hospital wards. A comparative study.	A growth medium infusate was used to make flushes to syringes	The pharmacy operator achieved lower contamination rates than nurses.	The airborne and surface environmental contamination was monitored during session. Prepared syringes were incubated and contaminants identified
Bertoglio et al 2013	Ward based catheter flushing and locking	To compare the use of pre-filled syringe with manually filled syringes for flushing for reducing catheter-related bloodstream infections (CRBSI). A retrospective observational cohort study.	Saline syringes	The CRBSI rate was 2,7 % in the pre-filled syringe group and 6,3 % in the manually filled syringe group	Blood samples and bacterial identification
Buerke et al 2010	Ward-based (CT unit) contrast agent administration via automatic injectors	To compare single- and multiple use of syringes in automatic injectors to evaluate microbiologic contamination and time benefit. A comparative study.	CT contrast agent and saline	No bacterial contamination of prefilled single use contrast or saline syringes. Contamination of two saline syringes with multiple use of saline syringes	Surface and palm samples: replicate organism detection and counting agar plates and Contrast or saline sample: injection to tryptic soy broth and after incubation streaking to agar plates
Chemaly et al 2011	Manufactured syringes	To evaluate contaminated pre-filled heparin and saline syringes clinical impact in patients. A multicenter case series study.	Pre-filled heparin and saline syringes	Patients got clinical symptoms	Blood culture and bacterial identification of automated methods
Crill et al 2010	Pharmacy based delivery method	To evaluate microbial contamination associated with different methods of neonatal intravenous fat emulsion preparation and delivery. A comparative study.	Intravenous fat emulsion infusions	The number of contaminated samples did not significantly differ between these different methods	Each sample was inoculated into a blood culture bottle, incubated and then microorganism were identified
De Giorgi et al 2010	Ward based preparation and administration	To analyze safety risks in injectable medications and to assess the potential impact and pharmaco-economic	Gentamicin, morphine and dopamine	Thirty-one failure modes identified and the most critical failure mode was microbial contamination	

		aspects of safety tools. A prospective study.			
De Smet et al 2012	Ward based administration and catheter flushing	To study if Burkholderia cepacia bloodstream infections may result from the use of Ringer lactate solution as multiple-dose vial for catheter flushing. A prospective surveillance study. A simulation and observation study. A comparative study.	Ringer lactate solution as multiple-dose vial for catheter flushing	Three patients died from burkholderia cepacia bloodstream infection.	Blood cultures are sampled when signs of Systemic Inflammatory Response are present. Blood is cultured and incubated and organisms identified. Also environmental sampling were taken.
Dias et al 2008	Pharmacy based preparations	To management of patients with CVC infection due to contaminated infusate and to define the amount of patients to get the infection through this route. A descriptive study.	Heparin catheter-lock solution	48 patient get this contaminated solution and 32 of these patients get infection due to <i>P.putida</i> , the same bacilli isolated from heparin solution	Blood culture and bacterial identification of automated methods
Dolan et al 2010		To provide practise guidance for health care facilities on essential safe injection, infusion, and vial practices that should be implemented in such settings. Good practices for safe infusion.		List of aseptic techniques to environmental conditions, IV solutions, flushing, syringes, vials, blood glucose monitoring devices and HCW	
Gargiulo et al 2012	Ward prepared and admistered infusions	To evaluate the possibility that anaesthetists are administering potentielly pathogenic micro-organisms to their patients. A prospective microbiological and observational study.	Sterile water for injection and sodium chloride powder	Organisms were isolated from 13 % of bags, 5 % of syringes and 35 % of needles	The samples were filtered and filter mambrane placed onto agar plate and then incubation ja identification. Also needles analysed in sterile tryptic soy broth. Environment contamination investigated by agar plates and organisms were identified.
Gershman et al 2008	Pharmacy prepared syringes	To summarize the findings of those bloodstream infections after exposure to contaminated flush solutions. A comparative study.	Heparinized Saline Flush syringes	There were for example fever and chills among patients who get contaminated flushing solutions	Cultures of blood samples, of sections and of syringes were performed at microbiology laboratories
Gorski 2010	Home infusion therapy with central venous access device	To bring out techniques that prevent CVAD-infections.		Infection related to CVADs in home care patient are preventable	

		Recommendations for best practise in home infusion therapy.			
Isanhart et al 2008	Media-fill test in an academic laboratory setting	To evaluate aseptic skills of pharmacy student near the beginning and end of course. A comparative study.	A low- and a medium-risk media fill test to prepare syringes	The overall contamination rate at the baseline was 4,2 % versus 0 % at the final assesment.	Prepared syringes were sealed and incubated and then evaluated for presence of microbial contamination (solution turbidity)
Kerenyi et al 2011	Ward prepared infusions	To examine contamination rates of syringes with different drugs used to administer intravenous medications. A comparative study.	The investigated medications were cordarone, furosemide, noradrenaline, adrenaline, bupivacaine+sufentanil, metoprolol, insulin, propofol, potassium chloride and morphin	The overall contamination rate was 16 % and insulin had higher contamination rates	Culturing to different agar plates, incubation and identification and also determined antibiotic sensitivities
Rangel-Frausto et al 2010	Ward based administration	To compare closed and open infusion container for prevention of bloodstream infections. An open label, prospective cohort, active healthcare – associated infection surveillance.	Baxter Viaflex and commercially available open infusion containers	The CLAB rate was significantly higher during the open versus the closed container period.	Blood culturing and microorganism identification with standard laboratory methods
Sigward et al 2012	Pharmacy based preparation	To assess aseptic technique of pharmacy operators. A comparative study.	Simulated media-fill test (vials, syringes, minibags)	The overall operator failure rate in aseptic simulation test was 40 % and 2,3 % of the 300 preparations were contaminated	Filters incubated at agar plate and micro-organisms were identified with Vitek-2 identification system
Stucki et al 2009	Pharmacy and ward prepared media-fill testing	To evaluate the direct influence of environmental cleanliness and risk manipulations on prepared syringes. A comparative study.	Syringes filled with tryptic soy broth and air	None prepared in cleanroom contained microorganisms, 6 % were contaminated in the operating room and 16 % were contaminated in the ward. These results correlate with the airborne particulate.	Air particulate contamination determined with a discrete particle counter
Yoshida et al 2008	Both pharmacy and ward prepared infusions	To evaluate the risk factors for catheter-related bloodstream infection with CV catheters. A comparative study.	Different preparations, no meaning to results	Positive cultues showed an overall incidence of 2.26 per 1000 device-days. Higher odds ratio of positive cultures showed ICU and CV catheter placement for more than 30 days	Blood samples and bacterial identification
Younger et al 2008	Ward based setting up and priming an intravenous	Step-by-step guide for the equipment required, correct			

	infusion	preparation of patient and the procedure in setting up intravenous infusion. Article is also concerned about infection risk in intravenous infusion. Nursing standards.			
Letters, editorials and commentaries (n=7)					
Agalloco et al 2009	Pharmacy based preparation (cleanroom hand filling)	Commentary on hand filling in cleanrooms which is soon history and we have to use new technologies to guarantee the quality			
Chemaly et al 2009	Pre-filled syringes	The letter to emphasize infection control teams role in identifying contaminated sources	Heparin and saline pre-filled syringes		
Cozanitis and Mäkelä 2008	Ward based administration	Letter to point out that using the same syringe for a number of could harm patients	Propofol	Infection from contaminated syringes has been reported overwhelmingly	
Gorski 2013	Ward and home based infusion preparation and administration	Editorial to address infection prevention, peripheral intravenous catheter use and alternative site infusions		Infusion nurses across the globe will continue to face challenges as new technology emerges	
Kuehn 2012	Commentary of unsafe injection practises on medical wards	The purpose is to campaign against unsafe injection practises and to increase the knowledge of risks to patients because of unsafe practices			
McCrea 2013	Single-use vials	Column to explore the federal policies regarding safe injection practices and the focus on compliance			
Nakataki et al 2013	Infusion set needles	Brief report to examine the incidence of beacterial contamination in infusion set needles	Electrolytes or parenteral nutrition	Bacterial contamination was detected in 5,7 % of the samples	Needle was plated in culture medium and then inoculated medium was added to agar plates, incubated and then identified