





Evidence assessment: Summary of a systematic review

Who is this summary for?

This evidence assessment is meant for clinicians, administrators of health facilities and decision makers.

Interventions to improve antibiotic prescribing practices for hospital inpatients

Key findings

- Antibiotic resistance is a major public health problem caused by innapropriate use of antibiotics
- Restrictive methods (limiting the use of anitbiotics) and persuasive methods (*advising/educating physicians on the use of antibiotics*) methods can improve prescription practices, and lead to a reduced number of infections in hospitals, deaths and length of stay.
- Restrictive methods have a larger effect than persuasive methods.
- The studies included in this review were conducted in multiple countries, and the applicability of the interventions tested is broad.

Background

Antibiotic resistance is a serious problem for individual patients and health care systems. Illnesses caused by resistant bacteria are more difficult to treat and lead to higher rates of morbidity and mortality, and longer hospital stays. Up to 50% of antibiotic use in hospitals may be inappropriate.

Question

What types of interventions can improve prescribing of antibiotics to hospital inpatients?

Prescription of antibiotics in Cameroon: The prescription of antibiotics in Cameroon is not always governed by standardized protocols. Due to the limited availability and use of culture and sensitivity (identifying which medication will work best), many antibiotics are used inappropriately. This is aggravated by their over-the-counter availability. In addition, it is unclear how well recommended guidelines on antibiotic use are respected in Cameroon.

	What the review authors searched for	What the review authors found
Studies	Randomized and Controlled Clinical Trials	66 studies were included in the review. 43 were
	(RCT/CCT); controlled Before and After studies	ITSs, 13 were RCTs, 6 were CBAs, 2 were
	(CBA) and Interrupted Time series (ITS) on antibiotic prescribing to hospital inpatients.	CCTs, 1 was a Cluster CCT and 1 was a Cluster RCT.
Participants	Health care professionals who prescribe	The interventions identified were delivered to
i ai cicipantis	antibiotics to hospital inpatients receiving acute	pharmacists in 22 studies, a specialist physicia
	care.	in 17 studies, a multi-disciplinary antimicrobia
		management team in 11 studies, change i
		antibiotic policy in 7 studies, physicians in th
		targeted department in 4 studies, by computer in
		4 studies and written feedback in 2 studies.
Interventions	A: Persuasive interventions:	Persuasive interventions:
	1. Dissemination of educational materials in	Educational outreach (22 interventions)
	printed form or via educational meetings; 2. Reminders;	dissemination of educational material (
	3. Audit and feedback;	interventions); reminders (8 interventions); audi and feedback (9 interventions).
	4. Educational outreach (academic detailing or	Restrictive interventions:
	review and recommend change).	Compulsory order forms (5 studies); exper
	B: Restrictive interventions:	approval (9 studies); removal by restriction (8
	1. Compulsory order form - prescribers had to	studies); review and make change (4 studies).
	complete a form with clinical details to justify use	Structural interventions (8 studies)
	of the restricted antibiotics;	
	2. Expert approval - the prescription for a	
	restricted antibiotic had to be approved by third	
	party 3. Restriction by removal - for example by	
	removing restricted antibiotics from drug	
	cupboards;	
	4. Review and make change - a reviewer changed	
	the prescription	
	C: Structural interventions: Health system	
	changes	
Controls		
Controls	No controls specified	No controls specified
	Antibiotic prescribing process measures	Fifty-two studies provided data about dru
	Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 10
Controls Outcomes	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 10 about microbiological outcomes. Fifty one studie
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 10 about microbiological outcomes. Fifty one studie provided interpretable data about only on
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 1 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only i
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 1 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only i three studies and microbiological only in 1
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 1 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only i three studies and microbiological only in 1 studies. Fifteen studies provided data about
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 10 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about drug outcomes, 14 about clinical outcomes and 16 about microbiological outcomes. Fifty one studies provided interpretable data about only on outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in nine studies, drugs plus microbiological in fou
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	No controls specified Fifty-two studies provided data about drug outcomes, 14 about clinical outcomes and 16 about microbiological outcomes. Fifty one studies provided interpretable data about only one outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study Only one study provided data about all three
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about drug outcomes, 14 about clinical outcomes and 16 about microbiological outcomes. Fifty one studies provided interpretable data about only one outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about drug outcomes, 14 about clinical outcomes and 16 about microbiological outcomes. Fifty one studies provided interpretable data about only one outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study Only one study provided data about all three outcomes.
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about drug outcomes, 14 about clinical outcomes and 16 about microbiological outcomes. Fifty one studies provided interpretable data about only one outcome: drugs only in 38 studies, clinical only in three studies and microbiological only in 10 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical in nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study Only one study provided data about all three outcomes. A fourth outcome ("Financial") is restricted to studies that provided information about the cos
	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 1 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only i three studies and microbiological only in 1 studies. Fifteen studies provided data about more than one outcome: drugs plus clinical i nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study Only one study provided data about all thre outcomes. A fourth outcome ("Financial") is restricted t studies that provided information about the cos of developing or implementing the intervention i
Outcomes	 Antibiotic prescribing process measures (decision to treat, choice of drug, dose, route or duration of treatment); Clinical outcome measures (mortality, length of hospital stay); Microbial outcome measure (colonization or infection with Clostridium difficile or 	Fifty-two studies provided data about dru outcomes, 14 about clinical outcomes and 1 about microbiological outcomes. Fifty one studie provided interpretable data about only on outcome: drugs only in 38 studies, clinical only i three studies and microbiological only in 1 studies. Fifteen studies provided data abou more than one outcome: drugs plus clinical i nine studies, drugs plus microbiological in fou studies, clinical plus microbiological in one study Only one study provided data about all thre outcomes. A fourth outcome ("Financial") is restricted t studies that provided information about the cos

this review was limited by the small number of comparable studies. Long term effects were not covered. **Citation:** Davey P, Brown E, Charani E, Fenelon L, Gould IM, Holmes A, Ramsay CR, Wiffen PJ, Wilcox M. Interventions to improve antibiotic prescribing practices for hospital inpatients. Cochrane Database of Systematic Reviews 2013, Issue 4. Art. No.: CD003543. DOI: 10.1002/14651858.CD003543.pub3.

Summary of findings:

Patient or population: Healthcare professionals Settings: Secondary care (inpatients in acute, not long term care only) Intervention: Any intended to improve antibiotic prescribing Comparison: Usual care

Outcomes	Effect measure (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Restrictive versus P	ersuasive interventions		•
Appropriate prescribing of antibiotics	32% difference in effect size (restrictive-persuasive) at one month 95% Cl 2 to 61%	53 comparisons from 40 studies (all ITS) in 46 hospitals	Low ⊕⊕00
	No significant difference at 6, 12 or 24 months		Indirect comparison between studies that provide data about effect of either persuasive or restrictive interventions
Microbial outcomes	53% difference in effect size (restrictive-persuasive) at 6 months 95% Cl 31 to 75% No significant difference at	20 comparisons from 14 studies (all ITS) in 14 hospitals	Low ⊕⊕00
	12 or 24 months		Indirect comparison between studies that provide data about effect of either persuasive or restrictive interventions
Patient outcomes	Risk of mortality for intervention versus control 0.92 (95% CI 0.81 to 1.06)	11 comparisons from 11 studies (7 RCT, 3 cluster-RCT, 1 cluster CCT) in 20 hospitals with	Moderate ⊕⊕⊕0
		9,817 patients	High risk of bias especially around study design
	Difference (in days) in length of stay for intervention versus control -0.04 days (95% Cl - 0.34	6 comparisons from 6 studies (4 RCT, 2 cluster-RCT) in 8	Very Low ⊕000
	to 0. 25)	hospitals with 8,071 patients	Studies are very heterogeneous and have high risk of bias
Interventions inter	ded to increase effective a	intibiotic prescribin	ng for pneumonia
Patient outcomes	Risk of mortality for intervention versus control 0.89 (95% Cl 0.82 to 0.97)	4 comparisons from 4 studies (3 CBA, 1 RCT) in 104 hospitals with 22,526 patients	Low ⊕⊕00
			High risk of bias especially around study design

Abbreviations

CBA: controlled before and after; CCT: controlled clinical trial; CI: confidence interval; ITS: interrupted time series; RCT: randomized controlled trial

Applicability

Forty two studies were from the USA. The remaining 24 studies were from 10 countries: Australia (2), Brazil (1), Canada (4), Colombia (1), France (2), Netherlands (2), Norway (1), Spain (1), Thailand (2) and the United Kingdom (8). Even though none of these studies was conducted in Africa, some of these interventions can easily be applied in low resource settings.

Conclusions

A wide variety of interventions has been shown to be successful in changing antibiotic prescribing to hospital inpatients. Restrictive methods are more effective than persuasive methods.

Prepared by

Marius Vouking, Christine Danielle Evina, Violette Tamo, Lawrence Mbuagbaw, Pierre Ongolo-Zogo: Centre for the Development of Best Practices in Health, Yaoundé, Cameroon. November 2013