Additional File 2: Sheet used to extract data from controlled studies on hypertension care in China

Study cha	racteristics										
Code	Study	Year	Design	Year of study	Size	Follow up year	Setting	Region	Participants	Age	Male (%)
Interventi	on, descript	tion:									
Main components						Providers					_
Patient education	Provider education	Improved monitor	self-manage	Family support	Organiza- tional	Unclear	GP/ Doctor	Nurse	Other	Control	
Results	(50)										
Systolic BP (I Baseline	mean+/-SD) mi	mнg	Daniella e			For donation			En do obot		_
			Baseline Control			Endpoint Intervention			Endpoint Control		+
Intervention n	Mean	SD	Control	Mean	SD	n Mean		SD	n	Mean	SD
D' !' . DD	(00)										
Diastolic BP (mean+/-SD) m	ттнд	Baseline			For donators		-	For docation		
Intervention			Control			Endpoint Intervention			Endpoint Control		_
n	Mean	SD	n	Mean	SD	n	Mean	SD	n	Mean	SD
Regular treat	ment rate										
Baseline			Baseline			Endpoint			Endpoint		
Intervention			Control			Intervention			Control		
n	event	%	n	event	%	n	event	%	n	event	%
HBP Control	rate										
Baseline			Baseline			Endpoint			Endpoint		
Intervention			Control			Intervention			Control		
n	event	%	n	event	%	n	event	%	n	event	%

eBox 3. Method used to assess the quality of included studies

"Quality Assessment tool for Quantitative Studies"

(http://www.nccmt.ca/registry/view/eng/14.html) was developed by the Effective Public Health Practice Project (EPHPP) to the assessment of studies on public health interventions. We simplified this tool and used the following checklist to assess the quality of included studies in this review:

- Study design: RCT, CT, quasi-experimental
- Participant representativeness: Are study participants likely to be representative of the target population? (very likely; somewhat likely; not likely; can't tell)
- Comparability between groups within a study: Were there between group differences for important confounders reported in the paper? (Yes, No, Can't tell)
- Blinding: Was the outcome assessor blinded to the intervention status? (Yes, No, Not reported, Not applicable)
- Withdrawal/drop outs: rate of withdrawals /drop-outs during study period; or Not Reported, NA.
- Sample size: Sample size or power calculation? (Yes, Partially, No)
- Intervention contamination: Control participants also received some interventions?
 (Very likely, somewhat likely, unlikely; not applicable)
- Completeness of outcomes measured/reported: BP outcome (yes/no), Treatment compliance outcome (Yes/no), Hypertension control outcome (Yes/no), CVD outcome (yes/no)