## G.14.2 Management strategies for people living with dementia and co-existing physical long term conditions

• What are the optimal management strategies (including treatments) for people living with dementia with co-existing physical long term conditions?

## G.14.2.1 Hypertension

Quality assessment								No of patients		Quality
o of dies	Design Ris		Indirectness	Inconsistency	Imprecision	Other considerations	Telmisartan (n=10)	Amlodipine (n=10)	Summary of results	
nical pro	ogression of co	morbidity	& associated	symptoms						
an diffe	rence in systoli	ic BP at 6 n	nonths (PPAR	versus CCB)						
	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD 2.00 (-7.64, 11.64)	Very low
an diffe	rence in diasto	lic BP at 6	months (PPAI	R versus CCB)						
	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD -2.00 (-8.20, 4.20)	Very low
an diffe	rence in pulse i	rate at 6 mo	onths (PPAR v	versus CCB)						
	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD 2.00 (-1.61, 5.61)	Very low
ical ou	ıtcomes, includ	ing cogniti	ve, functional	, behavioural a	bility					
an diffe	rence in MMSE	at 6 month	ns (PPAR vers	us CCB)						
	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD 0.00 (-3.10, 3.10)	Very low
an diffe	rence in ADAS-	Cog at 6 m	onths (PPAR	versus CCB)						
	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD -1.10 (-6.32, 4.12)	Very low
an diffe	rence in WMS-F	R (logical- ı	memory) at 6	months (PPAR	versus CCB)					
. –	Randomised open label trial	Serious <sup>1</sup>	Not serious	N/A	Very serious <sup>2</sup>	None	10	10	MD 3.00 (-0.18, 6.18)	Very low
ne 2)	Randomised	Serious <sup>1</sup>	Not serious	•	Very	None	10	10		

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Quality assessment							No of p	atients	Effect estimate	Quality
No of studies	Design	Risk of bias	Indirectness	Inconsistency	Imprecision	Other considerations	Telmisartan (n=10)	Amlodipine (n=10)	Summary of results	

		No of patients		Effect estimate	Quality					
No of studies	Design	Risk of bias	Indirectness	Inconsistency	Imprecision	Other considerations	Relative- HBPM (n=60)	ABPM (n=60)	Summary of results	
Clinical p	progression of comorbidi	ty & associ	iated symptor	ns						
Mean diff	ference in systolic BP aft	er 3 days (I	R-HBPM versi	us 24-h ABPM)						
Plichart (2013)	Randomised open comparative cross over study	Serious <sup>1</sup>	Not serious	N/A	Serious <sup>2</sup>	None	60	60	MD 11.30 (4.61, 17.99)	Low
Mean diff	ference in diastolic BP af	ter 3 days (	R-HBPM vers	us 24-h ABPM)						
Plichart (2013)	Randomised open comparative cross over study	Serious <sup>1</sup>	Not serious	N/A	Serious <sup>2</sup>	None	60	60	MD 1.00 (-2.76, 4.76)	Low
Mean diff	ference in systolic BP aft	er 3 days (l	R-HBPM versi	us day ABPM)						
Plichart (2013)	Randomised open comparative cross over study	Serious <sup>1</sup>	Not serious	N/A	Serious <sup>2</sup>	None	60	60	MD 9.70 (3.08, 16.32)	Low
Mean diff	ference in diastolic BP af	ter 3 days (	(R-HBPM vers	us day ABPM)						
Plichart (2013)	Randomised open comparative cross over study	Serious <sup>1</sup>	Not serious	N/A	Serious <sup>2</sup>	None	60	60	MD 0.00 (-3.76, 3.76)	Low
	owngrade 1 level, crossover of hort follow up period, 3 days	comparative of	design							

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