

Evidence Update

Non-Communicable Diseases Series

Do beta-blockers prevent heart disease and strokes in people with high blood pressure?

In people with high blood pressure, there is no evidence that beta-blockers reduce the number of deaths. Beta-blockers reduce the risk of stroke but are less effective than calcium channel blockers (CCBs) or renin-angiotensin system (RAS) inhibitors.

Inclusion criteria

Studies:

Randomized controlled trials.

Participants:

Adults, excluding pregnant women, with hypertension (high blood pressure) as defined by the study authors.

Intervention:

Beta-blockers either alone or as a first-line drug in a stepped care approach.

Control: no treatment, placebo, or alternative antihypertensive drug.

Outcomes:

Primary: death.

Secondary: coronary heart disease; stroke.

Adverse events: any adverse events.

Results

- Thirteen trials including 91,561 participants met the inclusion criteria. Nine used adequate methods to conceal allocation. Atenolol was the beta-blocker used in 75% of the participants in this review.
- Risk of death was no different for beta-blockers compared to no treatment or placebo (23,613 participants, 4 trials), diuretics, or RAS inhibitors, but it was significantly higher compared to CCBs (relative risk 1.07, 95% confidence interval 1.00 to 1.14; 44,825 participants, 4 trials).
- The effect of beta-blockers on coronary heart disease was no different to placebo, diuretics, RAS inhibitors, or CCBs.
- Participants treated with beta-blockers had a lower risk of developing stroke compared with placebo (RR 0.80, 95% Cl 0.66 to 0.96; 23,613 participants, 4 trials), but the risk was higher compared to CCBs (RR 1.24, 95% Cl 1.11 to 1.40; 44,167 participants, 3 trials) and RAS inhibitors (RR 1.30, 95% Cl 1.11 to 1.53; 9951 participants, 2 trials).
- Participants on a beta-blockers were more likely to stop treatment due to adverse events than those on a diuretic (RR 1.86, 95% Cl 1.39 to 2.50; 11,566 participants, 3 trials) or a RAS inhibitor (RR 1.41, 95% Cl 1.29 to 1.54; 9951 participants, 2 trials), but there was no significant difference with CCBs.







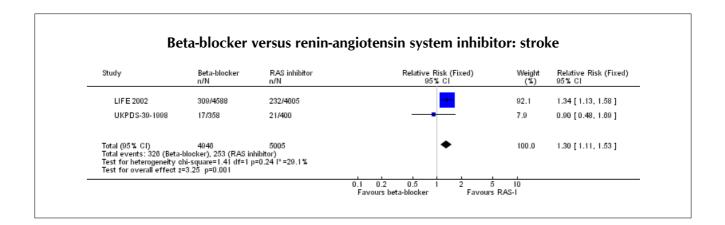
Adapted from Wiysonge CS, Bradley H, Mayosi BM, Maroney R, Mbewu A, Opie LH, Volmink J. Beta-blockers for hypertension. *Cochrane Database of Systematic Reviews* 2007, Issue 1. Art. No.: CD002003. DOI: 10.1002/14651858.CD002003.pub2. *Evidence Update* published in February 2008.

Produced by: the Effective Health Care Research Programme Consortium (www.liv.ac.uk/evidence), Liverpool School of Tropical Medicine, supported by the Department for International Development UK; and the Australasian Cochrane Centre. *Evidence Update* can be distributed free of charge.

Beta-blocker versus placebo or no treatment: death

Study	Beta-blocker n/N	Placebo n/N	Relative Risk (Fixed) 95% Cl	Weight (%)	Relative Risk (Fixed) 95% Cl
Coope 1986	60/419	69/465		11.7	0.97 [0.70, 1.33]
IPPPSH 1985	108/3185	114/3172		20.4	0.94 [0.73, 1.22]
MRC 1985	120/4403	253/8654		30.5	0.93 [0.75, 1.15]
MRC0A 1992	167/1102	315/2213	+	37.4	1.06 [0.90, 1.27]
Total (95% CI) Total events: 455 (Beta Test for heterogeneity Test for overall effect	chi-square=1.14 df=3	14504 p=0.77 I==0.0%	•	100.0	0.99 [0.88, 1.11]

Study	Beta-blocker n/N	CCB n/N	Relative Risk (Fixed) 95% Cl	Weight (ቕ)	Relative Risk (Fixed) 95% Cl
ASCOT 2005	422/9618	327/9639	—	63.8	1.29 [1.12, 1.49]
ELSA 2002	14/1157	9/1177		1.7	1.58 [0.69, 3.64]
INVEST 2003	201/11309	176/11267	-	34.4	1.14 [0.93, 1.39]
Total (95% CI) Total events: 637 (Bet Test for heterogeneity Test for overall effect	chi-square=1.37 df=2 p	22083 =0.50 ² =0.0%	•	100.0	1.24 [1.11, 1.40]



Authors' conclusions

Implications for practice:

The available evidence does not support the use of beta-blockers as first-line drugs for treating high blood pressure.

Implications for research:

More trials assessing the use of different subclasses of beta-blockers, compared with other antihypertensive drugs, are needed. The possible differential effects of beta-blockers on younger and older people should be assessed.

The Cochrane Database of Systematic Reviews is available from www.wiley.com, and free for eligible countries through www.healthinternetwork.org.