

- clonal expansion of fusidic acid-resistant *Staphylococcus aureus*: a cautionary tale. *Clin Infect Dis*. 2014;59:1451–4.
5. Cooke J. When antibiotics can be avoided in skin inflammation and bacterial colonization: a review of topical treatments. *Curr Opin Infect Dis*. 2014;27:25–9.
  6. Starship Clinical Guidelines. Skin infections: antibiotic choice. Starship Hospital, Auckland, New Zealand: Starship Hospital. [cited 2015 Jun 25]. Available from: <http://www.starship.org.nz>
  7. DermNet NZ. Staphylococcal skin infections. [cited 2015 Jun 25]. Available from: <http://dermnetnz.org/bacterial/staphylococci.html>
  8. Bryant L. Cephalosporins for people with penicillin allergy? *J Prim Health Care*. 2013;5(1):79–80.
  9. Gillies M, Ranakusuma A, Hoffmann T, Thorning S, McGuire T, Glasziou P, et al. Common harms from amoxicillin: a systematic review and meta-analysis of randomized placebo-controlled trials for any indication. *CMAJ*. 2015;187(1):E21–31.
  10. Eberl S, Renner B, Neubert A, Reisig M, Bachmakov I, König J, et al. Role of p-glycoprotein inhibition for drug interactions: evidence from in vitro and pharmacoepidemiological studies. *Clin Pharmacokinet*. 2007;46(12):1039–49.
  11. Baillargeon J, Holmes H, Lin Y, Raji M, Sharma G, Kuo Y. Concurrent use of warfarin and antibiotics and the risk of bleeding in older adults. *Am J Med*. 2012;125(2):183–9.
  12. Stöhlberger C, Finsterer J. Relevance of P-glycoprotein in stroke prevention with dabigatran, rivaroxaban, and apixaban. *Herz*. 2015;40(Suppl 2):140–5.
  13. Hughes J, Crowe A. Inhibition of p-glycoprotein-mediated efflux of digoxin and its metabolites by macrolide antibiotics. *J Pharmacol Sci*. 2010;113(4):315–24.
  14. Gomes T, Mamdani MM, Juurlink DN. Macrolide-induced digoxin toxicity: a population-based study. *Clin Pharmacol Ther*. 2009;86(4):383–6.
  15. van der Velden W, Huussen J, Ter Laak H, de Sévaux R. Colchicine-induced neuromyopathy in a patient with chronic renal failure: the role of clarithromycin. *Neth J Med*. 2008;66(5):204–6.
  16. Slimings C, Riley TV. Antibiotics and hospital-acquired *Clostridium difficile* infection: update of systematic review and meta-analysis. *J Antimicrob Chemother*. 2014;69(4):881–91.
  17. Gillespie D, Hood K, Bayer A, Carter B, Duncan D, Espinasse A, et al. Antibiotic prescribing and associated diarrhoea: a prospective cohort study of care home residents. *Age Ageing*. 2015. pii: afv072.

## Calcium intake and reducing blood pressure

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**THE PROBLEM:** Hypertension is a known public health problem that affects both the economically developed and developing world. It affects somewhere between 25 and 33% of the adult population.<sup>1</sup> Hypertension is one of the leading factors attributing to global mortality, and is the third highest risk factor for the global burden of disease.<sup>2</sup> The National High Blood Pressure Education Program in the US suggests that population strategies that aim to achieve a downward shift of the blood pressure distribution in the general population is an effective method to relieve some of this disease burden.<sup>3</sup> One potential population-based method could be dietary supplementation.

**CLINICAL BOTTOM LINE:** This systematic review shows that an increase in calcium intake will slightly reduce both systolic and diastolic blood pressure.<sup>4</sup> The effect was shown in a dose-response relationship, as well as being confirmed in multiple groups. Although the effect was small, it is based on high-quality evidence and, at the very least, the authors suggest that it should be an objective to make sure there is adequate calcium intake in the population. No adverse events were reported, but this would be an essential factor for any future research to monitor.

*Calcium supplementation: effect on systolic and diastolic blood pressure<sup>4</sup>*

	Success	Evidence	Harms
<b>Systolic blood pressure (SBP)</b>	Calcium significantly lowered SBP with a difference between the placebo group and the calcium supplementation group of -1.43 mm Hg (-2.15 to -0.72)  This effect showed a dose-response treatment effect and was largest in those taking >1500 mg/day and in those studies with patients with a mean age of <35 years of age	This was based on high-quality evidence from 16 individual studies containing 3048 participants in total	There were no reported adverse events
<b>Diastolic blood pressure (DBP)</b>	Calcium supplementation also significantly lowered DBP with a difference between the placebo group and the calcium supplementation group of -0.98 mm Hg (-1.46 to -0.50)  This effect showed a dose-response treatment effect and was largest in those taking >1500 mg/day, in men, and in those studies with patients with a mean age of <35 years of age.	This was based on high-quality evidence from 15 studies containing 2947 participants in total	

### References

1. Kearney PM, Whelton M, Reynolds K, Whelton PK, He J. Worldwide prevalence of hypertension: a systematic review. *J Hypertens*. 2004;22(1):11–9.
2. Ezzati M, Lopez AD, Rddgers A, Vander Hoom S, Murray CJ; Comparative Risk Assessment Collaborating Group. Selected major risk factors and global and regional burden of disease. *Lancet*. 2002;360(9343):1347–60.
3. Whelton PK, He J, Appel LJ, Cutler JA, Havas S, Kotchen TA et al.; National High Blood Pressure Education Program Coordinating Committee. Primary prevention of hypertension: clinical and public health advisory from the National High Blood Pressure Education Program. *JAMA*. 2002;288(15):1882–8.
4. Cormick G, Ciapponi A, Cafferata ML, Belizán JM. Calcium supplementation for prevention of primary hypertension. *Cochrane Database Syst Rev*. 2015;6:CD010037.

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