Media Muddies the Water on Treatment of Hypertension (HTN) Does MILD Hypertension need to be treated?

CBC news article: Blood pressure drug benefits overestimated for mild cases Link: http://www.cbc.ca/news/health/story/2012/10/02/blood-pressure-mild-drugs.html A report on CBC news is likely to raise a lot of questions from patients & health professionals on the treatment of HTN. As is common for media health stories, the facts are told to help make the story. RxFiles thought it would be useful to provide a heads up on issues raised that were valid versus those that were potentially misrepresented. This is not an exhaustive critical appraisal but may be of value in the follow up discussions you are likely to have.

Potentially Valid Potentially Misrepresented Need to confirm indication and The 2 studies in the news report were \Box The role of beta blockers post-MI has high quality evidence in RCT trials reassure patients when treatment not at all similar in populations, /meta-analysis over 2-4 years. Data is has a compelling indication (e.g. heart treatments or trial methodology. lacking regarding how long a benefit failure, atrial fibrillation, MI, etc.) with o Cochrane Review of "Pharmacotherapy for mild HTN"² evidence for outcome benefit. may persist. The 4yr observational was a systematic review & meta-It is reasonable to ask whether there REACH data raises the possibility of analysis that analysed the very is evidence for benefit of drug less benefit over time which should limited RCT evidence for the treatment when treating lower risk not be alarming. [While the RCT data treatment of mild HTN in primary population with mild HTN. shows benefit, it reflects the pre-2000 prevention. This review included Notes that mild HTN may be overera (pre-statin, revascularization, etc.)] drugs from multiple antihypertensive treated in otherwise low risk Observational trials are greatly classes, not just beta blockers. individuals. ["Mild HTN" is defined as a limited by potential confounding. o REACH Registry was a relatively low SBP between 140-159 & a DBP of o The trend in this observational quality observational trial looking at between 90-99. Current CHEP analysis of post-MI patients still beta blockers in patients with guideline BP threshold for initiating drug treatment is $\geq\!160^{~\text{SBP}}~\&\,\geq\!100^{~\text{DBP}}$ in favours beta blocker treatment. coronary artery disease.³ It did not ☐ Besides the limitations of an study treatment of "HTN" per se. those with no risk factors but $\geq 140^{SBP}$ observational trial, the study notes While BP targets for treatment of and 90^{DBP} if other risk factors present.]¹ that use of atenolol was prominent in otherwise uncomplicated HTN are Focuses on important outcomes the subjects. Beta blockers are not <140/90, the threshold at which to ("bigger question") such as MI, stroke generally 1st line for uncomplicated consider pharmacological treatment and death more than surrogate HTN⁵ & relative to other beta for low-risk patients is $\geq 160/100.^{1}$ outcomes such as blood pressure. blockers, there is evidence suggesting The findings in the Cochrane review Notes that drugs which have that atenolol may be less favorable.⁶ included only 8,912 patients with potential benefits may also have ☐ Single BP measurement is insufficient mild HTN from 4 RCTs who met the potential harms. The overall balance for mild HTN. Guidelines suggest 4-5 inclusion & exclusion criteria. Since of benefits & harms must be part of successive readings with proper these patients were by definition, low an individualized treatment decision. technique resting 5 minutes, arm @ heart level, etc. risk & only studied for 4-5 years, the Emphasizes essential & preferred role Pharmacist shown "counting pills" marginal/non-statistically significant of "lifestyle measures" in the which is stereotypical, but somewhat results are not surprising. treatment of mild HTN. unfortunate misrepresentation of [Mortality @4-5yrs, Tx vs Pl: 1.3% vs 1.5%; Cautions not to stop BP meds ↓0.2%; RR=0.85 (95% CI 0.63-1.15)] what many pharmacists do by way of suddenly as this may be harmful "Absence of evidence" (e.g. p>0.05) is drug therapy decision making, drug (especially for beta blockers). not the same as "not effective". monitoring & patient education. BP=blood pressure (units: mmHq) CI=confidence interval CV=cardiovascular DBP=diastolic BP HTN=hypertension PI=placebo RCT=randomized controlled trial RR=relative risk SBP=systolic BP Tx=treatment group

Take home points

- The limited evidence for antihypertensives in mild HTN is not new, and this is somewhat reflected in guidelines.
- Weigh the limited evidence, benefits, harms and patient values, and don't over treat mild HTN in low risk individuals. (Consider the threshold for treating and don't just rely on one BP reading. Do emphasize lifestyle measures in low risk.)
- Remember that treatment of moderate to high blood pressure does have evidence for overall benefit. Benefit is related to risk. In other words, patients with lower CV risk benefit less; those with higher risk, benefit more. As a result, greater caution & attention to the harms and side effects of medications is legitimate in deciding when & how to treat lower risk patients.
- Patients at higher risk of CV events should be reassured that lowering BP is associated with reduced risk of CV events.
- It's ok to reassess beta blocker use, but remember that in early post-MI especially 1st year and heart failure, they reduce mortality. (Do not stop beta blockers suddenly as this can cause serious withdrawal effects. To discontinue, taper over several weeks.)
- When treatment of uncomplicated HTN is indicated, preferred drug choices may include thiazide diuretics, ACEIs (or ARBs) or long-acting CCBs. Beta blockers are usually reserved for patients with compelling indications (heart failure, early post-MI, atrial fib.).
- Evidence is lacking to fully assess benefits/risks in low risk patients & to account for changes in post-MI practices e.g. revascularization.



Comment. There is a lot more that could be discussed. We could discuss the specifics of the trial/review data. We could discuss the concept of disease risk assessment. We could discuss the concept of how decisions based on the benefits and risks of treatment are more often clear cut for patients at higher risk, and how for those at lower risk, there is a lot more uncertainty that enters into the equation. We could discuss the changes in management of patients following an MI that might lower their risk to the point that the patients absolute risk is reduced, and the corresponding benefit of beta-blocker therapy would be reduced. We could discuss the potential role for thinking about subgroups within the broader group lumped in as "low risk". But this discussion has focused instead on a few "quick takes" after watching the news, knowing that we will be answering related questions on somewhat confusing information. We are sure there will be opportunity to revisit specific aspects of the information in the days to come. Now, off to some academic detailing...

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Patient decision aid (sample) - Hypertension: http://www.npc.nhs.uk/therapeutics/cardio/cd hyper/resources/pda hypertension.pdf
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References

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³ Bangalore S, Steg PHG, Deedwania P, et al. Beta blocker use and clinical outcomes in stable outpatients with and without coronary artery disease. JAMA 2012; 308:1340-1349.

⁴ Freemantle N, Cleland J, Young P, et al. Beta Blockade after myocardial infarction: <u>systematic review</u> and meta regression analysis. BMJ. 1999 Jun 26;318(7200):1730-7.

⁵ Wiysonge C, et al. Beta-blockers for hypertension. Cochrane Database Syst Rev. 2007 Jan 24;(1):CD002003.

⁶ Carlberg B, Samuelsson O, Lindholm LH. Atenolol in hypertension: is it a wise choice? Lancet. 2004 Nov 6;364(9446):1684-9.