

# RESEARCH

## Effect of intensive structured care on individual blood pressure targets in primary care: multicentre randomised controlled trial

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### Abstract

**Objective** To determine the effectiveness of intensive structured care to optimise blood pressure control based on individual absolute risk targets in primary care.

**Design** Pragmatic multicentre randomised controlled trial.

**Setting** General practices throughout Australia, except Northern Territory, 2009-11.

**Participants** Of 2185 patients from 119 general practices who were eligible for drug treatment for hypertension according to national guidelines 416 (19.0%) achieved their individual blood pressure target during a 28 day run-in period of monotherapy. After exclusions, 1562 participants not at target blood pressure (systolic 150 (SD 17) mm Hg, diastolic 88 (SD 11) mm Hg) were randomised (1:2 ratio) to usual care (n=524) or the intervention (n=1038).

**Intervention** Computer assisted clinical profiling and risk target setting (all participants) with intensified follow-up and stepwise drug titration

(initial angiotensin receptor blocker monotherapy or two forms of combination therapy using angiotensin receptor blockers) for those randomised to the intervention. The control group received usual care.

**Main outcome measures** The primary outcome was individual blood pressure target achieved at 26 weeks. Secondary outcomes were change in mean sitting systolic and diastolic blood pressure, absolute risk for cardiovascular disease within five years based on the Framingham risk score, and proportion and rate of adverse events.

**Results** On an intention to treat basis, there was an 8.8% absolute difference in individual blood pressure target achieved at 26 weeks in favour of the intervention group compared with usual care group (358/988 (36.2%) v 138/504 (27.4%)): adjusted relative risk 1.28 (95% confidence interval 1.10 to 1.49, P=0.0013). There was also a 9.5% absolute difference in favour of the intervention group for achieving the classic blood pressure target of  $\leq 140/90$  mm Hg (627/988 (63.5%) v 272/504 (54.0%)): adjusted relative risk 1.18 (1.07 to 1.29, P<0.001). The intervention group achieved a mean adjusted reduction in systolic blood

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Appendix I  
Appendix II  
Appendix III

pressure of 13.2 mm Hg (95% confidence interval –12.3 to –14.2 mm Hg) and diastolic blood pressure of 7.7 mm Hg (–7.1 to –8.3 mm Hg) v 10.1 mm Hg (–8.8 to 11.3 mm Hg) and 5.5 mm Hg (–4.7 to –6.2 mm Hg) in the usual care group ( $P<0.001$ ). Among 1141 participants in whom five year absolute cardiovascular risk scores were calculated from baseline to the 26 week follow-up, the reduction in risk scores was greater in the intervention group than usual care group (14.7% (SD 9.3%) to 10.9% (SD 8.0%); difference –3.7% (SD 4.5%) and 15.0% (SD 10.1%) to 12.4% (SD 9.4%); –2.6% (SD 4.5%): adjusted mean difference –1.13% (95% confidence interval –0.69% to –1.63%;  $P<0.001$ ). Owing to adverse events 82 (7.9%) participants in the intervention group and 10 (1.9%) in the usual care group had their drug treatment modified.

**Conclusions** In a primary care setting intensive structured care resulted in higher levels of blood pressure control, with clinically lower blood pressure and absolute risk of future cardiovascular events overall and with more people achieving their target blood pressure. An important gap in treatment remains though and applying intensive management and achieving currently advocated risk based blood pressure targets is challenging.

## Introduction

Although hypertension is a readily detectable and modifiable condition it is responsible for more deaths worldwide than any other cardiovascular risk factor, including tobacco use, obesity, and lipid disorders.<sup>1</sup> In the United States, of all modifiable risk factors hypertension has been associated with the greatest population attributable risk for all cause mortality (30%) and deaths from cardiovascular disease (40%).<sup>2</sup> Epidemiological studies suggest a relation between raised blood pressure and increased risk of cardiovascular events.<sup>3</sup> Large scale clinical trials examining the efficacy of a broad range of antihypertensives found a consistent and continuous reduction in cardiovascular risk (primary or secondary events and across all age groups) according to the baseline values and magnitude of change in both systolic and diastolic blood pressure.<sup>3</sup> Traditionally, the ideal blood pressure target has been  $\leq 140/90$  mm Hg for those aged less than 80 years and these are reflected in current guidelines from the National Institute for Health and Clinical Excellence in the United Kingdom.<sup>4</sup> The introduction of absolute risk profiling for primary prevention purposes<sup>4 5</sup> and more stringent, individualised blood pressure targets for higher risk groups such as people with type 2 diabetes or more advanced forms of cardiovascular disease<sup>6 7</sup> has provided clinicians in other regions of the world with a challenge in meeting these targets. Considering the recent debate on whether more stringent blood pressure targets fail to provide cardioprotection and perhaps even confer harm at blood pressure levels approaching 110/75 mm Hg<sup>8</sup> it is likely that more stringent blood pressure targets will be subject to further consideration. Regardless of recommended blood pressure targets, however, the translation of potential benefits from clinical trials to the real life management of hypertension (much of which occurs in primary care) is less than ideal. Many people prescribed antihypertensive treatment have a blood pressure  $>140/90$  mm Hg.<sup>9</sup> A key component of effective translation in the primary care setting seems to be the application of structured programmes to apply antihypertensive treatments by a rigorous stepped care approach<sup>10</sup> to overcome the common phenomenon of prescription resistance<sup>11</sup> (whereby clinicians are reluctant to up-titrate existing drug treatment or prescribe additional antihypertensive agents) and to achieve better individual blood pressure control. For example, nurse led interventions that comprise a structured care algorithm are associated with greater reductions in blood pressure levels than usual care overall.<sup>12</sup> Few studies, however, have specifically tested the application

of structured care in the setting of more stringent, risk based blood pressure targets in a large and diverse range of general practices—that is, small to large clinics and those with and without practice nurses to implement structured care.

As described elsewhere,<sup>13</sup> the Valsartan Intensified Primary care Reduction of Blood Pressure (VIPER-BP) study was a pragmatic, multicentre, randomised trial evaluating the clinical effectiveness and overall safety of a more intensive and structured approach to optimising blood pressure control in a group of people with persistently high blood pressure in a real world primary care setting.

In a typical cohort of primary care patients with suboptimal blood pressure (individualised target according to their risk profile<sup>6</sup>), we hypothesised that a greater proportion of those randomised to the intervention would achieve individualised blood pressure control at the end of 26 weeks compared with those randomised to usual care.

## Methods

The Australian universal health insurance scheme (Medicare) provides citizens with free access to public hospitals and reimbursed access (70% of services with no co-payment) to a widespread network of primary care or general practice clinics. A parallel system, the Pharmaceutical Benefits Scheme, provides subsidised drug treatment to all Australians, with capped payments for those with a chronic illness; most prescriptions for drugs being generated by general practitioners managing people with chronic conditions such as hypertension.

## Participants

Study participants were recruited by 260 general practitioners from 119 general practices distributed throughout every state and territory of Australia except the sparsely populated Northern Territory. People routinely managed by these doctors were potentially eligible to participate if they were aged 18 or more, had a diagnosis of hypertension requiring active drug treatment according to guidelines, and consented to participate. We excluded people who had a mean initial systolic blood pressure of 180 mm Hg or more while sitting, were prescribed three or more antihypertensives; had moderate to severe renal disease (clinical diagnosis or estimated glomerular filtration rate  $<60$  mL/min/1.73m<sup>2</sup>), or had contraindications to angiotensin receptor blockers, calcium channel blockers, or thiazide diuretics. Recruitment took place from July 2009 to December 2010, with follow-up completed in July 2011.

## Study design

This was an open label, two arm, randomised trial comparing an intensive blood pressure management strategy with usual care (control).<sup>13</sup> Randomisation was by computer generated, group assignment stratified according to nominated blood pressure target (three strata) and block randomisation (units of 12) per general practitioner. An independent data management team at Baker IDI coordinated a standardised protocol for randomising participants over the telephone. Participants were assigned to the groups using a prespecified ratio of 1:2 for usual care versus intervention (with a further ratio of 1:2 within the intervention arm to assign participants to a monotherapy strategy and to two possible combined treatment strategies). Quality control of the sites and general practitioners was achieved by visits before randomisation and at study end. The study was independently monitored by routine visits, and case report forms were verified against clinical records for 20% of patients. We

used standardised operating procedures for data queries and management. All queries were resolved by the time the study dataset was locked and no further changes were permitted. Figure 1 provides a graphical representation of the study timelines.

## Study entry

Eligible participants underwent baseline assessment, including determination of absolute cardiovascular risk.<sup>5</sup> On the basis of this assessment, we identified target risk factor measures (including blood pressure, lipid levels, exercise levels, smoking status, and body mass index) for each participant according to national guidelines.<sup>6</sup> A computer program developed by Baker IDI enabled clinical profiling in all participants and subsequent management in intervention patients (see appendix 1 in the supplementary file). Eligible participants began a standard 28 day run-in period of treatment with open label, oral valsartan 80 mg daily, with a mandatory blood pressure check 14 days after the start of treatment to determine the need for immediate (rescue) randomisation if systolic blood pressure was  $\geq 180$  mm Hg or clinically indicated. We then randomised into the study those who did not achieve their prespecified blood pressure target within this run-in period or were not specifically withdrawn or lost to follow-up (blinded allocation).

## Blood pressure measurement

Throughout the study we used a standardised protocol based on national guidelines to record blood pressure measurements.<sup>6</sup> After participants had rested for at least five minutes and while they were seated we obtained three blood pressure measurements separated by one minute intervals using an appropriate sized cuff and a calibrated, semiautomated oscillometric device approved by the Therapeutic Goods Administration of Australia. We defined aberrant measurements as those where the lowest reading was 10 mm Hg or more for systolic blood pressure or 5 mm Hg or more for diastolic blood pressure below the highest of the three seated readings. If this occurred, the participant was rested for three minutes and the procedure repeated. Given the size and nature of the study, it was impractical to obtain independent, blinded blood pressure levels. Nominated study staff at each site (general practitioner or practice nurse) verified the blood pressure measurements by an independent audit of the clinical details in the case records, outputs from the blood pressure monitor, and the computer decision tool.

## Usual care

For those assigned to usual care, we asked general practitioners to follow their usual pattern of clinic visits and treatment strategies (no restriction on prescribed drugs) to achieve the individualised blood pressure target and other prevention targets identified during the run-in phase; the only mandatory study visits were at weeks 6 (a typically brief 10-15 minute consultation) and 26 (final comprehensive evaluation). We explicitly acknowledged that this was an enhanced form of usual care given that the participants had already been subject to clinical profiling and had an identified blood pressure target and their doctors potentially had knowledge of the intervention (with the inherent problem of contamination).

## Study intervention

Those allocated to the intervention arm followed an intensive stepped programme of management, with mandatory visits to their doctor at weeks 6, 10, 14, and 18 after randomisation to review their blood pressure and to adjust their treatment if

needed according to prespecified algorithms (see appendix II in the supplementary file). Typically, these consultations lasted 10-15 minutes (about 60 minutes in total) in addition to the time taken for clinical profiling at baseline. Reflecting contemporary treatment options for the management of hypertension, different drug pathways were mandated (initial monotherapy with valsartan 160 mg or combined with hydrochlorothiazide or amlodipine, as single combined pills). Scheduling of visits and up-titration of drug treatment was guided by the same computer program developed by Baker IDI to facilitate the initial risk profiling and management. A final clinic visit for clinical re-evaluation was scheduled at 26 weeks.

## Study endpoints

The primary endpoint was the percentage of participants who achieved their individualised blood pressure target according to national guidelines<sup>6</sup> during 26 weeks of follow-up (using last recorded blood pressure for intention to treat analyses). These stringent blood pressure targets, which have been rarely examined for feasibility in a controlled trial, are  $\leq 125/75$  mm Hg if a patient has proteinuria,  $\leq 130/80$  mm Hg if end organ damage is present (including any form of cardiovascular disease, diabetes, or microalbuminuria), and the classic target of  $\leq 140/90$  mm Hg for those without evidence of the other two criteria.

Key secondary endpoints included change in mean sitting systolic and diastolic blood pressure and absolute risk for cardiovascular disease within five years based on the Framingham risk score<sup>5</sup> (where applicable). Secondary endpoints also included the type and rate of adverse events potentially attributable to antihypertensive treatment, and serious adverse events including all cause mortality and fatal and non-fatal cardiovascular events—for example, acute myocardial infarction, stroke, and heart failure. Several other secondary endpoints will be the subject of future or expanded reports, including changes in quality of life, depression, self care behaviours, and evidence of end organ damage (including electrocardiographic evidence of left ventricular hypertrophy and newly detected proteinuria).

## Study power

We largely met the study targets for overall recruitment and number of participants entering the run-in phase. Based on the combination of a strong potential for interventional contamination, balanced against the need to establish clinically significant differences between the two groups, the study was initially powered to detect a minimum absolute difference of 7% between the groups for the primary endpoint.<sup>13</sup> As proportionately more patients achieved blood pressure control during the study run-in and thus the number of randomised participants was reduced, we recalculated that a total randomised cohort of more than 1500 participants would still provide sufficient study power to detect a minimum 8% difference between groups with more than 85% study power (two sided  $\alpha$  level of 0.05 and adjusting for 2:1 randomisation for study intervention versus usual care).<sup>13</sup>

## Statistical analyses

The study statistician (on behalf of the clinical safety and efficacy committee) independently analysed the data according to a prospectively designed statistical analysis plan. If blood pressure values were missing at 26 weeks, we carried forward the last recorded blood pressure measurement after randomisation for primary endpoint analysis. Baseline and outcome data were analysed using SPSS for Windows version 19.0. Continuous data are presented as means (standard



deviations) or medians (interquartile ranges). Categorical data are presented as percentages.

We carried out efficacy analyses on an intention to treat population, consisting of all participants randomised to the intervention arm or to the usual care arm and who had at least one recorded blood pressure measurement after randomisation. Analyses were based on the treatment group to which the participant was randomised. We compared the primary endpoint measure (blood pressure control at week 26) between the study groups using a log binomial generalised linear model with stratification status at randomisation as a covariate. Stratification status was fit as a categorical variable with the three blood pressure target groups ( $\leq 125/75$  mm Hg,  $\leq 130/80$  mm Hg, and  $\leq 140/90$  mm Hg) based on the participant's clinical profile. Change in systolic and diastolic blood pressure from baseline to 26 weeks were each analysed by analysis of covariance, with treatment group and stratification status as factors and mean baseline blood pressure as a covariate. We also describe the proportion and rate of adverse events (per participant).

## Results

Overall, 2337 participants (1381 men (59.1%), mean age 58 (SD 12) years) with hypertension from 119 general practices were enrolled into the study; 2185 participants from 114 practices subsequently started the run-in phase of initial treatment (valsartan 80 mg/day for 14–28 days, fig 2). (The figure in appendix II in the supplementary file shows a simplified flow chart of study activity and interventions according to group randomisation, including the initial phase of clinical profiling of all participants who entered the run-in phase of the study, and the two study arms after randomisation.) Of those who entered the run-in phase, 1329 (60.8%) had pre-existing hypertension and had been treated for a median of 5 (interquartile range 2–10) years. Previous treatment, which was stopped on entry to the study, comprised monotherapy with either angiotensin converting enzyme inhibitors or angiotensin receptor blockers ( $n=907$ , 68.2%), a calcium channel antagonist ( $n=233$ , 17.5%), or combination antihypertensive treatment ( $n=259$ , 19.5%). The initial blood pressure target for these 2185 participants was  $\leq 125/75$  mm Hg in 330 (15.1%),  $\leq 130/80$  mm Hg in 993 (45.4%), and  $\leq 140/90$  mm Hg in 862 (39.5%). During the run-in phase, 416 (19.0%) participants reached their initial prespecified blood pressure target; comprising 20 (6.7%) of those with a preliminary blood pressure target of  $\leq 125/75$  mm Hg, 128 (12.9%) with  $\leq 130/80$  mm Hg, and 268 (31.1%) with  $\leq 140/90$  mm Hg. A further 207 (9.5%) participants withdrew from the study (56 (2.6%) experienced an adverse event) and were not randomised (fig 2).

Table 1 shows the baseline characteristics of the 1562 participants randomised to the usual care group ( $n=524$ , 33.5%) or intervention group ( $n=1038$ , 66.5%); 360 (23.0%) participants in the intervention group were randomised to the monotherapy arm and 678 (43.4%) to the combination therapy arm. The proportion of patients treated for hypertension before the study run-in phase (40% with angiotensin converting enzyme inhibitors, 20% with an angiotensin receptor blocker, and 21% with combination therapy including a calcium antagonist or diuretic) had increased to 66.9%, with little difference between the groups. The groups were well matched for clinical and personal characteristics and prespecified blood pressure targets. At randomisation, 219 (14.0% with equal proportions in each group) were assigned to a blood pressure target that differed from the recommended (automated) clinical profile; these reflected general practitioners' discretion for a mostly higher

blood pressure target than recommended. An endpoint blood pressure was recorded in 504/524 participants in the usual care group and 988/1038 in the intervention group, at any time point after randomisation.

## Patterns of blood pressure management

Drug treatment after randomisation was recorded in 501 participants (95.6%) in the usual care arm and 991 (95.5%) in the intervention arm. In the intervention group, attendance (within 14 days) at scheduled visits varied from 79.9–91.0%; the highest attendance being at six weeks (compared with 91.4% in the usual care group, table 2). Overall, participants in the usual care group had 854 clinic reviews (mean 1.7 visits) after randomisation compared with 3477 in the intervention group (mean 3.5 visits). Figures in appendix II in the supplementary file show the (complex) pattern of prescribing in the usual care group and the composite treatment groups comprising the intervention group. At randomisation, most participants in the intervention arm were prescribed their per protocol treatment: 334/360 (92.8%) monotherapy (valsartan 160 mg daily), 228/250 (91.2%) hydrochlorothiazide combination therapy (valsartan 80 mg + hydrochlorothiazide 12.5 mg daily), and 398/417 (95.4%) amlodipine combination (valsartan 80 mg + amlodipine 5 mg daily); 11 participants in the combination arm (all early study withdrawals) did not have their combination treatment selected at randomisation. In the next six weeks, 108 (32%), 118 (52%), and 207 (52%) of these participants, respectively, had progressed to the next level treatment as per study protocol. At baseline in the usual care group, 367 (70.0%) participants were still receiving valsartan 80 mg/day and 67 (12.8%) were receiving the increased dose of 160 mg/day. By study completion (based on last prescribed treatment) a greater proportion of participants in the usual care group ( $n=227$ ) than intervention group ( $n=154$ ) were prescribed monotherapy (43.3 v 14.8%) and fewer were prescribed combination (217 (41.4%) v 609 (58.7%)) or triple (21 (4.1%) v 155 (14.9%)) antihypertensive therapy. Alternatively, prescription resistance with deviation from the management protocol was evident in 360 participants (34.7%): most deviations (>90%) involved general practitioners not up-titrating antihypertensive treatment to achieve one of the two lower blood pressure targets.

## Primary endpoint: individualised blood pressure control

On an intention to treat basis, there was an 8.8% absolute difference in the primary endpoint in favour of the intervention group, with 358/988 (36.2%) participants in the intervention arm achieving their individual blood pressure target compared with 138/504 (27.4%) in the usual care arm: adjusted relative risk 1.28 (95% confidence interval 1.10 to 1.49,  $P=0.0013$ ). Table 2 shows absolute blood pressure values (with change in blood pressure from baseline) and overall pattern of blood pressure control for both groups during progressive visits. Overall, 627/988 (63.5%) participants in the intervention group compared with 272/504 (54.0%) in the usual care group achieved the classic blood pressure target of  $\leq 140/90$  mm Hg. This represents an absolute difference of 9.5% in favour of the intervention group: relative risk 1.18 (95% confidence interval 1.07 to 1.29,  $P<0.001$ ).

## Secondary endpoint: change in blood pressure

Clinically important falls in mean and systolic blood pressure were observed in both groups during the 26 weeks of follow-up,

with a lower diastolic and systolic blood pressure value recorded after randomisation in 311 (59.4%) participants in the usual care group compared with 719 (69.3%) in the intervention group. Individualised control of blood pressure showed an inverse gradient; lower rates of achieving the blood pressure target were associated with more stringent targets ( $P<0.001$ ) but better rates of control throughout in the intervention group. During study follow-up, mean blood pressure changed (randomisation to endpoint blood pressure) from 150 (SD 17)/88 (SD 11) mm Hg to 136 (SD 15)/81 (SD 10) mm Hg in the intervention group compared with 149 (SD 17)/87 (SD 11) mm Hg to 139 (SD 15)/82 (SD 11) mm Hg in the usual care group (table 2); adjusted mean reduction in blood pressure 13.2 (95% confidence interval 12.3 to 14.2) mm Hg and 7.7 (7.1 to 8.3) mm Hg compared with 10.1 (8.8 to 11.3) mm Hg and 5.5 (4.7 to 6.2) mm Hg in favour of intervention ( $P<0.001$ ). In those who completed the 26 weeks of follow-up (186 (SD 20) days) the equivalent reduction in blood pressure was also in favour of the intervention ( $P<0.001$ ): adjusted mean reduction in blood pressure 14.6 (13.7 to 15.6) mm Hg and 8.2 (7.6 to 8.8) mm Hg compared with 10.4 (9.1 to 11.7) mm Hg and 5.6 (4.7 to 6.4) mm Hg. The intervention was associated with greater levels of individual blood pressure control in all three target groups (fig 3). Overall, those assigned the combination treatment arm of the intervention group had the highest rate of individual blood pressure control (37.9% for the endpoint and 42.1% at week 26). Similarly, an analysis of change in blood pressure from baseline to 26 weeks ( $n=1323$ ) according to target group showed consistent benefits of the intervention (8-12% reduction in systolic blood pressure and 7-14% reduction in diastolic blood pressure) compared with usual care (6-8% and 5-9%, respectively), with the most gains in blood pressure reduction achieved through the combination treatment arm overall (see appendix II in the supplementary file).

## Change in absolute cardiovascular risk

Among the 1141 participants in whom absolute cardiovascular risk profiles were calculated from baseline to 26 weeks' follow-up (that is, complete data on profiling both before and after randomisation), the intervention group had a significantly greater reduction in risk scores (14.7% (SD 9.3%) to 10.9% (SD 8.0%);  $-3.7\%$  (SD 4.5%)) than the usual care group (15.0% (SD 10.1%) to 12.4% (SD 9.4%);  $-2.6\%$  (SD 4.5%)); the adjusted mean reduction in risk being in favour of the intervention group ( $-1.13\%$ , 95% confidence interval  $-0.69\%$  to  $-1.63\%$ ;  $P<0.001$ ). Overall, the proportion of high risk participants ( $>15\%$  absolute risk of an event) in the intervention group decreased from 41.2% to 25.6% and in the usual care group from 40.2% to 30.1%.

## Safety profile

A total of 1044 adverse events were recorded in 592 (27.1%) participants during the study run-in phase, with 424 (40.6%) classified as related to treatment by the treating doctor. Overall, 2184 adverse events were recorded after randomisation, with 2152 occurring among the 1492 participants who had at least six weeks' follow-up (see appendix III in the supplementary file for a summary of the adverse events). This included 196 (57.8%) participants allocated to the monotherapy arm and 355 (54.7%) to the combination therapy arm; this compared with 246 (48.8%) participants in the usual care group. Of the adverse events after randomisation, 543 (24.9%) were classified as related to treatment, with 422 (0.41/participant) occurring in the intervention arm and 121 (0.23/participant) in the usual care arm (table 3). The three most common adverse events after

randomisation attributed to study treatment were dizziness (including postural hypotension, 86 events), peripheral oedema (79 events), and fatigue or lethargy (46 events). The rate of dizziness (0.07 v 0.03 adverse events/participant) and oedema (predominantly peripheral oedema, 0.06 v 0.03 adverse events/participants) was highest in the intervention group overall, and most common in the combination arm. One (0.06%) participant randomised to the intervention group died due to adenocarcinoma. Post-randomisation, 24 (4.6%) participants in the usual care group experienced 35 serious adverse events compared with 37 (3.7%) participants and 57 serious adverse events in the intervention group (6.7 v 5.6 events/100 patients). In the usual care and intervention groups, respectively, this included six and eight events involving admission to hospital for chest pain/angina pectoris, one (in each group) for acute myocardial infarction, and four (in each group) for other cardiovascular events. Overall, due to an adverse event drug treatment was altered in 82 (7.9%) intervention participants compared with 10 (1.9%) usual care participants (1.1% v 0.2%, respectively, were withdrawn by the general practitioner or voluntarily withdrew).

## Discussion

A strategy comprising automated risk profiling plus standardised guideline based, stepwise drug treatment (initial monotherapy using an angiotensin receptor blocker or two forms of combination therapy) and intensified follow-up and treatment titration, resulted in more participants (absolute difference 8.8%) with persistent hypertension achieving their individual blood pressure target at 26 weeks than participants receiving usual care. Importantly, two thirds of the participants had a history of persistent hypertension despite receiving a typical range of antihypertensive treatments, and the remainder were unresponsive to valsartan monotherapy; one of the most commonly prescribed antihypertensive agents globally. The study population had an average five year risk of a major cardiovascular event of over 15% and therefore represented the more difficult end of the spectrum of people with hypertension managed in primary care. The detailed clinical profiling at study entry and general practitioners' access to the computer assisted treatment protocol may explain the large, clinically significant, reductions in blood pressure in both groups (mean change  $>13/7$  mm Hg in the intervention group and  $>10/5$  mm Hg in the control group). The combined application of more intensive follow-up and drug treatment is reflected in greater use of combination therapies (prescribed as single pills) and progressive reductions in blood pressure between six and 26 weeks in the intervention group. These incremental benefits in blood pressure control largely influenced more favourable absolute cardiovascular risk scores in participants assigned to intervention, with just over one third in this group achieving their individual blood pressure target and around two thirds the classic blood pressure target of  $\leq 140/90$  mm Hg; the same advocated by NICE in the United Kingdom.<sup>4</sup> If further adjustment is made for those who initially responded to more intensive management and monotherapy using an angiotensin receptor blocker at study run-in (apportioning equal success and failure to both treatment groups), this figure increases from 64% to 72%. An important trade-off for what could be considered to be modest absolute differences in absolute blood pressure levels between groups was the small increase in the proportion of participants who withdrew from the study (1.1% v 0.2%) and more frequent (7.9% v 1.9%) modification of treatment due to adverse events in the intervention group.

## Strengths and limitations of this study

The intervention directly addressed the difficult challenge of optimising blood pressure control in a real world general practice setting using commonly prescribed forms of antihypertensive agents. This is a major concern for countries with aging populations, where hypertension related heart and cerebrovascular disease is a growing problem.<sup>14</sup> By necessity, this pragmatic study used an open label design (with no blinding of treatment pathways or blood pressure measurements possible) with scheduled visits that varied according to clinic schedules and prescribed drug treatment and relied on the mainstream healthcare system in Australia. The broader application and possible impact of the intervention (particularly in other healthcare systems) therefore needs to be interpreted cautiously. However, we do have preliminary projections (data not shown) based on national primary care data in Australia, that even the modest absolute differences in blood pressure between groups would have a potentially large impact (as reflected in the primary endpoint and absolute risk differences) among the thousands of Australians who don't achieve their blood pressure targets each year. The selection of valsartan based treatments, rather than an open choice of agents, was predicated on the ability to standardise drug treatment in the intervention arm with precise up-titration pathways and to determine the potential impact of monotherapy versus combination therapy on short (six weeks) and longer term (26 weeks) blood pressure control. Certainly these agents targeting the renin-angiotensin system represent some of the most commonly applied in clinical practice internationally, and the treatment regimen is broadly consistent with the NICE guidelines (phase 3) for people with hypertension who do not respond to initial monotherapy.<sup>4</sup> We deliberately chose individual randomisation rather than a cluster design to increase the robustness of our findings for two reasons. Firstly, we wanted to avoid potential imbalances between the groups when using a larger unit of randomisation (that is, a general practice). Secondly, we wanted to expose participants with the same general practitioner to the two study groups (and indeed treatment arms within the intervention group) to truly determine if the programme would alter blood pressure management at the individual level. Although this resulted in almost inevitable cross contamination of more intensive management in the usual care group (with clinically significant reductions in blood pressure in both groups), we were still able to show that the intervention was associated with incremental benefits in blood pressure control (8.8% absolute difference in the primary endpoint) tempered by more adverse events requiring adjustment of treatment. Alternatively, unblinded blood pressures in the clinic reported by general practitioners (albeit using a standardised protocol and automated monitors) may have resulted in an element of bias, the recruitment of participants with so called white coat hypertension,<sup>15</sup> or an observed regression to the mean (blood pressure) over time. Wherever possible we minimised these factors by implementing a 28 day standardised treatment and blood pressure monitoring run-in phase, applying a standardised protocol for blood pressure monitoring and undertaking independent clinical auditing for data verification. Whether these data would be reproduced if we had studied 24 hour ambulatory blood pressure or home blood pressure measurements is open to debate.<sup>16</sup> However, at this time current guidelines are based on clinic blood pressure measurements and the study reflects current clinical practice in most countries. We do not know as of yet if the reduction in absolute cardiovascular risk will be maintained or result in fewer cardiovascular events in the future. Notably, 4% of randomised participants did not have an endpoint blood pressure recorded

and were excluded from endpoint analyses. For these 70 participants, retrospective analyses where either the randomised blood pressure was brought forward (relative risk 1.27, 95% confidence interval 1.09 to 1.48) or the change in blood pressure was imputed according to study group and blood pressure target (1.26, 1.08 to 1.46) did not substantially alter the primary endpoint results ( $P < 0.001$  for both comparisons). The general practitioners were reluctant to try to meet the more aggressive guideline targets for blood pressure reduction and in almost 15% of cases opted for a more lenient target, in effect making the task of achieving control easier.

## Supporting the evidence in favour of structured primary care

Consistent with data from other countries such as the United States,<sup>2</sup> we have strong evidence from community based studies,<sup>9</sup> preliminary data from the pilot stages of the VIPER-BP study,<sup>13</sup> and data from the usual care group, that present approaches to the management of hypertension have meant that the "rule of halves" still applies—that is, close to 50% of those treated for hypertension in primary care remain hypertensive. Regardless of recent suggestions that lower blood pressure targets in high risk people should be reassessed,<sup>8</sup> there is still clear evidence from population cohorts<sup>2</sup> and meta-analyses of clinical trial data<sup>3</sup> that lowering blood pressure (even by the modest, incremental margins associated with more intensive management) will noticeably attenuate the rates of stroke and other cardiovascular disease. In support of a more aggressive approach to the management of hypertension, retrospective results from both the Valsartan Antihypertensive Long-term Use Evaluation trial<sup>17 18</sup> and Avoiding Cardiovascular Events in Combination Therapy in Patients Living with Hypertension<sup>19</sup> trial suggest there are advantages to achieving early blood pressure control and that application of combination treatments are more likely to achieve this goal. The intervention in the current study brought all these elements together, with more than double (81% v 40%) the proportion of intervention participants receiving combination therapy compared with monotherapy at 26 weeks. To our knowledge this study (incorporating a combined drug and disease management approach to the management of blood pressure facilitated by a standardised computer support program) is one of the largest studies of its kind. Although there have been several large hypertension trials with a major focus on the efficacy of different combinations of drug treatment<sup>20</sup> and specific studies of algorithm based strategies,<sup>10 12</sup> there are few large scale, multicentre studies that have specifically compared a structured and intensive care approach with usual care in achieving better blood pressure control based on more stringent risk based targets in patients with persistent hypertension managed in primary care. The results of the present study therefore strengthen the conclusions of a recently updated Cochrane review recommending a stepped and intensive approach to the management of hypertension in primary care; particularly in those participants who remain hypertensive despite initial treatment.<sup>10</sup> In particular, the results complement the findings of a similarly large scale (cluster randomised design) Canadian study<sup>21</sup> of a simplified treatment algorithm in primary care that showed a 12% absolute difference (65% v 53%) in favour of the study intervention. Notable differences were the low proportion of participants requiring more stringent blood pressure targets (about 15%) and the lack of initial standardised care that in the present study resulted in almost one in five participants achieving early blood pressure control during the study run-in phase. Thus our randomised cohort represented a more "challenging" group of patients with



hypertension. It could be argued that we have now reached a similar stage of developing the evidence in favour of more structured care (that is, beyond a purely drug related approach) to optimise blood pressure management in primary care to that of heart failure management programmes; where head to head trials of different forms of structured care are being undertaken<sup>22</sup> to refine (rather than to establish) a new level of best clinical practice.

## Clinical implications

Several important issues highlighted by this pragmatic study involving more than 250 general practitioners are relevant to clinical practice guidelines and health policy in Australia and other high income countries where hypertension remains a problem. Firstly, the initial application of standardised monotherapy (valsartan 80 mg/day) in a structured approach, irrespective of previous treatment, was immediately successful in controlling a large group of people with hypertension and should be considered when reviewing primary endpoint data. Given its size and direct relevance to other high income countries, the study is likely to influence future blood pressure guidelines, particularly informing debate around the attainment of lower blood pressure targets, whether these lower targets are truly achievable, and the role of prescription resistance<sup>11</sup> in limiting what is meant by “truly achievable.” Only a small proportion of participants in the most stringent blood pressure group achieved this target despite the intensive approach with three or more drugs applied in the intervention. Given the nature of the study, these data are likely to reflect real world blood pressure management, and any consideration of retaining more stringent blood pressure targets (particularly if low values are proved to be unnecessary in those with pre-existing cardiovascular disease<sup>8</sup>) has to consider the impact of at least aiming for lower blood pressure to achieve values  $\leq 140/90$  mm Hg. Both an overall reluctance by general practitioners to prescribe higher doses of combination or triple treatment and individual patient factors seem to have prevented more participants from reaching their individual blood pressure target and the classic target of  $\leq 140/90$  mm Hg. It is worth highlighting that in the Australian healthcare system, subsidised drug treatment ensures that costs are not usually a barrier to the application of more aggressive treatment.

Although the absolute difference in change in blood pressure between the groups might be considered modest, they not only contributed to more people achieving the primary endpoint in the intervention group but also favourably altered both the absolute risk of future cardiovascular events (a clinically meaningful 1% difference in mean scores between groups) and the relative risk reduction of future coronary artery disease (–6% difference in mean risk) and strokes (–11% difference) in favour of the intervention (data not shown). The expected trade-off between more intense blood pressure control and an increase in adverse events occurred, with more participants in the intervention group withdrawn on this basis (1.1% v 0.2%) or requiring active change to their treatment (7.9% v 1.9%). This needs to be carefully considered against the overall difference (8.8% in favour of the intervention group) in the primary endpoint. Overall, the number and frequency of adverse events was consistent with the nature and size of the patient cohort and the drugs being prescribed. Whereas there seems to be a gradient in blood pressure control according to the amount of drug treatment (with greater control achieved in those assigned to immediate combination therapy at randomisation), spending additional clinical time (>1 hour over six months) with participants probably also contributed. Such a conclusion is

consistent with previous systematic reviews and meta-analyses,<sup>10</sup> and the patient centred approach advocated by contemporary guidelines.<sup>4 6 7</sup> This observation is further strengthened by the initial response to the run-in phase of the study where information on non-drug measures was provided with only low dose monotherapy (19% blood pressure control) and substantial anecdotal evidence from participants who appreciated the “beyond pharmacological” approach to their management.

## Future challenges

Despite noticeable reductions in blood pressure in both arms of the VIPER-BP study, a relatively small proportion of participants achieved these more stringent blood pressure targets; the classic target of  $\leq 140/90$  mm Hg being far more attainable. Given the intensity of the intervention and facilitated pathways to higher doses of combination antihypertensive treatment, these data highlight critical questions around the application of more stringent blood pressure targets in high risk people—that is, those with established cardiovascular disease or renal disease. The results are particularly relevant to clinicians when clinical guidelines and health policies seek to apply expert recommendations that may not be truly achievable. As such, these data may prompt a re-evaluation of blood pressure targets to reflect real world limitations for doctors managing hypertension in the busy and often complex primary care setting. The results also point to major patient related barriers to eviscerating blood pressure targets as even with best practice drug prescription and clinical support most participants still did not achieve the primary endpoint of blood pressure recommended by current guidelines. Most importantly, however, we have shown that a structured blood pressure management approach (computer assisted) results in higher blood pressure control rates in a primary care setting. As always, the challenge will be to implement this strategy into wider clinical practice by further analysing what did and did not work in this and other similar studies and how best to deliver pragmatic tools to facilitate the optimal management of blood pressure in the primary care setting.

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**What is already known on this topic**

Hypertension is the leading risk factor for cardiovascular disease worldwide and one of the most common conditions managed in primary care

Despite effective drug treatments, a substantial proportion of affected people remain above their ideal blood pressure target

Systematic reviews suggest that additional gains in blood pressure control can be attained with a more intensive structured approach to management

**What this study adds**

Compared with usual care, an intensive, structured programme for blood pressure control in primary care was associated with an 8.8% increase in the proportion of participants achieving individual targets and 9.5% increase in those achieving the classic target of  $\leq 140/90$  mm Hg

Despite parallel reductions in absolute cardiovascular risk, achieving more stringent blood pressure targets remains a challenge

Modification of treatment owing to adverse events was more common in the intervention group than usual care group (7.9% v 1.9%)

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## Tables

**Table 1 | Baseline characteristics according to group randomisation (n=1562). Values are numbers (percentages) of participants unless stated otherwise**

Characteristics	Usual care group (n=524)	Intervention group		
		Total (n=1038)	Monotherapy arm (n=360)	Combination therapy arm (n=678)
Men	323 (61.6)	640 (61.7)	222 (61.7)	418 (61.7)
Mean (SD) age (years)	59.3 (12.4)	59.2 (11.8)	59.5 (12.3)	59.1 (11.6)
Clinical profile:				
Previous hypertension	353 (67.4)	692 (66.7)	253 (70.3)	439 (64.7)
Heart disease	38 (7.3)	93 (9)	35 (9.7)	58 (8.6)
Type 2 diabetes	106 (20.2)	195 (18.8)	69 (19.2)	126 (18.6)
Proteinuria	93 (17.7)	183 (17.6)	64 (17.8)	119 (17.6)
Microalbuminuria	127 (24.2)	242 (23.3)	73 (20.3)	169 (24.9)
Mean (SD) eGFR (mL/min/1.73m <sup>2</sup> )	87.8 (19.6)	88.2 (19.7)	86.7 (18.4)	88.9 (20.4)
All cardiovascular disease	38 (7.3)	93 (9)	35 (9.7)	58 (8.6)
No cardiovascular disease or end organ damage	132 (25.2)	305 (29.4)	104 (28.9)	201 (29.6)
ECG evidence of LVH	36 (6.9)	75 (7.2)	23 (6.4)	52 (7.7)
Blood pressure profile (mm Hg):				
Mean (SD) systolic	149.2 (16.6)	150.0 (16.9)	150.1 (17.0)	149.9 (16.8)
Mean (SD) diastolic	87.4 (11.4)	88.4 (10.8)	88.0 (10.6)	88.6 (10.9)
Target blood pressure (mm Hg):				
≤140/90	145 (27.7)	304 (29.3)	106 (29.4)	198 (29.2)
≤130/80	286 (54.6)	557 (53.7)	190 (52.8)	367 (54.1)
≤125/75	93 (17.7)	177 (17.1)	64 (17.8)	113 (16.7)

eGFR=estimated glomerular filtration rate; ECG=electrocardiography; LVH=left ventricular hypertrophy.

**Table 2| Mean systolic and diastolic blood pressure and absolute change from baseline by study visits. Values are means (standard deviations) unless stated otherwise**

Measures at study visits	Usual care group	Intervention group		
		All	Monotherapy arm	Combination therapy arm
Baseline:	n=524	n=1038	n=360	n=678
Systolic blood pressure	149.2 (16.6)	150.0 (16.9)	150.1 (88.0)	149.9 (16.8)
Diastolic blood pressure	87.4 (11.4)	88.4 (10.8)	17.0 (10.6)	88.6 (10.9)
No (%) achieving blood pressure target*	—	—	—	—
Week 6:	n=479	n=945	n=319	n=626
Systolic blood pressure	142.1 (15.2)	141.5 (16.0)	144.6 (16.8)	140.0 (15.3)
Diastolic blood pressure	83.9 (10.8)	83.7 (10.4)	84.9 (10.4)	83.1 (10.3)
Change in blood pressure from baseline*	-7.2 (-3.6)	-8.3 (-4.7)	-5.1 (-3.2)	-9.9 (-5.5)
No (%) achieving blood pressure target*	97 (20.3)	223 (23.6)	57 (17.9)	166 (26.5)
Week 10:		n=872	n=294	n=578
Systolic blood pressure	—	139.8 (16.3)	143.8 (17.7)	137.8 (15.2)
Diastolic blood pressure	—	82.71 (1.0)	84.3 (11.1)	81.9 (10.9)
Change in blood pressure from baseline*	—	-10.1 (-5.5)	-6.1 (-3.5)	-12.1 (-6.5)
No (%) achieving blood pressure target*	—	247 (28.3)	66 (22.5)	181 (31.3)
Week 14:		n=831	n=281	n=550
Systolic blood pressure	—	136.3 (14.9)	139.4 (15.7)	134.7 (14.3)
Diastolic blood pressure	—	80.410.4	81.3 (10.7)	80.0 (10.1)
Change in blood pressure from baseline*	—	-13.4 (-7.7)	-10.5 (-6.5)	-14.9 (-8.4)
No (%) achieving blood pressure target*	—	295 (35.5)	80 (28.5)	215 (39.1)
Week 18:		n=829	n=276	n=553
Systolic blood pressure	—	135.2 (14.8)	136.5 (15.3)	134.6 (14.6)
Diastolic blood pressure	—	79.9 (10.3)	79.8 (10.7)	79.9 (10.1)
Change in blood pressure from baseline*	—	-14.5 (-8.3)	13.2 (-7.9)	-15.1 (-8.5)
No (%) achieving blood pressure target*	—	330 (39.8)	104 (37.7)	226 (40.9)
Week 26:	n=466	n=857	n=289	n=568
Systolic blood pressure	138.8 (14.9)	134.6 (14.2)	136.5 (15.6)	133.6 (13.3)
Diastolic blood pressure	82.0 (10.5)	79.9 (10.0)	80.4 (10.5)	79.6 (9.7)
Change in blood pressure from baseline*	-10.5 (-5.3)	-15.2 (-8.5)	-10.6 (-7.6)	-16.1 (-8.9)
No (%) achieving blood pressure target*	129 (27.7)	343 (40.0)	104 (36.0)	239 (42.1)
Endpoint blood pressure†:	n=504	n=988	n=339	n=649
Systolic blood pressure	139.1 (15.0)	136.1 (15.3)	138.0 (17.0)	135.1 (14.2)
Diastolic blood pressure	82.3 (10.5)	80.6 (10.2)	81.3 (10.8)	80.2 (9.9)
Change in blood pressure from baseline*	-10.2 (-5.2)	-13.7 (-7.8)	-11.8 (-6.8)	-14.6 (-8.4)
No (%) achieving blood pressure target*	138 (27.4)	358 (36.2)	112 (33.0)	246 (37.9)

\*Calculated for those only with data recorded at each time point and on an intention to treat basis.

†Calculated from last recorded data after randomisation.



**Table 3| Number of adverse events by system organ class and relation to study treatment derived from before randomisation (n=2185) and endpoint blood pressure (n=1492) population**

System organ class*	Before randomisation (n=2185)		Usual care group (n=504)		Intervention group					
					All (n=988)		Monotherapy arm (n=339)		Combination therapy arm (n=649)	
	Unrelated	Related	Unrelated	Related	Unrelated	Related	Unrelated	Related	Unrelated	Related
Total No of adverse events	620	424	550	113	1078	411	356	121	722	290
Blood and lymphatic system	4 (0.6)	1 (0.2)	2 (0.4)	0 (0)	2 (0.2)	0 (0)	0 (0)	0 (0)	2 (0.3)	0 (0)
Cardiac	74 (11.9)	100 (23.6)	88 (16)	23 (20.4)	127 (11.8)	102 (24.8)	32 (9)	34 (28.1)	95 (13.2)	68 (23.4)
Congenital, familial, and genetic	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	1 (0.2)	0 (0)	0 (0)	0 (0)	1 (0.3)
Ear and labyrinth	7 (1.1)	4 (0.9)	11 (2)	1 (0.9)	20 (1.9)	5 (1.2)	5 (1.4)	4 (3.3)	15 (2.1)	1 (0.3)
Endocrine	0 (0)	0 (0)	2 (0.4)	0 (0)	1 (0.1)	0 (0)	0 (0)	0 (0)	1 (0.1)	0 (0)
Eye	14 (2.3)	2 (0.5)	15 (2.7)	1 (0.9)	25 (2.3)	8 (1.9)	11 (3.1)	2 (1.7)	14 (1.9)	6 (2.1)
Gastrointestinal	81 (13.1)	67 (15.8)	32 (5.8)	15 (13.3)	100 (9.3)	44 (10.7)	42 (11.8)	16 (13.2)	58 (8)	28 (9.7)
General disorders and administration site conditions	54 (8.7)	52 (12.3)	47 (8.5)	20 (17.7)	73 (6.8)	106 (25.8)	23 (6.5)	18 (14.9)	50 (6.9)	88 (30.3)
Hepatobiliary	1 (0.2)	0 (0)	1 (0.2)	0 (0)	3 (0.3)	0 (0)	1 (0.3)	0 (0)	2 (0.3)	0 (0)
Immune system	3 (0.5)	0 (0)	1 (0.2)	0 (0)	9 (0.8)	0 (0)	3 (0.8)	0 (0)	6 (0.8)	0 (0)
Infections and infestations	78 (12.6)	2 (0.5)	86 (15.6)	1 (0.9)	161 (14.9)	2 (0.5)	45 (12.6)	0 (0)	116 (16.1)	2 (0.7)
Injury, poisoning, and procedural complications	28 (4.5)	0 (0)	19 (3.5)	0 (0)	60 (5.6)	3 (0.7)	17 (4.8)	1 (0.8)	43 (6)	2 (0.7)
Investigations	13 (2.1)	2 (0.5)	8 (1.5)	1 (0.9)	22 (2)	8 (1.9)	8 (2.2)	2 (1.7)	14 (1.9)	6 (2.1)
Metabolism and nutrition	10 (1.6)	3 (0.7)	10 (1.8)	2 (1.8)	20 (1.9)	4 (1)	11 (3.1)	3 (2.5)	9 (1.2)	1 (0.3)
Musculoskeletal and connective tissue	49 (7.9)	29 (6.8)	58 (10.5)	9 (8)	106 (9.8)	26 (6.3)	38 (10.7)	7 (5.8)	68 (9.4)	19 (6.6)
Neoplasms benign, malignant, and unspecified (including cysts and polyps)	1 (0.2)	0 (0)	5 (0.9)	0 (0)	14 (1.3)	0 (0)	5 (1.4)	0 (0)	9 (1.2)	0 (0)
Nervous system	67 (10.8)	73 (17.2)	37 (6.7)	18 (15.9)	80 (7.4)	39 (9.5)	27 (7.6)	13 (10.7)	53 (7.3)	26 (9)
Psychiatric	21 (3.4)	26 (6.1)	15 (2.7)	5 (4.4)	30 (2.8)	10 (2.4)	17 (4.8)	3 (2.5)	13 (1.8)	7 (2.4)
Renal and urinary	12 (1.9)	3 (0.7)	11 (2)	0 (0)	14 (1.3)	6 (1.5)	2 (0.6)	3 (2.5)	12 (1.7)	3 (1)
Reproductive system and breast	3 (0.5)	2 (0.5)	6 (1.1)	3 (2.7)	14 (1.3)	4 (1)	4 (1.1)	2 (1.7)	10 (1.4)	2 (0.7)
Respiratory, thoracic, and mediastinal	44 (7.1)	19 (4.5)	57 (10.4)	6 (5.3)	96 (8.9)	13 (3.2)	35 (9.8)	6 (5)	61 (8.4)	7 (2.4)
Skin and subcutaneous tissue	28 (4.5)	19 (4.5)	21 (3.8)	0 (0)	66 (6.1)	14 (3.4)	21 (5.9)	3 (2.5)	45 (6.2)	11 (3.8)
Social circumstances	0 (0)	0 (0)	0 (0)	0 (0)	1 (0.1)	0 (0)	1 (0.3)	0 (0)	0 (0)	0 (0)
Surgical and medical procedures	13 (2.1)	0 (0)	10 (1.8)	1 (0.9)	16 (1.5)	0 (0)	4 (1.1)	0 (0)	12 (1.7)	0 (0)
Vascular	15 (2.4)	17 (4)	8 (1.5)	5 (4.4)	18 (1.7)	14 (3.4)	4 (1.1)	4 (3.3)	14 (1.9)	10 (3.4)
Unknown	0 (0)	3 (0.7)	0 (0)	3 (2.7)	0 (0)	2 (0.5)	0 (0)	0 (0)	0 (0)	2 (0.7)

41 adverse events with unknown incident date, not presented in table.

\*Defined as highest level according to Medical Dictionary for Regulatory Activities (MedDRA, [www.meddrasso.com/index.asp](http://www.meddrasso.com/index.asp)), which contains medical terminology used to classify information on adverse events associated with use of biopharmaceuticals and other medical products (for example, medical devices and vaccines).

Table 3 (continued)

System organ class*	Before randomisation (n=2185)		Usual care group (n=504)		Intervention group					
					All (n=988)		Monotherapy arm (n=339)		Combination therapy arm (n=649)	
	Unrelated	Related	Unrelated	Related	Unrelated	Related	Unrelated	Related	Unrelated	Related

Coding these data to a standard set of MedDRA terms allows health authorities and the biopharmaceutical industry to more readily exchange and analyse data related to the safe use of medical products.

## Figures

Timeline	Usual care	Intervention valsartan monotherapy	Intervention valsartan + hydrochlorothiazide	Intervention valsartan + amlodipine
Site initiation	<b>a</b>			
Clinical profiling	<b>b</b>			
Treatment run-in	<b>c</b>			
Randomisation	Random allocation according to individual blood pressure goal			
Baseline (time 0)	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
0 to 6 weeks	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
6 weeks	Initial evaluation of blood pressure			
6 to 10 weeks	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
10 to 14 weeks	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
14 to 18 weeks	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
18 to 26 weeks	<b>d</b>	<b>e</b> ①	<b>e</b> ②	<b>e</b> ③
26 weeks	Re-evaluation of blood pressure and other clinical variables to determine study endpoints			
Site closure	<b>f</b>			

**a** 30-60 minute general practitioner investigator site initiation visit

**b** All participants were subjected to clinical profiling to determine their individual blood pressure target and other risk targets using the Baker IDI computerised risk profiling tool

**c** Eligible participants started a 28 day run-in phase of valsartan 80 mg/day with an interim visit at 14 days to assess blood pressure levels and the need for withdrawal or rescue randomisation if required

**d** Usual primary care with no restriction on prescribed pharmacotherapy (see appendix I in supplementary file) or schedule of visits to achieve individualised blood pressure control (prespecified at randomisation according to initial clinical profiling)

**e** Intervention applied via a computer assisted algorithm to: a) determine if participants have achieved their individual blood pressure goal, b) initiate treatment up-titration as per predetermined treatment protocol (see below), and c) automate clinic follow-up visits at 6, 10, 14, 18, and 26 weeks

① Initially prescribed valsartan 160 mg/day with progressive stepped up-titration if not achieving individual blood pressure target to valsartan 320 mg/day then valsartan/hydrochlorothiazide 320 mg + 12.5 mg/day then valsartan/hydrochlorothiazide 320 mg + 25 mg/day then consideration of alternative/triple therapy to achieve individual blood pressure control (see appendix I in supplementary file)

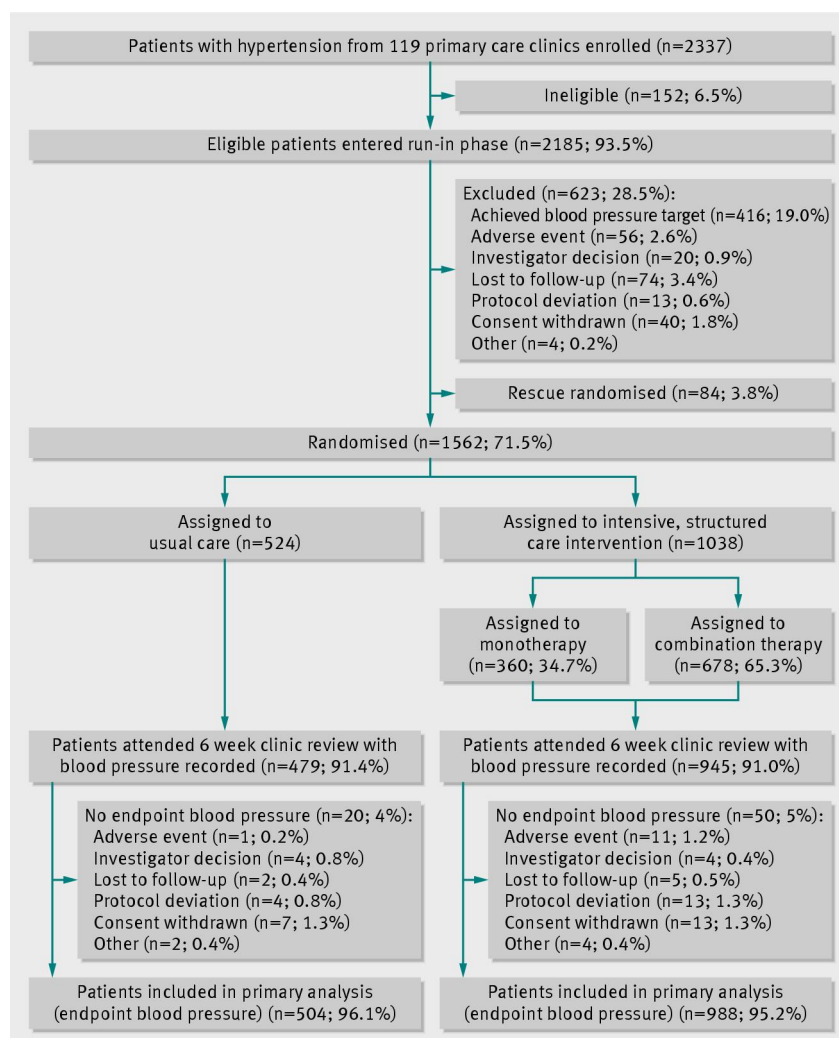
② Initially prescribed valsartan/hydrochlorothiazide 80 mg/day + 12.5 mg/day with progressive stepped up-titration if not achieving individual blood pressure target to valsartan/hydrochlorothiazide 160 mg + 12.5 mg/day then valsartan/hydrochlorothiazide 160 mg + 25 mg/day then consideration of alternative/triple therapy to achieve individual blood pressure control (see appendix I in supplementary file)

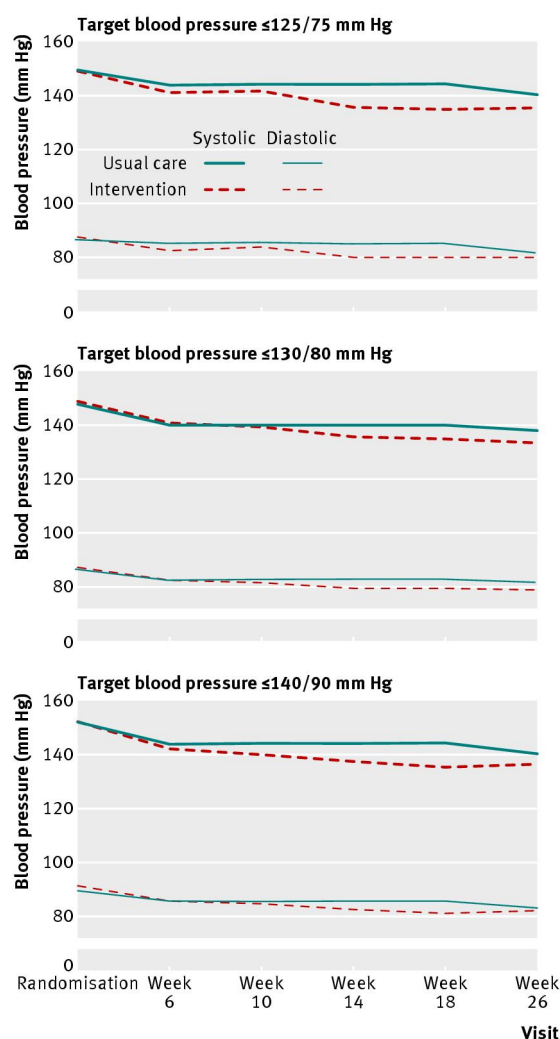
③ Initially prescribed valsartan/amlodipine 80 mg/day + 5 mg/day with progressive stepped up-titration if not achieving individual blood pressure target to valsartan/amlodipine 160 mg + 5 mg/day then valsartan/amlodipine 160 mg + 10 mg/day then consideration of alternative/triple therapy to achieve individual blood pressure control (see appendix I in supplementary file)

**f** Clinical audit (20% data verification) and site closure visit

**Fig 1** Summary of study timelines



**Fig 2** Flow of participants through study



**Fig 3** Change in systolic and diastolic blood pressure according to individual blood pressure target at randomisation (n=1492). On an adjusted basis, participants assigned to intervention were significantly more likely to achieve their target at the two higher blood pressure target levels