

Clinical trials of anti hypertensive agents for diabetes type 2 in patients with hypertension

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1 angiotensin receptor blocker

| Trial | Treatments | Patients | Trials design and methods |
|---|---|---|--|
| irbesartan vs placebo | | | |
| IDNT (irbesartan vs pbo) , 2001 n=579/569 follow-up: 2.6 years | Irbesartan 300 mg daily versus placebo | hypertensive patients with nephropathy due to type 2 diabetes | Parallel groups double blind Worldwide |
| IPDM (150mg) , 2001 n=195/201 follow-up: 2 years | irbesartan 150 mg daily versus placebo | hypertensive patients with type 2 diabetes and microalbuminuria | Parallel groups double-blind Worldwide |
| irbesartan vs amlodipine | | | |
| IDNT (irbesartan vs amlodipine) , 2001 n=579/567 follow-up: 2.6 years | Irbesartan 300 mg daily versus amlodipine 10 mg daily | hypertensive patients with nephropathy due to type 2 diabetes | Parallel groups double blind Worldwide |
| valsartan vs amlodipine | | | |
| NAGOYA HEART , 2011 <i>unpublished</i> [NCT00129233] n=575/575 follow-up: 3.2 y median | blood-pressure-lowering therapy based on valsartan; blood-pressure goal of <130/80 mm Hg versus blood-pressure-lowering therapy based on amlodipine; blood-pressure goal of <130/80 mm Hg | patients with hypertension with type 2 diabetes or impaired glucose tolerance | Parallel groups open Japan |
| losartan vs atenolol | | | |
| LIFE (diabetic subgroup) , 2002 n=586/609 follow-up: 4.7 years | losartan 50mg daily at step 1 versus atenolol 50mg daily at step 1 | patients with diabetes (subgroup) , hypertension, and signs of left-ventricular hypertrophy on electrocardiograms | Parallel groups double-blind USA, UK, Nordic countries |

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NAGOYA HEART, 2011:

Matsushita K, Muramatsu T, Kondo T, Maeda K, Shintani S, Murohara T Rationale and design of the NAGOYA HEART Study: comparison between valsartan and amlodipine regarding morbidity and mortality in patients with hypertension and glucose intolerance. J Cardiol 2010;56:111-7 [20409690] 10.1016/j.jjcc.2010.03.004

LIFE (diabetic subgroup), 2002:

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2 angiotensin-converting enzyme inhibitors

| Trial | Treatments | Patients | Trials design and methods |
|--|--|--|--|
| captopril or atenolol vs control | | | |
| UKPDS 38 , 1998 n=758/390 follow-up: 8.4y (median) | tight control of blood pressure aiming at a BP <150/85 (with the use of captopril or atenolol as main treatment, other treatment were added if the control criteria were not met) versus less tight control aiming at a blood pressure of <180/105 (avoiding treatment with ACE inhibitors or beta-blockers) | hypertensive patients with type 2 diabetes | Parallel groups open UK |
| ACE inhibitors vs placebo | | | |
| HOPE (diabetic subgroup) , 2000 n=1808/1759 follow-up: 4.5 years | ramipril 10 mg once per day orally versus placebo | patients with diabetes (sub group), aged 55 years or older, who had a previous cardiovascular event or at least one other cardiovascular risk factor, no clinical proteinuria, heart failure, or low ejection fraction | Factorial plan double-blind North, South america, Europe |
| captopril vs atenolol | | | |
| UKPDS 39 , 1998 n=400/358 follow-up: ND | captopril 25 mg/d aiming at a BP <150/85 versus atenolol 50mg/d aiming at a BP <150/85 | hypertensive patients with type 2 diabetes | Parallel groups open UK |
| ACE inhibitor vs calcium-channel blocker | | | |

continued...

| Trial | Treatments | Patients | Trials design and methods |
|--|--|---|--|
| STOP-2 (ACEI vs CCB) (diabetic subgroup) , 2000 n=235/231 follow-up: 5.03y | ACE inhibitor versus calcium antagonists | diabetic (subgroup) elderly patients aged 70-84 years | open with blind assessment Sweden |
| lisinopril vs chlorthalidone | | | |
| ALLHAT (lisi vs chlor, diabetic subgroup) , 2002 n=2431/4498 follow-up: 4.9 y | lisinopril 10 to 40 mg/d versus chlorthalidone 12.5 to 25 mg/d | diabetic (subgroup) participants aged 55 years or older with hypertension | Parallel groups double-blind |
| captopril vs diuretic and/or beta-blockers | | | |
| CAPP (diabetic subgroup) , 1999 n=309/263 follow-up: 6.1 year | Captopril initial dose of 50 mg daily given in one or two doses versus thiazide diuretic or beta-blocker | Patients aged 25-66 years with a measured diastolic blood pressure of 100 mm Hg or more on two occasions; subgroup of diabetic patients | Parallel groups open with blinded assessment Sweden, Finland |
| ACE inhibitor vs diuretic or beta-blocker | | | |
| STOP-2 (ACEI, diabetic subgroup) , 2000 n=235/253 follow-up: 5.03y | ACE inhibitor versus conventional treatment (diuretic or beta-blocker) | diabetic (subgroup) elderly patients aged 70-84 years with hypertension | Parallel groups open with blind assessment Sweden |

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3 calcium-channel blockers

| Trial | Treatments | Patients | Trials design and methods |
|---|---|--|---|
| amlodipine vs placebo | | | |
| IDNT (amlodipine vs PBO) , 2001 n=567/569 follow-up: 2.6 years | Amlodipine 10 mg daily versus placebo | hypertensive patients with nephropathy due to type 2 diabetes | Parallel groups double-blind Worldwide |
| nitrendipine vs placebo | | | |
| Syst-Eur (diabetic subgroup) , 1999 n=252/240 follow-up: 2 years | Calcium-channel blocker versus placebo | subgroup of diabetic patients, age, >=60 years) with systolic blood pressure of 160 to 219 mm Hg and diastolic pressure below 95 mm Hg | Parallel groups double blind |
| benazepril + amlodipine vs benazepril + hydrochlorothiazide | | | |
| ACCOMPLISH (diabetic subgroup) , 2010 [NCT00170950] n=1432/1410 follow-up: 36 months | benazepril, combined with amlodipine versus benazepril, combined with hydrochlorothiazide | patients with diabetes (subgroup) and hypertension at high risk of cardiovascular and related events | Parallel groups double-blind US, Norway, Denmark, Finland |
| amlodipine vs chlorthalidone | | | |

continued...

| Trial | Treatments | Patients | Trials design and methods |
|--|---|---|---|
| ALLHAT (amlodipine vs chlor, diabetic subgroup) , 2002 n=2664/4498 follow-up: 4.9 y | amlodipine versus chlorthalidone | diabetic (subgroup) participants aged 55 years or older with hypertension | Parallel groups double-blind |
| nifedipine vs coamilofide | | | |
| INSIGHT (diabetic subgroup) , 2000 n=649/653 follow-up: 4 y | Nifedipine GITS 30 mg daily versus co-amilofide hydrochlorothiazide 25 mg plus amilofide 2.5 mg | diabetic (subgroup) patients aged 55-80 years with hypertension (blood pressure \geq 150/95 mm Hg, or \geq 160 mmHg systolic) | Parallel groups double-blind Europe, Israel |
| diltiazem vs diuretic and/or beta-blocker | | | |
| NORDIL (diabetic subgroup) , 2000 n=351/376 follow-up: 4.5 y | Diltiazem 180/360 mg diltiazem daily at step one versus thiazide diuretic or a beta-blocker at step one | diabetic patients (subgroup), aged 50-74 years who had diastolic blood pressure of 100 mm Hg or more | Parallel groups open Norway, Sweden |
| calcium-channel blocker vs diuretic or beta-blocker | | | |
| STOP-2 (CCB, diabetic subgroup) , 2000 n=231/253 follow-up: 5.03y | Calcium-channel blocker versus diuretic or beta-blocker | diabetic (subgroup) elderly patients aged 70-84 years | Parallel groups open with blind assessment Sweden |
| nisoldipine vs enalapril | | | |
| ABCD (hypertension) , 1998 n=235/235 follow-up: 5 y | nisoldipine (long acting) versus enalapril | patients with non-insulin-dependent diabetes and hypertension | Factorial plan Double blind USA |
| amlodipine vs fosinopril | | | |
| FACET , 1997 n=191/189 follow-up: 3.5 y | amlodipine (long acting) 10 mg daily versus fosinopril 20 mg daily | hypertensive patients with NIDDM | Parallel groups open Italy |

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4 diuretics

| Trial | Treatments | Patients | Trials design and methods |
|--|--|--|---------------------------------|
| chlorthalidone vs placebo | | | |
| SHEP (diabetic subgroup) , 1996 n=283/300 follow-up: 5 year | low dose of chlorthalidone (12.5-25.0 mg/d) with a step-up to atenolol (25.0-50.0 mg/d) or reserpine (0.05-0.10 mg/d) if needed versus placebo | men and women aged 60 years and older , non-insulin-treated diabetic (sub group) patients with isolated systolic hypertension (systolic BP \geq 160 mm Hg; diastolic BP, $<$ 90 mm Hg) | Parallel groups double-blind |

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5 intensive treatment

| Trial | Treatments | Patients | Trials design and methods |
|---|---|---|---|
| intensive vs usual | | | |
| ACCORD (blood pressure) , 2010 [NCT00000620] n=2363/2371 follow-up: 4.7 y | intensive blood-pressure control, targeting a systolic pressure of less than 120 mm Hg versus standard blood-pressure control | high-risk patients with type 2 diabetes, high HbA1c concentrations ($>$ 7.5%), and cardiovascular disease (or \geq 2 cardiovascular risk factors) | Factorial plan open United States, Canada |

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6 Treatment blood pressure target

| Trial | Treatments | Patients | Trials design and methods |
|---|---|---|---------------------------|
| more intensive blood pressure lowering strategie vs less intensive blood pressure lowering strategie | | | |
| ABCD target (H) , 2000 n=237/233 follow-up: 5 year | intensive treatment with a diastolic blood pressure goal of 75 mmHg versus moderate treatment with a diastolic blood pressure goal of 80-89 mmHg | diabetes patients with DBP \geq 90 mmHg | Parallel groups open |
| ABCD target (N) , 2002 n=237/243 follow-up: | intensive treatment (diastolic blood pressure decrease of 10 mmHg below baseline DBP) versus moderate treatment (diastolic blood pressure goal of 80-89 mmHg) | diabetes patients with diastolic blood pressure between 80 and 89mmHg | Parallel groups open |

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