Effect of anti-diabetic drugs in primary prevention of cardiovascular disease in type 2 diabetes and prediabetes: a systematic review and meta-analysis of randomized trials

Online-only supplemental materials

Supplemental Methods.

Supplemental Results.

Supplemental List. List of citations of all eligible trials. Two eligible trials, NCT00121966 and NCT02273050, reported the results by different anti-diabetic drugs. These 2 trials were divided as separate studies in the quantitative synthesis.

Supplemental Table S1: Search strategy on Medline (Ovid).

Supplemental Table S2: Characteristics of included trials.

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Supplemental Table S4. Detailed results of meta-analysis of anti-diabetic drugs in preventing major cardiovascular events.

Supplemental Figure 1. Flow-diagram for literature search and screening.

Supplemental Figure 2. Funnel plot of anti-diabetic drugs in preventing major cardiovascular events.

Supplemental References.

Supplemental Methods

This systematic review was reported in compliance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA). The research protocol was registered on the PROSPERO registry (CRD42020196192, 30 July 2020).

Study eligibility

Eligible studies included those where diet, exercise, lifestyle advice, or any other non-pharmacological intervention was used alongside drug therapy, providing the only difference between comparison arms was the presence or absence of anti-diabetic drug treatment. Trials were included if at least one of the following macrovascular outcomes were reported: fatal or non-fatal myocardial infarction, fatal or non-fatal stroke, cardiovascular death, unstable angina, heart failure, and revascularization.

Search strategy

Relevant RCTs reporting management of diabetes and prediabetes were sought from database inception up to 20 September 2022 from Medline, Embase, the Cochrane library, Clinicaltrials.gov registry, and International Clinical Trials Registry Platform (ICTRP) of RCTs. Searches included keywords and MeSH terms for type 2 diabetes mellitus, impaired fasting glucose, impaired glucose tolerance, non-insulin anti-diabetes drugs and filters for RCTs (see Supplement Table 1 for search details). Hand-searching of the bibliographies of review articles and eligible trials was also conducted.

Screening

Initial screening of identified citations was undertaken by a member of the research group. Any potentially eligible trial was retained and further independently assessed by two researchers. Disagreements were resolved through discussion or consultation with a third experienced researcher.

Data extraction

Data were extracted independently by two researchers and entered into a structured extraction form. The following data were extracted from each eligible trial: 1) details of the citation (author, publication year, trial registration number); 2) study design, treatment details, and outcome measures; 3) patients' characteristics (age, body mass index [BMI], baseline glucose measures,

etc.); 4) main results about treatment effect. Discrepancy in extracted data was resolved through discussion or by consulting a third researcher, if necessary. Data not reported in original publications were sought from the online appendices, clinical trials registry websites, or other publications derived from the same trial.

Risk of bias assessment

The risk of bias was assessed independently by two reviewers with disagreements settled through discussions. The Cochrane Collaboration's tool for assessing risk of bias was used to assess six domains: sequence generation, concealment of allocation, blinding, incomplete outcome data, and selective reporting.²

Statistical analysis

Heterogeneity was assessed using the I² statistic, which describes the percentage of variability in effect estimates that was due to clinical and/or methodological factors rather than sampling error.³ Heterogeneity was considered substantial if I² was greater than 50%. Publication bias was examined using the funnel plot and Egger's test.³ All statistical analyses were conducted by using R (R version 4.3.1, R Foundation for Statistical Computing, Vienna, Austria, 2023, https://www.R-project.org/). A two-sided significance level of 0.05 was adopted for all analyses except for the statistical test for publication bias, in which a two-sided P value less than 0.1 was indicative of statistical significance.

Supplemental Results

Flow-diagram for literature search and screening is shown in Supplement Figure 1. Full citations of eligible trials are listed in the Supplement List. Characteristics of eligible trials are summarized in Supplement Table 2. The majority of patients (99.2%) were over-weight or obese. All-cause mortality, myocardial infarction, stroke, and the broader composite cardiovascular outcome were reported in 68, 90, 83, and 129 trials, respectively.

Overall, 40 trials evaluated the effects of various dipeptidyl peptidase 4 (DPP4) inhibitors, 6 evaluated metformin, 6 evaluated thiazolidinediones, 29 evaluated sodium glucose cotransporter 2 (SGLT2) inhibitors, 31 evaluated glucagon-like peptide 1 (GLP1) receptor agonists, and the other 20 trials evaluated alpha-glucosidase inhibitors, sulfonylureas, meglitinides, or combination of different drugs. In 98 trials (74.2%), some participants were already on certain anti-diabetic medications (background treatments) at baseline. Intervention treatment details are shown in Supplement Table 2.

Detailed results regarding risk of bias of eligible trials are shown in Supplement Table 3. All studies mentioned that they had allocated patients randomly to groups, although 33.3% of eligible trials did not report the details of randomization methods. A placebo was used in 94.7% of eligible trials. More than 80% of eligible trials had low risk of bias in the domains of "incomplete outcome data" and "selective reporting". No obvious evidence for publication bias was identified through visual inspection of the funnel plot (Supplement Figure 2) or Egger's test (P = 0.630).

Supplemental List. List of citations of all eligible trials. Two eligible trials, NCT00121966 and NCT02273050, reported the results by different anti-diabetic drugs. These 2 trials were divided as separate studies in the quantitative synthesis.

- 1. Coniff RF, Shapiro JA, Seaton TB, Bray GA. Multicenter, placebo-controlled trial comparing acarbose (BAY g 5421) with placebo, tolbutamide, and tolbutamide-plus-acarbose in non-insulin-dependent diabetes mellitus. *American Journal of Medicine* 1995; **98** (5): 443-51.
- 2. UK Prospective Diabetes Study (UKPDS) Group. Effect of intensive blood-glucose control with metformin on complications in overweight patients with type 2 diabetes (UKPDS 34). *The Lancet* 1998; **352**(9131): 854-65.
- 3. Johnston PS, Lebovitz HE, Coniff RF, Simonson DC, Raskin P, Munera CL. Advantages of α-glucosidase inhibition as monotherapy in elderly type 2 diabetic patients. *The Journal of Clinical Endocrinology & Metabolism* 1998; **83**(5): 1515-22.
- 4. Chiasson JL, Josse RG, Gomis R, Hanefeld M, Karasik A, Laakso M. Acarbose Treatment and the Risk of Cardiovascular Disease and Hypertension in Patients with Impaired Glucose Tolerance: The STOP-NIDDM Trial. *Journal of the American Medical Association* 2003; **290** (4): 486-94.
- 5. Buse JB, Henry RR, Han J, Kim DD, Fineman MS, Baron AD. Effects of exenatide (exendin-4) on glycemic control over 30 weeks in sulfonylurea-treated patients with type 2 diabetes. *Diabetes care* 2004; **27**(11): 2628-35.
- 6. Ratner R, Goldberg R, Haffner S, et al. Impact of intensive lifestyle and metformin therapy on cardiovascular disease risk factors in the diabetes prevention program. *Diabetes Care* 2005; **28**(4): 888-94.
- 7. Investigators. DT. Effect of rosiglitazone on the frequency of diabetes in patients with impaired glucose tolerance or impaired fasting glucose: a randomised controlled trial. *The Lancet* 2006; **368**(9541): 1096-105.
- 8. Ramachandran A, Snehalatha C, Mary S, Mukesh B, Bhaskar AD, Vijay V. The Indian Diabetes Prevention Programme shows that lifestyle modification and metformin prevent type 2 diabetes in Asian Indian subjects with impaired glucose tolerance (IDPP-1). *Diabetologia* 2006; **49** (2): 289-97.
- 9. Bosi E, Camisasca RP, Collober C, Rochotte E, Garber AJ. Effects of vildagliptin on glucose control over 24 weeks in patients with type 2 diabetes inadequately controlled with metformin. *Diabetes Care* 2007; **30**(4): 890-5.
- 10. Goldstein BJ, Feinglos MN, Lunceford JK, Johnson J, Williams-Herman DE. Effect of initial combination therapy with sitagliptin, a dipeptidyl peptidase-4 inhibitor, and metformin on glycemic control in patients with type 2 diabetes. *Diabetes Care* 2007; **30 (8)**: 1979-87.
- 11. Hermansen K, Kipnes M, Luo E, et al. Efficacy and safety of the dipeptidyl

- peptidase-4 inhibitor, sitagliptin, in patients with type 2 diabetes mellitus inadequately controlled on glimepiride alone or on glimepiride and metformin. *Diabetes Obes Metab* 2007; **9**(5): 733-45.
- 12. Mita T, Watada H, Shimizu T, et al. Nateglinide reduces carotid intima-media thickening in type 2 diabetic patients under good glycemic control. *Arteriosclerosis, Thrombosis, and Vascular Biology* 2007; **27** (11): 2456-62.
- 13. DeFronzo RA, Hissa MN, Garber AJ, et al. The efficacy and safety of saxagliptin when added to metformin therapy in patients with inadequately controlled type 2 diabetes with metformin alone. *Diabetes Care* 2009; **32**(9): 1649-55.
- 14. Hollander P, Li J, Allen E, Chen R, Investigators CV. Saxagliptin added to a thiazolidinedione improves glycemic control in patients with type 2 diabetes and inadequate control on thiazolidinedione alone. *J Clin Endocrinol Metab* 2009; **94**(12): 4810-9.
- 15. Kawamori R, Tajima N, Iwamoto Y, Kashiwagi A, Shimamoto K, Kaku K. Alpha-glucosidase inhibitor for the prevention of type 2 diabetes mellitus: a randomised, double-blind trial in Japanese subjects with impaired glucose tolerance. *Nihon rinsho Japanese journal of clinical medicine*, 2009. (accessed.
- 16. Kooy A, de Jager J, Lehert P, et al. Long-term effects of metformin on metabolism and microvascular and macrovascular disease in patients with type 2 diabetes mellitus. *Archives of internal medicine* 2009; **169**(6): 616-25.
- 17. Pratley RE, Kipnes MS, Fleck PR, Wilson C, Mekki Q, Alogliptin Study G. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor alogliptin in patients with type 2 diabetes inadequately controlled by glyburide monotherapy. *Diabetes Obes Metab* 2009; **11**(2): 167-76.
- 18. Pratley RE, Reusch JE, Fleck PR, Wilson CA, Mekki Q, Alogliptin Study G. Efficacy and safety of the dipeptidyl peptidase-4 inhibitor alogliptin added to pioglitazone in patients with type 2 diabetes: a randomized, double-blind, placebo-controlled study. *Curr Med Res Opin* 2009; **25**(10): 2361-71.
- 19. Ramachandran A, Snehalatha C, Mary S, et al. Pioglitazone does not enhance the effectiveness of lifestyle modification in preventing conversion of impaired glucose tolerance to diabetes in Asian Indians: Results of the Indian Diabetes Prevention Programme-2 (IDPP-2). *Diabetologia* 2009; **52**(6): 1019-26.
- 20. Rosenstock J, Aguilar-Salinas C, Klein E, Nepal S, List J, Chen R. Effect of saxagliptin monotherapy in treatment-naive patients with type 2 diabetes. *Current medical research and opinion* 2009; **25**(10): 2401-11.
- 21. Rosenstock J, Rendell MS, Gross JL, Fleck PR, Wilson CA, Mekki Q. Alogliptin added to insulin therapy in patients with type 2 diabetes reduces HbA(1C) without causing weight gain or increased hypoglycaemia. *Diabetes Obes Metab* 2009; 11(12): 1145-52.

- 22. Bailey CJ, Gross JL, Pieters A, Bastien A, List JF. Effect of dapagliflozin in patients with type 2 diabetes who have inadequate glycaemic control with metformin: a randomised, double-blind, placebo-controlled trial. *The Lancet* 2010; **375**(9733): 2223-33.
- 23. Lindblad U, Lindberg G, Mansson NO, et al. Can sulphonylurea addition to lifestyle changes help to delay diabetes development in subjects with impaired fasting glucose? The Nepi ANtidiabetes StudY (NANSY). *Diabetes, obesity & metabolism*, 2010. (accessed.
- 24. Liutkus J, Rosas Guzman J, Norwood P, et al. A placebo-controlled trial of exenatide twice-daily added to thiazolidinediones alone or in combination with metformin. *Diabetes Obes Metab* 2010; **12**(12): 1058-65.
- 25. Truitt KE, Goldberg RB, Rosenstock J, et al. A 26-week, placebo- and pioglitazone-controlled, dose-ranging study of rivoglitazone, a novel thiazolidinedione for the treatment of type 2 diabetes. *Current Medical Research and Opinion*, 2010. (accessed.
- 26. Vilsbøll T, Rosenstock J, Yki-Järvinen H, et al. Efficacy and safety of sitagliptin when added to insulin therapy in patients with type 2 diabetes. *Diabetes, Obesity and Metabolism* 2010; **12**(2): 167-77.
- 27. Zinman B, Harris SB, Neuman J, et al. Low-dose combination therapy with rosiglitazone and metformin to prevent type 2 diabetes mellitus (CANOE trial): a double-blind randomised controlled study. *The Lancet* 2010; **376**(9735): 103-11.
- 28. Barzilai N, Guo H, Mahoney EM, et al. Efficacy and tolerability of sitagliptin monotherapy in elderly patients with type 2 diabetes: a randomized, double-blind, placebo-controlled trial. *Current Medical Research and Opinion* 2011; **27**(5): 1049-58.
- 29. Chacra AR, Tan GH, Ravichandran S, List J, Chen R, Investigators CV. Safety and efficacy of saxagliptin in combination with submaximal sulphonylurea versus up-titrated sulphonylurea over 76 weeks. *Diab Vasc Dis Res* 2011; **8**(2): 150-9.
- 30. DeFronzo RA, Tripathy D, Schwenke DC, et al. Pioglitazone for diabetes prevention in impaired glucose tolerance. *The New England Journal of Medicine* 2011; **364**(12): 1104-15.
- 31. Del Prato S, Barnett A, Huisman H, Neubacher D, Woerle HJ, Dugi K. Effect of linagliptin monotherapy on glycaemic control and markers of β-cell function in patients with inadequately controlled type 2 diabetes: a randomized controlled trial. *Diabetes, Obesity and Metabolism* 2011; **13**(3): 258-67.
- 32. Gomis R, Espadero RM, Jones R, Woerle H, Dugi K. Efficacy and safety of initial combination therapy with linagliptin and pioglitazone in patients with inadequately controlled type 2 diabetes: a randomized, double-blind, placebocontrolled study. *Diabetes, Obesity and Metabolism* 2011; **13**(7): 653-61.

- 33. Gram J, Henriksen JE, Grodum E, et al. Pharmacological treatment of the pathogenetic defects in type 2 diabetes: the randomized multicenter South Danish Diabetes Study. *Diabetes Care* 2011; **34**(1): 27-33.
- 34. Nowicki M, Rychlik I, Haller H, et al. Long-term treatment with the dipeptidyl peptidase-4 inhibitor saxagliptin in patients with type 2 diabetes mellitus and renal impairment: a randomised controlled 52-week efficacy and safety study. *Int J Clin Pract* 2011; **65**(12): 1230-9.
- 35. Olansky L, Reasner C, Seck T, et al. A treatment strategy implementing combination therapy with sitagliptin and metformin results in superior glycaemic control versus metformin monotherapy due to a low rate of addition of antihyperglycaemic agents. *Diabetes, Obesity and Metabolism* 2011; **13**(9): 841-9.
- 36. Owens DR, Swallow R, Dugi KA, Woerle HJ. Efficacy and safety of linagliptin in persons with type 2 diabetes inadequately controlled by a combination of metformin and sulphonylurea: a 24-week randomized study. *Diabet Med* 2011; **28**(11): 1352-61.
- 37. Strojek K, Yoon KH, Hruba V, Elze M, Langkilde AM, Parikh S. Effect of dapagliflozin in patients with type 2 diabetes who have inadequate glycaemic control with glimepiride: a randomized, 24-week, double-blind, placebo-controlled trial. *Diabetes Obes Metab* 2011; **13**(10): 928-38.
- 38. Taskinen MR, Rosenstock J, Tamminen I, et al. Safety and efficacy of linagliptin as add-on therapy to metformin in patients with type 2 diabetes: a randomized, double-blind, placebo-controlled study. *Diabetes, Obesity and Metabolism* 2011; **13**(1): 65-74.
- 39. Chou HS, Truitt KE, Moberly JB, et al. A 26-week, placebo- and pioglitazone-controlled monotherapy study of rivoglitazone in subjects with type 2 diabetes mellitus. *Diabetes, Obesity and Metabolism* 2012; **14** (11): 1000-9.
- 40. DeFronzo RA, Burant CF, Fleck P, Wilson C, Mekki Q, Pratley RE. Efficacy and tolerability of the DPP-4 inhibitor alogliptin combined with pioglitazone, in metformin-treated patients with type 2 diabetes. *J Clin Endocrinol Metab* 2012; **97**(5): 1615-22.
- 41. Haak T, Meinicke T, Jones R, Weber S, Eynatten MV, Woerle HJ. Initial combination of linagliptin and metformin improves glycaemic control in type 2 diabetes: A randomized, double-blind, placebo-controlled study. *Diabetes, Obesity and Metabolism* 2012; **14** (6): 565-74.
- 42. Pan CY, Yang W, Tou C, Gause-Nilsson I, Zhao J. Efficacy and safety of saxagliptin in drug-naive Asian patients with type 2 diabetes mellitus: A randomized controlled trial. *Diabetes/Metabolism Research and Reviews* 2012; **28** (3): 268-75.
- 43. Rosenstock J, Vico M, Wei L, Salsali A, List JF. Effects of dapagliflozin, an

- SGLT2 inhibitor, on HbA(1c), body weight, and hypoglycemia risk in patients with type 2 diabetes inadequately controlled on pioglitazone monotherapy. *Diabetes Care* 2012; **35**(7): 1473-8.
- 44. Wilding JP, Woo V, Soler NG, et al. Long-term efficacy of dapagliflozin in patients with type 2 diabetes mellitus receiving high doses of insulin: a randomized trial. *Annals of internal medicine* 2012; **156**(6): 405-15.
- 45. Ahren B, Leguizamo Dimas A, Miossec P, Saubadu S, Aronson R. Efficacy and safety of lixisenatide once-daily morning or evening injections in type 2 diabetes inadequately controlled on metformin (GetGoal-M). *Diabetes Care* 2013; **36**(9): 2543-50.
- 46. Barnett AH, Charbonnel B, Li J, Donovan M, Fleming D, Iqbal N. Saxagliptin add-on therapy to insulin with or without metformin for type 2 diabetes mellitus: 52-week safety and efficacy. *Clin Drug Investig* 2013; **33**(10): 707-17.
- 47. Barnett AH, Huisman H, Jones R, von Eynatten M, Patel S, Woerle H-J. Linagliptin for patients aged 70 years or older with type 2 diabetes inadequately controlled with common antidiabetes treatments: a randomised, double-blind, placebo-controlled trial. *The Lancet* 2013; **382**(9902): 1413-23.
- 48. Dobs AS, Goldstein BJ, Aschner P, et al. Efficacy and safety of sitagliptin added to ongoing metformin and rosiglitazone combination therapy in a randomized placebo-controlled 54-week trial in patients with type 2 diabetes. *J Diabetes* 2013; 5(1): 68-79.
- 49. Haring HU, Merker L, Seewaldt-Becker E, et al. Empagliflozin as add-on to metformin plus sulfonylurea in patients with type 2 diabetes: a 24-week, randomized, double-blind, placebo-controlled trial. *Diabetes Care* 2013; **36**(11): 3396-404.
- 50. Lavalle-Gonzalez FJ, Januszewicz A, Davidson J, et al. Efficacy and safety of canagliflozin compared with placebo and sitagliptin in patients with type 2 diabetes on background metformin monotherapy: a randomised trial. *Diabetologia* 2013; **56**(12): 2582-92.
- 51. McGill JB, Sloan L, Newman J, et al. Long-term efficacy and safety of linagliptin in patients with type 2 diabetes and severe renal impairment: a 1-year, randomized, double-blind, placebo-controlled study. *Diabetes Care* 2013; **36**(2): 237-44.
- 52. Pinget M, Goldenberg R, Niemoeller E, Muehlen-Bartmer I, Guo H, Aronson R. Efficacy and safety of lixisenatide once daily versus placebo in type 2 diabetes insufficiently controlled on pioglitazone (GetGoal-P). *Diabetes Obes Metab* 2013; **15**(11): 1000-7.
- 53. Riddle MC, Aronson R, Home P, et al. Adding once-daily lixisenatide for type 2 diabetes inadequately controlled by established basal insulin: a 24-week, randomized, placebo-controlled comparison (GetGoal-L). *Diabetes Care* 2013; **36**(9): 2489-96.

- 54. Riddle MC, Forst T, Aronson R, et al. Adding once-daily lixisenatide for type 2 diabetes inadequately controlled with newly initiated and continuously titrated basal insulin glargine: a 24-week, randomized, placebo-controlled study (GetGoal-Duo 1). *Diabetes Care* 2013; **36**(9): 2497-503.
- 55. Roden M, Weng J, Eilbracht J, et al. Empagliflozin monotherapy with sitagliptin as an active comparator in patients with type 2 diabetes: A randomised, double-blind, placebo-controlled, phase 3 trial. *The Lancet Diabetes and Endocrinology*, 2013. (accessed.
- 56. Stenlof K, Cefalu WT, Kim KA, et al. Efficacy and safety of canagliflozin monotherapy in subjects with type 2 diabetes mellitus inadequately controlled with diet and exercise. *Diabetes, Obesity and Metabolism*, 2013. (accessed.
- 57. Wilding JP, Charpentier G, Hollander P, et al. Efficacy and safety of canagliflozin in patients with type 2 diabetes mellitus inadequately controlled with metformin and sulphonylurea: a randomised trial. *Int J Clin Pract* 2013; **67**(12): 1267-82.
- 58. Yki-Jarvinen H, Rosenstock J, Duran-Garcia S, et al. Effects of adding linagliptin to basal insulin regimen for inadequately controlled type 2 diabetes: a >/=52-week randomized, double-blind study. *Diabetes Care* 2013; **36**(12): 3875-81.
- 59. Ahren B, Johnson SL, Stewart M, et al. HARMONY 3: 104-week randomized, double-blind, placebo- and active-controlled trial assessing the efficacy and safety of albiglutide compared with placebo, sitagliptin, and glimepiride in patients with type 2 diabetes taking metformin. *Diabetes Care* 2014; **37**(8): 2141-8.
- 60. Bajaj M, Gilman R, Patel S, Kempthorne-Rawson J, Lewis-D'Agostino D, Woerle HJ. Linagliptin improved glycaemic control without weight gain or hypoglycaemia in patients with type 2 diabetes inadequately controlled by a combination of metformin and pioglitazone: a 24-week randomized, double-blind study. *Diabet Med* 2014; **31**(12): 1505-14.
- 61. Barnett AH, Mithal A, Manassie J, et al. Efficacy and safety of empagliflozin added to existing antidiabetes treatment in patients with type 2 diabetes and chronic kidney disease: a randomised, double-blind, placebo-controlled trial. *Lancet Diabetes Endocrinol* 2014; **2**(5): 369-84.
- 62. Bolli GB, Munteanu M, Dotsenko S, et al. Efficacy and safety of lixisenatide once daily vs. placebo in people with Type 2 diabetes insufficiently controlled on metformin (GetGoal-F1). *Diabet Med* 2014; **31**(2): 176-84.
- 63. Buse JB, Han J, Miller S, MacConell L, Pencek R, Wintle M. Addition of exenatide BID to insulin glargine: a post-hoc analysis of the effect on glycemia and weight across a range of insulin titration. *Curr Med Res Opin* 2014; **30**(7): 1209-18.
- 64. Ferdinand KC, White WB, Calhoun DA, et al. Effects of the once-weekly glucagon-like peptide-1 receptor agonist dulaglutide on ambulatory blood

- pressure and heart rate in patients with type 2 diabetes mellitus. *Hypertension* 2014; **64**(4): 731-7.
- 65. Forst T, Guthrie R, Goldenberg R, et al. Efficacy and safety of canagliflozin over 52 weeks in patients with type 2 diabetes on background metformin and pioglitazone. *Diabetes, Obesity and Metabolism* 2014; **16**(5): 467-77.
- 66. Ji L, Ma J, Li H, et al. Dapagliflozin as monotherapy in drug-naive Asian patients with type 2 diabetes mellitus: a randomized, blinded, prospective phase III study. *Clin Ther* 2014; **36**(1): 84-100 e9.
- 67. Kohan DE, Fioretto P, Tang W, List JF. Long-term study of patients with type 2 diabetes and moderate renal impairment shows that dapagliflozin reduces weight and blood pressure but does not improve glycemic control. *Kidney Int* 2014; **85**(4): 962-71.
- 68. McGill JB, Barnett AH, Lewin AJ, et al. Linagliptin added to sulphonylurea in uncontrolled type 2 diabetes patients with moderate-to-severe renal impairment. *Diab Vasc Dis Res* 2014; **11**(1): 34-40.
- 69. Pratley RE, Fleck P, Wilson C. Efficacy and safety of initial combination therapy with alogliptin plus metformin versus either as monotherapy in drug-naive patients with type 2 diabetes: A randomized, double-blind, 6-month study. *Diabetes, obesity & metabolism* 2014; **16**(7): 613-21.
- 70. Reusch J, Stewart M, Perkins C, et al. Efficacy and safety of once-weekly glucagon-like peptide 1 receptor agonist albiglutide (HARMONY 1 trial): 52-week primary endpoint results from a randomized, double-blind, placebo-controlled trial in patients with type 2 diabetes mellitus not controlled on pioglitazone, with or without metformin. *Diabetes, Obesity and Metabolism* 2014; 16(12): 1257-64.
- 71. Rosenstock J, Hanefeld M, Shamanna P, et al. Beneficial effects of once-daily lixisenatide on overall and postprandial glycemic levels without significant excess of hypoglycemia in type 2 diabetes inadequately controlled on a sulfonylurea with or without metformin (GetGoal-S). *J Diabetes Complications* 2014; **28**(3): 386-92.
- 72. Rosenstock J, Jelaska A, Frappin G, et al. Improved glucose control with weight loss, lower insulin doses, and no increased hypoglycemia with empagliflozin added to titrated multiple daily injections of insulin in obese inadequately controlled type 2 diabetes. *Diabetes Care* 2014; **37**(7): 1815-23.
- 73. Thrasher J, Daniels K, Patel S, Whetteckey J, Woerle HJ. Efficacy and Safety of Linagliptin in Black/African American Patients with Type 2 Diabetes: A 6-month, Randomized, Double-blind, Placebo-controlled Study. *Endocr Pract* 2014; **20**(5): 412-20.
- 74. Weir MR, Kline I, Xie J, Edwards R, Usiskin K. Effect of canagliflozin on serum electrolytes in patients with type 2 diabetes in relation to estimated glomerular

- filtration rate (eGFR). Curr Med Res Opin 2014; 30(9): 1759-68.
- 75. Wysham C, Blevins T, Arakaki R, et al. Efficacy and safety of dulaglutide added onto pioglitazone and metformin versus exenatide in type 2 diabetes in a randomized controlled trial (AWARD-1). *Diabetes Care* 2014; **37**(8): 2159-67.
- 76. Yu Pan C, Han P, Liu X, et al. Lixisenatide treatment improves glycaemic control in Asian patients with type 2 diabetes mellitus inadequately controlled on metformin with or without sulfonylurea: a randomized, double-blind, placebo-controlled, 24-week trial (GetGoal-M-Asia). *Diabetes/Metabolism Research and Reviews* 2014; **30**(8): 726-35.
- 77. Ahmann A, Rodbard HW, Rosenstock J, et al. Efficacy and safety of liraglutide versus placebo added to basal insulin analogues (with or without metformin) in patients with type 2 diabetes: a randomized, placebo-controlled trial. *Diabetes Obes Metab* 2015; **17**(11): 1056-64.
- 78. Chen Y, Ning G, Wang C, et al. Efficacy and safety of linagliptin monotherapy in Asian patients with inadequately controlled type 2 diabetes mellitus: A multinational, 24-week, randomized, clinical trial. *Journal of Diabetes Investigation*, 2015. (accessed.
- 79. Davies MJ, Bergenstal R, Bode B, et al. Efficacy of Liraglutide for Weight Loss Among Patients With Type 2 Diabetes: The SCALE Diabetes Randomized Clinical Trial. *Jama* 2015; **314**(7): 687-99.
- 80. Haering HU, Merker L, Christiansen AV, et al. Empagliflozin as add-on to metformin plus sulphonylurea in patients with type 2 diabetes. *Diabetes Res Clin Pract* 2015; **110**(1): 82-90.
- 81. Home P, Shamanna P, Stewart M, et al. Efficacy and tolerability of albiglutide versus placebo or pioglitazone over 1 year in people with type 2 diabetes currently taking metformin and glimepiride: HARMONY 5. *Diabetes, Obesity and Metabolism* 2015; **17**(2): 179-87.
- 82. Mathieu C, Shankar RR, Lorber D, et al. A Randomized Clinical Trial to Evaluate the Efficacy and Safety of Co-Administration of Sitagliptin with Intensively Titrated Insulin Glargine. *Diabetes Ther* 2015; **6**(2): 127-42.
- 83. Matthaei S, Catrinoiu D, Celinski A, et al. Randomized, Double-Blind Trial of Triple Therapy With Saxagliptin Add-on to Dapagliflozin Plus Metformin in Patients With Type 2 Diabetes. *Diabetes Care* 2015; **38**(11): 2018-24.
- 84. Pi-Sunyer X, Astrup A, Fujioka K, et al. A randomized, controlled trial of 3.0 mg of liraglutide in weight management. *The New England Journal of Medicine* 2015; **373**(1): 11-22.
- 85. Rosenstock J, Jelaska A, Zeller C, et al. Impact of empagliflozin added on to basal insulin in type 2 diabetes inadequately controlled on basal insulin: a 78-week randomized, double-blind, placebo-controlled trial. *Diabetes Obes Metab* 2015;

- **17**(10): 936-48.
- 86. Yang H, Min K, Park S, et al. A randomized, placebo-controlled, double-blind, phase 3 trial to evaluate the efficacy and safety of anagliptin in drug-naive patients with type 2 diabetes. *Endocrine journal*, 2015. (accessed.
- 87. Aroda VR, Rosenstock J, Wysham C, et al. Efficacy and Safety of LixiLan, a Titratable Fixed-Ratio Combination of Insulin Glargine Plus Lixisenatide in Type 2 Diabetes Inadequately Controlled on Basal Insulin and Metformin: The LixiLan-L Randomized Trial. *Diabetes Care* 2016; **39**(11): 1972-80.
- 88. Davies MJ, Bain SC, Atkin SL, et al. Efficacy and Safety of Liraglutide Versus Placebo as Add-on to Glucose-Lowering Therapy in Patients With Type 2 Diabetes and Moderate Renal Impairment (LIRA-RENAL): A Randomized Clinical Trial. *Diabetes Care* 2016; **39**(2): 222-30.
- 89. Dungan K, Weitgasser R, Perez Manghi F, et al. A 24-week study to evaluate the efficacy and safety of once-weekly dulaglutide added on to glimepiride in type 2 diabetes (AWARD-8). *Diabetes, Obesity and Metabolism* 2016; **18**(5): 475-82.
- 90. Ji L, Han P, Wang X, et al. Randomized clinical trial of the safety and efficacy of sitagliptin and metformin co-administered to Chinese patients with type 2 diabetes mellitus. *Journal of Diabetes Investigation* 2016; 7(5): 727-36.
- 91. Nauck M, Stewart M, Perkins C, et al. Efficacy and safety of once-weekly GLP-1 receptor agonist albiglutide (HARMONY 2): 52 week primary endpoint results from a randomised, placebo-controlled trial in patients with type 2 diabetes mellitus inadequately controlled with diet and exercise. *Diabetologia*, 2016. (accessed.
- 92. Oyama J, Murohara T, Kitakaze M, et al. The Effect of Sitagliptin on Carotid Artery Atherosclerosis in Type 2 Diabetes: The PROLOGUE Randomized Controlled Trial. *PLoS Med* 2016; **13**(6): e1002051.
- 93. Rodbard H, Lingvay I, Reed J, et al. Efficacy and safety of semaglutide onceweekly vs placebo as add-on to basal insulin alone or in combination with metformin in subjects with type 2 diabetes (SUSTAIN 5). Diabetologia; 2016: SPRINGER 233 SPRING ST, NEW YORK, NY 10013 USA; 2016. p. S364-S5.
- 94. Seino Y, Kaneko S, Fukuda S, et al. Combination therapy with liraglutide and insulin in Japanese patients with type 2 diabetes: A 36-week, randomized, double-blind, parallel-group trial. *J Diabetes Investig* 2016; 7(4): 565-73.
- 95. Takeda. Efficacy and Safety of Alogliptin and Metformin Fixed-Dose Combination in Participants With Type 2 Diabetes (NCT01890122). 2016. https://clinicaltrials.gov/ct2/show/study/NCT01890122 (accessed 11 March 2020).
- 96. Chacra A, Gantz I, Mendizabal G, et al. A randomised, double-blind, trial of the safety and efficacy of omarigliptin (a once-weekly DPP-4 inhibitor) in subjects

- with type 2 diabetes and renal impairment. Int J Clin Pract 2017; 71(6).
- 97. Gadde KM, Vetter ML, Iqbal N, Hardy E, Ohman P, investigators D-N-s. Efficacy and safety of autoinjected exenatide once-weekly suspension versus sitagliptin or placebo with metformin in patients with type 2 diabetes: The DURATION-NEO-2 randomized clinical study. *Diabetes Obes Metab* 2017; **19**(7): 979-88.
- 98. Groop PH, Cooper ME, Perkovic V, et al. Linagliptin and its effects on hyperglycaemia and albuminuria in patients with type 2 diabetes and renal dysfunction: the randomized MARLINA-T2D trial. *Diabetes Obes Metab* 2017; **19**(11): 1610-9.
- 99. Januzzi JL, Jr., Butler J, Jarolim P, et al. Effects of Canagliflozin on Cardiovascular Biomarkers in Older Adults With Type 2 Diabetes. *J Am Coll Cardiol* 2017; **70**(6): 704-12.
- 100. Neal B, Perkovic V, Mahaffey KW, et al. Canagliflozin and Cardiovascular and Renal Events in Type 2 Diabetes. *N Engl J Med* 2017; **377**(7): 644-57.
- 101. Shankar RR, Inzucchi SE, Scarabello V, et al. A randomized clinical trial evaluating the efficacy and safety of the once-weekly dipeptidyl peptidase-4 inhibitor omarigliptin in patients with type 2 diabetes inadequately controlled on metformin monotherapy. *Curr Med Res Opin* 2017; **33**(10): 1853-60.
- 102. Softeland E, Meier JJ, Vangen B, Toorawa R, Maldonado-Lutomirsky M, Broedl UC. Empagliflozin as Add-on Therapy in Patients With Type 2 Diabetes Inadequately Controlled With Linagliptin and Metformin: A 24-Week Randomized, Double-Blind, Parallel-Group Trial. *Diabetes Care* 2017; **40**(2): 201-9.
- 103. Sorli C, Harashima S-i, Tsoukas GM, et al. Efficacy and safety of once-weekly semaglutide monotherapy versus placebo in patients with type 2 diabetes (SUSTAIN 1): a double-blind, randomised, placebo-controlled, parallel-group, multinational, multicentre phase 3a trial. *Lancet Diabetes Endocrinol* 2017; **5**(4): 251-60.
- 104. Chen Y, Liu X, Li Q, et al. Saxagliptin add-on therapy in Chinese patients with type 2 diabetes inadequately controlled by insulin with or without metformin: Results from the SUPER study, a randomized, double-blind, placebo-controlled trial. *Diabetes Obes Metab* 2018; **20**(4): 1044-9.
- 105. Dagogo-Jack S, Liu J, Eldor R, et al. Efficacy and safety of the addition of ertugliflozin in patients with type 2 diabetes mellitus inadequately controlled with metformin and sitagliptin: The VERTIS SITA2 placebo-controlled randomized study. *Diabetes Obes Metab* 2018; **20**(3): 530-40.
- 106. Dou J, Ma J, Liu J, et al. Efficacy and safety of saxagliptin in combination with metformin as initial therapy in Chinese patients with type 2 diabetes: Results from the START study, a multicentre, randomized, double-blind, active-controlled, phase 3 trial. *Diabetes Obes Metab* 2018; **20**(3): 590-8.

- 107. Grunberger G, Camp S, Johnson J, et al. Ertugliflozin in Patients with Stage 3 Chronic Kidney Disease and Type 2 Diabetes Mellitus: The VERTIS RENAL Randomized Study. *Diabetes Ther* 2018; **9**(1): 49-66.
- 108. Guja C, Frias JP, Somogyi A, et al. Effect of exenatide QW or placebo, both added to titrated insulin glargine, in uncontrolled type 2 diabetes: The DURATION-7 randomized study. *Diabetes Obes Metab* 2018; **20**(7): 1602-14.
- 109. Miller S, Krumins T, Zhou H, et al. Ertugliflozin and Sitagliptin Co-initiation in Patients with Type 2 Diabetes: The VERTIS SITA Randomized Study. *Diabetes Ther* 2018; **9**(1): 253-68.
- 110. Yang W, Ma J, Li Y, et al. Dapagliflozin as add-on therapy in Asian patients with type 2 diabetes inadequately controlled on insulin with or without oral antihyperglycemic drugs: A randomized controlled trial. *J Diabetes* 2018; **10**(7): 589-99.
- 111. Aroda VR, Rosenstock J, Terauchi Y, et al. PIONEER 1: Randomized Clinical Trial of the Efficacy and Safety of Oral Semaglutide Monotherapy in Comparison With Placebo in Patients With Type 2 Diabetes. *Diabetes Care* 2019; **42**(9): 1724-32.
- 112. Pratley R, Amod A, Hoff ST, et al. Oral semaglutide versus subcutaneous liraglutide and placebo in type 2 diabetes (PIONEER 4): a randomised, doubleblind, phase 3a trial. *Lancet (London, England)* 2019; **394**(10192): 39-50.
- 113. Wiviott SD, Raz I, Bonaca MP, et al. Dapagliflozin and Cardiovascular Outcomes in Type 2 Diabetes. *N Engl J Med* 2019; **380**(4): 347-57.
- 114. Zinman B, Aroda VR, Buse JB, et al. Efficacy, safety, and tolerability of oral semaglutide versus placebo added to insulin with or without metformin in patients with type 2 diabetes: the PIONEER 8 trial. *Diabetes care* 2019; **42**(12): 2262-71.
- 115. Zinman B, Bhosekar V, Busch R, et al. Semaglutide once weekly as add-on to SGLT-2 inhibitor therapy in type 2 diabetes (SUSTAIN 9): a randomised, placebo-controlled trial. *Lancet Diabetes Endocrinol* 2019; **7**(5): 356-67.
- 116. Blonde L, Belousova L, Fainberg U, et al. Liraglutide as add-on to sodium-glucose co-transporter-2 inhibitors in patients with inadequately controlled type 2 diabetes: LIRA-ADD2SGLT2i, a 26-week, randomized, double-blind, placebo-controlled trial. *Diabetes, Obesity & Metabolism* 2020; **22**(6): 929-37.
- 117. Cho YM, Deerochanawong C, Seekaew S, et al. Efficacy and safety of gemigliptin as add-on therapy to insulin, with or without metformin, in patients with type 2 diabetes mellitus (ZEUS II study). *Diabetes, obesity & metabolism* 2020; **22**(1): 123-7.
- 118. Halvorsen YD, Lock JP, Freeman MW. Cardiovascular and glycemic benefits of bexagliflozin as an adjunct to metformin for the treatment of type 2 diabetes in adults: a 24-week, randomized, double-blind, placebo-controlled trial. *Diabetes*

- 2020; 69.
- 119. Sone H, Kaneko T, Shiki K, et al. Efficacy and safety of empagliflozin as add-on to insulin in Japanese patients with type 2 diabetes: a randomized, double-blind, placebo-controlled trial. *Diabetes, obesity & metabolism* 2020; **22**(3): 417-26.
- 120. Cherney DZI, Ferrannini E, Umpierrez GE, et al. Efficacy and safety of sotagliflozin in patients with type 2 diabetes and severe renal impairment. *Diabetes, Obesity and Metabolism* 2021; **23**(12): 2632-42.
- 121. Cioffi G, Giorda CB, Lucci D, et al. Effects of linagliptin on left ventricular DYsfunction in patients with type 2 DiAbetes and concentric left ventricular geometry: results of the DYDA 2 trial. *European journal of preventive cardiology* 2021; **28**(1): 8-17.
- 122. Davies M, Færch L, Jeppesen OK, et al. Semaglutide 2·4 mg once a week in adults with overweight or obesity, and type 2 diabetes (STEP 2): a randomised, double-blind, double-dummy, placebo-controlled, phase 3 trial. *Lancet (London, England)* 2021; **397**(10278): 971-84.
- 123. Ji L, Li L, Ma J, et al. Efficacy and safety of teneligliptin added to metformin in Chinese patients with type 2 diabetes mellitus inadequately controlled with metformin: A phase 3, randomized, double-blind, placebo-controlled study. *Endocrinol* 2021; 4(2): e00222.
- 124. Kaku K, Kadowaki T, Seino Y, et al. Efficacy and safety of ipragliflozin in Japanese patients with type 2 diabetes and inadequate glycaemic control on sitagliptin. *Diabetes, Obesity & Metabolism* 2021; **23**(9): 2099-108.
- 125. Nahra R, Wang T, Gadde KM, et al. Effects of Cotadutide on Metabolic and Hepatic Parameters in Adults With Overweight or Obesity and Type 2 Diabetes: A 54-Week Randomized Phase 2b Study. *Diabetes Care* 2021; **44**(6): 1433-42.
- 126. Rosenstock J, Wysham C, Frías JP, et al. Efficacy and safety of a novel dual GIP and GLP-1 receptor agonist tirzepatide in patients with type 2 diabetes (SURPASS-1): a double-blind, randomised, phase 3 trial. *Lancet (London, England)* 2021; **398**(10295): 143-55.
- 127. Weng J, Zeng L, Zhang Y, et al. Henagliflozin as add-on therapy to metformin in patients with type 2 diabetes inadequately controlled with metformin: a multicentre, randomized, double-blind, placebo-controlled, phase 3 trial. *Diabetes, obesity & metabolism* 2021; 23(8): 1754-64.
- 128. Akasaka H, Sugimoto K, Shintani A, et al. Effects of ipragliflozin on left ventricular diastolic function in patients with type 2 diabetes and heart failure with preserved ejection fraction: the EXCEED randomized controlled multicenter study. *Geriatrics & gerontology international* 2022; **22**(4): 298-304.
- 129. Dahl D, Onishi Y, Norwood P, et al. Effect of Subcutaneous Tirzepatide vs Placebo Added to Titrated Insulin Glargine on Glycemic Control in Patients With

- Type 2 Diabetes: The SURPASS-5 Randomized Clinical Trial. *Jama* 2022; **327**(6): 534-45.
- 130. Wada T, Mori-Anai K, Takahashi A, et al. Effect of canagliflozin on the decline of estimated glomerular filtration rate in chronic kidney disease patients with type 2 diabetes mellitus: a multicenter, randomized, double-blind, placebo-controlled, parallel-group, phase 3 study in Japan. *Journal of diabetes investigation* 2022.

Supplemental Table S1: Search strategy on Medline (Ovid).

#	Searches
1	exp diabetes mellitus, type 2/
2	diabet*.ab.
3	non-insulin dependent.ab.
4	type-2.ab.
5	type II.ab.
6	type 2.ab.
7	4 or 5 or 6
8	2 and 7
9	1 or 8
10	exp thiazolidinediones/
11	exp glipizide/
12	exp glyburide/
13	exp metformin/
14	exp acarbose/
15	thiazolidinedione*.ab.
16	pioglitazone.ab.
17	rosiglitazone.ab.
18	sulfonylurea*.ab.
19	sulphonylurea*.ab.
20	glipizide.ab.
21	glyburide.ab.
22	glimepiride.ab.
23	biguanide*.ab.
24	metformin.ab.
25	insulin secretagogues.ab.
26	meglitinide*.ab.
27	repaglinide.ab.
28	nateglinide.ab.

29 alpha-glucosidase inhibitors.ab. 30 alpha-glucosidase inhibitor.ab. 31 acarbose.ab. 32 exp Dipeptidyl-Peptidase IV Inhibitors/ 33 sitagliptin*.ab. 34 saxagliptin*.ab. 35 dpp-4.ab. 36 dpp-iv.ab.	
31 acarbose.ab. 32 exp Dipeptidyl-Peptidase IV Inhibitors/ 33 sitagliptin*.ab. 34 saxagliptin*.ab. 35 dpp-4.ab.	
32 exp Dipeptidyl-Peptidase IV Inhibitors/ 33 sitagliptin*.ab. 34 saxagliptin*.ab. 35 dpp-4.ab.	
33 sitagliptin*.ab. 34 saxagliptin*.ab. 35 dpp-4.ab.	
34 saxagliptin*.ab. 35 dpp-4.ab.	
35 dpp-4.ab.	
36 dpp-iv.ab.	
37 exp bromocriptine/	
38 bromocriptine.ab.	
39 colesevelam.ab.	
40 exp Glucagon-Like Peptide 1/	
41 liraglutide.ab.	
42 exenatide.ab.	
43	
44 clinical trial.pt.	
45 randomized.ab.	
46 placebo.ab.	
47 randomly.ab.	
48 trial.ti.	
49 exp Clinical Trial/	
50 44 or 45 or 46 or 47 or 48 or 49	
51 Animals/	
52 Humans/	
53 51 and 52	
54 51 not 53	
55 50 not 54	
56 50 not 54	
57 9 and 43	

#	Searches
58	56 and 57
59	letter.pt.
60	comment.pt.
61	editorial.pt.
62	59 or 60 or 61
63	58 not 62

Supplemental Table S2: Characteristics of included trials.

Trial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Type 2 diabetes								
Coniff 1995	255	24	55.9	29.9	6.9	1.0	Not reported	Acarbose and tolbutamide
Johnston 1998	364	56	67.8	29.7	8.4	0.6	Not reported	Miglitol and glyburide
UK Prospective Diabetes Study (UKPDS) Group 1998, UKPDS 34	753	558	53.0	31.7	7.2	0.6	2.3%	Metformin
Buse 2004, Exenatide-113	377	30	55.3	33.3	8.6	0.8	1.4%	Exenatide
Bosi 2007, NCT00099892	544	24	54.2	32.7	8.4	0.9	0.0%	Vildagliptin
Goldstein 2007, SITAGLIPTIN 036, NCT00103857	1091	24	53.5	32.1	8.8	1.4	1.2%	Sitagliptin and metformin
Hermansen 2007, NCT00106704	441	52	56.0	31.0	8.3	0.7	0.9%	Sitagliptin
Mita 2007	78	52	61.6	23.6	6.0	0.3	Not reported	Nateglinide
DeFronzo 2009, SAXAGLIPTIN 014, NCT00121667	743	24	54.6	31.4	8.1	0.8	3.6%	Saxagliptin
Hollander 2009, CV181-013, NCT00295633	565	24	54.1	30.0	8.3	0.5	1.2%	Saxagliptin
Kooy 2009, HOME, NCT00375388	390	224	61.5	30.0	7.9	0.4	4.1%	Metformin
Pratley 2009, Alogliptin Study 009, NCT00286494	493	26	55.4	32.8	8.0	0.5	0.0%	Alogliptin
Pratley 2009, NCT00286468	500	26	56.6	30.1	8.1	0.5	0.0%	Alogliptin
Rosenstock 2009, NCT00121641	401	24	53.5	31.7	7.9	0.7	2.3%	Saxagliptin
Rosenstock 2009, NCT00286429	390	26	55.4	32.5	9.3	0.5	1.5%	Alogliptin
Bailey 2010, NCT00528879	546	24	53.9	31.5	8.1	0.4	6.3%	Dapagliflozin
Liutkus 2010, NCT00603239	165	26	54.7	33.7	8.2	0.7	0.0%	Exenatide
Truitt 2010, NCT00143520	441	26	55.7	32.8	8.1	0.8	Not reported	Rivoglitazone and pioglitazone
Vilsboll 2010, NCT00395343	641	24	57.8	31.0	8.7	0.6	1.4%	Sitagliptin
Barzilai 2011, NCT00305604	206	24	71.9	31.0	7.7	0.7	Not reported	Sitagliptin
Chacra 2011, CV181040, NCT00313313	768	24	55.1	29.1	8.4	0.7	4.9%	Saxagliptin

Trial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Del Prato 2011, NCT00621140	503	24	55.7	29.0	8.0	0.7	Not reported	Linagliptin
Gomis 2011, NCT00641043	389	24	57.5	29.0	8.6	0.5	Not reported	Linagliptin
Gram 2011, South Danish (metformin), NCT00121966	371	104	56.3	33.9	8.6	0.6	0.5%	Metformin
Gram 2011, South Danish (rosiglitazone), NCT00121966	371	104	56.3	33.9	8.6	0.6	0.3%	Rosiglitazone
Nowicki 2011, D1680C00007, NCT00614939	170	52	66.5	30.7	8.3	0.7	2.4%	Saxagliptin
Olansky 2011, NCT00482729	1246	44	49.7	33.3	9.9	0.5	0.4%	Sitagliptin
Owens 2011, NCT00602472	1055	24	58.1	28.3	8.1	0.6	0.0%	Linagliptin
Strojek 2011, NCT00680745	596	24	59.8	29.8	8.1	0.5	1.5%	Dapagliflozin
Taskinen 2011, NCT00601250	700	24	56.5	29.9	8.1	0.6	0.0%	Linagliptin
Chou 2012, NCT00484198	1912	26	55.1	29.8	7.7	0.8	0.0%	Rivoglitazone and pioglitazone
DeFronzo 2012, NCT00328627	1554	26	54.4	31.2	8.5	1.3	1.6%	Alogliptin
Haak 2012, NCT00798161	791	24	55.3	29.1	8.7	1.1	0.0%	Metformin and linagliptin
Pan 2012, NCT00698932	568	24	51.4	25.9	8.1	0.5	0.0%	Saxagliptin
Rosenstock 2012, NCT00683878	420	24	53.5	Not reported	8.4	0.5	0.0%	Dapagliflozin
Wilding 2012, Dapagliflozin 006, NCT00673231	807	24	59.3	33.1	8.5	0.5	1.1%	Dapagliflozin
Ahren 2013, GetGoal-M, NCT00712673	680	24	54.7	32.9	8.1	0.5	2.6%	Lixisenatide
Barnett 2013, NCT00757588	455	54	57.2	32.3	8.7	0.4	0.6%	Saxagliptin
Barnett 2013, NCT01084005	241	24	74.9	29.7	7.8	0.6	0.0%	Linagliptin
Dobs 2013, NCT00350779	278	54	54.5	30.3	8.8	0.8	0.0%	Sitagliptin
Haring 2013, EMPA-REG METSU, NCT01159600	666	24	57.1	28.2	8.1	0.6	1.9%	Empagliflozin
Lavalle-Gonzalez 2013, CANTATA-D, NCT01106677	1284	26	55.4	31.8	7.9	0.7	0.0%	Canagliflozin and sitagliptin
McGill 2013	133	52	64.4	32.0	8.2	0.7	4.6%	Linagliptin
Pinget 2013, GetGoal-P, NCT00763815	484	132	55.8	33.9	8.1	0.6	0.2%	Lixisenatide

Trial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Riddle 2013, GetGoal-Duo 1 and GetGoal-L, NCT00975286	446	24	56.0	31.8	7.6	0.3	1.9%	Lixisenatide
Riddle 2013, GetGoal-L, NCT00715624	495	24	57.0	32.1	8.4	0.4	3.9%	Lixisenatide
Roden 2013, EMPA-REG MONO, NCT01177813	899	24	55.0	28.4	7.9	0.8	9.5%	Empagliflozin and sitagliptin
Stenlof 2013, CANTATA-M, NCT01081834	584	26	55.4	31.6	8.0	1.0	Not reported	Canagliflozin
Wildling 2013, CANTATA-MSU, NCT01106625	469	52	56.8	33.1	8.1	0.9	0.6%	Canagliflozin
Yki-Jarvinen 2013, NCT00954447	1261	54	60.0	31.0	8.3	0.7	0.6%	Linagliptin
Ahren 2014, HARMONY 3, NCT00838903	1012	164	54.5	32.6	8.1	0.5	0.9%	Albiglutide, sitagliptin, and glimepiride
Bajaj 2014, NCT00996658	272	24	53.8	28.2	8.4	0.6	2.4%	Linagliptin
Barnett 2014, EMPA-REG RENAL, NCT01164501	738	52	63.9	30.6	8.1	0.2	0.9%	Empagliflozin
Bolli 2014, GetGoal-F1, NCT00763451	482	112	56.1	32.5	8.0	0.5	0.9%	Lixisenatide
Buse 2014, NCT00765817	259	30	59.0	33.5	8.4	0.7	1.4%	Exenatide
Ferdinand 2014, NCT01149421	755	26	56.5	33.0	7.9	0.9	0.8%	Dulaglutide
Forst 2014, CANTATA-MP, NCT01106690	342	26	57.4	32.5	7.9	0.7	Not reported	Canagliflozin
Ji 2014, NCT01095653	393	24	51.4	25.6	8.3	0.8	1.6%	Dapagliflozin
Kohan 2014, NCT00663260	252	104	67.0	Not reported	8.4	0.1	3.0%	Dapagliflozin
McGill 2014, NCT00800683	133	52	64.4	32.0	8.2	0.7	9.2%	Linagliptin
Pan 2014, GetGoal-M-Asia, NCT01169779	390	24	54.8	26.9	7.9	0.4	0.0%	Lixisenatide
Pratley 2014, NCT01023581	784	26	45.7	30.7	8.4	1.0	0.0%	Metformin and alogliptin
Reusch 2014, HARMONY 1, NCT00849056	301	156	55.0	34.1	8.1	0.8	0.2%	Albiglutide
Rosenstock 2014, EMPA-REG MDI, NCT01306214	563	52	56.7	34.8	8.3	0.4	0.5%	Empagliflozin
Rosenstock 2014, GetGoal-S, NCT00713830	859	52	57.3	30.2	8.3	0.7	1.1%	Lixisenatide
Thrasher 2014, NCT01194830	226	24	53.9	32.7	8.7	0.6	1.8%	Linagliptin
Weir 2014, NCT01064414	269	26	68.5	Not reported	Not reported	0.4	2.2%	Canagliflozin

Trial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Wysham 2014, AWARD-1, NCT01064687	976	26	55.6	33.3	8.1	0.8	0.0%	Dulaglutide and exenatide
Ahmann 2015, NN2211-3917, NCT01617434	450	26	58.4	32.3	8.3	1.2	0.0%	Liraglutide
Chen 2015, NCT01214239	300	24	54.4	25.4	8.0	0.5	2.2%	Linagliptin
Davies 2015, SCALE Diabetes, NCT01272232	846	56	54.9	37.2	7.9	0.9	0.9%	Liraglutide
Haering 2015, EMPA-REG EXTEND METSU, NCT01289990	666	76	57.1	28.2	8.1	0.7	0.9%	Empagliflozin
Home 2015, HARMONY 5, NCT00839527	663	56	55.2	32.2	8.2	1.0	2.4%	Albiglutide and pioglitazone
Mathieu 2015, NCT01462266	658	24	58.8	32.1	8.8	0.4	1.3%	Sitagliptin
Matthaei 2015, NCT01619059	315	52	54.6	31.4	7.9	0.4	0.6%	Saxagliptin
Rosenstock 2015, NCT01011868	494	82	58.8	32.2	8.2	0.5	0.4%	Empagliflozin
Yang 2015	109	24	56.2	25.0	7.1	0.7	Not reported	Anagliptin
Aroda 2016, LixiLan-L, NCT02058160	736	30	60.0	31.1	8.1	0.5	0.0%	Lixisenatide
Davies 2016, LIRA-RENAL, NCT01620489	277	26	67.2	33.9	8.0	0.7	1.5%	Liraglutide
Dungan 2016, AWARD-8, NCT01769378	299	24	57.8	31.2	8.4	1.3	0.0%	Dulaglutide
Ji 2016, NCT01076088	744	24	52.7	25.8	8.7	0.9	Not reported	Metformin and sitagliptin
Nauck 2016, HARMONY 2, NCT00849017	309	52	52.9	33.5	8.1	0.9	2.9%	Albiglutide
Oyama 2016, PROLOGUE, UMIN000004490	463	104	69.3	25.1	7.0	0.2	0.4%	Sitagliptin
Rodbard 2016, SUSTAIN-5, NCT02305381	397	30	58.8	Not reported	8.4	1.6	0.0%	Semaglutide
Seino 2016, NCT01572740	257	36	60.5	25.6	8.8	0.8	2.2%	Liraglutide
Takeda 2016, NCT01890122	647	28	53.6	26.3	Not reported	-0.9	Not reported	Alogliptin and metformin
Chacra 2017, NCT01698775	213	24	65.2	30.1	8.3	0.3	2.0%	Omarigliptin
Gadde 2017, DURATION-NEO-2, NCT01652729	364	28	53.7	31.8	8.5	0.6	3.1%	Exenatide and sitagliptin
Groop 2017, MARLINA-T2D, NCT01792518	360	24	60.6	28.4	7.8	0.6	0.0%	Linagliptin
Januzzi 2017, NCT01106651	666	104	63.7	31.6	7.7	0.6	0.7%	Canagliflozin

rial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Neal 2017, CANVAS (primary prevention), NCT01989754	3486	126	62.7	32.3	8.3	0.6	1.5%	Canagliflozin
Shankar 2017, NCT01755156	402	24	57.2	32.5	8.1	0.6	Not reported	Omarigliptin
Softeland 2017, NCT01734785	327	24	55.2	30.2	8.0	0.7	Not reported	Empagliflozin
Sorli 2017, SUSTAIN-1, NCT02054897	387	35	53.7	32.9	8.1	1.5	0.0%	Semaglutide
Chen 2018, SUPER, NCT02104804	462	24	59.1	26.2	8.5	0.6	0.0%	Saxagliptin
Dagogo-Jack 2018, VERTIS SITA2, NCT02036515	462	54	59.1	30.8	8.0	0.8	0.6%	Ertugliflozin
Dou 2018, START (Metformin), NCT02273050	423	24	50.1	26.6	9.4	0.9	1.0%	Metformin
Dou 2018, START (Saxagliptin), NCT02273050	417	24	50.1	26.6	9.4	0.2	2.1%	Saxagliptin
Grunberger 2018, VERTIS RENAL, NCT01986855	467	54	67.3	32.5	8.2	0.1	1.3%	Ertugliflozin
Guja 2018, DURATION-7, NCT02229383	461	28	57.7	33.7	8.5	0.7	0.8%	Exenatide
Miller 2018, VERTIS SITA, NCT02226003	291	28	55.6	32.2	8.9	1.2	1.9%	Ertugliflozin and sitagliptin
Yang 2018, NCT02096705	272	24	57.5	26.5	8.5	0.9	3.3%	Dapagliflozin
Aroda 2019, PIONEER 1, NCT02906930	703	31	55.0	31.8	8.0	0.9	0.9%	Semaglutide
Pratley 2019, PIONEER 4, NCT02863419	711	57	56.0	33.0	8.0	0.9	1.3%	Semaglutide and liraglutide
Wiviott 2019, DECLARE-TIMI 58 (primary prevention), NCT01730534	10186	219	64.7	32.0	8.3	0.2	1.2%	Dapagliflozin
Zinman 2019, PIONEER 8, NCT03021187	731	57	61.0	31.0	8.2	0.9	2.0%	Semaglutide
Zinman 2019, SUSTAIN 9, NCT03086330	302	30	57.0	31.9	8.0	1.4	Not reported	Semaglutide
Blonde 2020, LIRA-ADD2SGLT2i, NCT02964247	303	27	55.2	32.2	8.0	0.7	Not reported	Liraglutide
Cho 2020, ZEUS II, NCT02831361	283	24	60.4	26.7	8.4	0.7	0.0%	Gemigliptin
Halvorsen 2020, NCT03259789	317	26	55.8	29.8	Not reported	0.5	0.0%	Bexagliflozin
Sone 2020, NCT02589639	266	52	58.7	26.9	8.8	0.9	0.0%	Empagliflozin
Cherney 2021, SOTA-CKD4, NCT03242018	277	56	67.4	31.6	8.3	0.5	11.0%	Sotagliflozin
Cioffi 2021, DYDA 2, NCT02851745	188	48	69.0	30.0	6.4	0.2	1.1%	Linagliptin

Trial (first author, publication, name of trial, registration)	Sample size	Follow- up (week)*	Age (years, mean or median)	BMI (kg/m2, mean or median)	Baseline HbA _{1c} (%, mean)	Reduction in HbA _{1c} (%, mean)	One-year baseline CVD risk (%)	Drug used in the intervention group(s)
Davies 2021, STEP 2, NCT03552757	1210	75	55.0	35.7	8.1	1.2	0.5%	Semaglutide
Ji 2021, NCT02924064	246	24	55.3	26.1	7.9	0.7	1.8%	Teneligliptin
Kaku 2021, MK-0431J-843, NCT02577003	143	26	60.5	25.8	8.1	0.8	0.0%	Ipragliflozin
Nahra 2021, NCT03235050	834	54	56.8	35.0	8.1	0.7	0.9%	Cotadutide and liraglutide
Rosenstock 2021, SURPASS-1, NCT03954834	478	44	54.1	31.9	7.9	2.0	1.0%	Tirzepatide
Weng 2021, NCT04390295	483	24	54.8	25.5	8.5	0.8	0.0%	Henagliflozin
Akasaka 2022, EXCEED, UMIN000027095	68	24	71.1	25.3	8.0	0.1	6.8%	Ipragliflozin
Dahl 2022, SURPASS-5, NCT04039503	475	74	60.7	33.4	8.3	1.4	0.6%	Tirzepatide
Wada 2022, NCT03436693	308	104	62.5	26.9	7.8	0.2	1.0%	Canagliflozin
Intermediate hyperglycemia								
Chiasson 2003, STOP-NIDDM	1429	170	54.5	30.9	Not reported	Not reported	0.8%	Acarbose
DPP Research Group 2005, DPP	2155	155	50.6	34.1	5.9	-0.1	Not reported	Metformin
DREAM Trial Investigators 2006, DREAM, NCT00095654	5269	156	54.7	30.9	Not reported	Not reported	0.3%	Rosiglitazone
Ramachandran 2006, IDPP-1	398	129	45.9	25.8	6.2	Not reported	Not reported	Metformin
Kawamori 2009, Voglibose Ph-3 Study, UMIN000001109	1780	207	55.7	25.8	Not reported	Not reported	Not reported	Voglibose
Ramachandran 2009, IDPP-2, NCT00276497	407	155	45.3	26.1	5.8	0.1	Not reported	Pioglitazone
Zinman 2010, CANOE, NCT00116932	207	201	52.5	31.7	Not reported	Not reported	Not reported	Metformin and rosiglitazone
DeFronzo 2011, ACT NOW, NCT00220961	602	125	52.3	34.5	5.5	0.2	0.1%	Pioglitazone
Lindblad 2011, NANSY	274	191	60.0	29.7	4.9	Not reported	Not reported	Glimepiride
Pi-Sunyer 2015, SCALE Obesity and Prediabetes, NCT01272219	3731	56	45.1	38.3	5.6	0.2	0.1%	Liraglutide

Abbreviations: CVD, cardiovascular disease; BMI, body mass index; HbA_{1c} , hemoglobin A_{1c} .

*In studies reporting \leq 56 weeks of follow-up, this value represents the planned duration of treatment. In studies reporting \geq 56 weeks of follow-up this value represents the mean or median duration.

Supplemental Table S3: Risk of bias of included trials.

Type 2 diabetes Coniff 1995 Unclear Low Low Low Unclear Johnston 1998 UK Prospective Diabetes Study (UKPDS) Group 1998, UKPDS 34 Buse 2004, Exenatide-113 Unclear	reporting
Johnston 1998 UK Prospective Diabetes Study (UKPDS) Group 1998, UKPDS 34 Buse 2004, Exenatide-113 Unclear	
UK Prospective Diabetes Study (UKPDS) Group 1998, UKPDS 34 Buse 2004, Exenatide-113 Unclear Unclear Unclear Unclear Unclear Unclear Unclear High Goldstein 2007, NCT00099892 Unclear	Unclear
1998, UKPDS 34 Buse 2004, Exenatide-113 Unclear	Unclear
Bosi 2007, NCT00099892 Unclear Unclear Unclear Unclear High Goldstein 2007, SITAGLIPTIN 036,	Unclear
Goldstein 2007, SITAGLIPTIN 036,	Unclear
' Inclear Low Low Low Low	Unclear
NC100103837	Low
Hermansen 2007, NCT00106704 Low Low Low Unclear Low	Low
Mita 2007 Low Unclear Unclear Low Low	Unclear
DeFronzo 2009, SAXAGLIPTIN 014, NCT00121667 Low Low Unclear Unclear High	Low
Hollander 2009, CV181-013, NCT00295633 Low Low Unclear Low Unclear	Low
Kooy 2009, HOME, NCT00375388 Unclear Unclear Unclear Unclear Low	Low
Pratley 2009, Alogliptin Study 009, NCT00286494 Low Low Unclear Unclear Low	Low
Pratley 2009, NCT00286468 Unclear Unclear Low High High	Low
Rosenstock 2009, NCT00121641 Unclear Low Low Low Low	Low
Rosenstock 2009, NCT00286429 Low Low Unclear Unclear High	Low
Bailey 2010, NCT00528879 Low Low Low Low Low	Low
Liutkus 2010, NCT00603239 Unclear Unclear Unclear Unclear Low	Low
Truitt 2010, NCT00143520 Unclear Low Low Low Low	Low
Vilsboll 2010, NCT00395343 Low Unclear Unclear Low	

Barzilai 2011, NCT00305604	Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Del Prato 2011, NCT00621140	Barzilai 2011, NCT00305604	Low	Low	Low	Low	Low	Low
Comis 2011, NCT00641043 Cow Low Unclear Unclear Low Low Low Cram 2011, South Danish (metformin), NCT00121966 Unclear Unclear Unclear Unclear Unclear Unclear Low Low Low Crow	Chacra 2011, CV181040, NCT00313313	Low	Low	Low	Unclear	Low	Low
Gram 2011, South Danish (metformin), NCT00121966 Unclear Unclear Unclear Unclear Low Low NCT00121966 Unclear	Del Prato 2011, NCT00621140	Low	Low	Low	Low	Low	Low
NCT00121966 Gram 2011, South Danish (rosiglitazone), NCT00121966 Nowicki 2011, D1680C00007, NCT00614939 Low Low Unclear Unclear Unclear Unclear High Low Olansky 2011, NCT00482729 Unclear Unclear Unclear Unclear Unclear High Low Owens 2011, NCT00602472 Unclear Unclear Unclear Unclear Unclear Low Low Low Low Strojek 2011, NCT00680745 Low Unclear Unclear Unclear Unclear Unclear Low Low Low Low Chou 2012, NCT006484198 Unclear Uncl	Gomis 2011, NCT00641043	Low	Low	Unclear	Unclear	Low	Low
NCT00121966 Unclear Unclear Unclear Unclear Low Low Olansky 2011, NCT00482729 Unclear Unclear Unclear Unclear Unclear High Low Owens 2011, NCT00602472 Unclear Unclear Unclear Unclear Unclear Low	NCT00121966	Unclear	Unclear	Unclear	Unclear	Low	Low
Olansky 2011, NCT00482729 Unclear Unclear Low Unclear High Low Owens 2011, NCT00602472 Unclear Unclear Unclear Unclear Low Low Strojek 2011, NCT00680745 Low Unclear Low Low Low Taskinen 2011, NCT00601250 Unclear Unclear Unclear Unclear Unclear Low Chou 2012, NCT00484198 Unclear Low Low DeFronzo 2012, NCT00328627 Unclear Unclear Unclear Low Unclear		Unclear	Unclear	Unclear	Unclear	Low	Low
Owens 2011, NCT00602472 Unclear Unclear Unclear Low Low Low Strojek 2011, NCT00680745 Low Unclear Low Low Low Low Taskinen 2011, NCT00601250 Unclear Unclear Unclear Unclear Unclear Low	Nowicki 2011, D1680C00007, NCT00614939	Low	Low	Unclear	Unclear	High	Low
Strojek 2011, NCT00680745 Low Unclear Low Low Low Low Taskinen 2011, NCT00601250 Unclear Unclear Unclear Unclear Unclear Low Low Low DeFronzo 2012, NCT00328627 Unclear Unclear Low Unclear Unclear Unclear Haak 2012, NCT00798161 Unclear Low Low Low Low Dew Low Low Pan 2012, NCT00698932 Low Low Low Low Unclear U	Olansky 2011, NCT00482729	Unclear	Unclear	Low	Unclear	High	Low
Taskinen 2011, NCT00601250 Unclear Unclear Unclear Low Low Low Chou 2012, NCT00484198 Unclear Low Low Low Low Low Low Low Low Low Unclear Low Unclear Low Low <td>Owens 2011, NCT00602472</td> <td>Unclear</td> <td>Unclear</td> <td>Unclear</td> <td>Unclear</td> <td>Low</td> <td>Low</td>	Owens 2011, NCT00602472	Unclear	Unclear	Unclear	Unclear	Low	Low
Chou 2012, NCT00484198 Unclear Low Low Low Low Low Low DeFronzo 2012, NCT00328627 Unclear Unclear Low Haak 2012, NCT00798161 Unclear Low	Strojek 2011, NCT00680745	Low	Unclear	Low	Low	Low	Low
DeFronzo 2012, NCT00328627 Unclear Unclear Unclear Low Unclear Low Low Low Low Low Low Low Lo	Taskinen 2011, NCT00601250	Unclear	Unclear	Unclear	Unclear	Low	Low
Haak 2012, NCT00798161 Unclear Low Low Low Low Low Low Pan 2012, NCT00698932 Low Low Low Low Low Rosenstock 2012, NCT00683878 Unclear Unclear Unclear Unclear Low Wilding 2012, Dapagliflozin 006, NCT00673231 Low Low Low Ahren 2013, GetGoal-M, NCT00712673 Unclear	Chou 2012, NCT00484198	Unclear	Low	Low	Low	Low	Low
Pan 2012, NCT00698932 Low Low Low Low Low Low Low Low Wilding 2012, NCT00683878 Unclear Unclear Unclear Unclear Low Low Low Ahren 2013, GetGoal-M, NCT00712673 Unclear Unclear Unclear Unclear Unclear Unclear Low Low Low Barnett 2013, NCT00757588 Low Low Low High Low Low Low Low Dobs 2013, NCT00350779 Low Unclear Low Low High Low Low High Low	DeFronzo 2012, NCT00328627	Unclear	Unclear	Low	Unclear	Unclear	Low
Rosenstock 2012, NCT00683878 Unclear Low	Haak 2012, NCT00798161	Unclear	Low	Low	Low	Low	Low
Wilding 2012, Dapagliflozin 006, NCT00673231 Low	Pan 2012, NCT00698932	Low	Low	Low	Low	Low	Low
Ahren 2013, GetGoal-M, NCT00712673 Unclear Unclear Unclear Unclear Unclear Unclear Low Low Low Low Low Low Low Lo	Rosenstock 2012, NCT00683878	Unclear	Unclear	Unclear	Unclear	Low	Low
Barnett 2013, NCT00757588 Low Low High Low Low Low Dobs 2013, NCT01084005 Low Low Low Low Low Low High Low	Wilding 2012, Dapagliflozin 006, NCT00673231	Low	Low	Low	Low	Low	Low
Barnett 2013, NCT01084005 Low Low Low Low Low Low Dobs 2013, NCT00350779 Low Unclear Low Low High Low	Ahren 2013, GetGoal-M, NCT00712673	Unclear	Unclear	Unclear	Unclear	Low	Low
Dobs 2013, NCT00350779 Low Unclear Low Low High Low	Barnett 2013, NCT00757588	Low	Low	High	Low	Low	Low
	Barnett 2013, NCT01084005	Low	Low	Low	Low	Low	Low
	Dobs 2013, NCT00350779	Low	Unclear	Low	Low	High	Low
Training 2015, EIVIT A-NEG IVIE 150, NC 101159000 Low Low Unicidal Unicidal Low Low	Haring 2013, EMPA-REG METSU, NCT01159600	Low	Low	Unclear	Unclear	Low	Low

Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Lavalle-Gonzalez 2013, CANTATA-D, NCT01106677	Low	Unclear	Low	Low	Low	Low
McGill 2013	Unclear	Unclear	Low	Low	Low	Unclear
Pinget 2013, GetGoal-P, NCT00763815	Low	Low	Low	Low	Low	Low
Riddle 2013, GetGoal-Duo 1 and GetGoal-L, NCT00975286	Low	Low	Unclear	Low	Low	Low
Riddle 2013, GetGoal-L, NCT00715624	Low	Low	Low	Low	Unclear	Low
Roden 2013, EMPA-REG MONO, NCT01177813	Low	Low	Low	Low	Low	Low
Stenlof 2013, CANTATA-M, NCT01081834	Unclear	Low	Low	Low	Low	Low
Wildling 2013, CANTATA-MSU, NCT01106625	Low	Low	Low	Low	Low	Low
Yki-Jarvinen 2013, NCT00954447	Low	Low	Unclear	Low	Low	Low
Ahren 2014, HARMONY 3, NCT00838903	Unclear	Unclear	Low	Low	Low	Low
Bajaj 2014, NCT00996658	Low	Low	Unclear	Low	High	Low
Barnett 2014, EMPA-REG RENAL, NCT01164501	Low	Low	Low	Low	Low	Low
Bolli 2014, GetGoal-F1, NCT00763451	Unclear	Unclear	Low	Low	Low	Low
Buse 2014, NCT00765817	Low	Low	Low	Low	Low	Low
Ferdinand 2014, NCT01149421	Low	Unclear	Unclear	High	Low	Low
Forst 2014, CANTATA-MP, NCT01106690	Low	Low	Low	Low	Low	Low
Ji 2014, NCT01095653	Low	Low	Low	Low	Low	Low
Kohan 2014, NCT00663260	Unclear	Unclear	Unclear	Unclear	High	Low
McGill 2014, NCT00800683	Unclear	Unclear	Low	Low	Low	Low
Pan 2014, GetGoal-M-Asia, NCT01169779	Low	Low	Unclear	Unclear	Low	Low
Pratley 2014, NCT01023581	Unclear	Low	Low	Low	Low	Low
Reusch 2014, HARMONY 1, NCT00849056	Low	Low	Low	Low	High	Low

Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Rosenstock 2014, EMPA-REG MDI, NCT01306214	Low	Low	Unclear	Unclear	Low	Low
Rosenstock 2014, GetGoal-S, NCT00713830	Unclear	Unclear	Unclear	Unclear	Low	Low
Thrasher 2014, NCT01194830	Low	Unclear	Unclear	Unclear	Low	High
Weir 2014, NCT01064414	Unclear	Unclear	Low	Low	Unclear	Low
Wysham 2014, AWARD-1, NCT01064687	Low	Low	Low	Unclear	Low	Low
Ahmann 2015, NN2211-3917, NCT01617434	Unclear	Unclear	Low	Low	Unclear	Low
Chen 2015, NCT01214239	Low	Low	Low	Low	Low	Low
Davies 2015, SCALE Diabetes, NCT01272232	Low	Low	Low	Unclear	Low	Low
Haering 2015, EMPA-REG EXTEND METSU, NCT01289990	Unclear	Unclear	Unclear	Unclear	Low	Low
Home 2015, HARMONY 5, NCT00839527	Low	Low	Unclear	Low	Low	Low
Mathieu 2015, NCT01462266	Unclear	Unclear	Unclear	Unclear	Unclear	Low
Matthaei 2015, NCT01619059	Low	Low	Unclear	Unclear	Low	Low
Rosenstock 2015, NCT01011868	Low	Low	Unclear	Unclear	Low	Low
Yang 2015	Unclear	Low	Low	Low	Low	Unclear
Aroda 2016, LixiLan-L, NCT02058160	Low	Low	High	High	Low	Low
Davies 2016, LIRA-RENAL, NCT01620489	Low	Low	Low	Low	High	Low
Dungan 2016, AWARD-8, NCT01769378	Unclear	Unclear	Unclear	Unclear	Low	Low
Ji 2016, NCT01076088	Low	Low	Low	Low	Low	Low
Nauck 2016, HARMONY 2, NCT00849017	Unclear	Low	Low	Low	Low	Low
Oyama 2016, PROLOGUE, UMIN000004490	Low	Low	High	Low	Low	Low
Rodbard 2016, SUSTAIN-5, NCT02305381	Unclear	Unclear	Unclear	Unclear	Unclear	Low
Seino 2016, NCT01572740	Low	Low	Unclear	Unclear	Low	Low

Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Takeda 2016, NCT01890122	Unclear	Unclear	Unclear	Unclear	Low	Low
Chacra 2017, NCT01698775	Low	Low	Unclear	Unclear	Low	Low
Gadde 2017, DURATION-NEO-2, NCT01652729	Low	Low	Low	Low	High	Low
Groop 2017, MARLINA-T2D, NCT01792518	Low	Low	Low	Low	Low	Low
Januzzi 2017, NCT01106651	Low	Low	Low	Low	Low	Low
Neal 2017, CANVAS (primary prevention), NCT01989754	Low	Low	Low	Low	Low	Low
Shankar 2017, NCT01755156	Unclear	Unclear	Unclear	Unclear	Low	Low
Softeland 2017, NCT01734785	Low	Low	Low	Low	Low	Low
Sorli 2017, SUSTAIN-1, NCT02054897	Low	Low	Low	Low	Low	Low
Chen 2018, SUPER, NCT02104804	Unclear	Unclear	Unclear	Unclear	Low	Low
Dagogo-Jack 2018, VERTIS SITA2, NCT02036515	Low	Unclear	Low	Low	Low	Low
Dou 2018, START (Metformin), NCT02273050	Low	Low	Unclear	Unclear	Low	Low
Dou 2018, START (Saxagliptin), NCT02273050	Low	Low	Unclear	Unclear	Low	Low
Grunberger 2018, VERTIS RENAL, NCT01986855	Low	Low	Low	Low	Unclear	Low
Guja 2018, DURATION-7, NCT02229383	Low	Low	Low	Low	Low	Low
Miller 2018, VERTIS SITA, NCT02226003	Low	Low	Low	Low	Low	Low
Yang 2018, NCT02096705	Low	Low	Unclear	Unclear	Low	Low
Aroda 2019, PIONEER 1, NCT02906930	Low	Low	Low	Low	Low	Low
Pratley 2019, PIONEER 4, NCT02863419	Low	Low	Low	Low	Low	Low
Wiviott 2019, DECLARE-TIMI 58 (primary prevention), NCT01730534	Low	Low	Low	Low	Low	Low
Zinman 2019, PIONEER 8, NCT03021187	Low	Low	Low	Low	Low	Low

Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Zinman 2019, SUSTAIN 9, NCT03086330	Low	Low	Low	Low	Low	Low
Blonde 2020, LIRA-ADD2SGLT2i, NCT02964247	Low	Low	Low	Low	Low	Low
Cho 2020, ZEUS II, NCT02831361	Low	Low	Low	Low	Low	Low
Halvorsen 2020, NCT03259789	Low	Low	Low	Low	Low	Low
Sone 2020, NCT02589639	Low	Low	Unclear	Unclear	Low	Low
Cherney 2021, SOTA-CKD4, NCT03242018	Unclear	Unclear	Low	Low	Low	Low
Cioffi 2021, DYDA 2, NCT02851745	Low	Low	Low	Low	Low	Low
Davies 2021, STEP 2, NCT03552757	Low	Low	Low	Low	Low	Low
Ji 2021, NCT02924064	Low	Low	Low	Low	Low	Low
Kaku 2021, MK-0431J-843, NCT02577003	Low	Low	Low	Unclear	Low	Low
Nahra 2021, NCT03235050	Low	Low	Low	Low	Low	Low
Rosenstock 2021, SURPASS-1, NCT03954834	Low	Low	Low	Low	Low	Low
Weng 2021, NCT04390295	Low	Low	Low	Low	Low	Low
Akasaka 2022, EXCEED, UMIN000027095	Low	Unclear	Low	Low	Low	Low
Dahl 2022, SURPASS-5, NCT04039503	Low	Low	Low	Low	Low	Low
Wada 2022, NCT03436693	Unclear	Unclear	Unclear	Unclear	Low	Low
Intermediate hyperglycemia						
Chiasson 2003, STOP-NIDDM	Low	Low	Low	Low	Low	Low
DPP Research Group 2005, DPP	Unclear	Low	Low	Low	Unclear	Low
DREAM Trial Investigators 2006, DREAM, NCT00095654	Low	Low	Low	Low	Low	Low
Ramachandran 2006, IDPP-1	Unclear	Unclear	Unclear	Unclear	Low	Unclear
Kawamori 2009, Voglibose Ph-3 Study, UMIN00001109	Low	Low	Low	Low	Low	Low

Trial (first author, publication, name of trial, registration)	Random sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessment	Incomplete outcome data	Selective reporting
Ramachandran 2009, IDPP-2, NCT00276497	High	Low	Low	Low	Low	Low
Zinman 2010, CANOE, NCT00116932	Low	Low	Low	Low	Low	Low
DeFronzo 2011, ACT NOW, NCT00220961	Low	Low	Low	Low	Low	Low
Lindblad 2011, NANSY	Unclear	Low	Low	Low	Unclear	Unclear
Pi-Sunyer 2015, SCALE Obesity and Prediabetes, NCT01272219	Unclear	Low	Low	Low	Low	Low

Supplemental Table S4. Detailed results of meta-analysis of anti-diabetic drugs in preventing major cardiovascular events.

Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
UK Prospective Diabetes Study (UKPDS) Group 1998, UKPDS 34	52	342	103	411	0.5	0.4	0.8	7.3%
Buse 2004, Exenatide-113	1	254	1	123	0.5	0.0	7.8	0.3%
Bosi 2007, NCT00099892	1	362	0	182	1.5	0.1	37.4	0.3%
Goldstein 2007, SITAGLIPTIN 036, NCT00103857	4	915	1	176	0.8	0.1	6.9	0.5%
Hermansen 2007, NCT00106704	2	222	2	219	1.0	0.1	7.1	0.7%
DeFronzo 2009, SAXAGLIPTIN 014, NCT00121667	5	564	3	179	0.5	0.1	2.2	1.2%
Hollander 2009, CV181-013, NCT00295633	4	381	1	184	1.9	0.2	17.5	0.5%
Kooy 2009, HOME, NCT00375388	37	196	34	194	1.1	0.7	1.8	5.4%
Pratley 2009, Alogliptin Study 009, NCT00286494	3	396	0	97	1.7	0.1	33.9	0.3%
Pratley 2009, NCT00286468	1	401	0	99	0.7	0.0	18.4	0.3%
Rosenstock 2009, NCT00121641	3	306	1	95	0.9	0.1	9.1	0.5%
Rosenstock 2009, NCT00286429	0	260	1	130	0.2	0.0	4.1	0.3%
Bailey 2010, NCT00528879	3	409	4	137	0.2	0.1	1.1	1.1%
Liutkus 2010, NCT00603239	1	111	0	54	1.5	0.1	36.9	0.3%
Vilsboll 2010, NCT00395343	1	322	2	319	0.5	0.0	5.5	0.5%
Chacra 2011, CV181040, NCT00313313	3	501	6	267	0.3	0.1	1.1	1.3%
Gram 2011, South Danish (metformin), NCT00121966	3	184	2	187	1.5	0.3	9.3	0.8%
Gram 2011, South Danish (rosiglitazone), NCT00121966	4	187	1	184	4.0	0.4	36.1	0.5%
Nowicki 2011, D1680C00007, NCT00614939	4	85	2	85	2.0	0.4	11.5	0.9%

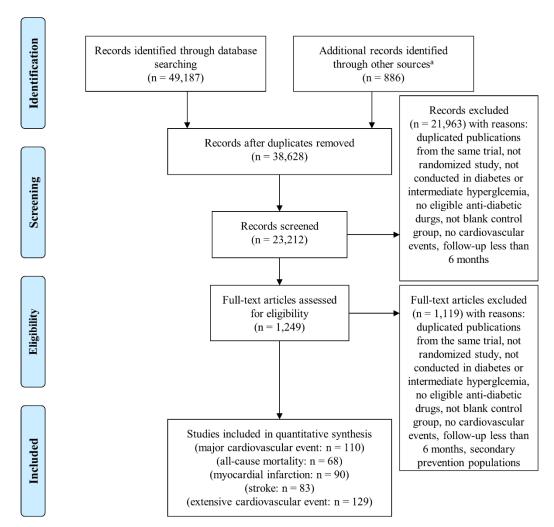
Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
Olansky 2011, NCT00482729	2	625	2	621	1.0	0.1	7.1	0.7%
Owens 2011, NCT00602472	1	792	0	263	1.0	0.0	24.6	0.3%
Strojek 2011, NCT00680745	3	450	1	146	1.0	0.1	9.4	0.5%
Taskinen 2011, NCT00601250	1	523	0	177	1.0	0.0	25.1	0.3%
Chou 2012, NCT00484198	3	1775	0	137	0.5	0.0	10.6	0.3%
DeFronzo 2012, NCT00328627	3	1425	1	129	0.3	0.0	2.6	0.5%
Haak 2012, NCT00798161	2	719	0	72	0.5	0.0	10.6	0.3%
Pan 2012, NCT00698932	2	284	0	284	5.0	0.2	105.4	0.3%
Rosenstock 2012, NCT00683878	1	281	0	139	1.5	0.1	36.9	0.3%
Wilding 2012, Dapagliflozin 006, NCT00673231	4	610	1	197	1.3	0.1	11.6	0.5%
Ahren 2013, GetGoal-M, NCT00712673	2	510	2	170	0.3	0.0	2.4	0.7%
Barnett 2013, NCT00757588	2	304	1	151	1.0	0.1	11.0	0.5%
Barnett 2013, NCT01084005	2	162	0	79	2.5	0.1	52.2	0.3%
Dobs 2013, NCT00350779	3	181	0	97	3.8	0.2	74.8	0.3%
Haring 2013, EMPA-REG METSU, NCT01159600	2	441	2	225	0.5	0.1	3.6	0.7%
Lavalle-Gonzalez 2013, CANTATA-D, NCT01106677	1	1101	0	183	0.5	0.0	12.3	0.3%
McGill 2013	5	68	3	65	1.6	0.4	7.2	1.1%
Pi-Sunyer 2015, SCALE Obesity and Prediabetes, NCT01272219	2	2487	2	1244	0.5	0.1	3.6	0.7%
Riddle 2013, GetGoal-Duo 1 and GetGoal-L, NCT00975286	1	223	2	223	0.5	0.0	5.5	0.5%
Riddle 2013, GetGoal-L, NCT00715624	2	328	3	167	0.3	0.1	2.0	0.8%
Roden 2013, EMPA-REG MONO, NCT01177813	12	671	10	228	0.4	0.2	0.9	2.9%

Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
Wildling 2013, CANTATA-MSU, NCT01106625	1	313	1	156	0.5	0.0	8.0	0.3%
Yki-Jarvinen 2013, NCT00954447	6	631	4	630	1.5	0.4	5.3	1.5%
Ahren 2014, HARMONY 3, NCT00838903	11	911	3	101	0.4	0.1	1.5	1.4%
Bajaj 2014, NCT00996658	0	183	1	89	0.2	0.0	4.0	0.3%
Barnett 2014, EMPA-REG RENAL, NCT01164501	3	419	3	319	0.8	0.2	3.8	1.0%
Bolli 2014, GetGoal-F1, NCT00763451	6	322	3	160	1.0	0.2	4.0	1.3%
Buse 2014, NCT00765817	0	137	1	122	0.3	0.0	7.3	0.3%
Ferdinand 2014, NCT01149421	1	505	1	250	0.5	0.0	7.9	0.3%
Ji 2014, NCT01095653	2	261	1	132	1.0	0.1	11.3	0.5%
Kohan 2014, NCT00663260	6	168	5	84	0.6	0.2	2.0	1.6%
McGill 2014, NCT00800683	5	68	6	65	0.8	0.2	2.7	1.6%
Pan 2014, GetGoal-M-Asia, NCT01169779	2	196	0	194	5.0	0.2	104.8	0.3%
Pratley 2014, NCT01023581	3	675	0	109	1.1	0.1	22.2	0.3%
Reusch 2014, HARMONY 1, NCT00849056	2	150	1	151	2.0	0.2	22.6	0.5%
Rosenstock 2014, EMPA-REG MDI, NCT01306214	2	375	1	188	1.0	0.1	11.1	0.5%
Rosenstock 2014, GetGoal-S, NCT00713830	7	574	3	285	1.2	0.3	4.5	1.3%
Thrasher 2014, NCT01194830	0	106	1	120	0.4	0.0	9.3	0.3%
Weir 2014, NCT01064414	0	179	1	90	0.2	0.0	4.1	0.3%
Wysham 2014, AWARD-1, NCT01064687	2	835	0	141	0.8	0.0	17.8	0.3%
Ahmann 2015, NN2211-3917, NCT01617434	1	225	0	225	3.0	0.1	74.4	0.3%
Chen 2015, NCT01214239	1	201	1	99	0.5	0.0	7.9	0.3%
Davies 2015, SCALE Diabetes, NCT01272232	5	634	2	212	0.8	0.2	4.3	0.9%

Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
Haering 2015, EMPA-REG EXTEND METSU, NCT01289990	4	441	3	225	0.7	0.2	3.1	1.1%
Home 2015, HARMONY 5, NCT00839527	7	548	3	115	0.5	0.1	1.9	1.3%
Mathieu 2015, NCT01462266	1	329	2	329	0.5	0.0	5.5	0.5%
Matthaei 2015, NCT01619059	0	153	1	162	0.4	0.0	8.7	0.3%
Rosenstock 2015, NCT01011868	7	324	1	170	3.7	0.5	30.6	0.6%
Aroda 2016, LixiLan-L, NCT02058160	3	367	0	369	7.1	0.4	137.9	0.3%
Davies 2016, LIRA-RENAL, NCT01620489	4	140	1	137	4.0	0.4	36.3	0.5%
Dungan 2016, AWARD-8, NCT01769378	1	239	0	60	0.8	0.0	18.9	0.3%
Nauck 2016, HARMONY 2, NCT00849017	1	204	3	105	0.2	0.0	1.6	0.5%
Oyama 2016, PROLOGUE, UMIN000004490	3	232	2	231	1.5	0.2	9.1	0.8%
Rodbard 2016, SUSTAIN-5, NCT02305381	1	264	0	133	1.5	0.1	37.6	0.3%
Seino 2016, NCT01572740	0	127	2	130	0.2	0.0	4.2	0.3%
Chacra 2017, NCT01698775	1	107	1	106	1.0	0.1	16.0	0.3%
Gadde 2017, DURATION-NEO-2, NCT01652729	1	303	1	61	0.2	0.0	3.2	0.3%
Groop 2017, MARLINA-T2D, NCT01792518	2	182	0	178	4.9	0.2	103.7	0.3%
Januzzi 2017, NCT01106651	9	450	3	216	1.4	0.4	5.4	1.4%
Neal 2017, CANVAS (primary prevention), NCT01989754	78	2039	54	1447	1.0	0.7	1.5	7.5%
Sorli 2017, SUSTAIN-1, NCT02054897	1	258	0	129	1.5	0.1	37.3	0.3%
Chen 2018, SUPER, NCT02104804	5	232	0	230	11.1	0.6	202.7	0.3%
Dagogo-Jack 2018, VERTIS SITA2, NCT02036515	1	309	1	153	0.5	0.0	7.9	0.3%
Dou 2018, START (Metformin), NCT02273050	0	210	1	213	0.3	0.0	8.3	0.3%
Dou 2018, START (Saxagliptin), NCT02273050	0	210	2	207	0.2	0.0	4.1	0.3%

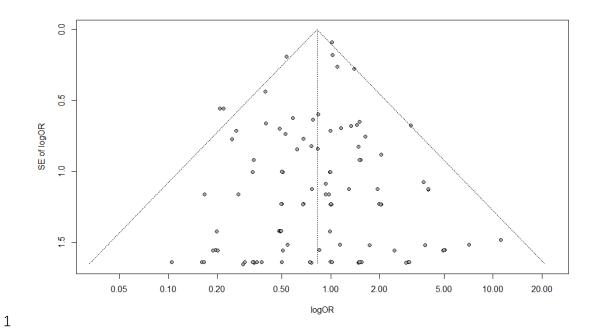
Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
Grunberger 2018, VERTIS RENAL, NCT01986855	6	313	2	154	1.5	0.3	7.4	1.0%
Guja 2018, DURATION-7, NCT02229383	0	231	1	230	0.3	0.0	8.2	0.3%
Miller 2018, VERTIS SITA, NCT02226003	0	194	1	97	0.2	0.0	4.1	0.3%
Yang 2018, NCT02096705	0	139	2	133	0.2	0.0	4.0	0.3%
Aroda 2019, PIONEER 1, NCT02906930	2	525	1	178	0.7	0.1	7.5	0.5%
Pratley 2019, PIONEER 4, NCT02863419	5	569	2	142	0.6	0.1	3.2	0.9%
Wiviott 2019, DECLARE-TIMI 58 (primary prevention), NCT01730534	273	5108	266	5078	1.0	0.9	1.2	10.1%
Zinman 2019, PIONEER 8, NCT03021187	10	547	4	184	0.8	0.3	2.7	1.7%
Cho 2020, ZEUS II, NCT02831361	1	188	0	95	1.5	0.1	37.9	0.3%
Halvorsen 2020, NCT03259789	1	158	0	159	3.0	0.1	75.1	0.3%
Sone 2020, NCT02589639	1	176	0	90	1.5	0.1	38.4	0.3%
Cherney 2021, SOTA-CKD4, NCT03242018	5	184	11	93	0.2	0.1	0.6	1.9%
Cioffi 2021, DYDA 2, NCT02851745	0	93	1	95	0.3	0.0	8.4	0.3%
Davies 2021, STEP 2, NCT03552757	8	807	3	403	1.3	0.4	5.1	1.4%
Ji 2021, NCT02924064	2	122	1	124	2.1	0.2	22.9	0.5%
Kaku 2021, MK-0431J-843, NCT02577003	1	73	0	70	2.9	0.1	72.8	0.3%
Nahra 2021, NCT03235050	6	722	1	112	0.9	0.1	7.8	0.6%
Rosenstock 2021, SURPASS-1, NCT03954834	0	363	1	115	0.1	0.0	2.6	0.3%
Weng 2021, NCT04390295	1	322	0	161	1.5	0.1	37.2	0.3%
Akasaka 2022, EXCEED, UMIN000027095	0	36	1	32	0.3	0.0	7.3	0.3%
Dahl 2022, SURPASS-5, NCT04039503	2	355	1	120	0.7	0.1	7.5	0.5%
Wada 2022, NCT03436693	9	154	3	154	3.1	0.8	11.8	1.4%
Chiasson 2003, STOP-NIDDM	4	714	18	715	0.2	0.1	0.6	1.9%

Trial (first author, publication, name of trial, registration)	Event number, interventi on group	Total number, interventi on group	Event number, control group	Total number, control group	Odds ratio	Lower bound of 95% CI	Upper bound of 95% CI	Weight (random- effects)
DREAM Trial Investigators 2006, DREAM, NCT00095654	32	2635	23	2634	1.4	0.8	2.4	5.2%
DeFronzo 2011, ACT NOW, NCT00220961	2	303	1	299	2.0	0.2	22.0	0.5%
Pinget 2013, GetGoal-P, NCT00763815	1	323	1	161	0.5	0.0	8.0	0.3%



^aReference list of potential relevant systematic reviews.

Supplemental Figure 1 Flow-diagram for literature search and screening.



Supplemental Figure 2 Funnel plot of anti-diabetic drugs in preventing major cardiovascular events.

1 Supplemental References

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- Moher D, Liberati A, Tetzlaff J, Altman DG; PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. BMJ. 2009 Jul 21;339:b2535. doi: 10.1136/bmj.b2535. PMID: 19622551; PMCID: PMC2714657.
- Higgins JP, Altman DG, Gøtzsche PC, Jüni P, Moher D, Oxman AD, Savovic J, Schulz KF, Weeks L, Sterne JA; Cochrane Bias Methods Group; Cochrane Statistical Methods Group. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. BMJ. 2011 Oct 18;343:d5928. doi: 10.1136/bmj.d5928. PMID: 22008217; PMCID: PMC3196245.
- Higgins, J. P., & Green, S. (Eds.). (2008). Cochrane handbook for systematic reviews of interventions.