

maintained treatment effect from long-term evidence in other T2DM populations. Responder were primarily defined using a composite end-point that based on an HbA1c  $\leq 7.0\%$  threshold AND no weight gain AND no documented symptomatic hypoglycemia. The NNT was calculated to determine the average number of patients needed to be treated in order to gain one additional successful responder. Sensitivity analyses were performed to examine the robustness of results. **Results:** For the primary composite end-point, cost per responder results were 136,290 CNY for lixisenatide group, 231,487 CNY for Basal-Plus group, and 222,424 CNY for Basal-Bolus group. The NNT analysis showed that there was approximately one additional responder for every 7.65 and 8.74 patients treated with lixisenatide combined with basal insulin compared to Basal-Plus and Basal-Bolus, respectively. The sensitivity analysis proved the robustness of results. **Conclusions:** Lixisenatide combined with basal insulin is a cost-effective treatment alternative compared with Basal-Plus and Basal-Bolus for T2DM patients inadequately controlled by basal insulin in China.

**PDB62  
INCIDENCE AND COST OF DIABETES RELATED EVENTS IN  
AN ADULT TYPE 1 DIABETES POPULATION FROM A U.S.  
PAYOR PERSPECTIVE**

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**Objectives:** Approximately  $\frac{3}{4}$  of adults with type 1 diabetes mellitus (T1DM) do not meet glycemic goals and therefore are at risk of developing diabetes-related com-



**PDB64  
ASSOCIATION BETWEEN CARDIOVASCULAR DISEASE  
RISK FACTORS AND HYPOGLYCAEMIC EVENTS IN TYPE 2  
DIABETES USING THE IQVIA CORE DIABETES MODEL**

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**Objectives:** There is clinical evidence that having a hypoglycemic event increases the risk of cardiovascular disease and mortality. In version 9.5 of the IQVIA Core Diabetes Model (CDM), an association relative risk factor (RR) of developing Cardiovascular Disease (CVD) and CVD mortality events related to non-severe (NSHE) and severe (SHE) hypoglycemic events over an adjustable duration were included. CVD conditions are Myocardial Infarction (MI), Stroke, Congestive Heart Failure (CHF) and angina. The current study compared the model outcomes obtained with and without applying a variable RR. **Methods:** The observational EDGE study comparing metformin + vildagliptin (M+V) versus metformin + sulphonylurea (M+S) was used as a base case. Basal insulin rescue therapy was applied to both arms when an HbA1c level of 7.5% was reached. Mean age at study start was 57.8 years and duration of diabetes 6 years. NSHE and SHE rate were applied as 13.94 per 100 patient-years (100 pt/year), 2.46 (100 pt/year) for M+S and 1.785, 0.315 (100 pt/year) for M+V. A RR of 1.1 was applied for all CVD conditions over a five-year period. UK 2018 costs were applied; costs and health benefits were discounted at 3.5%. **Results:** Using RR increase of 1.1 for CVD and mortality due to SHE and NSHE, changed life expectancy (LE) and QALY with -1.68 and -1.55% (LY: from 11.961 to 11.760; QALY: 8.335 to 8.91) for M+V and -1.55 and -1.50% (LY: from 11.959 to 11.716; QALY: 8.217 to 8.052) for M+S. LEER decreased with -77% (66.336 to 15.086). Small increase in life expectancy CVD



