

## Background and objective

- Methodological concerns in previous cost-effectiveness analyses (CEAs) of glucagon-like peptide-1 receptor agonists (GLP-1RAs) vs. long-acting insulins (LAIs) (e.g., using trial data as input) limit the applicability of study findings to real-world patients with type 2 diabetes (T2D).
- We conducted a **model-based CEA** of GLP-1RAs vs. LAIs from a healthcare sector perspective using **real-world, population-based data of Taiwanese T2D patients** to inform decision-makings.

## Methods

- Population: T2D patients initiating a GLP-1RA or LAI
- Model settings: Markov model (**Figure 1**), a 10-year simulation horizon and 3% annual discount rate on both costs and quality-adjusted life years (QALYs).
- Cost and health utility parameters:
  - Derived from **published literature of Taiwanese T2D patients**
  - Adjusted by patient characteristics** of the overall study cohort and subgroup populations in base-case and subgroup analyses, respectively.
- Transition probabilities:
  - LAI group: **cumulative risks of clinical events** measured from analyzing **Taiwan’s National Health Insurance Research Database (NHIRD) 2013-2019**
  - GLP-1RA group: modified cumulative risks of LAI group with the **relative hazards of clinical events** associated with using GLP-1RAs versus LAIs (measured from **propensity-score matched pairs of GLP-1RA and LAI users** in NHIRD)
  - Separate analyses conducted in subgroup populations (i.e., with/without prior cardiovascular diseases [CVDs] or chronic kidney diseases [CKDs])

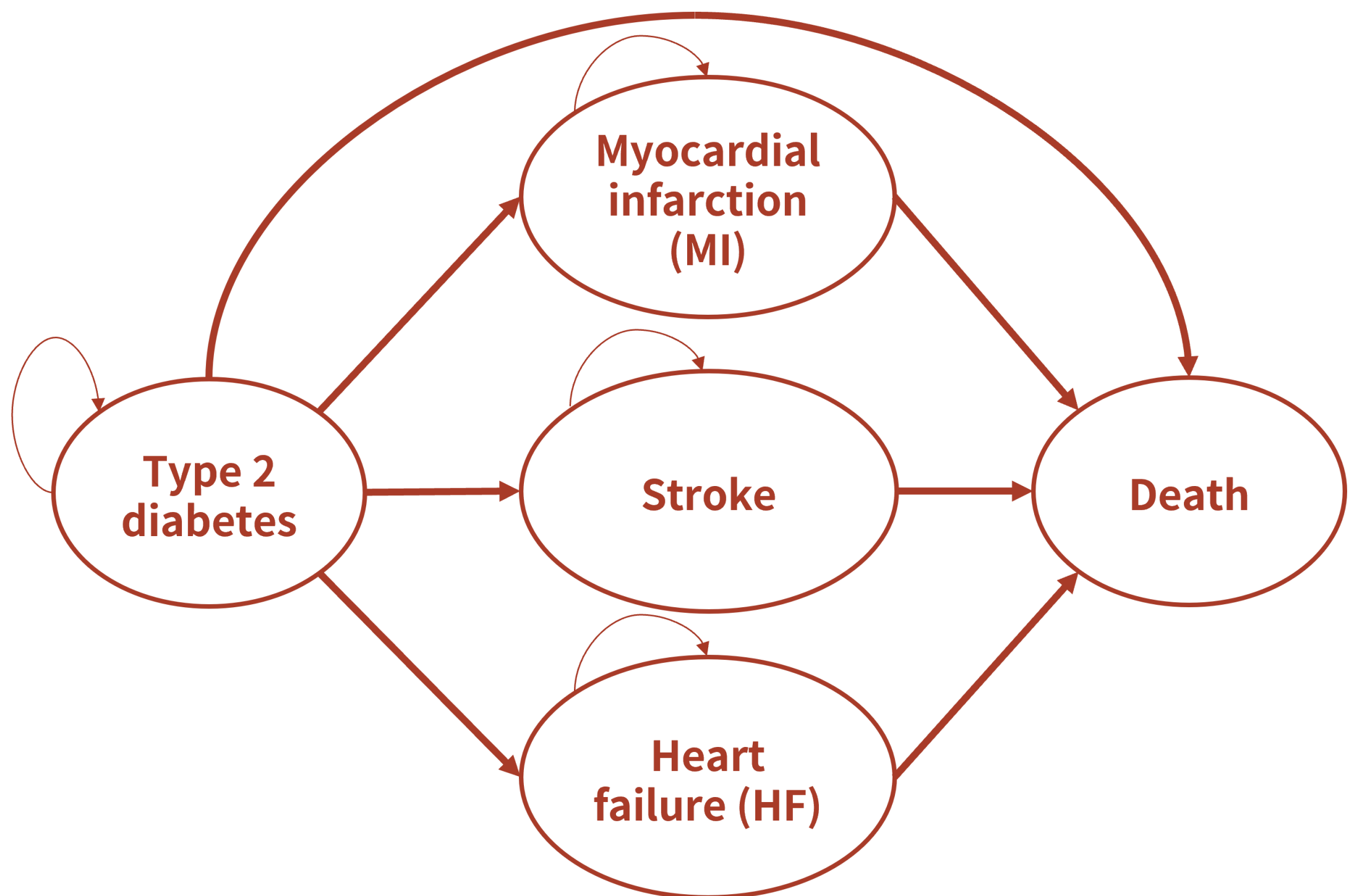


Figure 1: Model structure

## Results

- In the base-case analysis, using GLP-1RAs versus LAIs yielded an **incremental cost-effectiveness ratio (ICER) of 6,053 USD per QALY gained**, which was considered as **highly cost-effective** against the pre-defined willingness-to-pay (WTP) threshold of 33,011 USD (one-time gross domestic product per capita in Taiwan in 2021) (**Table 1**).

Table 1: ICER estimates of GLP-1RAs versus LAIs

|                            | ΔQALYs       | Δcosts (USD) | ICERs       |
|----------------------------|--------------|--------------|-------------|
| Base-case (overall cohort) | 0.804        | 4,866        | 6,053       |
| <b>Patients with CVDs</b>  | <b>0.996</b> | <b>-673</b>  | <b>-675</b> |
| Patients without CVDs      | 0.656        | 5,965        | 9,093       |
| Patients with CKDs         | 1.181        | 1,978        | 1,675       |
| Patients without CKDs      | 0.704        | 5,392        | 7,659       |

- Scenario analyses (e.g., varied simulation horizons and treatment effects of GLP-1RAs) showed consistent findings.
- Economic benefit of GLP-1RA therapy was revealed across patient subgroups, especially among those with prior CVDs or CKDs (**Table 1 and Figure 2**).

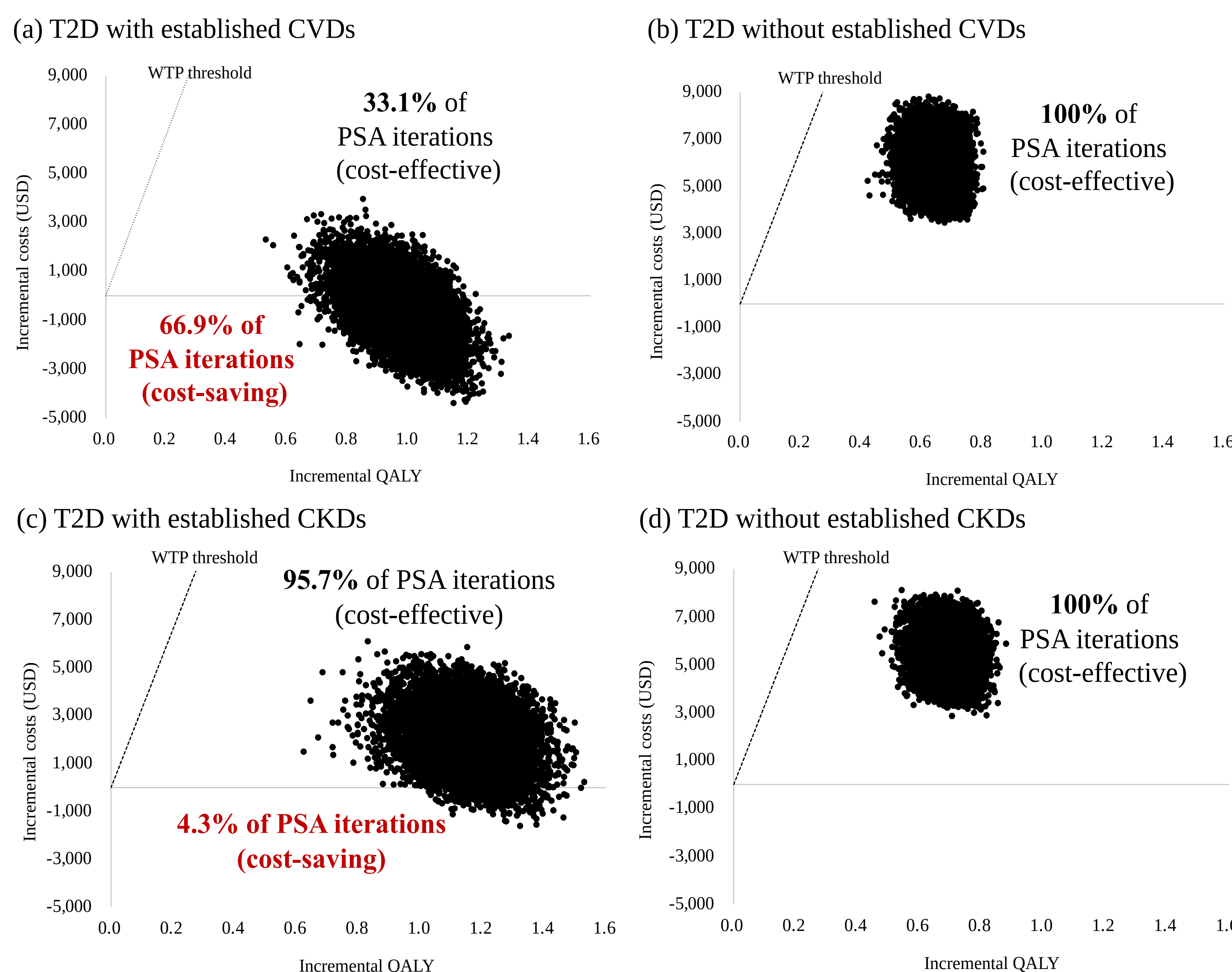


Figure 2: Results of probabilistic sensitivity analysis (PSA)

## Conclusions

Using GLP-1RAs versus LAIs would be highly cost-effective for T2D patients requiring injectable therapy. Remarkable economic benefit of GLP-1RA therapy among patients with CVDs or CKDs supports rational treatment decisions and healthcare resource optimization for these patients.