

The IHE Diabetes Cohort Model (IHE-DCM)

The IHE Diabetes Cohort Model (IHE-DCM) is a cohort-level, cost-effectiveness model developed for the evaluation of interventions in diabetes care. IHE-DCM can be used to support early-stage decision making, market access, value story dissemination, post-marketing evidence needs and analyses of long-term cost and health implications of public health issues.

Transparent, flexible, and easy to use

IHE-DCM is designed to meet strict HTA requirements for transparency in diabetes modelling while remaining easy to use and quick to run. Implemented in Microsoft® Excel, the model requires no external software and uses a familiar interface that supports review, adaptation, and clear communication of assumptions and results. This makes it well suited for engagement with clinical, payer, and policy stakeholders. Two model versions are available within the IHE-DCM framework: IHE-DCM-T1 for type 1 diabetes and IHE-DCM-T2 for type 2 diabetes.

Validation and real-world use

The validity and practical relevance of the model are supported by extensive evaluation through external validations and comparison with well-established microsimulation models [1-5]. The validation studies found that IHE-DCM produced results that were consistent with real-world data and with other established models. The model has also been accepted by HTA agencies in Sweden, Norway, Canada, and Australia, supporting its use in healthcare decision-making across different healthcare systems [6-12].

Scientific publications from Sweden, Denmark, Canada, and China further demonstrate the international use of IHE-DCM [13-24]. The model has been used to evaluate a broad range of interventions, including pharmaceutical treatments, insulin pumps, continuous glucose monitoring, and pediatric screening for type 1 diabetes. The practical relevance of the model is also reflected in licensing by external stakeholders, including life science companies and HTA agencies [25]. Overall, this evidence supports the credibility and practical value of the model for healthcare decision-making in different settings.

Accessing the model

IHE-DCM is available through global or country-specific license agreements covering IHE-DCM-T1, IHE-DCM-T2, or both. Each license includes supporting documentation, including a technical report and user guide, together with technical support. Additional model training can be scheduled on request. The model can also be accessed as part of specific projects conducted by IHE.

USER FRIENDLY



- Available through licensing
- Fully Excel-based
- Intuitive interface
- Short run time
- Can be shared with HTA agencies

ACCEPTED & VALIDATED



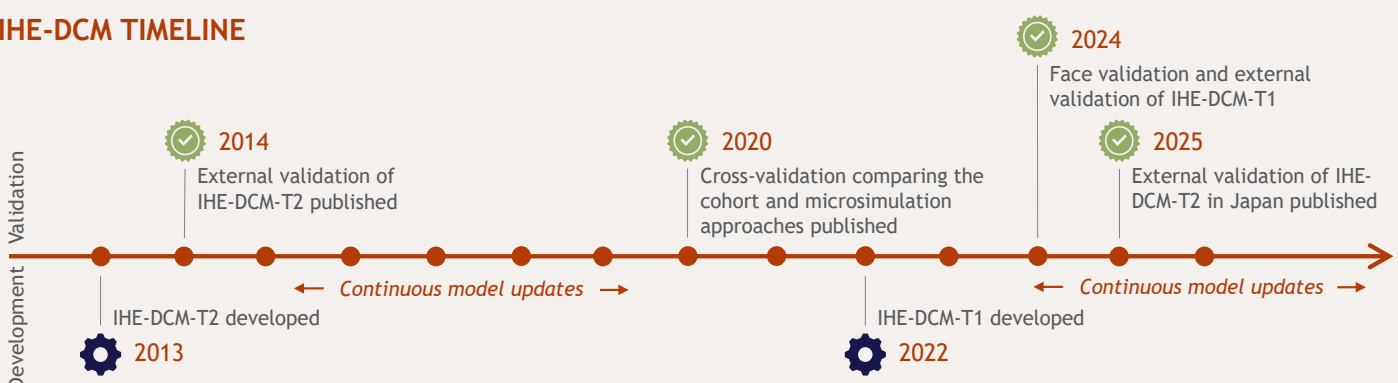
- Externally validated in several independent studies
- Accepted for use by HTA agencies in multiple countries
- Transparent model structure to support review

FLEXIBLE MODEL DESIGN

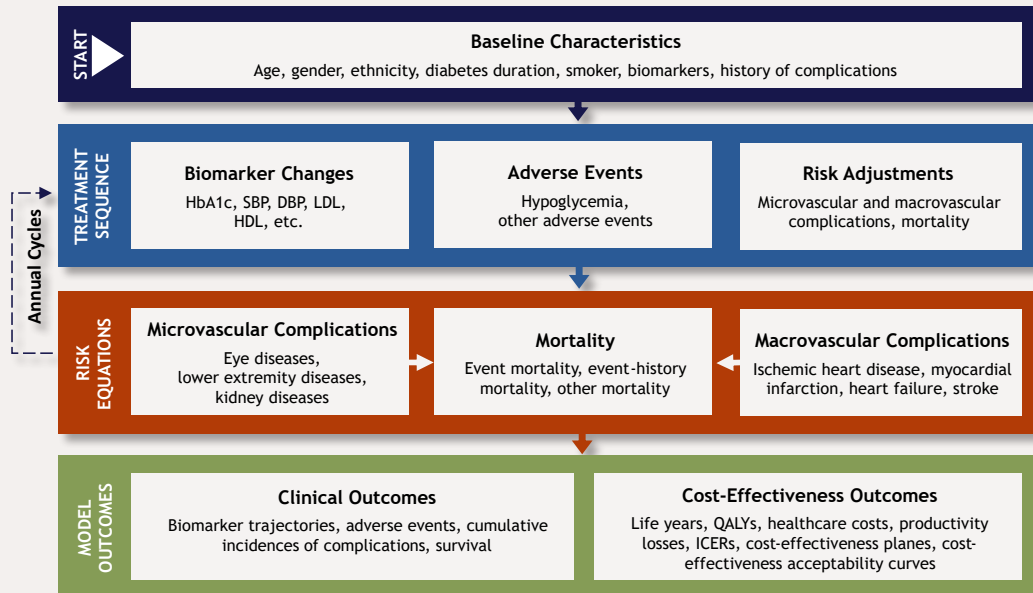


- Separate versions for type 1 and type 2 diabetes
- Designed to reflect different treatment options
- Supports comparison of multiple treatments in a single analysis
- Continuously developed and updated

IHE-DCM TIMELINE



IHE-DCM MODEL STRUCTURE



Risk prediction equations

Type 2 diabetes

- UKPDS-OM1 (UK)
- UKPDS-OM2 (UK)
- NDR (Sweden)
- FDS (Australia)
- JJRE (Japan)
- NIH (US; Eastman)

Type 1 diabetes

- DCCT/EDIC (US)
- NDR (Sweden)

Model description

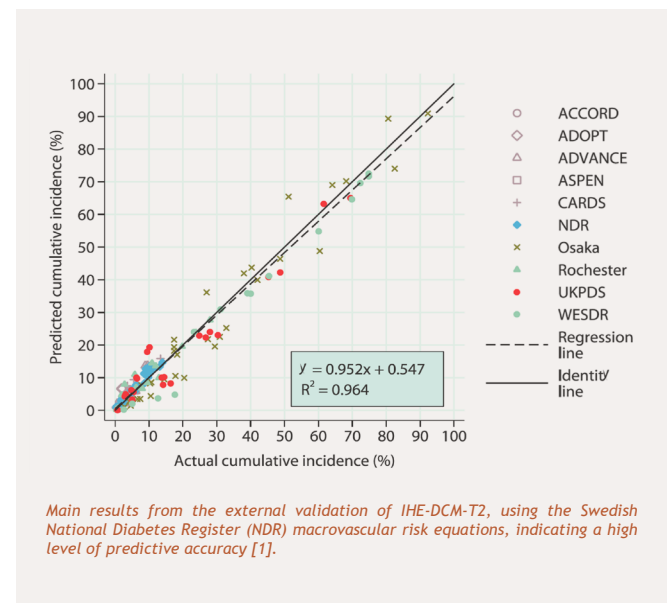
IHE-DCM is a cohort-level Markov cost-effectiveness model. It supports deterministic and probabilistic sensitivity analyses and operates with a one-year cycle length. The model is flexible, allowing most parameters to be user-defined.

The model includes key diabetes-related complications and mortality. Complications are chronic and generally progressive, and multiple complications can occur simultaneously. Transition probabilities are derived from established risk prediction equations for both type 1 and type 2 diabetes, based on evidence from multiple large cohort studies across various country settings.

Within this framework, the cohort is defined by characteristics such as age, diabetes duration, biomarker values, and baseline complications. Biomarkers evolve over time according to user-defined trends and treatment effects, while complications develop and progress. A flexible treatment sequence permits modelling of a wide range of treatment profiles.

The model estimates health outcomes, including life-years and quality-adjusted life years (QALYs), adjusted for complications and adverse events. It incorporates costs for treatments, adverse events, and complications, and can optionally include indirect costs such as productivity losses.

Cost-effectiveness is summarized using incremental cost-effectiveness ratios (ICERs) and through the probabilistic sensitivity analysis (PSA) functionality, results can be presented as cost-effectiveness planes and cost-effectiveness acceptability curves (CEACs).



- Lundqvist, A., et al., Validation of the IHE Cohort Model of Type 2 Diabetes and the Impact of Choice of Macrovascular Risk Equations. *PLoS One*, 2014. 9(10): p. e110235.
- Willis, M., et al., Comparing the Cohort and Micro-Simulation Modeling Approaches in Cost-Effectiveness Modeling of Type 2 Diabetes Mellitus: A Case Study of the IHE Diabetes Cohort Model and the Economics and Health Outcomes Model of T2DM. *Pharmacoeconomics*, 2020. 38(9): p. 953-969.
- Nilsson, K., et al., Validation of the IHE type 2 diabetes cohort model in the Japanese clinical setting. *Journal of Medical Economics*, 2025. 28(1): p. 944-963.
- K. Nilsson, S.P., A. Fridhammar., IHE Type 1 Diabetes Cohort Model (IHE-DCM-T1). Technical Report, Version 1.2.1. 2025.
- Altunkaya, J., et al., Examining the Impact of Structural Uncertainty Across 10 Type 2 Diabetes Models: Results From the 2022 Mount Hood Challenge. *Value in Health*, 2024. 27(10): p. 1338-1347.
- The Swedish Dental and Pharmaceutical Benefits Agency (TLV). [Title in English: Lyxumia is included in the pharmaceutical benefits scheme with restrictions. 2015 [2018-01-12]
- The Swedish Dental and Pharmaceutical Benefits Agency (TLV). Underlag för beslut om subvention - Ozempic (semaglutide). 2018: <https://www.tlv.se>.
- The Swedish Dental and Pharmaceutical Benefits Agency (TLV). Underlag för beslut om subvention - Xultophy. 2015: <https://www.tlv.se>.
- Norwegian Medicines Agency (NoMA). [Title in English: Rapid assessment for pre-approved reimbursement - Ozempic (semaglutide) for the treatment of type 2 diabetes mellitus]. 2019.
- Norwegian Medicines Agency (NoMA). [Title in English: Rapid health technology assessment for pre-approved reimbursement - Rybelsus (semaglutide) for the treatment of type 2 diabetes mellitus]. 2019: https://www.dmp.no/globalassets/documents/offentlig-finansiering-og-pris/metodevurderinger/rybelsus_dmiil_2021.pdf
- CADTH, CADTH common drug review: pharmacoeconomic review report: semaglutide (Ozempic). 2019: <https://www.cadth.ca/sites/default/files/cdr/pharmacoeconomic/sr0594-ozempic-pharmacoeconomic-review-report.pdf>.
- PBS. Public Summary Document - November 2019 PBAC Meeting. 2019; Available from: <https://www.pbs.gov.au/industry/listing/elements/pbac-meetings/psd/2019-11/files/semaglutide-psd-november-2019.docx>.
- Kiadalini, A.A., et al., Cost-Utility Analysis of Glucagon-Like Peptide-1 Agonists Compared with Dipeptidyl Peptidase-4 Inhibitors or Neutral Protamine Hagedorn Basal Insulin as Add-On to Metformin in Type 2 Diabetes in Sweden. *Diabetes Ther*, 2014.
- Steen Carlsson, K. and U. Persson, Cost-effectiveness of add-on treatments to metformin in a Swedish setting: liraglutide vs sulphonylurea or sitagliptin. *J Med Econ*, 2014. 17(9): p. 658-69.
- Ericsson, Å. and A. Lundqvist, Cost Effectiveness of Insulin Degludec Plus Liraglutide (IDegLira) in a Fixed Combination for Uncontrolled Type 2 Diabetes Mellitus in Sweden. *Applied Health Economics and Health Policy*, 2017. 15(2): p. 237-248.
- Ericsson, A. and A. Fridhammar, Cost-effectiveness of once-weekly semaglutide versus dulaglutide and lixisenatide in patients with type 2 diabetes with inadequate glycaemic control in Sweden. *J Med Econ*, 2019. 22(10): p. 997-1005.
- Johansen, P., et al., Cost Effectiveness of Once-Weekly Semaglutide Versus Once-Weekly Dulaglutide in the Treatment of Type 2 Diabetes in Canada. *Pharmacoecon Open*, 2019. 3(4): p. 537-550.
- Ericsson, A., et al., Cost-effectiveness of liraglutide versus lixisenatide as add-on therapies to basal insulin in type 2 diabetes. *PLoS One*, 2018. 13(2): p. e0191953.
- Lindvig, A., et al., The economic burden of poor glycaemic control associated with therapeutic inertia in patients with type 2 diabetes in Denmark. *Curr Med Res Opin*, 2021. 37(6): p. 949-956.
- Eliasson, B., et al., Long-Term Cost-Effectiveness of Oral Semaglutide Versus Empagliflozin and Sitagliptin for the Treatment of Type 2 Diabetes in the Swedish Setting. *Pharmacoecon Open*, 2022. 6(3): p. 343-354.
- Nilsson, K., et al., Model-based predictions on health benefits and budget impact of implementing empagliflozin in people with type 2 diabetes and established cardiovascular disease. *Diabetes Obes Metab*, 2023. 25(3): p. 748-757.
- Stafford, S., et al., Cost-Effectiveness of Once-Weekly Semaglutide 1 mg versus Canagliflozin 300 mg in Patients with Type 2 Diabetes Mellitus in a Canadian Setting. *Appl Health Econ Health Policy*, 2022. 20(4): p. 543-555.
- Ruan, Z., et al., Long-Term Cost-Effectiveness Analysis of Once-Weekly Semaglutide versus Dulaglutide in Patients with Type 2 Diabetes with Inadequate Glycemic Control in China. *Diabetes Ther*, 2022. 13(10): p. 1737-1753.
- Liu, L., et al., Long-Term Cost-Effectiveness of Subcutaneous Once-Weekly Semaglutide Versus Polyethylene Glycol Lixisenatide for Treatment of Type 2 Diabetes Mellitus in China. *Diabetes Ther*, 2023. 14(1): p. 93-107.
- The Swedish Dental and Pharmaceutical Benefits Agency (TLV). Utvärdering av avancerade insulinpumpar för vuxna med diabetes typ 1. 2025; Available from: <https://www.tlv.se>.