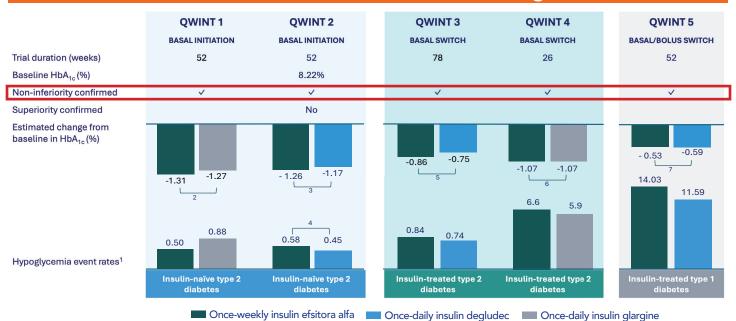






Clinical Data Overview Phase 3 QWINT Program



^{1.} Events per participant-year of exposure; 2. 95% CI, -0.22 to 0.04; 3. 95% CI, 0.94 to 1.78; 4. 95% CI, -0.22 to 0.061; 5. 95% CI, -0.19 to 0.12; 6. 95% CI, -0.22 to 0; 7. 95% CI, -0.075% to 0.19%. ClinicalTrials gov identifier. NCT05662332. ClinicalTrials gov identifier: NCT05662332. ClinicalTrials gov identifier: NCT056262568. QWINT 2. https://www.clinicaltrials.gov/ci2/show/NCT05525400; ClinicalTrials gov identifier: NCT055427606. QWINT 3. https://www.clinicaltrials.gov/ci2/show/NCT05525400; ClinicalTrials gov identifier: NCT05642766. QWINT 4. https://www.clinicaltrials.gov/ci2/show/NCT05427640; ClinicalTrials gov identifier: NCT05462766. QWINT 4. https://www.clinicaltrials.gov/ci2/show/NCT0542766. QWINT 4. https://www.clinicaltrials.gov/ci2/show/NCT

Clinical Data Overview Phase 3 ONWARDS Program



In people with type 2 diabetes: No statistical difference in estimated hypoglycaemia events

Once-weekly insulin icodec Once-daily insulin glargine U100 Once-daily insulin degludec Once-daily basal insulins

*Statistically significant. 1 Severe or clinically significant hypoglycaemia events (blood glucose <3 mmo/L) per patient year, included for end of trial/end main phase in-trial. 2 Duration refers to trial main phase.

ONWARDS 1: QW insulin icodec vs QD insulin glargine U100 both with non-insulin anti-diabetic treatment in insulin-naive people with T2D; ONWARDS 2: QW insulin icodec vs QD insulin degludec in people with T2D switching from a QD insulin; ONWARDS 3: QW insulin icodec vs QD insulin degludec in insulin-naive people with T2D; ONWARDS 4: QW insulin icodec vs QD insulin degludec both with mealtime insulin in people with T2D treated with basal and bolus insulin; ONWARDS 5: QW insulin icodec vs QD basal insulin with an app providing dosing recommendation in insulin-naive people with T2D; ONWARDS 6: QW insulin icodec vs QD insulin degludec both with mealtime insulin in people with T1D T1D: Type 1 diabetes; T2D: Type 2 diabetes. Note: Overview refer to primary end-points in main phases of trials