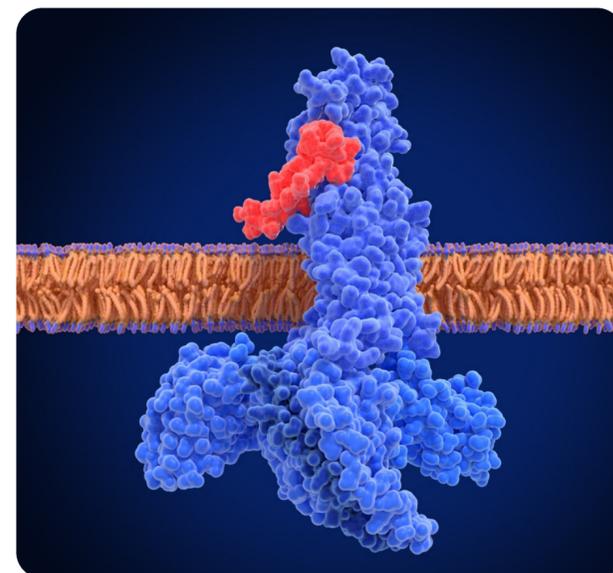


From Insulins to GLP-1 Receptors:

Preparing for the Next Wave
of Off-Patent Molecules



Supporting You at Every Stage of your Drug Development Journey

Preclinical

- Discovery
- Chemistry
- Bioanalysis
- Toxicology

Biopharma

- Clinical Bioanalysis solution for Bio Therapeutics
- Non-Clinical Characterization Solutions - Discovery Biology, Bioprocess, and Analytical Characterization

Clinical Studies

- Phase I Healthy Volunteer study for Antidiabetic drug
- Type 2 Diabetic Phase II/Phase III Trials
- Phase IV & PMS Studies
- Healthy Volunteer Bioequivalence (BE)/ Bioavailability (BA) studies involving Glucose Clamp & GLP1

Bioanalytical Solutions

- Bioanalytical Method Development and Validations for NCEs & Generics

State-of-the-Art Infrastructure & Expertise for Endocrinology Drug Development

Operational Capabilities

- Experience in Recruiting 400+ subjects
- 38 Endocrinologists across 20 sites
- GCP Trained Study Staff
- Trained doctor and team in Advanced Cardiac Life Support (ACLS) and Basic Life Support (BLS)
- Utilized 1322 clamps in 15 studies
- Experience ranging from 8 to 36 hours per clamp
- Capable of performing 4 to 6 clamps per day for efficient study completion
- Trained senior staff specialized in Endocrinology studies with a PHD in Medicine and Pharmacology.
- Tertiary care tie-up with Sterling hospital for advanced medical care

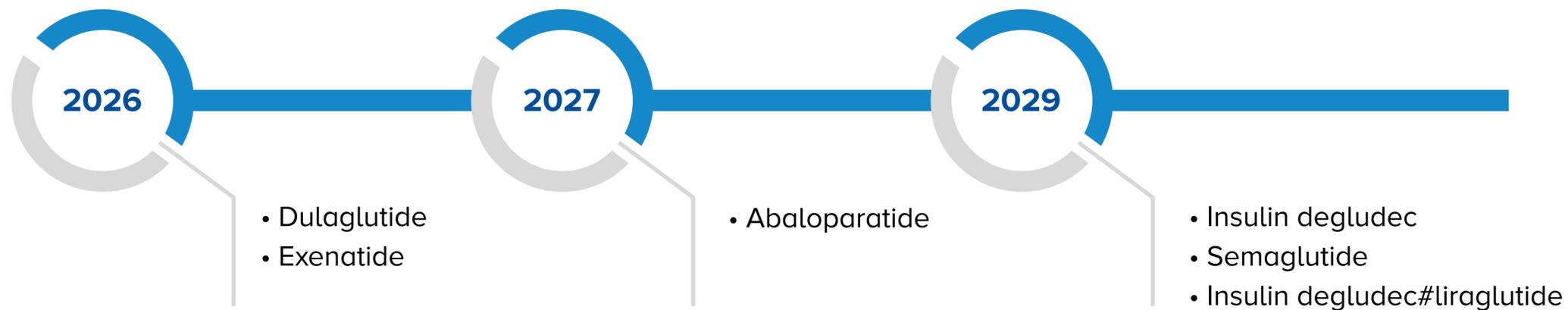
Advanced Facilities

- Yellow Spring Instruments (YSI) for accurate blood glucose assessment
- 12 state-of-the-art beds with backup generators
- 30-bed Phase-I capacity across two units.
- Special care area with cardiac monitor, defibrillator, ECG machine, suction machine, oxygen cylinder, cardiac arrest kit, and anaphylaxis kit.
- Resuscitation centre with a fully equipped resuscitation trolley and emergency medications

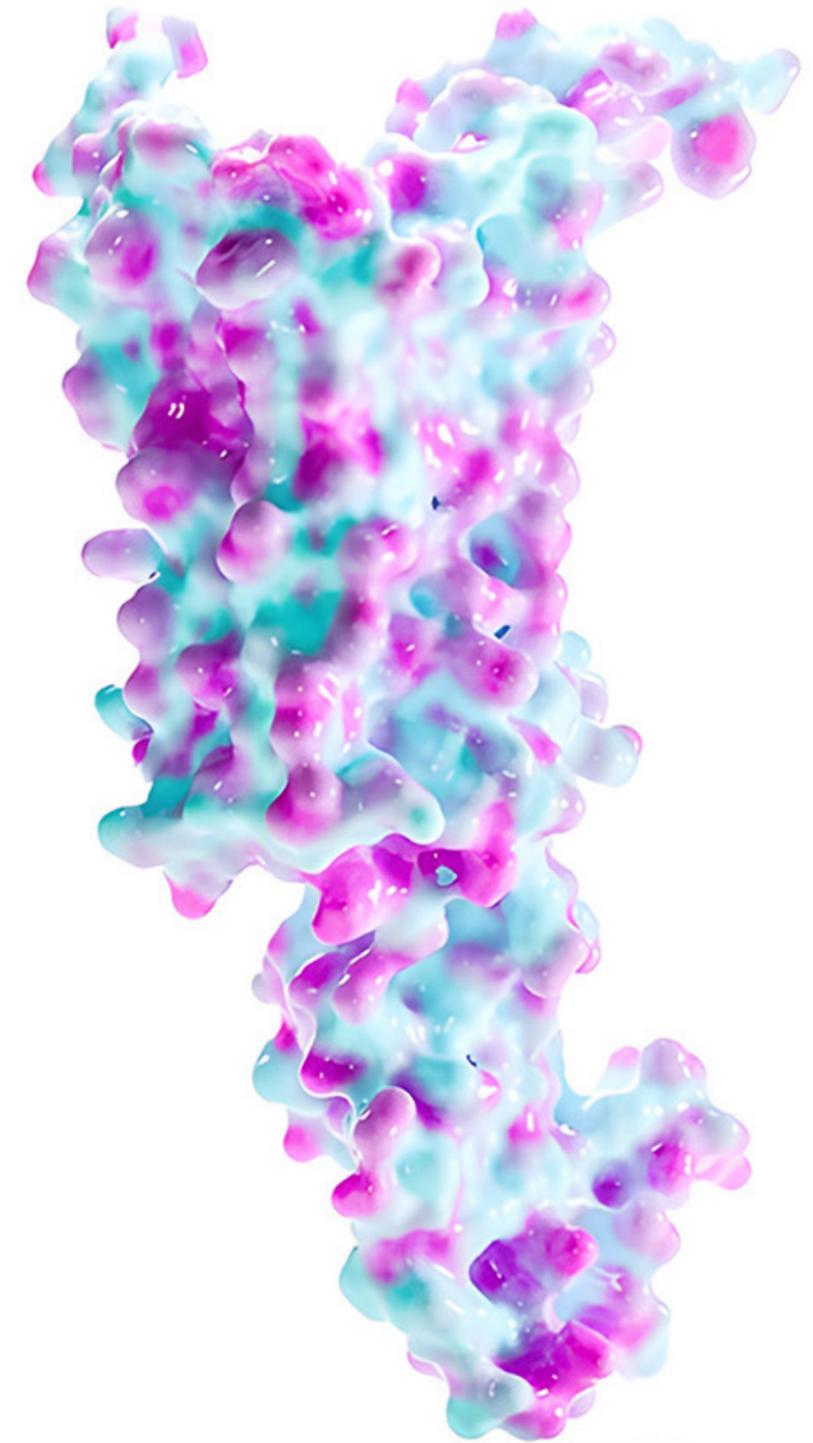
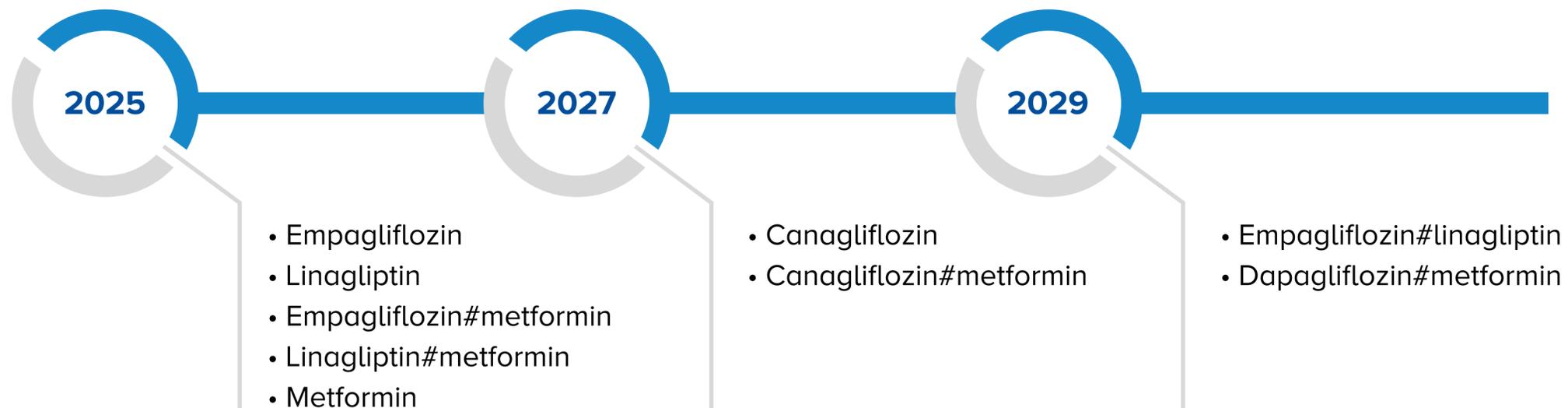


Stay Ahead of the Curve: Veeda's Solution for Molecules going Off-Patent

Biotherapeutics Going off Patent

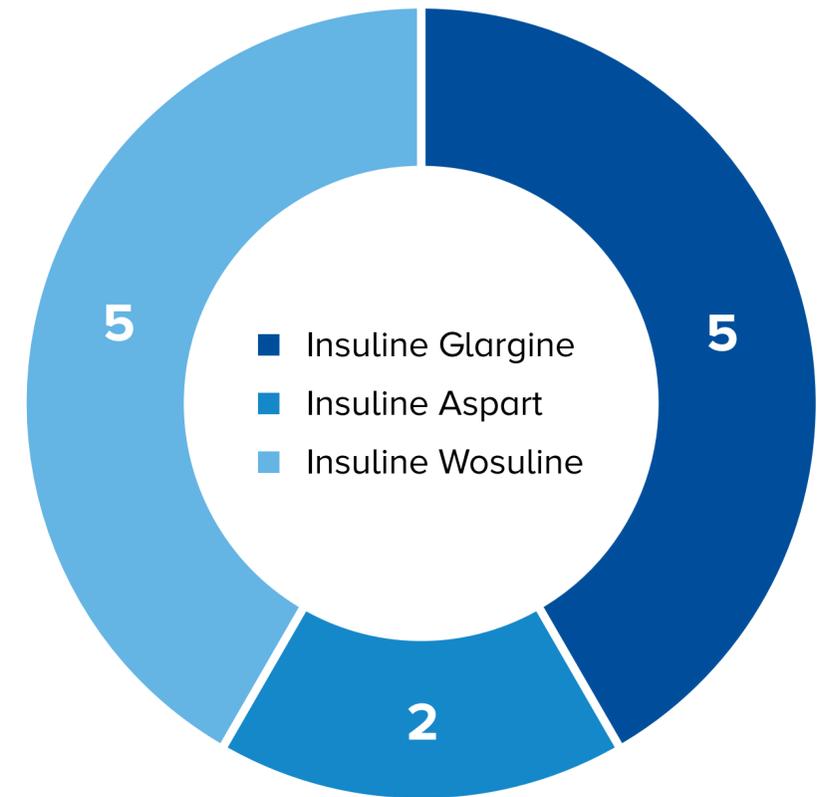


Molecules Requiring Healthy Volunteer Generic Bioequivalence (BE) Studies:



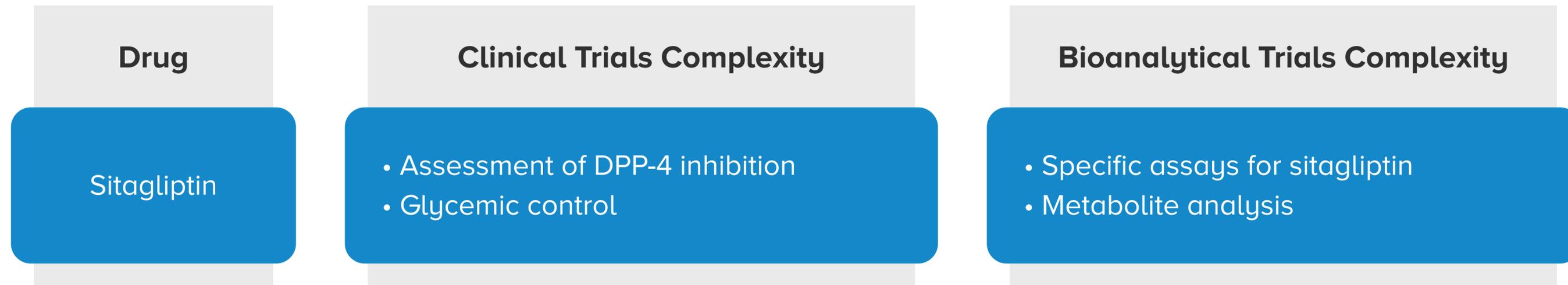
Diverse Insulin Portfolio: Successfully executed 12 Different Studies

Insulin Product	Clinical Trials Complexity	Bioanalytical Trials Complexity
Insulin Glargine	<ul style="list-style-type: none">• Individualized titration• Hypoglycemia management	<ul style="list-style-type: none">• Sensitive long-acting assays• Metabolite analysis
Insulin Aspart	<ul style="list-style-type: none">• Postprandial glycemic control• Injection site rotation	<ul style="list-style-type: none">• Rapid-acting assays• Injection site impact
Insulin Wosulin	<ul style="list-style-type: none">• Individual patient response• Hypoglycemia risk assessment	<ul style="list-style-type: none">• Development of Wosulin-specific assays• Pharmacokinetic/pharmacodynamic differences



Insights into Sitagliptin: Diabetes Care with DPP-4 Inhibitors

- DPP-4 inhibitors like Sitagliptin prolong the action of GLP-1 and GIP, resulting in increased insulin secretion and decreased glucagon secretion. This mechanism ultimately helps to lower blood sugar levels in patients with type 2 diabetes.



Completed Oral Antidiabetic Glucose Clamp Studies with 20 Subjects

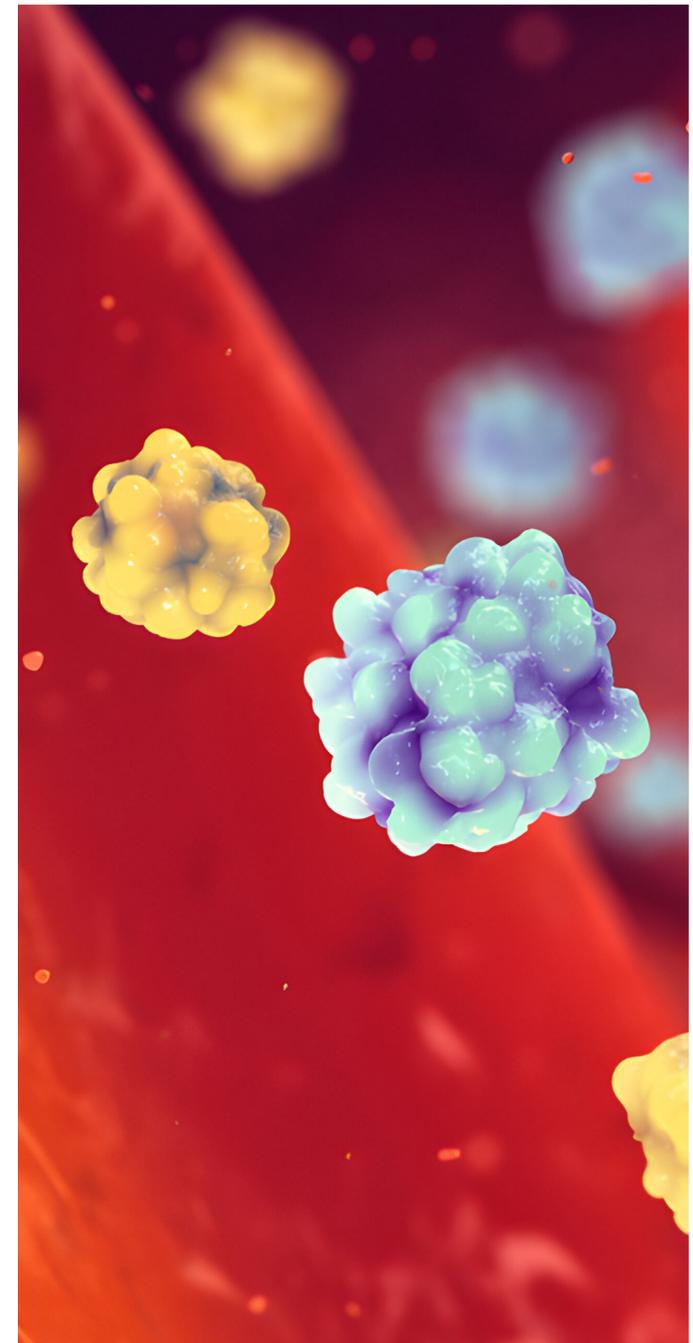
Biosimilar Experience across Different Diabetic Molecules

- Specialized in clinical and bioanalytical evaluation, with expertise in immunogenicity assessment using advanced techniques such as HPLC, ELISA and LC-MS/MS
- Biosimilar development includes rigorous analytical techniques such as intact mass analysis, peptide mapping, and amino acid sequencing by focusing on peptides like Insulin, & C-Peptide
- Advanced methods like capillary electrophoresis to assess molecular heterogeneity and confirm the equivalence of our biosimilar

Completed Healthy Volunteer GLP-1 Studies for -

Liraglutide

Semaglutide



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Excellence
In Everything