



Partners in creating a healthier tomorrow

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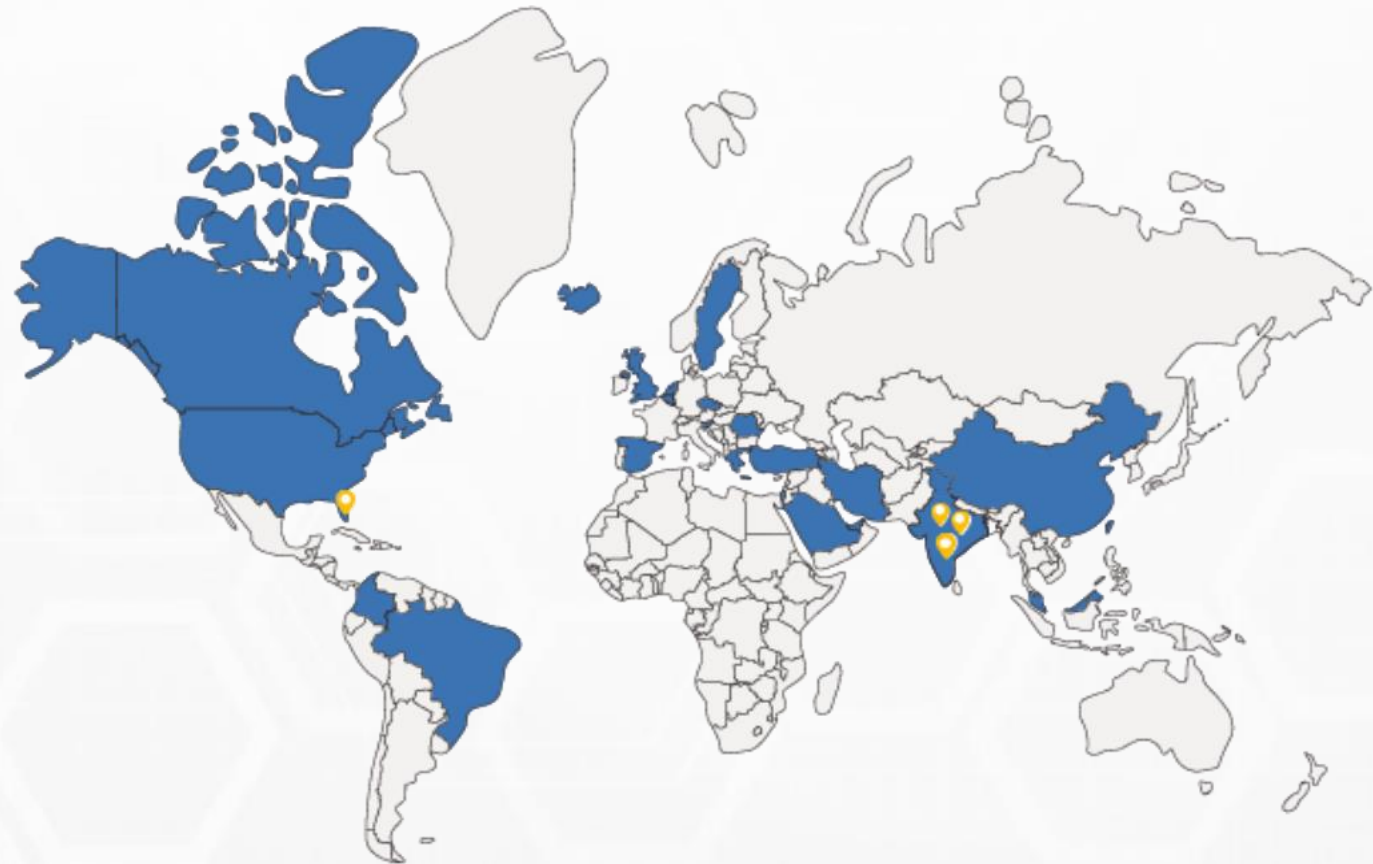
Veeda Group Capabilities




Veeda Group

- Veeda Clinical Research Limited (“Veeda”) together with its subsidiary, Bionees India Private Limited (“Bionees”), (together referred to as the “Veeda Group”) offers a comprehensive portfolio of clinical, preclinical and bio/analytical services to support innovator, biosimilar and generic drug development programs of our global clientele
- We are an independent, institutional investors owned, Board governed and professionally managed contract research group offering scientific leadership, global quality management systems and long term operational and financial stability through a continuing investment in our people, processes, systems, infrastructure and technology and a deep commitment to quality
- Together, we serve clients globally in the following industries:
 - Pharmaceutical and Biopharmaceutical
 - Agrochemical and Industrial Chemicals
 - Herbal/ Nutraceuticals
 - Medical Devices

Our Global Foot Print



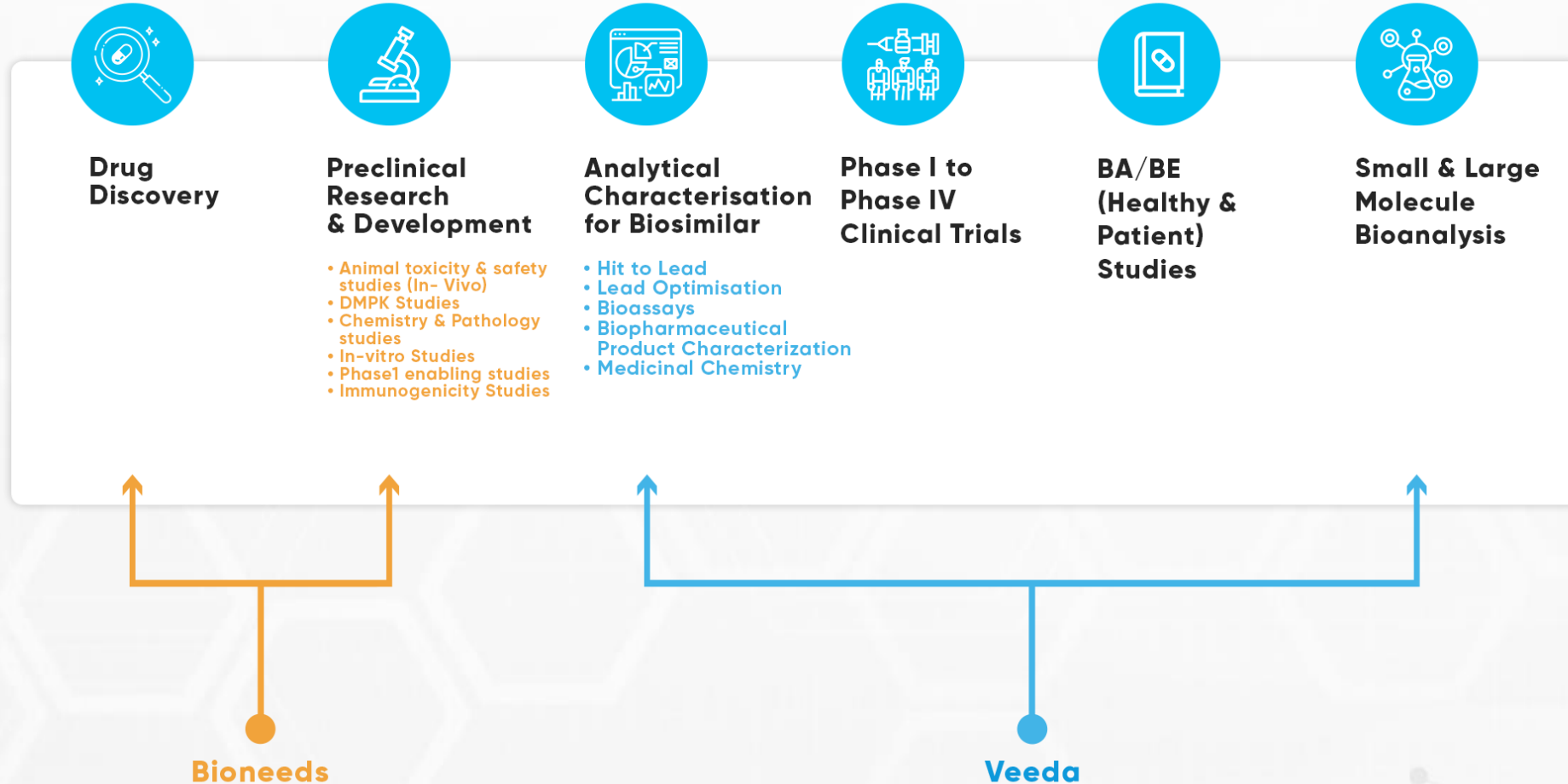
 Serving clients across these geographies

 Veeda's Team Presence

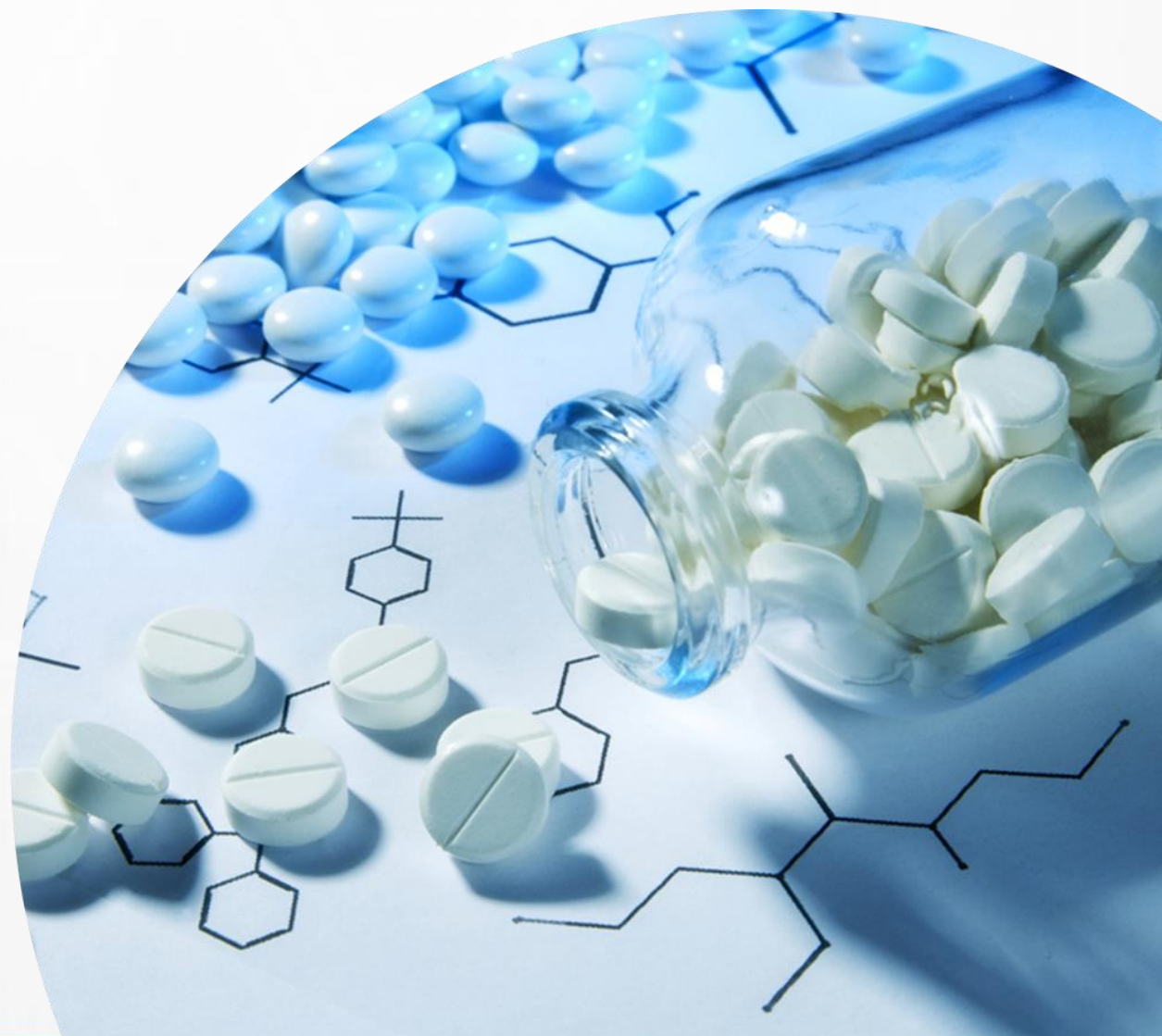
Drug Development Services Overview



Drug Development Journey



Complex Generics Overview



What are Complex Generics?

- The U.S. Food and Drug Administration (FDA) defines a generic drug as one that is identical – or bioequivalent – to a brand name drug in dosage form, safety, strength, route of administration, quality, performance characteristics and intended use.
- A simple generic is a copy of a small molecule reference drug and is chemically identical to its branded counterpart.
- A complex generic is a generic that could have a complex active ingredient, complex formulation, complex route of delivery, or complex drug device combinations.

Source: U.S. FDA

The European Medicines Agency (EMA) defines a generic drug as a medicine that is developed to be the same as a medicine that has already been authorized. It contains the same active substances and is used at the same doses to treat the same diseases as the reference drug.

EMA refers to complex generics as “hybrid medicines,” whose “authorization depends partly on the results of tests on the reference medicine and partly on new data from clinical trials.”

Source: EMA

It is challenging, time-consuming and expensive to develop complex generics and demonstrate the equivalence, safety and efficacy of the therapy.

Complex Products

- **Complex active ingredients**
 - E.g., Complex mixtures of APIs, polymeric compounds, peptides
- **Complex formulations**
 - E.g., Liposomes, suspensions, emulsions, gels
- **Complex routes of delivery**
 - E.g., Locally acting such as ophthalmic, otic, dermatological and inhalational drugs
- **Complex dosage forms**
 - E.g., Long acting Injectables and implantable
- **Complex drug-device combinations**
 - E.g., Metered Dose Inhalers, nasal sprays and transdermal
 - Other products where complexity or uncertainty concerning the approval pathway or other alternative approach would benefit from early scientific engagement

Key Considerations

- **Regulatory requirements**
 - Product specific guidance (PSG) from the FDA Office of Generic Drugs (OGD), European Medicines Agency's (EMA)
 - Interactions with regulatory agencies and precedents: trial parameters, data end points
- **Study design and planning**
 - Protocol, study design, location(s), population
 - Right collaboration for clinical studies
 - Approval from local regulatory bodies
- **Compliance**
 - Quality execution
 - Cost and time

Veeda Can Support

- **Study design and planning**
 - Clinical study design, development of study protocol
 - Approval from local regulatory bodies and Ethics committee
- **Compliance**
 - Cost effective quality execution in timely manner
- **Experts at Handling Complex Generic Studies**
 - Inhalation studies
 - Glucose Clamp Studies
 - Complex dosage formulations – Long Acting Injections (LAI), Rectal dosage forms
 - Robust and high-sensitive analytical instruments
 - Biosimilar studies – 06 studies

Veeda's Drug Development Capabilities - Generic Drugs

End-to-End BA/BE study development and execution (pilot and pivotal) towards ANDA submission for different regulatory authorities like USFDA, EMA, ANVISA, Health Canada, WHO, MHRAUK, CDSCO and many more.

Toxicity testing for special products, Impurity synthesis & LCMS characterization, Invitro microbial kill rate study, generic drug stability testing

**Generic
Drugs
Capabilities**

505 (b)(2) method development and submission for branded generics, orphan drugs, prodrugs, and Drug Efficacy Study Implementation (DESI) drugs.

Diverse Therapeutic Areas Of Expertise



Cardiology



Rheumatology



Dermatology



Ophthalmology



Gynecology



Gastroenterology



ENT



Oncology



Psychiatry



Respiratory



Endocrinology

Team Experience Across various Therapeutic Areas and Indications

Sr. No.	Area	Indication	Regulatory Submissions
1	Psychiatry	Major Depressive Disorder, Schizophrenia, Bipolar disorder, Bipolar I depression	USFDA, EMA and DCGI
2	Medical Devices	CAD, Arrhythmia, Heart failure, Uncontrolled hypertensions,	USFDA & DCGI
3	Cardiology	Hypertension, Ischemic cardiomyopathy, CVD, ACS	USFDA, EMA and DCGI
4	Endocrinology	DM-I, DM-II, Diabetic nephropathy	USFDA, EMA and DCGI
5	Oncology	Advanced Ovarian Cancer, Metastatic breast cancer, Renal Cell Carcinoma, Multiple Myeloma, Colorectal Cancer, Solid Tumors / Lymphoma, NSCLC, Cervix Cancer,	USFDA, EMA, ENVISA and DCGI
6	Respiratory	Asthma, COPD	USFDA & DCGI
7	Dermatology	Atopic dermatitis, Oral lichen planus, Dermatomycoses	DCGI
8	Nephrology	CKD, Urinary tract infection and pyelonephritis	USFDA & DCGI
9	Gastroenterology	Arsenic Poisoning, GERD, Constipation, Ulcerative Colitis	USFDA & DCGI
10	Infectious diseases	Bacterial Infection, Skin Infection, Hepatitis B Infection	USFDA & DCGI
11	Ophthalmology	Chronic Open Angle Glaucoma, Ocular Hypertension	USFDA & DCGI
12	Neurology	Epilepsy, Seizures	DCGI
13	Vaccine	Rabies, Leishmaniasis & serious fungal infections	DCGI
14	Orthopaedic	Psoriasis and Rheumatoid Arthritis& Osteoporosis	USFDA & DCGI

Clinical Research



Infrastructure

- **VEDANT**

Clinical,
Bio-analytical facility

- **SATYAMEV CORPORATE PARK**

Corporate Office
Bio-analytical facility

- **SHIVALIK**

Dedicated Clinical facility

- **MEHSANA**

Clinical and
Screening facility

- **SKYLAR**

Common screening
facility for both Shivalik
and Vedant

- **INSIGNIA**

Dedicated
Bio-analytical facility

- **ARCHIVES**

Archival area in each facility.
Separate archival facility at Changodar

Spread across **16** clinics

Shivalik

170 Beds +

7 Special care beds +

12 Intensively monitored
beds to conduct Phase I
study

Vedant

226 Beds +

8 Special care beds +
18 Intensively monitored
beds to conduct Phase I
study

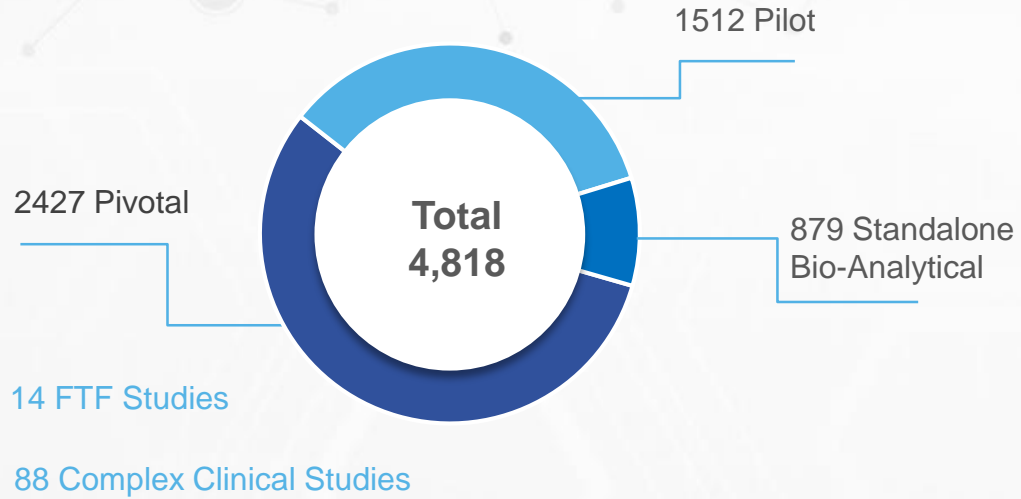


Mehsana

162 Beds +

7 Special care beds

Experience







75 Special Studies

*Both Pilot and Pivotal BA/BE

- 16 Glucose Clamps studies (1322 clamps)
- 36 Inhalation Studies
- 8 Suppositories
- 15 Patches Studies
- 27 Phase – I Studies
- 1 Phase – II Study

Volunteer Database (More than 84,947)

- Male Volunteers —  > 62,179
- Female Volunteers —  > 5,659
- Elderly Males —  > 13,322
- Post - Menopausal Females —  > 3,787

Glucose Clamp



Veeda's Glucose Clamp Studies Overview

Veeda's proficiency in glucose clamp studies

- 1322 Clamp utilized in 16 studies
- Clamp experience ranging from 8 hours to 36 hours
- Capability of performing 4 to 6 clamps per day
- Risk Mitigation and Management Strategies (RMMS)
- Robust database of healthy volunteers with previous history of participation in clamp studies while adhering to mandatory compliance instructions and other study related restrictions

Veeda's clinical facilities that support glucose clamp studies

- YSI (Yellow Spring Instruments), for accurate assessment of blood glucose values
- 12 state-of-art beds in Phase I unit with stature lifts that have back-up generators
- Well-equipped special care area to handle medical emergencies with provisions such as cardiac monitor, defibrillator, ECG machine, suction machine, oxygen cylinder, cardiac arrest kit, and anaphylaxis kit
- Resuscitation centre with resuscitation trolley and all necessary and movable emergency medications and equipment.
- In-house ambulance
- Tertiary care tie up with Sterling hospital, Gujarat, a 280 bedded multi-specialty hospital for sophisticated and advanced emergency medical care
- Fire and chemical hazard systems with standard operating procedures (SOPs) and well-maintained equipment

Veeda's Glucose Experience

Veeda have experience in 16 Glucose clamp studies

Veeda's Experience in handling Glucose Clamp Studies

Drug synopsis	No. of subjects
Wosulin N (Insulatard as reference)	14
Wosulin N (Novolin as reference)	18
Wosulin N (Insulatard as reference)	12
Wosulin N (two batches of Novolin as reference)	18
MK-0431 (Sitagliptin 100mg) Tablets	12
Insulin Glargine 100IU/ml	4
Insulin Glargine 100IU/ml	18
TRC150094 50mg Tablets	20
Insulin Glargine	40
Insulin Aspart	71
Insulin Wosulin	60
WCK 9444 G01	20
Sun_Phase 1 AS012_150/300/600/1200/2400mg_Cohort 1A&B, Cohort 2A, MAD, 100 mg	48
Insulin Glargine injection 100 IU/MI	48
Insulin Aspart Mix 30 100 U/mL	60
Insulin Glargine	48

Inhalation



Inhalation Overview

Inhalation: Infrastructure

- **State-of-the-art Negative Pressure Rooms**
- **Advantages:**
 - Provides uniform environment with relatively consistent temperature, humidity, air flow, oxygen content and other major environmental factors for respiratory dosing
 - Eliminates any chances of cross contamination from one dosed subject to another during dosing procedure
 - Better regulatory acceptance due to assured well controlled dosing procedure
- **Specifications:**
 - Change room 1 and 2 at 25 Pa (capacity of 3 persons at a time in each room)
 - Dosing room 1 and 2 at 10 Pa (capacity of 3 persons at a time in each room)
 - Ensure that the movement of air between these rooms is unidirectional, from change rooms to dosing room
 - ACPH (Air Cycles Per Hour) time of 25 cycles, gap of 4 minutes between two consecutive dosing is appropriate

Inhalation: Training of Volunteers

- **Training of volunteers on placebo inhalers, aerosol inhalation monitors (AIM), in-check Dial meters and 2-dose devices:**
 - To educate on the proper inhalation technique devoid of any leakage - Exhalation followed by inhalation: full inhalation at rate of 70-90 l/min for DPI and consistent inhalation at 30 l/min to 60 l/min for pMDI.
 - To understand uniform inhalation rate.
 - For precise interpretation of inhalation flow with respect to time.
 - For interpretation of inhalation volumes, turbulent flow, and acceleration rates.
 - For understanding the co-ordination between pMDI actuation and inhalation.

Leverage Veeda's Inhalation Capabilities

Dedicated team with proper understanding of Inhalation studies and its challenges, for clinical development phase

State-of-the-art Negative Pressure Rooms

- Change room 1 and 2 at 25 Pa
- Dosing room 1 and 2 at 10 Pa (capacity of 3 persons at a time in each room)

Training of volunteers on placebo inhalers, aerosol inhalation monitors (AIM), in-check Dial meters and 2-dose devices to ensure precise dosing and study compliance

Database of trained Volunteers - hence increasing study compliance

High-end sensitive equipment which ensure measurement of LLOQ – upto 0.1 pg, hence addressing dosing and PK requirements

Inhalation: Experience

Completed 37 studies with more than 1500 volunteers

Type of studies	Number of studies	Number of volunteers
Pressurized metered-dose inhalers (pMDIs)	16	851
Dry powder inhalers (DPIs)	16	557
Nasal sprays	3	86
Activated charcoal suspension studies	2*	48

* Part of pMDI/DPIs study

Inhalation: Bioanalytical Capabilities



Drug Name	Therapeutic Class	Matrix	Anti-Coagulant	Equipment	LLOQ	ULOQ
Fluticasone Propionate	Corticosteroid, Asthma	Human Plasma	K ₂ EDTA	LCMS-8060	0.80 pg/mL	500 pg/mL
Formeterol	Asthma	Human Plasma	K ₃ EDTA	LCMS-8060	0.4 pg/mL	200 pg/mL
Tiotropium	Anticholinergic	Human Plasma	K3EDTA	LCMS-8060	0.20 pg/mL	100 pg/mL
Budesonide	Glucocorticoid	Human Plasma	K3EDTA	LCMS-8050	10 pg/mL	8000 pg/mL
Salmeterol	Asthma	Human Plasma	K3EDTA	LCMS-8050	1.0 Pg/mL	500 Pg/mL
Mometasone	Corticosteroid, Asthma	Human Plasma	K3EDTA	LCMS-8060	0.20 pg/mL	30.0 pg/mL
Ipratropium	Asthma	Human Plasma	K3EDTA	LCMS-8060	0.60 pg/mL	180 pg/mL
Formoterol(Sensitive)	Asthma	Human Plasma	K ₃ EDTA	LCMS-8060	0.2 pg/mL	100 pg/mL
Becloamethasone Dipropeonate + Beclomethasone 17-monopropionate	Corticosteroid, Asthma	Human Plasma	K ₃ EDTA	LCMS-8050	10.0pg/mL	5000pg/mL
Fluticasone Furoate + Vilanterol	Corticosteroid, Asthma	Human Plasma	K ₂ EDTA	Sciex 6500+	0.5pg/mL 1.0pg/mL	125pg/mL 500pg/mL
Budesonide(Sensitive)	Glucocorticoid	Human Plasma	K3EDTA	Sciex 6500+	3.0 pg/mL	8000 pg/mL

Inhalation - Patient trial capabilities

- Team Experience in Conducting two clinical endpoint phase III studies in moderate asthma. Around 650 patients were enrolled in these two studies & More than 60 active sites were involved
- Veeda Experience: We have conducted start-up phase till DCGI approval for one pilot study in COPD recently for Tiotropium Bromide inhalation spray, 2.5 mcg BASE/INH
- Database of more than 40 pulmonologists including Centralized spirometry in COPD, moderate asthma patients and respective hospital sites are having registered IRBs
- 25 sites possess experience in conducting clinical end point studies while around 15 sites have experience of doing patient PK studies
- A good selection of vetted and authorised providers of clinical services (i.e. temp controlled shipments for IMP and biological samples, GMP compliant repackaging services for IMP, NABL accredited lab for pathology service, translating agencies for ICD translation in regional languages)

Long Acting Injectable (LAI)



LAI: experience

- Till date Veeda CR has completed **16** BA/BE studies involving Long Acting Injectables (LAIs)
- Experience in understanding the challenges, clinical development, study design, and execution of LAI antipsychotic drugs like
 - Aripiprazole depot injection
 - Olanzapine modified release injection
 - Paliperidone palmitate modified release injection
 - Risperidone modified release injection
 - Leuprolide acetate injection

Drug synopsis	No. of subejcts	No. of Periods
Methylprednisolone Acetate Injectable Suspension USP 80 mg/ mL	54	2
Methylprednisolone Acetate Injectable Suspension USP 80 mg/ mL	24	2
Methylprednisolone Acetate Injectable Suspension USP 80 mg/ mL	24	2
Methylprednisolone Acetate Injectable Suspension, USP 80 mg/ml	180	1
Octreotide acetate injectable suspension 30 mg	48	1
Octreotide Acetate Injectable Suspension 30 mg	32	1
MPA 80 mg/ml Injection	120	3
Fulvestrant intramuscular (IM) Injection 250 mg/5 mL	12	1
Fulvestrant intramuscular (IM) Injection 250 mg/5 mL	48	1
Fulvestrant IM Injection 50 mg/mL	48	1
Fulvestrant Inj.	214	1
Naltrexone ER Injectable Suspension, 380 mg	300	1
Paliperidone Palmitate Prolonged-Release Injectable Suspension for injection 25 mg, G-01	290	1
Pegfilgrastim (6 mg/0.6 mL) in Single-dose prefilled syringe G-01_Part-01	426	3
Filgrastim 5 µg/kg SC Injection	201	2
ADL-018(XOLAIRrecombinant DNA-derived humanized IgG1 monoclonal antibody)	306	1

Normal injectable : experience

Drug synopsis	No. of subejects	No. of Periods
Phytonadione Injection 10 mg/1 ml	69	
Iron Sucrose injection 20mg-mL-Fasting	256	Total 08 Group
Iron Sucrose Injection 20mg/mL Fasting	48	
Iron Sucrose Injection 20 mg/mL fasting	25	
Phytonadione Injectable Emulsion, USP 10 mg/mL fasting	15	
Phytonadione Injectable 1 mg/0.5 mL fasting study	54	
Iron sucrose Injection 100 mg/5 ml- Fasting(Gr-01)	180	Total 06 Group of 30 each
Phytonadione Injectable Emulsion, USP 1 mg/0.5 mL	48	
Phytonadione Injectable Emulsion, USP 10 mg/1 mL	48	Two group
Phytonadione Injectable Emulsion, USP 1 mg/0.5 mL	52	
Paliperidone Palmitate Prolonged-Release Injectable Suspension (25 mg)	290	Total 11 Group
Phytonadione Injection 1 mg/0.5 ml	18	
Phytonadione injectable emulsion 1 mg/mL	32	

505 B2 Studies



Bridging Approaches to support 505(b)(2) Applications



Clinical Studies

- Single & Multiple dose BA / BE
- Dose proportionality
- Pharmacokinetic / Pharmacodynamic
- Food effect
- Safety / Efficacy studies
- Drug drug Interaction.
- Single ascending dose / Multiple ascending dose.

Pre Clinical Studies

- Pre clinical

In Vitro Studies

- In vivo – Bio waiver
- In vitro dose dumping studies
- In vitro PD studies

505(b)(2) Veeda experience

Veeda CR has been a partner in supporting 505(b)(2) applications with ~51 studies experience with various clients.

505(b)(2)	Test	RLD	Design
Salt change	Drug hemitartrate. Tablets	Drug mesylate Tablets	Single dose BE
Change in formulation & dosage form	Drug 300mg ER tablets	Drug 150 mg IR capsules (2x150mg)	comparative BA
Change in formulation & strength	Drug sublingual tablets 0.6 mg	Drug Tablets 1mg	comparative BA
Change in formulation	Drug ODT 2 mg	Drug Tablets (2 mg)	Single dose BE

505(b)(2) Veeda experience

505(b)(2)	Test	RLD	Design
FDC	Fixed dose Combination of statin and cholesterol-lowering Agent	Individual Formulations of statin + cholesterol-lowering Agent	Single dose BE
FDC	Fixed dose Combination of statin and cholesterol-lowering Agent	Individual Formulations of statin + cholesterol-lowering Agent	Single dose BE
Change in formulation	Statin Drug oral suspension 20mg/5ml (total dose - 80 mg)	Drug tablets	Single dose BE
Change in formulation	Drug 20 mg Soluble Tablets	Drug Tablets 2.0 mg (2.0 mg X 10)	Comparative PK Study
Strength change	Drug 600 mg PR tab	Drug XR tablets 200 mg (3 tablets X 200 mg)	Multiple dose BE

Veeda Expertise – 505(b)2

- Our team is experienced in study design, having carried out more than 45 investigations, and employs a tailored strategy to satisfy each study's particular requirements
- Our team can help navigate the complex Regulatory pathways and ensure timely approvals
- Our strong team of Bioanalytical scientists can develop complex validated methods within 4-6 weeks

Patch Studies



Transdermal Patches

Veeda has experience in conducting Transdermal patch studies Bioequivalence (BE) with pharmacokinetic (PK) endpoints and adhesion study, Skin irritation and sensitization study (Proof of Procedure). We have successfully completed **15** Transdermal Patch Clinical Studies (5 Pivotal study, 2 Pilot study, and 7 PK endpoint and Adhesion trials).

Drug synopsis	No. of subjects	Formulation
Iodex (Tingle) topical products containing counterirritants	80	Topical Cream
Doxepin 5% Topical Cream	48	Topical Cream
Ivermectin Cream 1 %	80	Topical Cream
Estradiol Transdermal System USP, 0.1 mg/day	64	Transdermal system
Rotigotine ER Transdermal System	24	Transdermal system
Methylphenidate ER Transdermal System	18	Transdermal system
Estradiol 0.1 mg/day Transdermal System USP	64	Transdermal system
Ethinyl estradiol and Norelgestromin Transdermal System	36	Transdermal system
Rivastigmine 9.5 mg/24 hours Transdermal System	60	Transdermal system
Ethinyl estradiol and Norelgestromin Transdermal System, 0.035/0.15 mg per	66	Transdermal system
Rivastigmine Transdermal System, 9.5 mg/24 hours	12	Transdermal system
Scopolamine Transdermal System, 1mg/72 hours	36	Transdermal system
Ivermectin Cream 1%	18	Topical cream
Lidocaine and Prilocaine cream (Lidocaine 2.5% and Prilocaine 2.5%)	16	Topical cream
Diclofenac gel 2.32%	128	Topical gel

Rectal Dosage Form



Rectal Dosage Form

Veeda has an experience of successfully completing 6 studies (2 Pilot & 4 Pivotal)

Drug synopsis	Sample size (Subjects recruited)	No. of subjects completed/evaluable
Artesunate Rectal Capsules 100mg	120	98
Mesalamine 1 gm Rectal Suppository	36	32
Mesalamine 1 gm Rectal Suppository	86	72
Artesunate Rectal Capsules 100mg	72	69
Budesonide rectal foam 2mg/actuation	18	18
Diazepam Rectal Gel 10 mg delivery system (5 mg / mL)	28	28

Veeda's proficiency in Rectal dosage studies

- Subject Selection: Apart from General screening, per rectal examination and proctoscopy done by General Surgeon before examination thereby ensuring appropriate subject selection
- Rectal drug administration Procedure: Well-documented administration procedures in protocol/study manuals
- Monitoring Subject Compliance

Phase 1



Phase 1

Total Phase I Studies Experience: 26

Completed: 25

Ongoing: 1

- Total 30 bedded Phase-I capacity, spread across two units.
- Well developed **12 bedded Phase 1 unit** to support Phase 1 studies.
- Additional **18 bedded Phase 1 unit**; operational since February 2021.
- Team of scientists having **in-depth knowledge and experience** of handling Phase 1 studies.

Type of Studies	No. of Studies
NAV	7
NAV+Patient	1
SAD Studies	6
MAD Studies	3
Safety and Immunogenicity study with Multiple-dose regime (with only one dose)	1
Drug Interaction Studies	3
Glucose Clamp Studies	1
Food Effect Studies	2
MD+Safety+Tolrability+PK+PD+Patients	1
Relative BA	1

Phase 1 Experience

Drug synopsis	No. of subjects	No. of Periods
Valsartan 320mg gastro retentive (GR) extended release (XL) tablets	6	Three
Paracetamol 1000mg	16 (M) +12 (F)	Four
IPV-Open Label Phase 1 Clinical Study for Evaluation of Safety and Immunogenicity of Sabin based Inactivated Polio Vaccine in Healthy Adult Human Male Subjects	12+12	Three
MK-0822 25mg tablets-An Open-Label, 2-Period, Fixed-Sequence Study to Assess the Effects of Multiple Oral Doses of Diltiazem, a Moderate CYP3A4 Inhibitor, on the Single-Dose Pharmacokinetics of MK-0822 in Healthy Volunteers	12	Two
TRC4186-An Open-label, Food and Formulation Effect, Single-dose Study (Part A) and a Double-blind, Placebo-controlled, Randomized, Multiple-dose, Dose-ascending Study (Part B) to Evaluate the Safety, Tolerability and Pharmacokinetics of TRC4186 Administered Orally in Healthy Male Subjects	24	Three
DRL-16536-A randomized, double-blind, placebo controlled, parallel group, multiple dose study to evaluate the safety, tolerability and pharmacokinetics of DRL-16536 following oral once daily dose administration, in healthy, adult, male, human volunteers	12	-
TRC4186 -MAD Gp 03	9	Single
Erythropoietin 0.5/1.5mcg/kg-GP01-A Randomised, Double-Blind, Placebo-Controlled, Single Dose Study to Assess Safety, Pharmacokinetics and Pharmacodynamics of Polysialylated Erythropoietin Administered Subcutaneously to Healthy, Adult, Male Subjects	16	Single
E3410548	30	Five in one
Valsartan 160mg	39	Two
MK-0859 (Treatment A)-A Study to Evaluate the Effect of Multiple Oral Doses of Diltiazem on Single-Dose Pharmacokinetics of MK-0859 in Healthy Volunteers	10	Two

Phase 1 Experience

Drug synopsis	No. of subejcts	No. of Periods
SPA 100 (aliskerin/amlodipine 300/10 mg tablets)	36	Two
Aliskiren (SPP100)/ amlodipine/ hydrochlorothiazide 300/10/25 mg tablet	36	Two
SPP100 (aliskiren) 300 mg Tablets	8	NA
AZD+D28:N281656 Oral solution-A Randomized, Single-Blind, Placebo-Controlled, Phase I Study to Assess the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics after Multiple Oral Doses of AZD1656 in Subjects with T2DM Treated with Metformin	6	NA
0.8 mg recombinant salmon calcitonin/200 mg 5-CNAC + Paracetamol 1 G + Ibuprofen 600 mg	28	Four
ADV-1002401 oral solution-A First in Human, Placebo-Controlled, Randomized, Double-Blind, Rising Single Dose Study of ADV-1002401 to Evaluate Safety, Tolerability, Pharmacokinetics, and Pharmacodynamics in Healthy, Adult Volunteers and Adult Type-II Diabetic Volunteers	6	one
P3914 Tablets, SAD study, G01, Part A-Randomized, Double-blind, Placebo-controlled Phase I-Ib Study of P3914 to Evaluate the Safety, Tolerability, Food effect & Pharmacokinetics in Healthy Male Subjects and Efficacy & Safety of P3914 in Patients With Acute Dental Pain	6	One
SMRX 11 Injection, SAD FIM G01-Open Label, Placebo-Controlled, Single Ascending-Dose, Phase I Safety Study of SMRX 11 (Clot Specific Streptokinase) to Determine Pharmacokinetics and Tolerability in Healthy Male Subjects	4	One
Granulocyte Colony Stimulating Factor (Gr 01)	6	2
Sun_Phase 1 AS012 , Cohort 2A, MAD, 100 mg, Set-03	3	3
AXA1665	16	4+2
Insulin Glargine injection 100 IU/MI	48	4
MKP 10241 Suspension Cohort 1	72	One
AB1001 Topical Gel (Cohort 1)	15	NA
Pharmacokinetic Interactions Between HRF-10071 and Tenofovir alafenamide/Emtricitabine	18	3

Infrastructure

Scale and Range

- **54 LC-MS/MS machines**
 - Insignia (31), Vedant (16) and Satyamev(07)
 - API 6500/5500/4000/4500/3200/3000/2000
 - Shimadzu 8060/8050/8040
 - Quattro Premier
- **2 ICP-OES**
- **Watson LIMS**
- **BSL-2 Laboratory**

Storage Capacity



Plasma Sample:

- 41 Deep freezers of -80°C (1 M samples capacity) and 12 Deep freezers of -20°C (0.15 M samples capacity)
- 01 Cold Room -20C (0.3 M samples capacity)



IP Storage:

- 6 Walking type stability chambers with overall capacity to store 74,000 Ltr for retention at room temperature
- 5 Humidity chambers with overall capacity of 4,200 Ltr
- 4 Pharmaceutical refrigerators having storage capacity of 11,350 Ltr at 2-8 °C

Experience

Capabilities

Total available Bioanalytical methods are more than 1221

995 + 20

Generics +
Pharmacodynamics/
Immunogenicity

108

Complex
Generics

98

NCEs

Salient Features

- Average processing capacity of 1,00,000 samples per month
- Central Bioanalytical Laboratory for global Phase II/ Phase III trials

Types of Methods

- Capability to develop methods with lowest quantification level- up to 0.1 pg
- Methods developed for:
 - Endogenous molecules
 - Amino Acids (Multiple analysis in single injection)
 - Hormones
 - Steroids
 - Inhalation formulation
 - Elemental Bioanalysis (Other matrix- Urine)
 - Immunogenicity
 - Large molecules/ECLIA/ELISA
 - Chiral and Liposomal
- Tissue distribution studies.

Complex Methods Experience

- **Iron Sucrose:** For Transferrin bound iron the serum samples are filtered through SPE cartridges to remove free and formulation bound iron while the filtrate contains TBI which is further analyzed by ICP OES.
- **Peptides (small molecules) by LCMSMS:** sensitivity and extraction issues
 - Desmopressin
 - Leuprolide
 - Octreotide
- **Biomarker analysis - α 1 Acid Glycoprotein – AAG:** Method HPLC-UV, large molecule (biomarker) validated method for clinical support.

Large Molecules



Large Molecules: Bioanalytical Experience

- Veeda has recently developed and validated below large molecules as per current EMEA guidance using commercially available kits by ELISA technique of G-CSF, Insulin Aspart, C – Peptide and PTH
- Immunogenicity validation study sample analysis was done for Denosumab, Teriparatide and Romiplostim
- Developed these large molecules using validated methods study samples were analysed as below:

Sr. No.	Analyte	No. Samples Analyzed	No. of Samples Analyzed for ISR	% of ISR Samples within Acceptance
1	G-CSF	2142	158	98.70%
2	Insulin Aspart	2139	158	94.90%
3	C- Peptide	2400	176	98.20%
4	PTH	340	34	88.33%

Large Molecules: Bioanalytical Experience

- Completed Enoxaparin PD end point and Immunogenicity studies (03 studies for US FDA, 01 study each for EU and ANVISA)
- Pharmacodynamic endpoint-Anti-Xa detection and Anti- 1Ia detection using Chromogenic method
- Immunogenicity-Anti heparin PF4 screening assay & Total Ig GAM screening assay using ELISA technique
- TFPI detection using ELISA technique, Linearity range: 0.050 IU/mL to 0.200 IU/mL
- Heparin Clotting assay Coagulation method, Linearity range: 0.100 IU/mL to 1.00 IU/mL
- IgG, IgA&IgM Screening assay using ELISA technique
- Total no of sample Analyzed PK-14784, PD-3866 & Immunogenicity - 3783 samples

Large Molecules by LC/MS/MS:

Human Insulin: Insulin Glargine Method Under development by LCMSMS, approach is to have intact molecule analyzed in MS (SH-8060) [As Multiple molecule MRM).

- Proposed Linearity Range - 50 pg/mL-10000 pg/mL/
- The other analogs of insulin (Glargine, aspart are also planned to develop).
- The other Insulin analogues like Insulin Aspart is also in the pipeline.
- Insulin Aspart- validated method available for Insulin Aspart. Linearity range- 100 to 5000 pg/mL.
- Other Methods – planned, under validation- Insulin glargine + Metabolites (M1 and M2)

Recognitions



Recognitions

Celebrating
19 YEARS
of excellence in Clinical Research

Organization	Award Category
	Best Clinical Research Organization - India
	Clinical Trial Company of the Year
	Bharat Udhdyog Ratan Award in Clinical Research

Organization	Award Category
	Top CLRO Company
	Best Quality Clinical Research Services in India



Organization	Award Category
	National Excellence Award
	Best Pharmaceutical CRO
Health & Safety Awards	Best Clinical Research- India
	Best Clinical Research- India
	Mark of Excellence
	Indian Clinical Research company of the year

Organization	Award Category
	Best Quality Clinical Research Organization in India
	Best Quality Clinical Research Organization in India
	Indian Clinical Research company of the year

Organization	Award Category
 Excellence in Science & BRINGING TECHNOLOGY. ENABLING SCIENCE.	MS Excellence in BABE Services, Largest Indian CRO

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Partners in creating a healthier tomorrow

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