

30 years of Putting Science to Work



A Global CRO/CDMO

- Integrated Drug Discovery, Development and Manufacturing service provider
- Small Molecules and Biologics, ADCs, Oligonucleotides
- Listed on Indian Stock Exchanges (NSE and BSE)



Scientific Ecosystem

- 2 Mn sq. ft. world-class R&D and Manufacturing infrastructure
- 5200+ qualified scientists
- Ongoing \$510Mn (423.15 Mn Euro) investment program
- Highly effective supply chain practices



IP Position

- IP assigned to clients
- Strong track record of Data Integrity and Security
- Over 400 patent assignments by clients recognizing Syngene



Marquee Clients

- 400+ active clients last FY
- Partnering with large / mid-size / emerging BioPharma (EBP) and other industries
- Clients concentrated in US, Europe & Japan
- Track record of working with diverse industry sectors



Quality Focus

- Quality driven organization
- Excellent track record of compliance with global regulators
- US FDA, EMA, ANVISA and PMDA approved, GLP Certified, Ministry of Health of Russian Federation, AAAI AC & CAP accredited/certified facilities
- 15+ regulatory and 160+ client audits in the last 3 years



Track Record

- Collaborations and partnerships to deliver numerous clinical candidates
- Delivery history for integrated CMC programs towards FIH and beyond



Integrated Drug Discovery

Across multiple therapeutic areas and modalities to drive innovation and speed

Therapeutic Areas

- Oncology
- Immunology
- Neuroscience
- Infectious diseases
- Cardiovascular and metabolism
- Inflammation
- Rare/orphan diseases

Modalities

- **Small molecules**
- **Peptides**
- Oligonucleotides
- **Antibodies**
- Antibody drug conjugates
- **PROTACs**
- CAR-Ts

Drug discovery at a global scale

1600

600,000

scientists, with 95%+ Ms/Ph.D.

sq. ft. of lab space

200+

global clients

Efficiency through co-localization of team members and capabilities

- IDD programs for 8 clients over 6 therapeutic areas
- Syngene-Synvent for therapeutic discovery on behalf of clients
- High rate of repeat business and return-clients



Realize Your Next

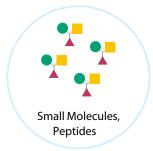
Drug candidate through our Chemistry, Biology, Safety/Tox and Computational and Data Sciences

- 95% on time delivery rate for known target compounds, >90% for unknown targets
- 25,000 reagents and 8,000 unique scaffolds in stock
- LCMS lab with ability to perform 96000 analyses/year and access to NMR lab to run 25000 samples/ year
- High-throughput mass based preparative HPLC lab for purifications with current capacity of 30000 compounds/year
- Completed ~40 PROTACs libraries (>98% success rate)
- 5000+ compounds in HT ADME assays / year
- 2500+ rodent (n=3) PK studies / year
- ADME studies turnaround in ~10 calendar days with >95% TAT adherence
- Computational and Data Sciences support across multiple stages of the discovery paradigm
- Effective use of CADD, cheminformatics and AI for multiparametric optimization



Our Discovery Toolbox

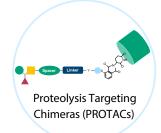
We have the capabilities to apply the most appropriate modality for target validation and therapeutic discovery

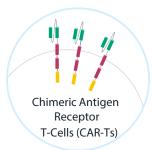














Our End-to-End Discovery Platform

Our end-to-end platform allows us to offer integrated as well as standalone services across the discovery continuum

		Target Identification and Validation	Hit Identification	Hit to Lead	Lead optimization	IND enabling
SynVent (Integrated Drug Discovery)	Biology	Pathway analysis Omics In silico target ID/validation	In vitro assays: Biochemical Orthogonal HTS Formats	<i>In vitro</i> biology / pharmacology In vitro ADME	<i>In vivo</i> PK/PD, Efficacy	Later translational PK/PD/Efficacy
		Cell en				
	Chemistry	Library Synthesis Hit validation Resynthesis Series Qualification Prioritization	HTS/DEL/ Fragments/ Virtual Screening	Optimization	Tox-suitable Formulation (maximize exposure)	Route scouting Scale up
			Medicinal chemistry Synthetic chemistry			Pre-clinical formulation
	Safety assessment			In silico	Early screening: In vitro tox, Genotox, hERG, Ion channels, Safety Pharm (in vivo)	DRF Tox (R/NR) GLP Tox (R/NR) Safety Pharmacology Bioanalysis/TK
	Computational and Data Sciences	Integrative omics analysis, Target assessment, Disease Models	Virtual Screening, Cheminformatics, Focused libraries, Structure/Ligand/Fragment based design, Predictive models, multiparameter optimization, biologics engineering			Quantitative systems models, Translational informatics, Patient response

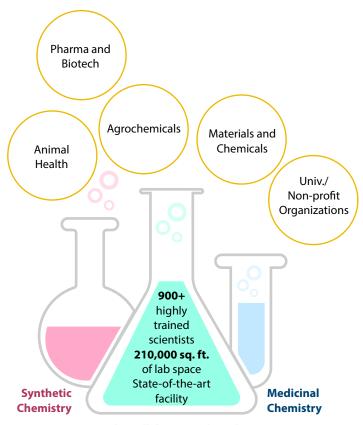
We deliver our services (both Integrated & Functional) via a range of proven collaboration models



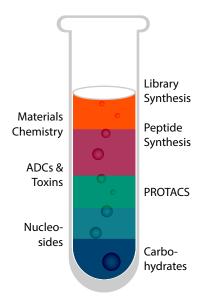


Discovery Chemistry at Syngene

Discovery chemistry at Syngene is delivered by 900+ highly qualified scientists, backed by strong analytical chemistry support, robust IT infrastructure and ELN, as well as efficient data and compound management



Integrated & collaborative drug discovery Seamless co-ordination between Chemistry, Biology, DMPK and animal studies



Analytical facilities supported by a team of 80 people Strong IT support for ELN, data and compound management



Discovery Biology at Syngene

The Discovery Biology team supports drug discovery across multi-modal platforms including small and large molecules as well as cell and gene therapy, across multiple therapy areas. Our diverse team of scientists solve complex biological problems in an integrated fashion, embracing scientific excellence, operational efficiency and commercial competitiveness.



650 +scientists



110,000

sq. ft lab space State of the art infrastructure

















Syngene Safety Assessment Services

Full range of in vivo and in vitro toxicology services for comprehensive non-clinical development of drugs

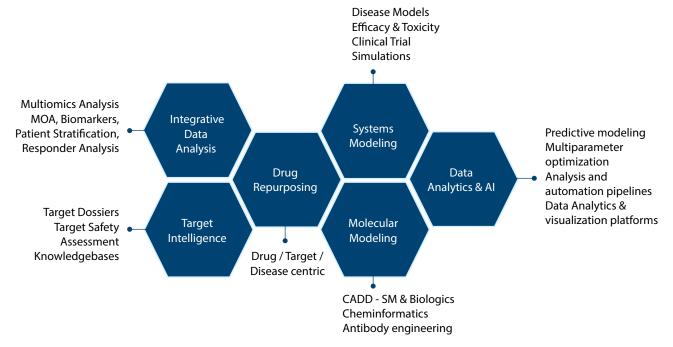
Exploratory studies	GLP studies	Phase I	Phase II	Phase III	Specialty studies
Diamental in the	Analytical and bioanalytical studies to support safety tox program	Repeat dose studies & Genetic toxicology	Sub-chronic toxicity study		In silico • Derek & Sarah Nexus
Pharmacokinetics	Repeat Dose Toxicology (2 Species)				In vitro Cytotoxicity Skin Absorption Skin
(MTD/Dose Escalation/Short- term repeat dose toxicology)					
	Genotoxicity (In vitro/ In vivo)	Repeat dose studies & Genetic toxicology	Reproductive Toxicology (Male fertility/ Pre and Postnatal development)		Skin Irritation Skin Corrosion Eye Irritation Phototoxicity
	Toxicokinetics		•		<i>In vivo</i> • Impurity
Genotoxicity screening (in vitro)					Qualification • Medical Device Testing - ISO10993
(III VILIO)	Core Battery Safety Pharmacology		Chronic Toxicology & Carcinogenicity		Dermal Tox - Rat & Rabbit
Safety					 Local tolerance/ irritation - Rat & Rabbit GPMT/LLNA Intravenous infusion - Rat Phototoxicity
Pharmacology: CNS, Respiratory, CV	Reproductive (Seg I-III) & Juvenile Toxicology				



Computational and Data Sciences Capabilities

Syngene's Computational and Data Sciences capabilities enable informed decisions at all stages of the drug discovery continuum











About Syngene

Syngene International Ltd. (BSE: 539268, NSE: SYNGENE, ISIN: INE398R01022) is an integrated research, development and manufacturing services company serving the global pharmaceutical, biotechnology, nutrition, animal health, consumer goods and specialty chemical sectors. Syngene's more than 5200+ scientists offer both skills and the capacity to deliver great science, robust data management and IP security and quality manufacturing at speed to improve time-to-market and lower the cost of innovation. With a combination of dedicated research facilities for Amgen, Baxter and Bristol-Myers Squibb, as well as 2 Mn sq. ft of specialist discovery, development and manufacturing facilities, Syngene works with biotech companies pursuing leading-edge science as well as multinationals, including GSK and Merck KGaA.

For more details, visit www.syngeneintl.com or write to us at bdc@syngeneintl.com







