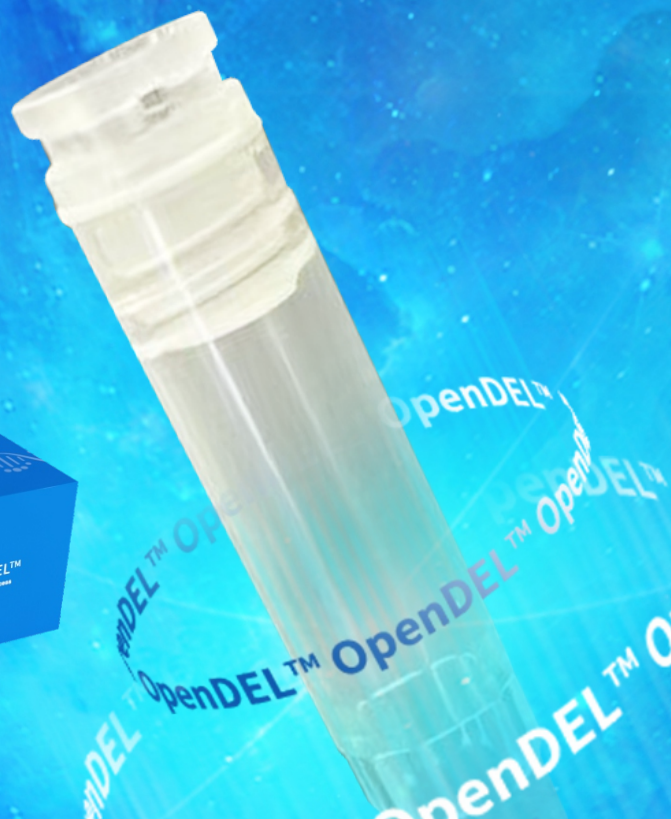




HitGen OpenDEL™

Affordable Access to Billions of
Novel Molecules & Data for AI/ML
in Drug Discovery Research





OpenDEL™ – Fully Transparent Open Access

▶ Access to

- Small Molecule Structures
- Building Blocks
- Scaffolds
- DNA Codons
- Synthetic Schemes
- Selection Manual

▶ No Structure Disclosure Fee

▶ No Compound IP License Fee

OpenDEL™ to Help Your Research

▶ For AI/ML:

- Post-selection DEL data for AI/ML

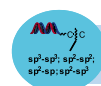
▶ For Drug Discovery:

- Rapid and efficient target ligandability evaluation
- Novel Hit discovery

■ Content of OpenDEL™

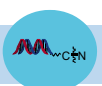
- ▶ 3 Billion Compounds
- ▶ 50 Encoded Libraries
- ▶ 10 2-Cycle Libraries
- ▶ 40 3-Cycle Libraries
- ▶ No 4-Cycle Libraries
- ▶ Physical Materials
- ▶ Selection Manual
- ▶ Synthetic Schemes
- ▶ Building Block lists
- ▶ DNA Codon lists

■ Chemistry Diversity



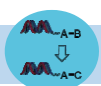
C-C Bond Formation

- Indole C3 alkylation
- Suzuki coupling
- Photoredox
- Sonogashira coupling



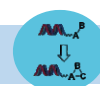
C-N Bond Formation

- Acylation
- Reductive Amination with Aldehyde
- Reductive Amination with Ketone
- Buchwald-Hartwig Cross Coupling
- SN_2/SN_A
- Sulfonylation



Functional Group Transformation

- NO_2 to Amine
- Azide to amine
- Amine to azide
- Halide to azide
- Halide to amine
- Halide to alkyne
- Halide to acid
- Halide to cyanide
- Aldehyde to alkyne
- Alkene to aldehyde
- Thioether oxidation



In-situ Heterocycle Formation

- Triazoles
- Imidazolindinone
- Pyridones
- 1,2,4-oxadiazole
- isoindolinone
- Benzimidazole
- Benzotriazole
- Indazolone
- 2-pyridinone



■ Building Block/Scaffold Diversity

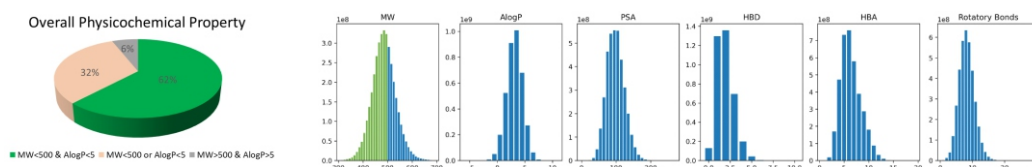
Mono-functional group BBs: >20,000

Bi-functional group BBs: >3,000

Novel scaffolds: >550

BBs: amines, acids, aldehydes, boronates, protected amino acids, free amino acids, amino esters, diamines, acid-aldehydes, acid-aryl-halides, etc.

■ Physicochemical Property



	Average MW (Da)	MW<500 & AlogP<5	MW<550	MW<500	MW<450
2-Cycle DELs	414	88%	99.7%	95%	74%
3-Cycle DELs	484	62%	91.5%	66%	23%

■ OpenDEL™ Service

HitGen provides full technical support and/or on-demand services for the entire drug discovery process



■ OpenDEL™ Access Model

	OpenDEL™ Kits	OpenDEL™ Selection Manual	On-DNA Reference Compound	OpenDEL™ Selection	Sequencing (NGS)	Data analysis & Hit Proposal	Off-DNA Synthesis	Binding/Functional Confirmation
OpenDEL™ Taster	4 selection samples ✓	✓				✓		
OpenDEL™ Standard	10 selection samples ✓	✓				✓		
OpenDEL™ Premium	10 selection samples ✓	✓	(If having) ✓	✓	✓	✓		
OpenDEL™ Premium Plus	10 selection samples ✓	✓	(If having) ✓	✓	✓	✓	Up to 10 compounds ✓	Up to 10 compounds ✓



Customer Testimonial

“

HitGen has created one of the most exciting new DEL products. Septerna was initially attracted to the openness and flexibility with the OpenDEL model. And we have now successfully identified functionally validated hits for multiple GPCR targets from the OpenDEL libraries. Most importantly, the HitGen team has been a very open and collaborative partner throughout the process.

”

By Septerna, A US-based biotechnology company discovering and advancing novel small molecule medicines targeting G Protein-Coupled Receptors

About HitGen Inc.

HitGen Inc. (SSE: 688222.SH), is a drug discovery research company with headquarters in Chengdu, China, and subsidiaries in Cambridge, UK and Houston, USA. HitGen has established leading technology platforms to enable the discovery and optimization of small molecules and nucleic acid drugs. Our key technology platforms include world-leading DNA-encoded library technology (DEL), fragment-based drug discovery and structure-based drug design technologies (FBDD/SBDD), as well as the emerging technology platforms for synthetic therapeutic oligonucleotide technology (STO), and targeted protein degradation technology (TPD). Through our diverse and flexible business models, we have built up collaboration partnership with several hundred biopharmaceutical research organizations worldwide. HitGen has multiple programmes from early discovery to clinical trial stage.



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