

**National Biotechnology Research Park**

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(Administrative Affairs)**

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(BioHub Taiwan)**

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# NATIONAL BIOTECHNOLOGY RESEARCH PARK INVESTMENT DIRECTORY AND RESOURCES IN BIOMEDICAL ECOSYSTEM



BioTReC



NBRP

No.1 December.2022



# Table Of Contents

<b>About NBRP</b>	<b>01</b>
<b>BioTReC, Academia Sinica</b>	<b>03</b>
- Biomedical Translation Research Center	
- BioHub Taiwan	
<b>Investment Opportunities</b>	<b>09</b>
- Resident Companies	
- Resident Companies in Shared Space	
<b>Cooperative Partnership</b>	<b>48</b>
- BioHub Taiwan Accelerator	

# ABOUT NBRP



To stimulate and promote valuable and effective biotechnology research, Academia Sinica in concordance with governmental development policy, has constructed the National Biotechnology Research Park (NBRP) in Nangang, Taipei; opened in October 2018.

NBRP not only aspires to be Taiwan's premier state-of-the-art biotechnology research park but also embraces the obligation to maintain natural environment. It is hoped that NBRP can serve as a model for innovative research and development, while upholding high standards for ecological preservation.



## INSIGHT & MISSION

NBRP is the 1<sup>st</sup> biomedical research park established in Taiwan. It aims to accelerate the development timelines for new drugs serving unmet medical needs, with the ultimate goal of enhancing human health and welfare.

As a coherent engine driving pharmaceutical research and development, NBRP has built a bio-corridor for academic research and pharma R&D to maximize the efficiency across value chains. NBRP focuses on the first part of journey toward biotechnological success by creating strong links between basic research and pre-clinical studies. Pre-clinical research results are then led to further product development and mass production.

## NBRP ECOSYSTEM

NBRP serves as a single enterprise and plays a pivotal role in facilitating innovative research by closely integrating the resources of four government institutions to create an efficient cluster effect. These institutions include the Biomedical Translation Research Center of Academia Sinica (BioTReC), the Development Center for Biotechnology of Ministry of Economic Affairs (DCB), the National Applied Research Laboratories National Laboratory Animal Center of National Science and Technology Council (NLAC), and the Taiwan Food and Drug Administration of Ministry of Health and Welfare (TFDA).



# Biomedical Translation Research Center

The Biomedical Translation Research Center (BioTReC) was founded in September 2019 at the National Biotechnology Research Park with the aspiration to support the development of biotechnology in Taiwan. The objectives of BioTReC align with government initiatives to accelerate bio-entrepreneurship from early discovery stages to commercialization.

The center consists of four divisions, including the Translational Medicine Division (TMeD) Innovation Incubation Center (BioHub Taiwan), Smart Medicine Division (SMeD), and Emerging Infectious Disease Division (EIDD). The establishment of BioTReC aims to forge a professional and innovative ecosystem that facilitates new drug discovery, as creation of a thriving biotechnology industry will ultimately enhance human health and welfare.

## 01 Translational Medicine Division (TMeD)

To enhance the international competitiveness of new drug development and medical applications from translational research to clinical use, TMeD placed strong focus in facilitating industry-oriented innovation and multi-disciplinary collaboration in translational medicine. In addition, TMeD provides resident companies and domestic researchers in academia, research, and industry with advanced instruments, equipment, and technical services for translational research on disease prevention, detection, diagnosis, and treatment.

## 02 Innovation Incubation Center (BioHub Taiwan)

BioHub Taiwan consolidates resources and provides one-stop service to support resident companies during R&D stage and subsequent product commercialization with the mission to foster the growth of domestic startups and entrepreneurs. BioHub Taiwan is committed in building an innovative biomedical ecosystem by bridging domestic / foreign R&D technologies and resources to promote development of innovative biomedical ecosystems. BioHub Taiwan also places strong emphasis in integrating interdisciplinary knowledge and skill sets into the NBRP Academy to cultivate business talents and provide business mentorship to bridge the gap between fundamental research and industrial applications.

## 03 Emerging Infectious Disease Division (EIDD)

EIDD was established with the mission to pre-deploy and reserve R&D capacities to immediately respond to outbreaks of emerging infectious diseases. Besides providing P2 and P3 facility services and rapidly develops identification/rapid testing technologies and therapeutic antibodies, drugs, and vaccines, as well as next-generation technologies to fight against serious infectious diseases; EIDD also collaborates with relevant units in industry, government, academia, and research that help to build up a comprehensive strategy for the prevention and treatment of critical infectious diseases.

## 04 Smart Medicine Division (SMeD)

SMeD aims to help accelerate the R&D of big data technology transfer in the smart healthcare sector, to advance precision medicine development, and to build an industry ecosystem to improve overall social interest. SMeD will integrate the Taiwan Biobank Project, Taiwan Precision Medicine Initiative (TPMI), Cancer Moonshot Project of Academia Sinica, and other relevant big data in biomedicine, and it will also establish a big data center.

## Best-in-class Core Facilities

01  
Medicinal Chemistry and Analytical Core Facility



02  
Taiwan Mouse Clinic



03  
Core Facilities for Translational Medicine



04  
RNA Technology Platform and Gene Manipulation Core

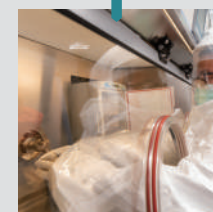
05  
Taiwan Biobank



06  
Human Therapeutic Antibody Development Platform



09  
Nucleic Acid Pilot (\*Under Planning)



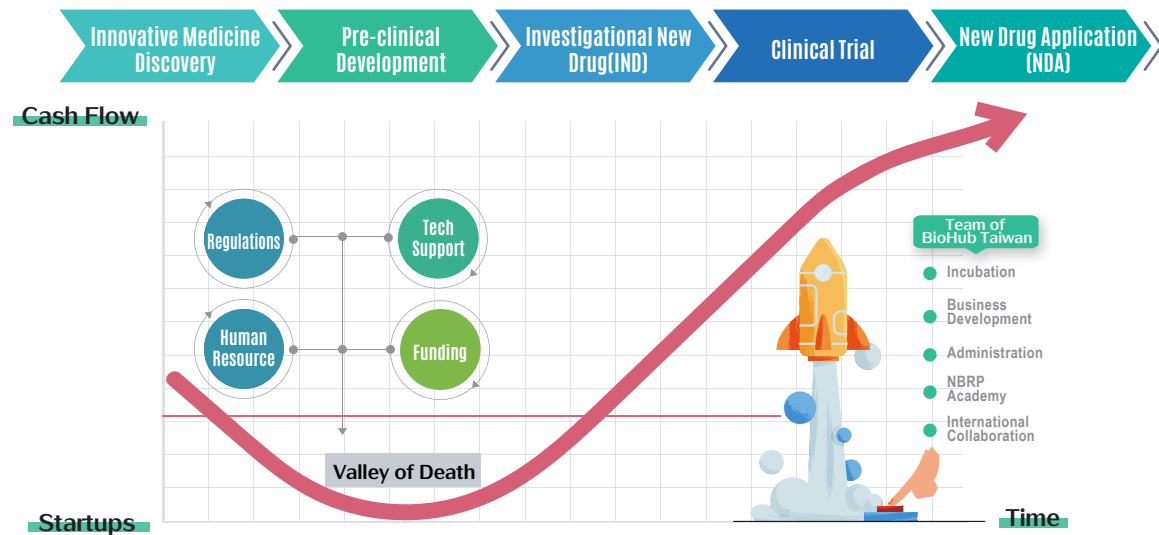
07  
Infectious Diseases Core Facility



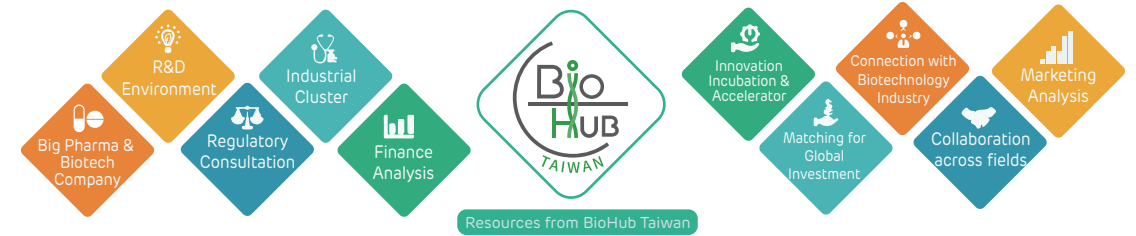
08  
Animal Imaging Facility



With the aspiration to support the development of biotechnology in Taiwan, BioHub Taiwan @NBRP was opened in Oct. 2018, in collaboration with different ministries in NBRP, to promote impactful fundamental research into commercialization. The Innovation Incubation Center of BioTReC, Academia Sinica was established in Sep. 2019 to coordinate and execute the mission of BioHub Taiwan.

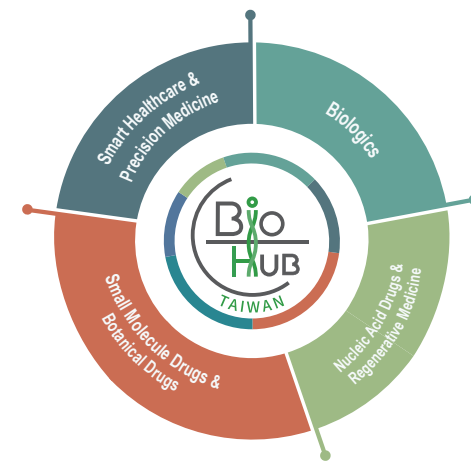


## One-stop Service for Startups



BioHub Taiwan aims to consolidate resources among different ministries, academics, and industries to provide one-stop service to support resident companies during R&D stage and subsequent product commercialization. With the mission to foster the growth of domestic startups and entrepreneurs, BioHub Taiwan is committed in building and promoting the development of innovative biomedical ecosystem by bridging domestic / foreign R&D technologies and resources. Its strategic focus yields the formation of new hotspot for biomedical industry as well as investment in Taiwan and the Asia-Pacific region.

## Resident Start-ups & Companies in NBRP



AcadeMab Biomedical	Ascendo Biotechnology	AIBIOS K.K.	CHO Pharma	Immunwork	OBI Pharma
Oneness	PapIVax	Pharmaessentia	Vacino Biomed	ZHENG YANG BIOMEDICAL	Zenith BioDesign Lab
Anbogen Therapeutics	AnHorn Medicines	ALPS Biotech	Agari Pharma	Excelsior Biopharma	Exland Biotechnology
Foresee Pharmaceuticals	Pharmasaga	SINEW PHARMA	Saint Therapeutics	Win Lux Biotech	*The One Biopharmaceutical
Acer	Bio Preventive Medicine	Horuseye Technology	i Analyzer Incorporation	JUN ZHI Biomedical	
Instant NanoBiosensors	LumiSTAR Biotechnology	LuminX Biotech	Molsentech	Nebulum Technologies	
NanoRay	SYNCELL	VITAE Biomedical			
Advanced Connection Technology	Apexcella Biomedical	DSignal BioTechnology	MELLO BIOTECH TAIWAN	Ochre Bio Taiwan	
Polaris Biopharmaceuticals	ROCK BIOMEDICAL	Topmunnity	*Elite BioPharma	*SMOBIO Technology	
BMCC	Taiwan BIO	VMIC	(*non-commercial service providers)		

\*Contract review in process  
Data as of 2nd September 2022

# NBRP Academy

To foster an entrepreneurial mindset and develop business skills for biotech start-ups, BioHub Taiwan places strong emphasis in integrating interdisciplinary knowledge and skill sets to facilitate establishment of NBRP Academy. With the aim of helping biotech talents succeed at different phases and stages, it consolidates the resources from the cooperative partner at NBRP. These institutions include the BioTReC of Academia Sinica, the DCB of Ministry of Economic Affairs, the Center for Drug Evaluation (CDE), the NLAC of National Science and Technology Council, Biotech Industrial Academy (BIA), and Amgen Taiwan.

NBRP Academy not only aims to cultivate business talents by constructing the training program on scientific innovation and entrepreneurship, but it also provides consultation services and guidance for startups to bridge the gap between fundamental research and industrial applications.

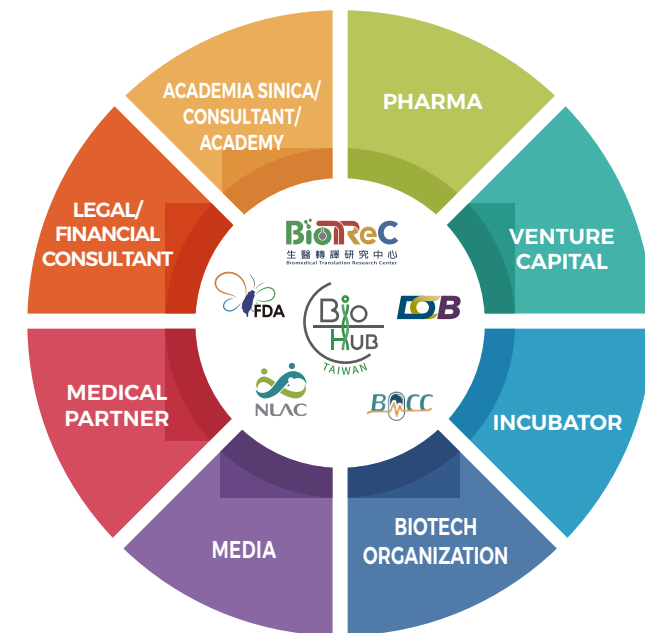


# BioHub Taiwan Accelerator



BioHub Taiwan Accelerator was found with the aim of building a competitive ecosystem for the biotech industry in Taiwan. It integrates resources among different academic, private, public and research sectors to assist with commercializing biotech discoveries and facilitating the growth and expansion of collaborations with innovative biotech companies.

It has partners coming from venture capital (VC), law firms, international pharmaceutical companies, and incubators; This innovative biomedical ecosystem can generate positive communication environment, by sharing resources and experiences, more companies can replicate the success model and create a win-win solution.



## Core curriculum

- Business Plan workshop - Industry, Venture Capital & Accounting Firm Experts in Taiwan
- IP & Market analysis workshop - Cortellis and Derwent Innovation
- Clinical Research Protocol workshop - Taiwan FDA & Center for Drug Evaluation

## Entrepreneurship counseling

- Entrepreneurship Workshops - Industry Experts
- Bio-Startup Seminar - Consultant & Industry Experts

## Experience sharing with successful entrepreneurs

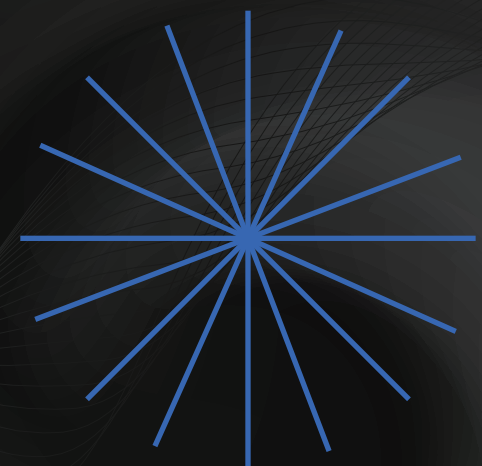
- International link & Professional mentoring - Industry Experts



NBRP BioHub Taiwan

Investment  
Opportunities

Resident Companies





# Biologics





**COMPANY DESCRIPTION**

Ascendo Biotechnology, Inc. ("ASCENDO") is an innovative biotech Cayman-registered company originated from Taiwan. The Company's mission is to develop cutting-edge approaches to revive the immunological momentum for patients with cancer and chronic viral infection to provide an opportunity to cure their diseases.

**BUSINESS DESCRIPTION**

ASCENDO focuses on the development of a first-in-the-class innate immune checkpoint inhibitor and vaccines derived from a very versatile platform from bench to bedside. The exit strategies include out-licensing, partnership, and M&A.

**TECHNOLOGY PLATFORM / PRODUCT**

**•ASD141**

- First-in-the-class innate immune checkpoint inhibitor
- Break the bottleneck of current cancer immunotherapy through preventing a novel regulatory ligand from binding to a common regulator of PRR pathways
- Significantly enrich the infiltration of functional APCs and neoantigen specific cytotoxic T in TME
- Very synergetic with approved effector cell level of immune checkpoint inhibitors such as anti PDI/ PDL1 or CTLA4 in animal models
- Can dramatically ameliorate the acute injection reactions when in combination with anti-CTLA4 in animal models
- Very good tolerability in animal pharmacology and PK studies
- Current stage: Preclinical development and plan to submit IND in one year.

**•NanoCherub®**

- A versatile Electro-Kinetic nanocomplex antigen adjuvant & delivery platform for therapeutic vaccine and preventive vaccine development

**•ASD253**

- Therapeutic vaccine for hepatitis B that can achieve hepatitis B Surface antigen seroconversion in 100% of test animals of both HDI mice model and AAV model.
- No liver function abnormality was observed in all the animals tested in HDI mice and AAV mice model.
- Current stage: pre-clinical testing

**•ASD254**

- RBD based subunit protein preventive vaccine of Covid-19
- Non-polarized Th1/Th2 immunogenicity
- Very long lasting protection and clean GLP toxicity data.

Founded in 2020, Academab Biomedical Inc. is a start-up company dedicated to the development of new antibody drugs in the medical field that requires urgent breakthroughs. As cancer is widely acknowledged as a leading cause of death across the globe, we take it upon ourselves to research on the cancer therapy as our first step. From there, we expanded to other life-threatening diseases in different fields. To this end, we have developed multiple in-house technology platforms to break fresh ground for new antibody drugs. Our mission is to improve life quality and human health by providing the first-in-class or best-in-class new antibody drugs.

With rapid breakthrough of biotechnology, drugs for cancer treatment have improved dramatically, evolving from small-molecule drugs to protein/antibody-based drugs, including antibody-drug conjugates (ADC) and CAR-T therapy. Having proprietary platforms and unparalleled technologies for the development of antibody, it can support the company's sustainability on new product pipeline, and/or to support our partners moving forward via services provided.

With comprehensive patent portfolio deployment, Academab continues to reinforce core capabilities for the R&D of antibody drugs to clinical. We are committed to the development of potential therapeutic antibody drugs, boosting our competitiveness, improving human health and wellbeing, and promoting the antibody derived biologics through global collaboration.

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CHO PHARMA, INC.  
醣基生醫股份有限公司



CHO PHARMA INC. was founded in 2013. As a pioneer in innovative glycochemistry and carbohydrate new drug development, we are devoted in providing better solutions for severe human diseases. Our core expertise is in the field of glycoengineering and the development of vaccine and antibody against novel carbohydrate antigens. Several pipeline products, which cover indications for human cancer, infection disease and autoimmune disease, have been created through our proprietary technologies CHOOptimax™ and the integrated anti-carbohydrate vaccine and antibody development platform. CHO-H01, our first product for human non-Hodgkin lymphoma, has been approved to conduct phase I/II clinical trials in both Taiwan and the US. More product candidates are expected to be launched for clinical trials in the near future. Another product, universal COVID-19 vaccine, is a new vaccine also designed by CHOOptimax™.



Immunwork Inc.  
免疫功坊股份有限公司



Immunwork, Inc. was founded in 2014. We focus on the research, development, and commercialization of novel drugs to fulfill unmet medical needs. Our drugs are created based on our proprietary technology platform for treating multiple types of cancer, diabetes, obesity, non-alcoholic steatohepatitis (NASH), pathological blood clots, and other selected severe clinical conditions. Our unique platform allows us to design and generate novel "T-E" drug molecules, which contain targeting (T) and effector (E) moieties. These designs would help to improve the efficacy and limit toxicity. The majority of our T-E drugs are based on "multi-arm linkers" and various "drug bundles". These drug candidates include antibody-drug conjugates (ADCs), antibody-radionuclide conjugates (ARCs), ultra-long-acting therapeutic peptides, and drugs in other modalities. Six of the drug candidates have been selected and progressed into the pre-clinical development stage. Meanwhile, we are looking forward to applying the multi-arm linker platform for designing additional novel drug candidates.

**1 Lead Antibodies / High Flexibility**

- Can apply to all kinds of human IgGs and human IgG Fc-fusion proteins produced from diverse systems, such as mammalian cells, insect cells and yeast.

**N-glycosylation Profile**

- Highly heterogeneous

**2 SAR Study / Glycan-based SAR Study**

- Candidates with single, well-defined N-glycan structure can be made for functional studies.

**Optimize Desired Functions or Discover New Functions**

- Common structures like G0(F), G2(F), G2S2(F), or customized structures

**3 Pharmacokinetics**

**GMP Manufacturing / Rapid and Well Defined Process**

- Modular platform technology to generate sub kilograms of functionally optimized product within 24 hours.

**High Yield & Purity**

- >95% single glycoform with limited loss during process.

**Cost- and Time-Efficient**

**4 Drug Product / Regulatory**

- Product has been approved for Phase I/II trials in US and Taiwan.

**Simplified CMC**

- No worry for batch variation caused by glycosylation.

**N-glycosylation profile**

- Homogeneous

Selected Products	Indications	Status
TE-1146, TE-1132, TE-8214	Oncology	Pre-clinical Development
TE-8105, TE-8024	Diabetes, obesity, NASH	Pre-clinical Development
TE-6168	Pathological blood clots	Pre-clinical Development

**TE-8105**  
GLP-1, Fatty acid bundle

**TE-1146**  
Anti-CD38 scFv-Fc, Lantidomide bundle

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Founded in 2002 as a 100% subsidiary of Optimer in Taiwan, OBI Pharma was publicly launched in 2012 and listed on the Taiwan Stock Exchange in 2015.

OBI Pharma has positioned itself as a new drug research and development company dedicated to developing innovative cancer therapies. Initially focusing on developing the Globo series of glycosphingolipids (Globo H, SSEA-4), OBI has transformed into a cancer pharmaceutical company with a diversified technology platform and multiple targets over the years. OBI aims to develop first-of-a-kind new cancer drugs and innovative therapies with novel therapeutic targets to meet the needs of patients who still lack effective medical treatments, improve their health and enhance their quality of life.

OBI's core competency lies in pioneering first-in-class new drugs. OBI has a complete Globo Series pipeline line, including active immunization (vaccines), passive immunotherapy (monoclonal antibodies), and conjugates of small molecule drugs with antibodies (ADCs); and also developing new drugs in the small molecule field, targeting cancers with high expression of Aldo-Keto reductase AKR1C3. In recent years, OBI has been actively developing new products targeting TROP2. In addition, OBI also has achieved good results in developing cell therapy and the COVID-19 vaccine with full potential for development.

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Oneness Biotech has been keen on employment training, new drug development and innovation of technology. Our pipelines cover Phases I, II, III, NDA and approval phases. With strong R&D team and capability for new drugs, we have set up raw material cultivation bases, API and finished dosage manufacturing plant according to GACP and PIC/S GMP standards for botanical drugs. The streamlined control ensures the effectiveness, safety, and consistency of our drug quality. Besides, we have developed antibody new drugs for severe asthma, atopic dermatitis, and other indications based on the established proprietary technologies including fully-human antibody library, a high-performance screening platform, a stable and high-yield antibody-producing cell line, and a cancer stem cell platform.

The first approved new drug: Fespixon® (for DFUs), a first-in-class macrophage-regulating topical new drug has been granted approval in Taiwan and Macao. It is also under NDA review by health authorities in China, South East Asian countries as well as India, and has been granted a Fast Track designation by the US FDA to accelerate the development for the US market. Oneness dedicates its R&D efforts to chronic dermatological and immunological disorders and has out-licensed its atopic dermatitis antibody new drug for US\$ 530 million to the global medical dermatology leader, LEO Pharma A/S in 2020. Through licensing or co-development, Oneness can provide patients with cost-effective treatments and achieve our founding mission of "developing new drugs and caring for life".

Research Code	Therapeutic Area	Indication	Pre-Clinical	Phase I	Phase II	Phase III	NDA	Market
ON101 (FESPIXON®)	Dermatology	Diabetic Foot Ulcer	Taiwan & Macau	[Progress bar]				
			China, Southeast Asia & India	[Progress bar]				
			US	[Progress bar]				
FB825	Dermatology	Atopic Dermatitis	US	[Progress bar]				
	Immunology	Allergic Asthma	Taiwan	[Progress bar]				
FB704A	Immunology	Asthma	Taiwan (US IND)	[Progress bar]				
OB318	Immunology	Hepatocellular Carcinoma	Taiwan (US IND)	[Progress bar]				
FB918	Immunology	Asthma	[Progress bar]	[Progress bar]				
SNS812	Infection	Pan-COVID	[Progress bar]	[Progress bar]				

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PharmaEssentia Corporation (TPEX: 6446) is a fully integrated biotech company based in Taiwan. The company engages in innovation, clinical development, production and manufacturing of new drugs, and global commercialization. PharmaEssentia utilized its self-developed PEG molecule and PEGylation technology platform to produce a long-acting protein drug with the advantages of prolonged efficacy time and the simplest compound. The innovative long-acting interferon P1101 (Ropeginterferon alfa-2b) was invented by PharmaEssentia. Since 2019, Ropeginterferon alfa-2b has been approved by the agencies for the treatment of Polycythemia Vera (PV) in European Union, Taiwan, Switzerland, Israel, South Korea. In 2021, Ropeginterferon alfa-2b was approved by US FDA for PV.

PharmaEssentia is currently conducting the multi-country and multi-center Phase III clinical trials of P1101 for the treatment of Essential Thrombocythemia (ET) globally. After completion of the Phase III trials, the company will submit new drug application (NDA) in various countries. The company's Taichung plant has completed commercial mass production and was GMP certified by Taiwan FDA, EMA, South Korea MFDS, and US FDA. In addition, the company's filled and finished plant was also GMP and GDP certified by Taiwan FDA.

Therapeutic Area	Candidate	Indication	Market	Preclinical	Early Phase	Pivotal	Regulatory	Approval	Partnership
Hematology	Ropeginterferon alfa-2b (P1101)	PV	EU, CH, IL	█	█	█	█	█	ADP ORBRIAN
			TW, KR	█	█	█	█		
			US	█	█	█	█		
			JP, CN	█	█	█	█		
			Global	█	█	█	█		
Hematology	Next generations of P1101	ET	Global	█	█	█	█	ADP ORBRIAN	
			Adult T-Cell Leukemia	JP, TW, CN	█	█	█		█
Infectious Disease	P1101 + Entry inhibitor	HDV	US, EU, TW	█	█	█	█	Atfenex	
Dermatology	KX01	Psoriasis	TW, CN, MO	█	█	█	█		
			SG, JP, KR	█	█	█	█		
Dermatology	KX01	Actinic keratosis	TW, JP, KR	█	█	█	█	Atfenex	
			Global	█	█	█	█		
Oncology	P1101 + anti PD-1	HCC	Global	█	█	█	█	Axis	
			PEG-IL2	Solid tumor	Global	█	█		█
			PEG-Gamma	Solid tumor	Global	█	█		█
			P1101 Combo	Solid tumors	Global	█	█		█
Oncology	TCRT	Solid tumors	US, TW	█	█	█	█	Axis	

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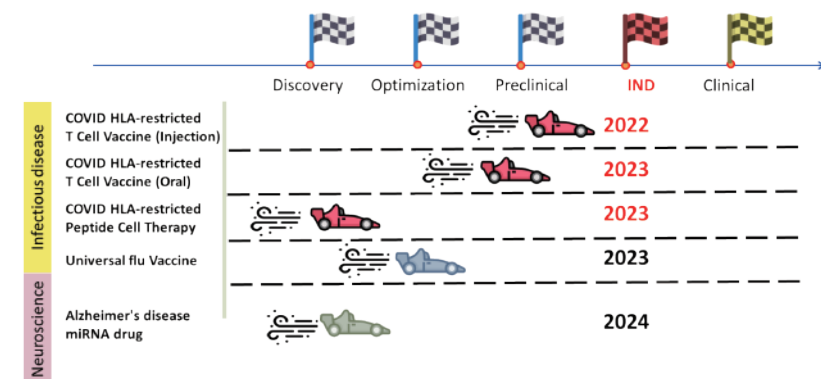


Established in July 2020, Vacino Biotech Co., Ltd. is committed to the development of nucleic acid drugs and vaccine products. With self-developed next-generation vaccine design AD tech technology, COVID-19 T-cell vaccines are put into practice to solve the short period neutralizing antibodies of B-cell vaccines with short protection. In addition, nucleic acid drug target delivery technology has been developed, which has been invested in the field of Alzheimer's treatment to increase the drug load of brain therapeutic drugs, and developed nucleic acid drugs with modular technology.

COVID-19 T-cell vaccine: Research and development of COVID-19 T-cell vaccine to produce T-cell protection against the world's major COVID-19 viruses, and through oral dosage forms, more people can get protection in the shortest possible time and build up herd immune protection. The vaccine does not require the use of adjuvants, achieves the best safety profile, and is suitable for the elderly and adolescents.

Nucleic acid drugs: Research and development of neural nucleic acid drugs to inhibit gene expression through miRNA to achieve the protective effect of brain function. Through targeted delivery technology, a new generation of nucleic acid drugs can easily increase the effective dose of drugs reaching the brain.

## PRODUCT PIPELINE



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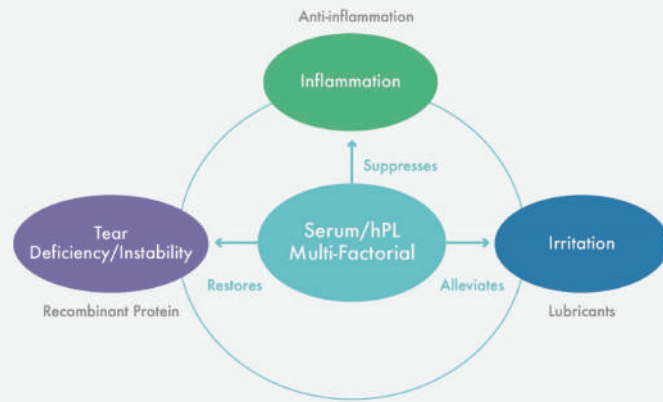
Zheng Yang Biomedical Technology Co., Ltd.  
正揚生醫科技股份有限公司



正揚生醫科技股份有限公司  
ZHENG YANG BIOMEDICAL TECHNOLOGY CO., LTD.

Zheng Yang Biomedical Technology Co., Ltd. (ZYBT) is a new drug development company, dedicated to the research and development of Elate Ocular™ dry eye biologics and new drugs. ZYBT was established in 2015 with cooperation with professors from EMORY University (Cambium Medical Technology, LCC) on cell culture supplement (FDhPL) patented product manufacturing, development and global sales rights. ZYBT cooperates with Cambium to develop new dry eye drugs and owns the global rights to manufacture its active ingredients, Aurarix™. The cell culture supplement technology extends to the treatment on dry eye. In the fourth quarter of 2017, Cambium obtained FDA approval to perform Phase I/II clinical trials, and completed Phase I/II clinical trials in 2019. Its clinical indicators and output results are very positive, current plan is to enter Phase II/III clinical trial application.

# Nucleic Acid Drugs & Regenerative Medicine →→



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Advanced Connection Technology Inc. not only has accumulated years of medical research development experience, but also connect several resources to achieve regenerative medicine, precision medicine and digital health fields; ACTT has a high standard GTP laboratory, professional cell culture specialist, to provide a variety of regenerative medicine development; Next Generation Sequencing laboratory brings all kinds of precision medicine research and development; With our SMO in the hospital, can provide resources for clinical trials and assist building data base With plenty of research experience and high level of digital connection can lower the cost and shorten the development path. Create the perfect clinical research and development services with full resources.

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DrSignal BioTechnology Co., Ltd. is a biotech-company which leads the world in building an automated cell production line and providing CDMO services as our main business model. Current relevant companies are either manufacturers developing and selling automated machines, or those providing cell manufacture OEM services with conventional GTP laboratories. Instead, DrSignal aims at the integration and standardization of the automated cell production with AI-assisted optimization, with our expertise in the biomedicine, hardware, and artificial intelligence.

## Contact US

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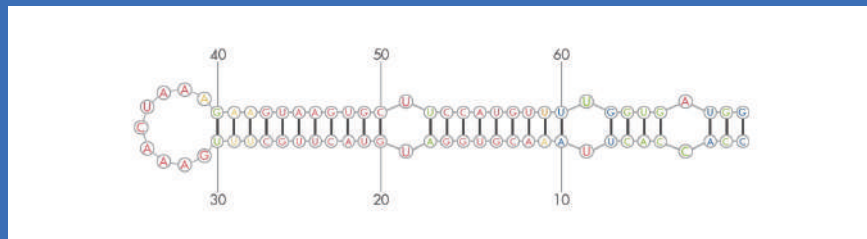
Mello Biotech is a research company specializes in development and commercialization of microRNA and stem cell technologies. Based on patented proprietary technologies in microRNA and stem cells, Mello Biotech currently focuses on the development of microRNA precursor for cancer treatment. Based on technical and commercial viability, initial indication is targeting "4th stage NSCLC patients not eligible for target therapy or immunotherapy". Current in-vivo study data demonstrates good efficacy and safety in orthotopic animal models. The company aims to validate efficacy and safety of treatment in human. Objectives include manufacturing, pre-clinical experiments, application for IND, and clinical trial phase I.

### Target Market Condition

- NSCLC: ~600,000 new patients annually in Japan, Europe & US.
- Indication: 4th stage NSCLC patients not eligible for target therapy or immunotherapy, second line after chemotherapy.
- Poor survival with current treatments – PFS in months.
- Current treatments are costly, toxic, undesirable side effects, and limited in efficacy.

### Key Advantages & Features

- Proven efficacy in animal study.
- Inherently Safe – Technology derived from stem cells, laboratory proven safe.
- Technically Feasible - Deliverable and stable with proprietary delivery technology.
- Commercially Feasible – Large patient population with urgent medical need and limited options.
- Unique MOA - Allows for combination therapy.
- Comprehensive IP Protection - global patents secured in fundamental technology, manufacturing & applications.



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Founded in 2021, Rock BioMedical, Inc. (Rock Bio) is a biopharmaceutical company seeking to combine both active immunization (eg vaccines) and passive immunization (eg antibodies) for the eradication of diseases based on our proprietary Glyco-engineered Universal Vaccine platform. The initial focus will be on the development of next generation COVID vaccine featuring the unique property of our glycan-depleted SARS-CoV-2 spike protein (SMG) to induce broad protection against immune evading variants. Built upon the same technology platform, the company's pipeline also includes novel therapeutic antibody and universal flu vaccine candidates that together will revolutionize the way such common respiratory tract infection can be managed in the future.

The company is founded upon a set of core technologies licensed from Academia Sinica. It has an R&D base located in National Biotechnology Research Park in Nangang Taipei and a subsidiary in Boston Massachusetts USA to manage its global CRO activities for clinical development and international partnership.

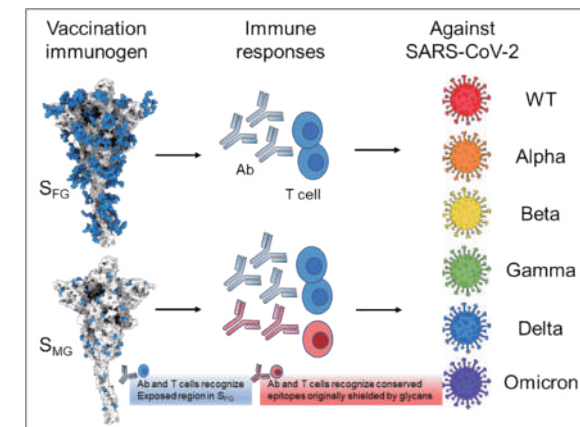


Image provided by Ma Lab

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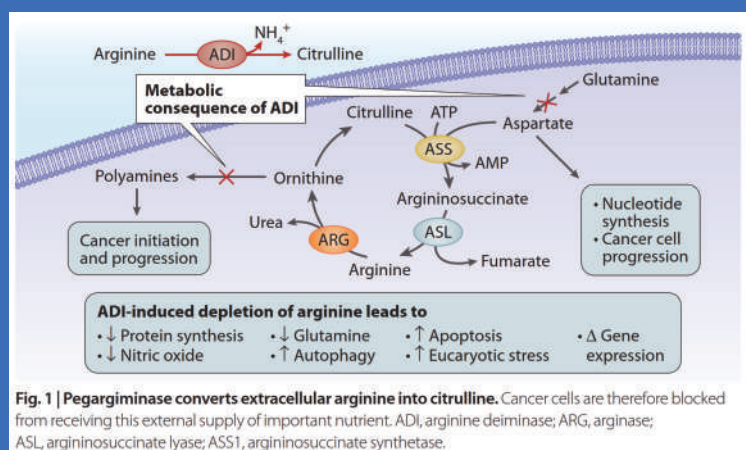


# Small Molecule Drugs & Botanical Drugs

Polaris Group is a multinational biotechnology company focused on developing novel anti-cancer therapies. Leading the way to new and better cancer treatments. Our lead drug candidate, Pegargiminase (ADI-PEG 20), is a biologic in late-stage clinical development for a wide range of cancers, including hepatocellular carcinoma, mesothelioma, pancreatic cancer, non-small cell lung cancer, melanoma, acute myeloid leukemia and others.

Polaris Group is involved in every stage of the drug development process. Our family of companies harnesses structure-based drug design technology to create novel oncology therapies, conducts clinical studies at top-tier cancer centers worldwide and operates cGMP production facilities in Northern California and China. Polaris Group is a Biotechnology CDMO provider for early-stage drug development.

Polaris Group announces positive top-line results from phase 2/3 ATOMIC study in patients with Malignant Pleural Mesothelioma to assess ADI-PEG 20 with Pemetrexed and Cisplatin. Given these results, Polaris Group will proceed with regulatory submissions in the United States in 2023. In 2022, ADI-PEG 20 received fast track designation from the FDA.



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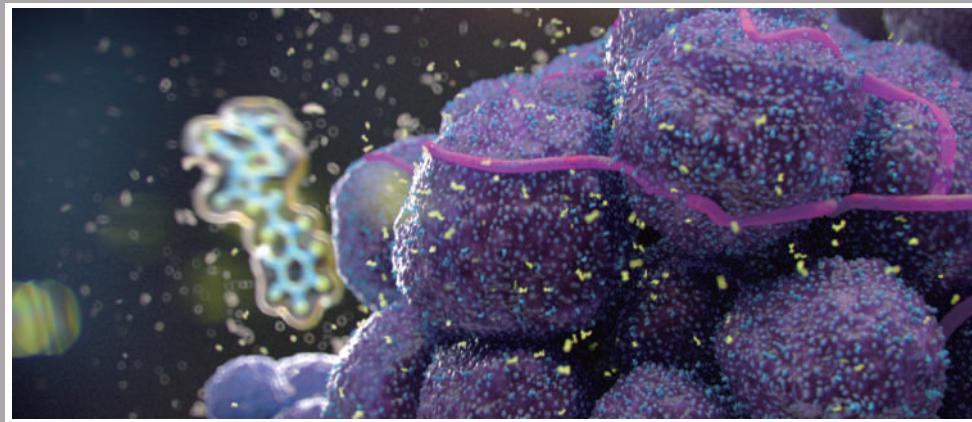




Anbogen Therapeutics (ABT) is a clinical stage biotech founded in 2019, based in Taipei, Taiwan. We focus on developing first/best-in-class precision oncology medicines. We target cancer patients harboring clinically validated driver mutations where we believe the unmet therapeutic needs are. There are two clinical-stage assets, ABT-101 and ABT-301 (HDAC-inhibitor), and several early research programs (e.g. ABT-202, a KRAS inhibitor) under development. The lead compound, ABT-101, is an oral TKI that targets actionable driver mutation, HER2 exon20 insertions, in NSCLC. With strong scientific knowledge and extensive experience in medicinal chemistry, drug discovery and clinical development, we are devoted to make impact to patients with cancer.



Founded in 2017, ALPS Biotech is a new drug research and developing company, dedicated to study biological functions of protein and natural product. In the short term, ALPS focuses on IP's generation including the early stage disease biomarkers discovery for developing in vitro diagnostic device (IVD), and immune therapy on oncology, as well as a variety of professional biotechnology service, such as protein/natural products separation, purification and identification technology. UP to now ALPS has filed many patents covered early diagnosis, better treatment for good quality of patient's life. Currently ALPS Biotech is looking for interested biotech or big pharma to co-development the IP's also.



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PRODUCT	TYPE	TARGET	DISEASE	STAGE	POTENTIAL MARKET (USD)
AL-121	Botanic drug	ACE-2	Anti- COVID-19	Pre-Clinical (In vitro hamster data)	5.6 billion
AL-126	NCE	NLRP3 Inflammasome	Huntington' s disease	Pre-Clinical (In vivo mouse data)	252 million
AL-101	Botanic drug	Novel mechanism	combine Radiation therapy	Pre-Clinical (In vivo mouse data)	2.9 billion
AL-102	Botanic drug	ACE protein	Anti- HBP	Pre-Clinical (In vivo testing)	13.5 billion
AL-801	mAb	GM2AP	Lung Cancer	Pre-Clinical	22.9 billion
AL-801DX	Diagnostic	GM2AP	Lung Cancer DX	Pre-Clinical (2 <sup>nd</sup> stage)	3.64 billion

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The research team of Agari Pharma consists of the experts from I-MEI FOODS CO., LTD and LOHAS BIOTECH DEVELOPMENT CORP. Base on the toxin and metal free agaricus blazei murill planted by unique AI-controlling planting approach, we further concentrated on developing botanical drugs from it.



AnHorn Medicines is a drug discovery platform company with a strong pipeline, offering therapies that target and degrade disease-causing proteins. Our mission is to develop new class of BIGPRO<sup>®</sup> (Bi-functional liGand induced PROteolysis) drugs under the foundation of Targeted Protein Degradation (TPD), to address a broad range of life-threatening and life-impairing diseases.

We offer effective drug candidates and partner with world leading pharmaceutical companies to carry project into real-world studies. We leverage on AnHorn's proprietary AIMCADD<sup>®</sup>, an AI-enabled platform, to turn innovation into new therapeutics - new opportunities to target difficult-to-treat and drug resisting diseases.

AnHorn's Drug Development Strategy:

- Focus on discovering BIGPRO<sup>®</sup> new drugs
- Go beyond next generation of targeted protein degradation by developing novel E3 ligase and its ligand
- Expand unexplored therapeutic opportunity in undruggable targets
- Establish a multilayer patent protection framework for projects

Program	Target	Indication	Discovery	Preclinical	Clinical Trial
AH-001	Androgen Receptor	Androgenetic Alopecia	Open for Partnering		
AH-003	Androgen Receptor	Prostate Cancer	Open for Partnering		
AR-V7 Program		Prostate Cancer	▶		
KRAS G12C/D/V, G13D Programs		KRAS-Mutated Cancers	▶		
Undisclosed Target Programs		Lymphoma & Various Cancers	▶		
Novel E3 Ligase Programs		Various Cancers	▶		

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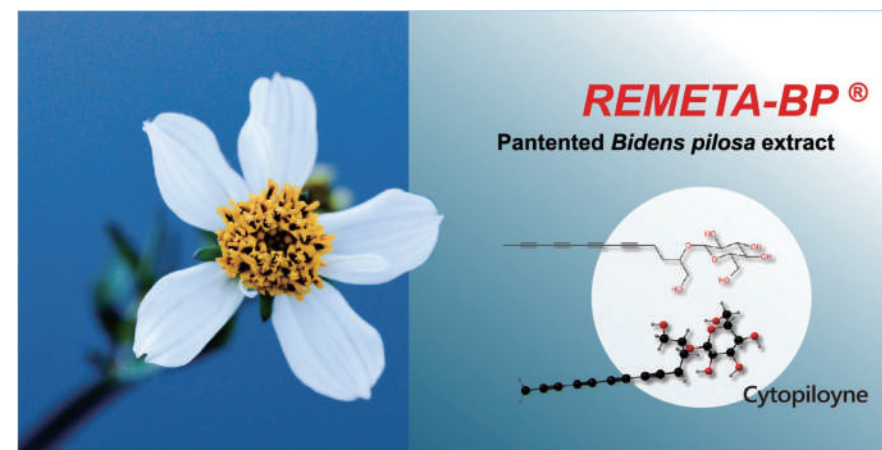


Excelsior group has been dedicated to specialty medicines and medical examinations in rare diseases for more than 2 decades. In 2016, Excelsior group established a TGF-beta research center and start up new drug development projects to screen new drug candidates of enhancers and inhibitors targeting at TGF-beta activities in various diseases, such as wound healing, cardiovascular diseases, autoimmune diseases, liver cirrhosis, pulmonary fibrosis, neurodegenerative diseases, and cancer treatments. In 2020, Excelsior group is expanding our team in the field of rare diseases and establish a new research center of specific diseases aim at genetic and metabolic disorders in National Biotechnology Research Park, NBRP. In NBRP, we are investing in precision medical testing, epidemiology study, new drug development, and novel therapeutic technologies in various rare diseases. Besides, we are also looking for more strategic alliances with Academia Sinica, National Health Research Institute and leading medical centers in Taiwan. Our aim is to establish a world-class laboratories in the research of TGF-beta and specific diseases and extend our services to Asian populations.



Exland biotechnology inc. manufactures, supplied, and exports functional plant extracts in Taiwan for the past 12 years. Our R&D team is staffed with the most experienced experts in traditional herbs and plants. We are proud to hold a multinational patent of our own extraction technology, Superdash<sup>®</sup> extraction, certified in Taiwan, USA, China, and Germany. Exland's Superdash<sup>®</sup> extraction technique yields the highest extraction efficiency resulting in higher concentration of active ingredients than our competitors. By utilizing water extraction, we are able to achieve zero organic solvent residues in our products, offering a healthier and higher quality choice to the market. Our mission strives towards the highest extraction excellence in functionality, taste, color and aroma. Given our background in medicine and nutrition, Exland has been ever invested in our R&D ability and close collaboration with our partners and clients to provide the best product for our consumers.

Selected Products	Indications	Status
EX042-002-05	Rare disease	Lead optimization
EX042-002-11	Rare cancer	Lead optimization
EX042-002-06	Tissue fibrosis	Prove of concept
EX042-004-09	Alzheimer's disease	Research



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Foresee is a Taiwan and US-based biopharmaceutical company listed on Taipei Exchange (TPEX : 6576) . Foresee's R&D activities consist of two areas, the proprietary stabilized injectable formulation (SIF) targeting specialty markets, and the first-in-class NCE programs targeting rare and severe diseases.

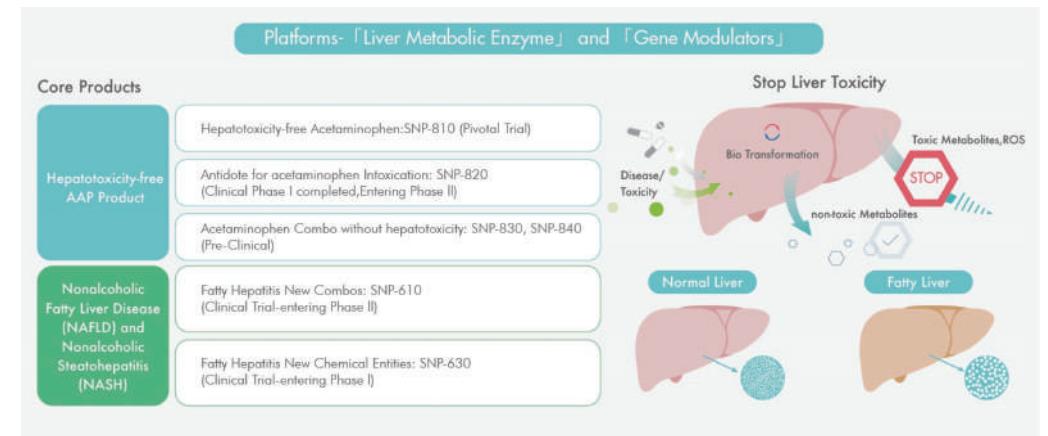
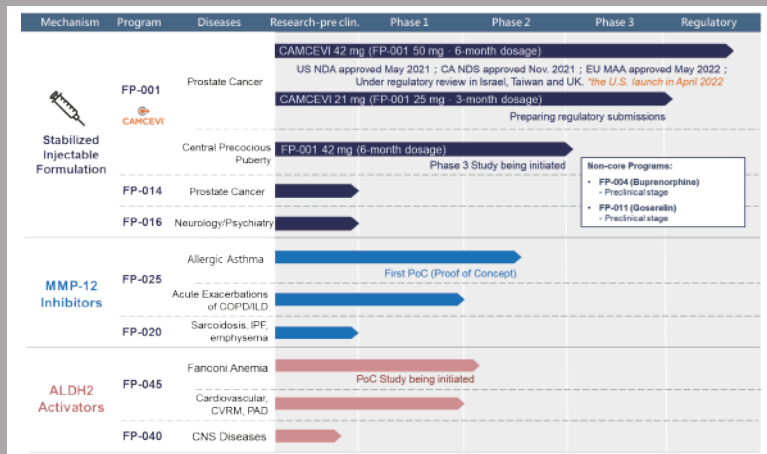
SIF technology : CAMCEVI<sup>®</sup> 42 mg (6-month depot injection formulation), indicated for the treatment of advanced prostate cancer, was the first commercial product developed from Foresee's SIF platform. CAMCEVI<sup>®</sup> 42 mg has been approved in the US, EU, Canada, and was launched in the US in April 2022.

NCE programs : FP-025, a first-in-class oral MMP-12 inhibitor targeting inflammatory and fibrotic diseases currently in a Phase 2 proof-of-concept study in allergic asthma. FP-045, a first-in-class oral ALDH2 activator, is in a Phase 1b/2 for Fanconi Anemia.

(The company currently develops six core products based on the two technology platforms of "Liver Metabolic Enzyme Activity" and "Gene Regulation". Two are for the treatment of non-alcoholic steatohepatitis (NASH), and three are new formulations and combo of hepatotoxicity-free painkiller Acetaminophen and one is an antidote for Acetaminophen poisoning.

New drugs for the treatment of NASH, of which SNP-610 has been approved by the US FDA and Taiwan's TFDA for Phase II clinical trials in May 2017, and is the first-in-class new drug in the market; SNP-630 is an optimized drug with a new ingredient NCE), a Phase I clinical trial is now underway.

Among the other three SNP-8 products, SNP-810 is a new non-hepatotoxic Acetaminophen analgesic drug, which has been approved for the US OTC monograph sales license and Taiwan's drug license. At the same time, it has applied for a US FDA pivotal trial. SNP-820 is an antidote for Acetaminophen poisoning and is applying for entry into clinical trials in Taiwan and the United States by way of application for orphan drugs. Compound SNP-830 and SNP-840 are patented new compound non-hepatotoxic analgesic drugs containing SNP-810 and analgesic and anesthetic ingredients. In the future, SNP-810 and non-addictive analgesics such as Sebacyol Dinalbuphine Ester will be developed into new therapeutic effects. The new compound drug is currently undergoing preparations for clinical trials of the new compound new drug.



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Saint Therapeutics (ST) is a clinical-stage biotechnology company focused on revolutionizing tissue repair and disrupting the billion-dollar scar reduction and chronic wound healing industries. Building upon over 20 years of research into how fetal skin heals rapidly and without scarring, ST is developing SLI-F06, a unique, patent-protected, first-in-class peptide drug to significantly reduce scar formation after surgery and to promote healing.

SLI-F06 has completed nonclinical efficacy and safety, human Phase 1, and preliminary Phase 2a studies of injectable liquid SLI-F06 for scar reduction during surgery. Both Phase 1 and 2a results show preliminary evidence of improvement in several individual POSAS parameters related to scarring on the SLI-F06 treated side in high tension.

The FMOD/SLI-F06 technology is also supported by 20 academic papers; \$14M in NIH/SBIR grant awards; 16 patents granted in USA, Canada, UK, France, Germany, China, Korea, and Taiwan; and 10 pending patents in various countries.

Formulation	Indication	Discovery	Preclinical	Phase 1	Phase 2	Phase 3	NDA
Liquid SLI-F06 - Injectable	Acute (1 <sup>st</sup> time) scar reduction	Lead					
	Preexisting scar revision surgery	Lead					
	Diabetic foot ulcer healing	Secondary					
Hydrogel SLI-F06 - Injectable - Implantable	Bioactive adjuvant to increase device (sutures, mesh) healing						
	Tissue healing (tendon)						
	Tissue healing (fascia, muscle)						
Topical SLI-F06	Diabetic foot ulcer healing						
	Burn injury						

## Contact us

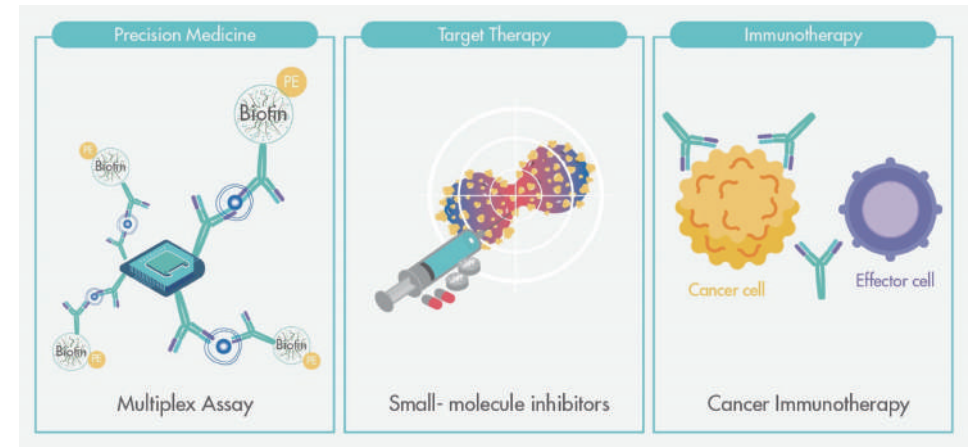
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At WinLux Biomedicine Technology Co., Ltd., a subsidiary of Chainwin Agriculture and Animal Technology Global Business Group, we strive to set the standard for quality, safety, and value in the discovery, research, and development in the healthcare industry. Our aim includes establishing diseases database, multiplex diagnostic platform, and devoting in new drug development. Throughout our work, WinLux is committed to being transparent, acting with integrity, protecting the environment, and respecting human rights.

Led by the Chairman Chen, Jin-Cai, WinLux reveals its core values from the basis of everything we do: Accountability, Innovation, Quality Foremost, and Life Respect. Consistent with our responsibility as one of the world's premier innovative biopharmaceutical companies, we shall keep accomplishing ultimate goals, "to make a difference for all who rely on us."



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# Smart Healthcare & Precision Medicine →→

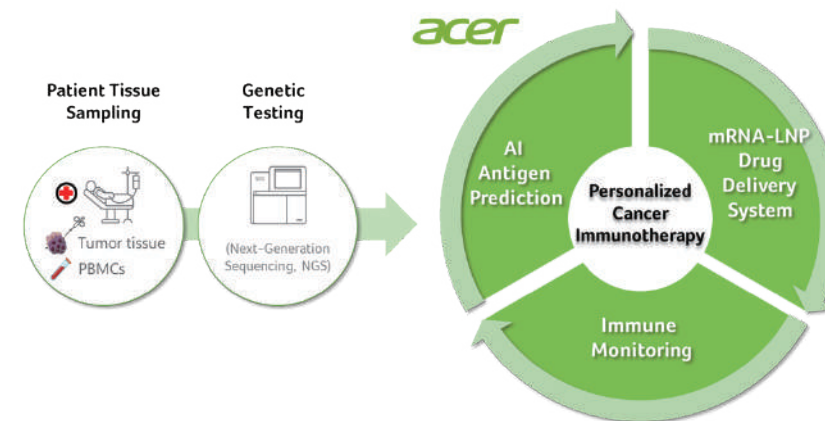


Acer Incorporated  
宏碁股份有限公司



The Bio-Medical team of Acer Value Lab focuses on the research and development of personalized cancer therapeutic vaccine that will induce immune responses to specifically recognize and destroy tumors in patients. We are currently developing three platforms for personalized neoantigen-based immunotherapy approaches: 1) Artificial Intelligence (AI) model for neoantigen prediction; 2) mRNA-lipid nanoparticle (LNP) system for payload delivery; 3) Immune monitoring platform for analyzing specific T cell subsets and effector functions to identify novel biomarkers of immunotherapy.

Our team is in collaboration with clinical and research institutions as well as technology platforms and we welcome partnerships to engage personalized treatment strategies.



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Bio Preventive Medicine Corp. (BPM), a leading company in renal biomarker area, focuses on translating clinically validated and IP-protected (>79 patents) biomarkers into precision diagnostic solutions for unmet clinical needs. The disease area we focus on includes diabetic kidney disease (DKD), renal injury, and oncology.

Human uPTM3-DKD (DNlite-IVD013) ELISA test is a urinary prognosis test for predicting/monitoring risk of kidney disease progression, and is perfect for routine clinical practice in any healthcare institutes and clinical labs. Besides precision management of DKD, it can also be applied for risk assessment of Acute Kidney Disease, and Graft Failure in kidney transplant patients.


Commercially, DNlite-IVD103 is CE-IVD marked, approved by Malaysia MDA, and also launched in Japan and US as RUO kit. Furthermore, several other worldwide regulatory approvals are in preparation. BPM is ISO17025 and ISO13485 accredited, and listed in Taiwan emerging stock market (code: 6810).



The company was founded in April 2021, with the goal of developing portable analytical instruments to meet the needs of real-time and on-the-spot measurement. In the initial stage, it is expected to be applied to biomedical, food safety, environmental safety, semiconductor, chemical and environmental protection industries. The medium-term development can be applied to customs, postal systems, airports for illegal drugs as well as explosives detection. In the long run, the second generation of products will be mini size similar to a smart phone.

The company's initial main products include handheld ion mobility spectrometer, portable liquid chromatographs, portable mass spectrometers, and portable liquid chromatography mass spectrometers.

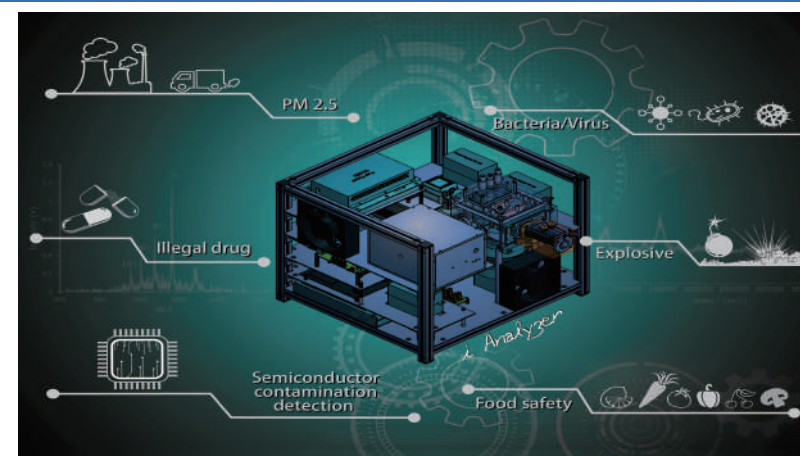
The key technologies provide portable and low-cost for the above instruments. The company will keep Taiwan as the R&D center, and establish overseas sales departments to enter the global market.



**DNlite IVD103** predicts renal function loss years in advance through non-invasive ELISA-based testing. A true diagnostic gold standard for DKD in clinics and labs.

**DNlite IVD103 Human uPTM3-DKD ELISA Test**

- ▶ Single and Unique Biomarker
- ▶ Predict Progressive Decliner
- ▶ Predicting & Monitoring Renal Function Loss
- ▶ Precision DKD Management
- ▶ Non-invasive Urinary ELISA Test



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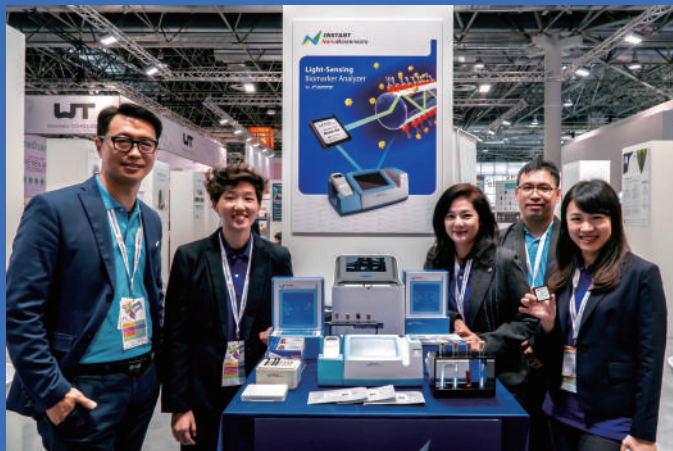




Instant NanoBiosensors is an innovative life science company which has developed the ultra-sensitive digital immunoassay platform based on patented FOPPR™ (fiber optical particle plasmon resonance) sensing technology and patented IN-Chip (auto-flowing microfluidic chip) technology for the life science research and in vitro diagnostics markets. The simple and ultra-sensitive blood-based biomarker platform that could pinpoint the Alzheimer's disease process well before any cognitive symptoms would allow physicians to give any potential disease-slowing therapy at a very early point in the disease process. Our goal is to offer clinical researchers and physicians testing solutions with higher precision, real-time and convenience to serve as the basis for early detection and disease monitoring. The additional beneficial features provided are portable ease of use and inexpensive sensor chips in a cost-effective manner to light up the hope for early detection of Alzheimer's disease in the future.

JUNZHI is a biotechnology team from Taiwan, and we have established research and development centers and Operations Center in National Biotechnology Research Park and Tainan Science Park respectively. The technologies authorized by many top research and development units, including Academia Sinica, National Tsing Hua University, National Yang-Ming University and Kaohsiung Medical University, etc. After many years of research and development, we developed the cancer screening service targeted on cancer of digestive system.

The technology of SAA MS screening and interpretation of MS spectra come from Academia Sinica. NTHU is responsible for the development of Magnetic Nanoparticles and antibody for extraction and purification of SAA. NYMU and Metal Industries Research & Development Centre help develop the bio-algorithms, making it more accurate. Last but not least, KMU work on clinical trials and the integration of clinical information to ensure that SAA TEST has high accuracy and also help practically in the medical field.



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LumiSTAR develops various platforms by packaging bioimaging and iPSC technologies together to make cellular response visualizable in real time. Our designed protein-based indicators with different dynamic ranges allow long-term observation and measurements, as they are non-toxic to the cells. Thus, drug efficacy and toxicity can be studied at early stages to shorten the process of product development.

**Applications:**

**1. iPSC High Content/Throughput Screening Platform:**

For Cell Therapy:

- Development of iPSC cell reprogramming and differentiation
- iPSC differentiated cells functional testing platform

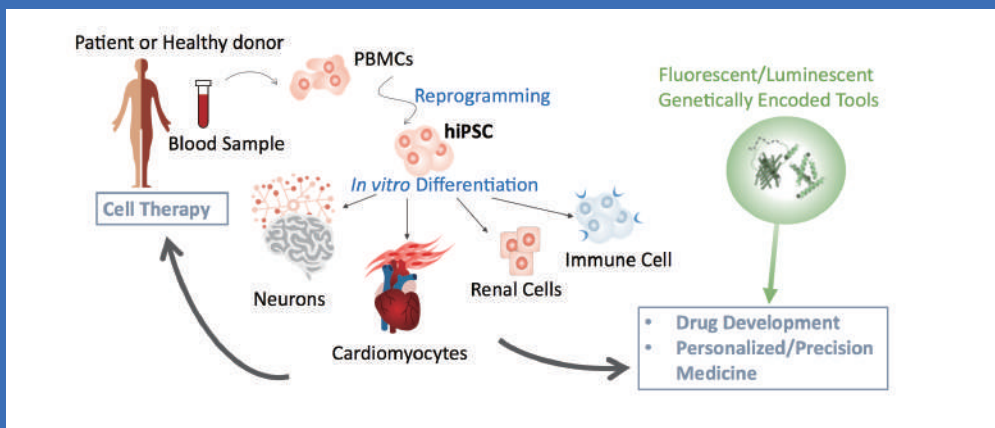
For Drug Development:

- Cardiac and neurodegenerative diseases models
- Drug efficacy study
- Toxicity evaluation at early stages
- Support phenotypic screening

**2. Pseudovirus High Throughput Imaging Screening Platform**

- Pseudovirus construction for various mutant strains
- Neutralizing antibody screening, vaccine validation and drug screening
- Customized service

**3. Personalized/Precision Medicine**



Contact us



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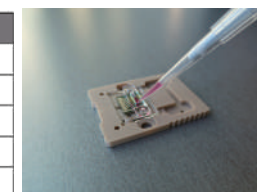
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Molsentech's Semiconductor-based Biosensor leverages ultra-high sensitivity to integrate the electrical charge of biomolecules with the technologies of surface modification and biomolecular detection to develop an assay that can be applied to detect not only RNA but also proteins for a variety of diseases, including COVID-19. Our biosensor assay combines the advantages the velocity of rapid screening and the high accuracy of traditional PCR assays, providing 5-minute testing time with 95% accuracy that makes it the best solution for early detection.

Molecular biosensing real-time analyzer, SENEDIA, is an electrical-based ultra-high sensitivity device capable of catching minimal charge change of electrical current. Compared with optical-based readers, the Semiconductor-based Biosensor powers SENEDIA to be able to detect ultra-low concentration of pathogens from minimal size of a wide range of samples (e.g. blood, saliva, urine, and etc.) to get detection results before symptoms appear, benefiting people's health on the front of disease prevention.

Products	Applications	Phases
Bio-FET Biosensor & Biomolecular Analyzer & Reagent	Covid19 RNA Rapid Test	Received EUA
	Respiratory Infectious Disease Detection	Pre-clinical
	Cancer Detection	Pre-clinical
	Pet Cancer Detection	Coming soon in 2023
	Microorganism Detection	Pre-clinical



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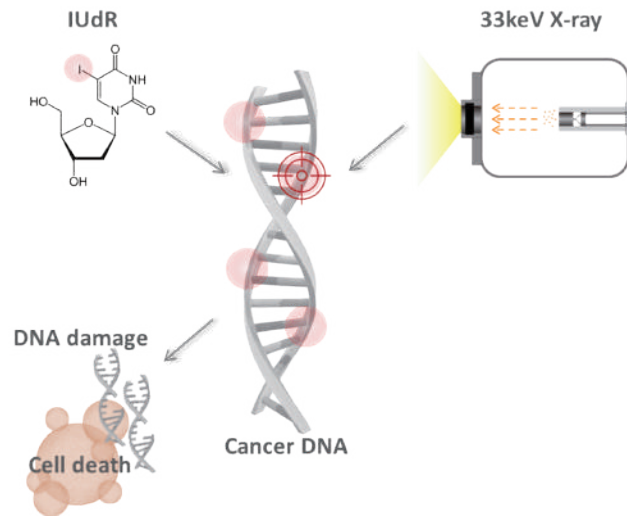
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NanoRay Biotech has stationed in National Biotechnology Research Park since 2020, devoting to the development of characteristic X-ray technology, applying to medical imaging and cancer therapy.

"Auger Molecular Therapy" is a novel cancer therapy technology. Based on the characteristic X-ray generated by the patented transmission type X-ray tube developed by NanoRay Biotech, Auger effect is efficiently induced in specific heavy atom that release enormous energy in nano-scale space of cancer DNA, inducing aggressive DNA damage and cancer cell death, while lowering the radiation dose received by normal tissues.

Auger Molecular Therapy has been approved for IND by US FDA. A phase 1 clinical trial will be administrated in 2023.



# Contact us



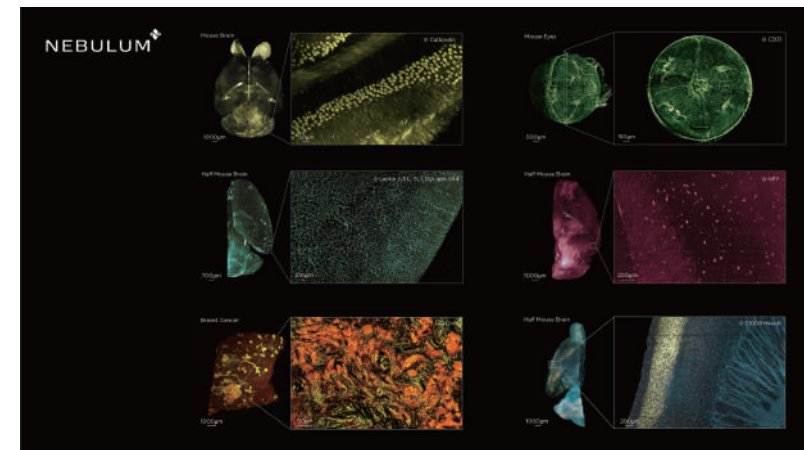
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Nebulum 3D Pathology Platform offers a one-stop solution for therapeutic approach optimization. We provide a universal, multi-dimensional visual analytic solution for scientists, clinical researchers and diagnosis professionals. This offers access and insights on biology studies, neuroscience research and pharmaceutical research for clinical diagnosis. Our 3D Whole Tissue Imagine Service will accelerate our client's research.

We offer a one-stop sample-to-data solution platform. With our proprietary tissue clearing, immunolabeling, lightsheet imaging and data processing technologies, our clients can gain access to the most up-to-date solutions and integrate the best available 3D tissue data into their research.



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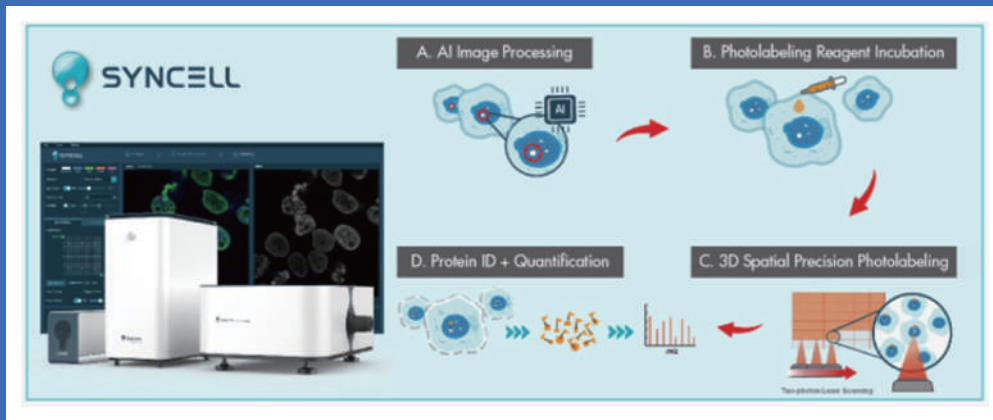
SYNCELL is building a novel spatial biology system, Microscoop™, the world's first in-cell "pickable" high-content microscope-based platform. The system leverages AI-based pattern-recognition algorithm to selectively harvest proteins and nucleic acids in situ from user-defined areas of interest. Using our proprietary subcellular photosensitive labeling reagents, the instrument can effectively "tag" and isolate both known and novel targets for a wide range of cell and tissue biology applications. It is uniquely suited for spatial proteomics and offers distinct capability for discovering novel protein expression to overcome the limitation of all current spatial proteomics systems that can only study known proteins. The system also has the potential to generate multi-omics data on the same cell or tissue sample to delineate new insights on novel biomarker functions that cannot be explored by current technologies.



VITAE Biomedical Co. Ltd. is an innovative screening technology company, providing valuable screening technologies for cancers and other diseases. We hold patents in multiple countries and seek to apply the results of our research to medical communities worldwide.

In response to the highly unmet needs for lung cancer screening, our featured reagent kits target a biomarker specific for lung cancer screening in patients' urine. Our breakthrough method is non-invasive, non-radiative, and made possible for early diagnosis. This technology provides excellent benefit-to-risk ratio to the public, serving the needs for frequent monitoring of high-risk individuals and lung cancer patients, and fulfilling a key screening element in regular health checks.

We are committed to continuously deliver our capabilities and services to the people, developing the most needed screening tools that enable accurate and speedy diagnosis with the hope of improving survival rates and life quality of patients.



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**Investment  
Opportunities**

NBRP BioHub Taiwan

**Resident Companies in  
Shared Space**







AIBIOS K.K.  
日商艾栢生技股份有限公司



We believe by using our unique knowledge can improve people's lives, and for the benefit of society, the economy, and the environment. With a strong passion, our founder and CEO Akio Sato found the company in 2015 Tokyo, Japan.

We're a group of people with passion, focus on create solutions that help people live healthier lives.

Today, our team in Japan, Taiwan, and United States, focuses on discovering and delivering transformational medicines and products in several key therapeutic areas including immunology, oncology, and neuroscience through our growing pipeline. Each day, we work to deliver sustainable solutions that improve the health of our business and the health of humankind.

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Apexcella Biomedical Inc.  
艾沛生醫股份有限公司



Apexcella Biomedical Inc. has developed unique three-stage apexNK cell expansion technology, which can produce mass apexNK cells with high purity and high cytotoxicity. Based on the technology, we are committed to develop the autologous or allogeneic apexNK cell therapies, which can provide cancer patients the next-generation, novel and effective or curative therapies.

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LuminX Biotech Co., Ltd.  
路明思生技股份有限公司



LuminX Biotech dedicates to accelerate the optimization of cell-therapy-products through specific understanding of therapeutic cells distribution in-vivo, long-term viability, as well as their biological fate either in preclinical or clinical setting and our technologies can transition a potential therapy from the research stage to advanced clinical and commercial applications to save lives.

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Ochre Bio Taiwan Ltd.  
赭石生醫有限公司



Ochre Bio, HQ'd in Oxford UK, develops genomic medicines that rejuvenate transplanted livers, ultimately with the goal of applying these therapies to treat fatty liver disease (NASH/NAFL) and other metabolic diseases. Ochre Bio Taiwan is Ochre's first overseas research site, focusing on Target Discovery and Target Validation. The site will employ single-cell and spatial sequencing, advanced genomics, and high-throughput screening to identify disease-causing genes, and develop combination therapies together with Oxford HQ. In Jan 2021, Ochre Bio completed the largest genomic atlas of the liver, and is currently progressing multiple GalNAc-siRNA candidates toward the clinic.

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Pharmasaga Co. Ltd.  
藥祇生醫股份有限公司



Pharmasaga is an Academia Sinica-derived biotech company, and focus on the development of a first-in-class diabetes drug, PS-001. PS-001 is an inhibitor of Pdia4 protein, which plays a critical role in diabetes development. Oral administration of PS-001 in diabetic mice can prevent the beta cell death through ROS decreasing, stabilize the blood sugar and reverse diabetes. The preclinical studies and CMC had been completed. The US IND of PS-001 had been approved, and the Phase I clinical trial will be launched.

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Topmunnity Therapeutics Taiwan Limited  
得勝醫學科技股份有限公司



Topmunnity Therapeutics Taiwan Ltd., established in 2019, hope to improve the efficiency and precision of drug treatment through innovative cancer therapy, and meet the needs of cancer patients who still lack effective treatment. Topmunnity focuses on the research and application of immunotherapy and hope to help prolong and improve the quality of patient's life.

The CAR-108 developed by Topmunnity is the CAR-T cell therapy (Chimeric antigen receptor T cells), the scFv can accurately identify glycosylated antigen on the tumors, the indication will be treating the solid tumors. In the cell-based experiments, it has been proven that CAR-108 can kill 80% of tumor cells effectively. In the in vivo animal study, the tumor was totally remission after the CAR-T treatment.

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Zenith BioDesign Lab Co., Ltd.  
禹智生物科技股份有限公司



Zenith BioDesign Lab(ZBL) has positioned itself upstream of the drug development industry chain through our antibody drug design platform technology and full-range capabilities for R&D. ZBL expects to facilitate drug development by combining the process and production strengths of CMOs or CDMOs in the biopharmaceutical industry. Synergistic collaborations built on our own patented ZBL-UmaskAb technology is one of our win-win business strategies.

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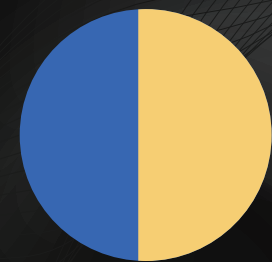
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**Biomedical  
Ecosystem  
Resources**

NBRP BioHub Taiwan

**BioHub Taiwan Accelerator**







We provide full range of legal services to local and foreign clients. As part of Baker McKenzie's global network of firms, we also deliver seamless legal service that is both global and local. With nearly 100 legal services professionals, including lawyers and patent engineers, the credentials of Baker McKenzie's lawyers in Taiwan speak for themselves.

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Famar International Patent and Trademark Offices integrates Famar Licensing Co., Ltd. to provide one-stop service. Except patent prosecution and trademark registration, Farmer provides professional services such as intellectual property management, technology authorization, technology evaluation and business model advice; assisting startup companies to establish their business mechanism. The clients we served include California Institute of Technology, Illumina, National Health Research Institutes, National Taiwan University, National Yang Ming Chiao Tung University, National Chung Hsing University, National Cheng Kung University, China Medical University, Council of Agriculture, Taipei and Taichung Veterans General Hospital, Yungshin Pharm Ind. Co. Ltd, Taiwan Advance Bio-Pharmaceutical Incorporation...etc. Our professional team is divided into two groups: the biochemical group (biotechnology, medicine, chemicals, materials, medical engineering) and the electromechanical group (electrical, electronics, semiconductor, optoelectronics, communications, machinery, general application items). The team is composed of experts with the experience of patent examination in Taiwan Intellectual Property Office and qualification as US or Taiwan patent attorney.

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GeneOnline is a leading biotech media and market intelligence platform offering in-depth healthcare and biopharma insights. Our Mission is to provide the latest global biotech/pharma trends covering exclusive KOL interviews, market updates, emerging technologies and more. We are the go-to option for any company seeking global exposure and partners in the biopharma sector.

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GeneOnline is a leading biotech media and market intelligence platform offering in-depth healthcare and biopharma insights. Our Mission is to provide the latest global biotech/pharma trends covering exclusive KOL interviews, market updates, emerging technologies and more. We are the go-to option for any company seeking global exposure and partners in the biopharma sector.

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Grant Thornton Taiwan  
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Grant Thornton Taiwan was founded in 1971. We are an ISO certified firm and have approximately 250 employees in Taiwan. We serve dynamic, growing companies that are on the move because we too are a dynamic, growing firm on the move. Having started out as a small practice in Taipei in 1971, we now have offices in Hsinchu, Taichung, and Kaohsiung, which provide geographic coverage that can meet the needs of most companies doing business in Taiwan. We understand the needs and aspirations of growing companies on a personal level, and we have over 50 years' experience helping companies achieve these aspirations. We measure client satisfaction on an annual basis as part of our ISO certification program. Grant Thornton Taiwan clients continuously rank as highly satisfied. This is partly thanks to our business structure, which is designed to react quickly to our clients' needs. At Grant Thornton Taiwan, we are empowered to make decisions locally, rather than having to move issues up through a complex chain of command. This allows us to consistently provide clients with solutions tailor-made to their circumstances and time restrictions. Grant Thornton Taiwan is structured to provide services to clients via different service divisions. Our service divisions include: Assurance, Bookkeeping, Corporate Finance, Company Registration/Liquidation, Legal Support Service, Payroll, Taxation, Transfer Pricing. Working with Grant Thornton Taiwan, you will feel safe in the knowledge that help and advice on the issues that inevitably arise in a complex regulatory environment are only a phone call away.

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IBMI / RBMP  
生策會 / 生策中心



Founded by the former president of the Legislative Yuan of Taiwan, Mr Jin-Pyng Wang, IBMI is an independent, not-for-profit organization voicing Taiwan-based health care industry, promoting interdisciplinary collaborations through its global platform, and creating policy dialogues between public and private sectors. On top of that, IBMI is also a trusted awarding and certification body to health care providers and an incubator to health care startups in areas of novel technologies, services and innovations.

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Pacific 8 Ventures



At Pacific 8 Ventures, we invest at the intersection of breakthrough technology platforms and Biology & Medicine. The team has a unique combination of venture capital, tech platforms, clinical medicine, biomedical engineering, and operational expertise. Through our support, we look to help build great companies that advance human health and well-being.

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Roche Products Ltd.  
羅氏大藥廠



Roche is a global pioneer in pharmaceuticals and diagnostics focused on advancing science to improve people's lives. The combined strengths of pharmaceuticals and diagnostics, as well as growing capabilities in the area of data-driven medical insights help Roche deliver truly personalized healthcare. Roche is working with partners across the healthcare sector to provide the best care for each person.

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Taiwania Capital  
台杉投資



Taiwania Capital is a venture capital firm that was founded in August 2017 by the National Development Fund of the Executive Yuan of Taiwan and private enterprises. Our mission is to establish a partnership with companies worldwide and boost Taiwan's economic growth. We invest in different stages of companies mainly in the tech and biotech fields. By linking technology and capital, Taiwania Capital aspires to also create a robust ecosystem in Taiwan that is rooted in entrepreneurial innovation.

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## Tiger Accelerator 台荷加速器



Based in Taiwan & the Netherlands, Tiger Accelerator serves as a bridge between entrepreneurship ecosystems in Asia and Europe. We've been facilitating valuable connections between startups, SMEs, corporates, investors, accelerators, incubators and public stakeholders in the two continents.

With our expertise and network in the two geographies, we link Asian and European innovators with the business resources they need for international expansion, support their international growth ambitions, as well as minimize the uncertainties and risks involved for entrepreneurs to enter unfamiliar markets.

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## Taiwan Research-based Biopharmaceutical Manufacturers Association (TRPMA) 台灣研發型生技新藥發展協會



TRPMA, established in 2012, is an industry association representing 56 Taiwanese R&D-based biopharmaceutical companies. Our mission is to build up a competitive bio pharma industry in Taiwan by consolidating public and private resources and facilitating product and business development. On behalf of Taiwan bioindustry, TRPMA serves as a window for promoting international partnership in the area of new drugs, vaccines, regenerative medicines and digital medical techs. We welcome your visit!

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## Taiwan Bio Industry Organization 台灣生物產業發展協會



Since its founding in 1989, the Taiwan Bio Industry Organization (Taiwan BIO) has been promoting the interests of its members from industry, the government and academia as well as promoting the development of the biomedical industries and their contribution to Taiwan's economy and people. The more than 100 organizations and 3,000 individuals that make up Taiwan BIO membership represent the entire life sciences industry spectrum, including from the biotechnology, pharmaceutical, diagnostics, medical devices and materials, agricultural biotechnology, and nutraceuticals sectors.

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