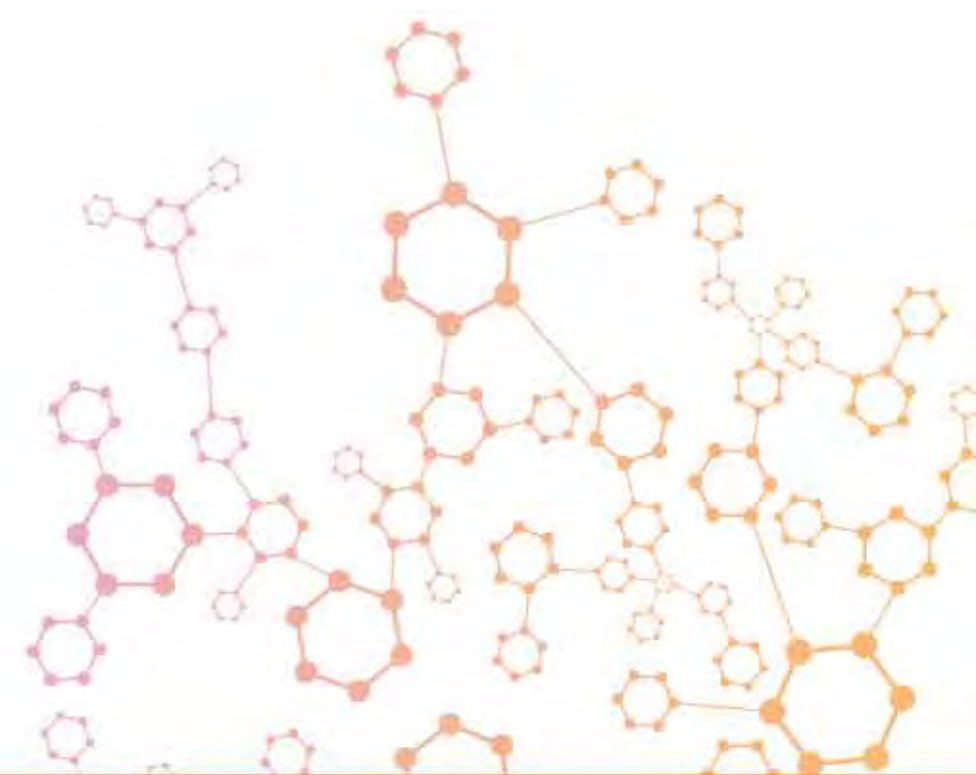




Taiwan

Pharmaceutical Alliance

State-of-the-art
Cost-effective Service
CMO/CDMO
Licensing In / Out
API / FDF / Biosimilar



About TPA

Taiwan Pharmaceutical Alliance (TPA) is comprised of the top Taiwanese pharmaceutical companies and was established by the Medical and Pharmaceutical Industry Technology and Development Center (PITDC) that is under the Industrial Development Bureau (IDB) as a part of the Ministry of Economic Affairs.

There are 15 members in the TPA currently. We have a collection of APIs, niche generics, 505(b)(2)s, biosimilars, and medical supplies. Almost all the therapeutic areas are covered by TPA while each member has their own expertise. The finished formulations contain various types of oral forms, semi-solids, injectables, eye drops, MDIs/DPIs/nasal sprays, patches, topicals, etc.

TPA also provides CMO/CDMO services. Just like any Taiwanese manufacturing facility, TPA members are PIC/S GMP accredited, most are certified by US FDA, EU EMA, and Japan PMDA. TPA's presence is throughout the world in many regions, such as the US, Europe, MENA, Japan, and ASEAN. Our high-quality products have been recognized by many international partners.

The vision of TPA is to collaborate with the international pharmaceutical community and to maintain long-term relationships with our partners. We are committed to bringing the best medicine to patients and to fulfilling the global unmet medical needs.

Therapy Area

- Cardiovascular
- Dermatology
- Oncology
- Respiratory
- Gastrointestinal
- Others
- Central Nervous System
- Ophthalmology
- Infectious Disease
- Women's Health
- Ear Nose Throat Disorders

Product

- Active Pharmaceutical Ingredients
- Various Finished Dosage Forms
- Traditional Chinese Medicine
- Medical Supplies (Rubber Stoppers)

Service

- Contract Manufacturing
- Contract Development and Manufacturing
- Contract Research





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In English Alphabetical Orders

















GLOBAL FOOTPRINT

| United States | Canada | Mexico | Japan | Korea | South East Asia | China | Middle East | Turkey |
| European Union | United Kingdom | Egypt | India | Australia | Taiwan |



Company name	Biologic/ Biosimilar	Oncology	Hormone	Ophthalmic	API	Packaging Material	Dietary Supplement	OTC
 Bora Pharmaceuticals	•	•		•		•		•
 Chung Mei Biopharma Co. LTD.							•	•
 Johnson Chemical Pharmaceutical Works Co., Ltd.							•	•
 Mithra Biotechnology Inc.	CRO Service							
 Pei Li Pharmaceutical Industrial Co., Ltd.		•	•					
 Prince Pharmaceutical Co., Ltd.					•		•	•
 Savior Lifetec Corporation					•			
 Sinphar Pharmaceutical Co., Ltd.		•	•	•		•	•	•
 Standard Chem. & Pharm. Co., Ltd.							•	
 Synmosa Biopharma Corporation			•		•		•	•
 Syn-Tech Chem. & Pharm. Co., Ltd.					•			
 Swiss Pharmaceutical Company Limited							•	•
 Taiwan Biotech Co., Ltd.				•	•		•	•
 Taiwan Tashen Biotech Co., Ltd.						•		
 UBI Pharma Inc.	•							•

Company name	Lyophilization	Vial / Cartridge	Liposomes / Microspheres	Tab.	OD Tab.	Soft Cap.	Cap.	Eye Drop	MDI / DPI / Spray	Suppository	Powder / Granule	Patch	Other product
 Bora Pharmaceuticals				•	•		•	•	•		•		
 Chung Mei Biopharma Co. LTD.				•	•		•		•		•		Ointment, emulsifiable paste, gel, and suspension.
 Johnson Chemical Pharmaceutical Works Co., Ltd.				•			•				•		
 Mithra Biotechnology Inc.													Biopharmaceutical Analysis, Pre-clinical/clinical Bioanalysis, Pharmaceutical Analysis, Clinical Research, Pharmacokinetics Study
 Pei Li Pharmaceutical Industrial Co., Ltd.				•	•	•	•			•			
 Prince Pharmaceutical Co., Ltd.													
 Savior Lifetec Corporation	•	•											
 Sinphar Pharmaceutical Co., Ltd.	•	•		•		•	•	•		•	•		
 Standard Chem. & Pharm. Co., Ltd.	•	•		•	•		•				•		
 Synmosa Biopharma Corporation				•	•		•		•		•		
 Syn-Tech Chem. & Pharm. Co., Ltd.											•		Solution , Crystal
 Swiss Pharmaceutical Company Limited		•		•	•		•				•		Liquid Semi-Solid
 Taiwan Biotech Co., Ltd.		•		•	•		•	•			•	•	Injection, Oral Solution, Blow-Fill-Seal, Cream, Ointment
 Taiwan Tashen Biotech Co., Ltd.	•	•					•				•		
 UBI Pharma Inc.	•	•	•	•			•				•		Ointment (CMO/US)

ALLIANCE MEMBERS



**Bora
Pharmaceuticals**
保瑞藥業股份有限公司

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GMP CERTIFIED

- Oral Solids: Taiwan FDA, USFDA, MHRA, GCC FDA, Jordan FDA, Health Canada, PMDA
- Semi-solids: US FDA, MHRA, Health Canada, PMDA
- Ophthalmic: US FDA, Taiwan FDA

**FOCUSED
THERAPEUTIC
AREA**

- ALL

INTRODUCTION Bora Pharmaceuticals is a leading international Contract Development and Manufacturing Organization (CDMO) with expertise in the production of complex oral solid dosage forms (tablets and capsules), liquids (solutions, suspensions, and nasal sprays), and semi-solids (creams and gels), Ophthalmic products (eye drops and ointment), and large molecule biologics(biosimilar) for both prescription (Rx) and over-the-counter (OTC) pharmaceutical products. We specialize in late-phase Clinical through Commercial manufacturing and packaging.

With three state-of-the-art cGMP manufacturing facilities located in Taiwan and Canada, we adhere to the highest international standards for manufacturing, packaging, research and development (R&D), and analytical testing. Our facilities are equipped to handle high potency compounds, solvents, flammables, as well as immediate-release (IR), sustained-release (SR), and extended-release (ER) products.

We have a global reach and serve over 100 markets worldwide, including the United States, Canada, European Union, Southeast Asia, Middle East, and South and Central Americas. All our sites comply with the Trade Agreements Act (TAA), and our packaging lines are fully serialized to ensure product traceability and security.

At Bora, we prioritize reliability, integrity, and innovation in every aspect of our operations. We are committed to delivering dependable, personalized service starting from proactive project management and technical support. Our goal is to consistently provide high-quality products and help you achieve your global outsourcing objectives.



**Chung Mei
Biopharma Co. LTD.**
中美生技醫藥股份有限公司

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GMP CERTIFIED • PIC/S GMP (oral liquid preparation: syrup, solutions, emulsions, suspensions ; solid preparation: lozenge, film-coated tablet, capsules, granules, tablet, capsule and pellet ; external liquid)
• ISO 22000 (tablet, capsule, powder, granule, etc.)
• Halal (tablet, capsule, powder, granule, etc.)

**FOCUSED
THERAPEUTIC
AREA**

• OTC drug

INTRODUCTION

Pharmaceutical is an ethical business and we respect every life with care. Chung Mei Enterprises Group established "Chung Mei Biopharma" in 2014. Begin in 1936, Chung Mei Pharmaceutical Co., Ltd. (CM) is primarily centered on over-the-counter pharmaceuticals. For 85 years, we relentlessly collaborate with leading pharmaceutical firms, research centers and biotech centers, local and foreign. We committed to fully integrated cutting-edge manufacturing techniques and with new concepts through constant research and development. Chung Mei services from professional manufacturing to end-user sales (OEM & ODM), and integrates the biotechnology medical supply and sales chains. We can produce oral liquid preparation (syrup, solutions, emulsions, suspensions.); solid preparation (lozenge, film-coated tablet, capsules, granules, tablet, capsule and pellet) ; external liquid, and health food (tablet, capsule, powder, granule, etc.) We have also passed the high standards of cGMP, PIC/S GMP, ISO 22000 and Halal certifications. We anticipate bringing Taiwan's high-quality medicine into the Chinese and Asia markets. Let all of Taiwanese be proud of the mark "MIT".



**Johnson Chemical
Pharmaceutical Works Co., Ltd.**
強生化學製藥廠股份有限公司

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GMP CERTIFIED • TFDA (Oral Solids)

**FOCUSED
THERAPEUTIC
AREA**

- CNS
- Cardiovascular
- Gastro-intestinal
- Respiratory
- Vitamins
- Infectious diseases

INTRODUCTION

Johnson Chemical Pharmaceutical Works Co., Ltd. (aka. JCP) is a solid dosage form manufacture company founded in 1966. Over the years, JCP grows by expanding its capacity with a wide array of products, among which its CNS products hold the leading position in local market.

In 2010, JCP became the 7th PIC/S GMP certified pharmaceutical manufacturer in Taiwan. In 2013, the company was approved and went public for OTC listing in GreTai Market.

Through collaboration with academia, JCP is currently working on niche generics such as controlled release medicines in treatments for hypolipidemic - improved bioavailability with a lipid lowering agent. JCP is a certified manufacturer with Accreditation of Foreign Manufacturer from Japan's MHLW. JCP continues to grow as an international pharmaceutical company of high caliber by leveraging innovation, differentiation, quality, talent and globalization.

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GMP CERTIFIED · CRO Service (N/A)

FOCUSED THERAPEUTIC AREA · Contract Research Organization

INTRODUCTION Mithra Biotechnology Inc., the first Contract Research Organization (CRO) in Taiwan since 1988 offers integrated and multi-dimensional CRO services including bioanalytical services for chemical drugs and biologics, full characterization plans for biologics, bioavailability/bioequivalence (BA/BE) studies, and early phase clinical trials to facilitate pharmaceutical development. With profound expertise in biological products diverse in monoclonal antibodies, growth factors, antibody drug conjugates (ADC) and other various proteins, the company's services include primary structure characterization, PTMs identification, disulfide bond assignment, carbohydrate structure analysis, variants analysis and peptide mapping for biosimilar comparison or lot-to-lot QC, and degradation/impurities profiling. The company has proven expertise and experience in meeting international standards to serve the fast growing pharmaceutical industry, as evident from the accredited GLP compliance laboratory and GMP QC testing laboratory working on both small molecule drugs and protein drugs. The laboratory is equipped with state-of-the-art analytical instruments for sensitive and high throughput drug quantification in various biological matrices. A great variety of comprehensive methods have been developed and validated for contract research, as well as SOPs been established and applied to ensure the result been delivered to its highest quality.

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GMP CERTIFIED · PIC/S GMP: 1. semi-solid dosage forms
2. solid dosage forms: coated tablet, tablet, granules, capsule, suppository, soft capsule
3. hormone products (semi-solid dosage forms, coated tablet, tablet, capsule)
· Japan PMDA: 1. coated tablet/ tablet
2. soft capsules
3. hormone related products.

FOCUSED THERAPEUTIC AREA · Oncology (Non-cytotoxic)
· Hormone, Obstetrics & Gynecology
· Genitourinary
· Metabolic Diseases
· Gastroenterology (GI)

INTRODUCTION Pei Li was established in 1976, focusing on research and development of high barrier generic drugs and healthcare products. Our facility has been passed Japan PMDA and Taiwan PIC/S GMP inspection. There are 6 products which have been approved in Japan. Among these products, three are hormone-related drugs and one is high potent soft capsules.

Pei Li's Specialty:

1. Specialize in developing and manufacturing soft gelatin capsule (soft gel) for poorly soluble compounds or highly potent active ingredients.
2. Specialize in developing and manufacturing highly potent pharmaceutical products. Meanwhile, Pei Li has a dedicated production facility for highly potent pharmaceutical products, which can prevent cross contamination between high potent products and general pharmaceutical products.



**Prince
Pharmaceutical Co., Ltd.**
王子製藥股份有限公司

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GMP CERTIFIED · PMDA
· TFDA

**FOCUSED
THERAPEUTIC
AREA** · API

INTRODUCTION Prince Pharmaceutical Co., Ltd. has been established in 1962. Now, we focus on two business scopes. One is APIs, the other one is Dietary Supplements.

With over 50-year experience, we are one of the world's leading producers of vitamin B1 derivatives active pharmaceutical ingredients, which has been successfully supplied to Taiwan, Japan and Europe etc.

Not only APIs of vitamin B1 derivatives, we also developed numbers of APIs such as Ufenamate, Tulobuterol, Aldioxa, Felbinac etc., many new projects of R&D are ongoing. Furthermore, in order to expand capacity of plant and fulfill new business, our new APIs plant which is located in Yulin started operation in 2020.

In business of Dietary Supplements, we provide professional OEM/ODM service for well-known domestic and foreign customers.

For years, Prince Pharma has always insisted on our corporate belief of "A company that improves people's health by providing the best pharmaceuticals" and treated customer satisfaction as our highest priority.



**Savior Lifetec
Corporation**
松瑞製藥股份有限公司

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GMP CERTIFIED · TFDA
· USFDA
· MHRA

**FOCUSED
THERAPEUTIC
AREA** · Antibiotic

INTRODUCTION Savior Lifetec Corporation (SLC) is a specialty injectable company based in Chunan, Taiwan. Our state of the art facilities are vertically integrated, capable of producing sterile API and finished dosage formulations. Our facilities comply with the highest quality standards, having passed inspections by authorities from the US, EU, and Japan. SLC also has dedicated state-of-the-art GLP compliant R&D centers for API research and pharmaceutical research.

SLC's expertise in sterile crystallization and the aseptic processing of sterile powder serves as the foundation for product innovation.



**Sinphar
Pharmaceutical Co., Ltd.**
杏輝藥品工業股份有限公司

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TEL +886-2-2760-3688

FAX +886-2-2760-0350

GMP CERTIFIED · PIC/S GMP
· PMDA

**FOCUSED
THERAPEUTIC
AREA** · Oncology
· Hormone

INTRODUCTION Sinphar Pharmaceuticals is a publicly traded company and a certified PIC/S GMP, ISO 9001, ISO 14001, ISO 22000, Food GMP and OHSAS 18001 pharmaceutical company in Taiwan.

"Safety, stability, validity, and convenience" are elements we constantly emphasis in our quality policy. Holding both strong R&D capability and stringent control over product quality, Sinphar has fully demonstrated trustworthy techniques and skills in food products, cosmetics, pharmaceuticals, and anti-cancer drugs in both own brands and contract manufacturing services.

Following years of development, Sinphar has gradually consolidated its R&D capabilities in cancer drugs and new drugs, allowing us to build Taiwan's first pharmaceutical R&D center that specializes in isolated, automated production of cancer injections and cytotoxic agents. For sales distribution channel, Sinphar was the first to establish a Sinphar Counter in 1997, adopting a shop-in-shop business model. Now there are approximately 1300 branches across Taiwan, making Sinphar Counter the largest specialized pharmacy in Taiwan.



**Standard Chem. &
Pharm. Co., Ltd.**
生達化學製藥股份有限公司

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FAX +886-6-633-2775

GMP CERTIFIED · PIC/S GMP
· US FDA (Oral Solids)
· PMDA (Oral Solids)
· KFDA (Oral Solids)

**FOCUSED
THERAPEUTIC
AREA** · Cardiovascular
· Digestive
· Diabetes
· Anti-microbial

INTRODUCTION Established in 1967, Standard Chem. & Pharm. Co., Ltd (SCP) is a renowned global pharmaceutical manufacturer with a strong international presence. We have gained widespread recognition for our commitment to excellence in the industry. Guided by our corporate philosophy of "Sincerity, Honesty, Excellence, and Innovation," SCP has emerged as a leading pharmaceutical company in Taiwan.

As the flagship company of the Standard Group, SCP offers a comprehensive product portfolio that encompasses Pharmaceuticals, APIs, and Health Supplements. This diverse range of offerings positions us as a fully integrated company, capable of meeting diverse market demands.

At SCP, we consistently surpass quality standards, consistently exceeding expectations. Our state-of-the-art manufacturing facilities have received numerous accolades from prestigious domestic and international regulatory agencies, including T-FDA, US-FDA, PMDA, KFDA, and TGA, among others. These recognitions serve as a testament to our unwavering commitment to maintaining the highest quality in our products.

Furthermore, SCP takes pride in our strong focus on customer satisfaction. We constantly strive to surpass partner expectations, ensuring mutually beneficial collaborations and solidifying our position as a leader in the Taiwanese pharmaceutical industry. By partnering with SCP, clients are guaranteed success in both the Taiwanese and international markets.

With decades of expertise and an unwavering commitment to innovation and excellence, Standard Chem. & Pharm. Co., Ltd stands as a trusted partner for pharmaceutical needs, driving progress in the global healthcare industry.

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GMP CERTIFIED • TFDA
• KFDA (Oral Solids, Hormone Oral Solids)

FOCUSED THERAPEUTIC AREA

- Cardiovascular
- Respiratory
- Hormones
- Urology
- Oncology
- CNS

INTRODUCTION Synmosa Biopharma Corporation, a listed company in Taiwan OTC stock market (TW.4114), was founded in 1980. We develop sustainable business ranging from R&D, manufacturing, contract manufacturing as well as sales and distribution for both pharmaceuticals and health supplements.

Through constant mergers and acquisitions of potential domestic candidates, we have built a series of PIC/S manufacturing plants that specialized in the manufacturing of metered-dose inhaler (MDI), hormone preparations, nasal sprays, effervescent tablet, granules and even ointment, shaping us into the leading pharmaceutical company in various therapeutic areas of expertise such as cardiovascular, respiratory, hormones, urology, oncology, and CNS. Until today, we are still continuously expanding and putting more and more efforts in developing innovative medications and brand-name drugs.

As for global business development, we show direct presence in China, as well as in Hong Kong where we allocated our own competent sales and marketing team; whereas for Northeast Asia and Southeast Asia market we have a long-established and stable collaborative business relationship with local partners. We believe in improving the quality of lives of those in need with the highest standard of pharmaceutical manufacturing technology as we move towards our vision of becoming the foremost biopharmaceutical corporation in Asia.

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GMP CERTIFIED • Taiwan FDA (powder, Solution, Crystal)
• US FDA (powder, Solution, Crystal)
• Japan PMDA (powder, Solution)
• EDQM (powder, Solution)
• Korea KFDA (powder, Crystal)
• Hungary FDA (powder, Crystal)
• Mexico FDA (Solution)

FOCUSED THERAPEUTIC AREA

- Cardiovascular
- Immunology
- Metabolic Diseases
- Psychiatry
- Respiratory

INTRODUCTION Syn-Tech was established in 1982 as a professional APIs manufacturer. In the past decades, all the employees in Syn-Tech followed the philosophy of honesty, sincerity, innovation and development to create a good reputed company in APIs industry.

We experienced in systems such as ISO and PIC/S GMP. We also participated all kind of audits from authorities and world brand companies. All these activities strengthened Syn-Tech became a high-quality producer. Until now, Syn-Tech still looks forward to improving its system and facility. Because we know medicine is for human life. Providing high quality products to cure people is the forever task in Syn-Tech.



**Swiss Pharmaceutical
Company Limited**
瑞士藥廠股份有限公司



**Taiwan
Biotech Co., Ltd.**
信東生技股份有限公司

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TEL +886-6-589-3966

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GMP CERTIFIED

- Solid Dosage Forms: (US FDA) Tablets, Granules, Capsules
- Semi-solid Dosage Forms: Ointment, cream, gel.
- Liquid Dosage Forms: Solution, syrup.
- Sterile Dosage Forms: 1. Injection Solution (Aseptic Process and Terminal Sterilization)
2. Sterile optic and nasal solution
- Cephalosporin Products (Dedicated Facility):
1. Sterile dosage forms: Powder for injection
2. Solid dosage forms: capsules and granules

FOCUSED THERAPEUTIC AREA

- Psychiatry and Central Nervous System
- Circulatory and Cardio-Vascular System
- Anti-Inflammatory and Analgesic
- Dermatologic and Topical Product
- Woman Health and Dietary Supplement
- Metabolism System
- Anti-Infectious
- Respiratory System
- Digestive System

INTRODUCTION Swiss Pharmaceutical was established in 1966 with the mission of developing and manufacturing quality pharmaceutical generics products. Throughout the past 55 years, Swiss operates with the mindset of continuous improvement to pursuit this mission. Headquartered in Tainan City and currently has 400+ outstanding staffs and annual revenue of USD 33 million in FY2020.

Swiss has gone through several transformation. The most recent key transformation started in 2007, when we acquired a 40,000 m2 property to build our new facility. The facility was PIC/S GMP certified in 2012 and has opened various of opportunities which eventually lead to our first US ANDA CMO project. Swiss was inspected in 2017 and 2019 by the US FDA.

For next step, we will continue to sharpen our skill in both product development and manufacturing and are seeking for partners who has the same mindset to form strategic alliance.

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GMP CERTIFIED

- Taiwan FDA (All)
- Japan PMDA (Sterile Injection, Sterile Eye Drops)
- ISO 13485 (Contact Lens Care Solution)

FOCUSED THERAPEUTIC AREA

- Ophthalmology
- Cardiovascular
- Central Nervous System
- Respiratory
- Metabolic Diseases
- Antimicrobial Agents

INTRODUCTION Taiwan Biotech is a PIC/S GMP pharmaceutical manufacturer in Taiwan. Established in 1945, we have more than 1,000 staffs now and have devoted over 78 years to develop and supply the superior generic drugs.

The products we offer cover a wide range of therapeutic categories focus on cardiovascular, CNS, diabetic, respiratory and eye care drugs. With the complete production capability, we provide the oral solids, injections, inhalation solution, eye drops, TDDS patch, creams and ointments as a one-stop shop to our customers.

Taiwan Biotech is specialized in the manufacture of aseptic Blow-Seal-Fill products, including the unit dose eye drops, contact lens care solution, inhalation solution, injection, oral solution and wound cleaning solution. Filling volume is available from 0.2ml to 20ml. With its profession in this field, Taiwan Biotech is a CMO partner for the top OTC and generic companies for over 15 years.

We provide the professional CMO/CDMO services and welcome the out-licensing and in-licensing business opportunities.



**Taiwan
Tasheh Biotech Co.,Ltd**
大協生化科技股份有限公司

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GMP CERTIFIED · ISO 9001:2015 & ISO 15378:2017
Pharmaceutical Packaging Rubber Components

FOCUSED THERAPEUTIC AREA · High-end Laminated Rubber Stoppers
· RTU & RTS Rubber Stoppers
· Pharmaceutical Packaging Related Rubber

INTRODUCTION Founded on 1975, with Taiwan's Ta Sheh Heng Yi Co., Ltd. as the parent company, Sheng Zou Group has spent more than 40 years researching, manufacturing and adapting to the global requirements necessary to supply vendors with the proper medical packaging materials. Current market demands include medicinal butyl rubber stoppers (for powder injection, freeze dried, fluid infusion, biological agent, oral liquid, etc.), high-end laminated rubber stoppers, RTS rubber stoppers and medical equipment-related rubber and plastic products like syringe plunger seals, IV injection discs, also medical cosmetic products like dropper plugs, inner plugs.

Currently, Sheng Zou Group has four factories, three are located in China, at Hefei, Suzhou, Kunshan, as well as manufacturing and R&D facilities in Hsinchu, Taiwan, and the Division in Suzhou is responsible for the global marketing. Sheng Zou Group continues to lead the industry with more than six billion pieces of rubber stoppers annually.



**UBI Pharma
Inc.**
聯亞藥業股份有限公司

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GMP CERTIFIED · Taiwan FDA (sterile injection, lyophilisate for injection, liquid solution, semi-solid, tablet, capsule, powder, biological medicinal product)
· US FDA (sterile Injection, lyophilisate for injection, semi-solid)
· Japan PMDA (sterile injection)

FOCUSED THERAPEUTIC AREA · All except high-potency, cytotoxic, sex hormone

INTRODUCTION UBI Pharma Inc. (UBIP) is one of the leading pharmaceutical companies in Taiwan. Established in 2014, UBIP specializes in developing and manufacturing high quality generics and fusion protein drugs, as well as providing contract development and manufacturing services to global clients.

Our facility is routinely inspected by TFDA, USFDA, and PMDA. We are capable of manufacturing finished dosage form of sterile (vial, ampoule and lyophilized) and non-sterile (tablet, capsule, liquid, powder, cream and ointment), and have been the contract manufacturer for multiple world renowned pharmaceutical companies for more than 20 years. We are also selected for manufacturing of government controlled drug and COVID-19 vaccine.

UBIP is continuing seeking global partners for providing high-quality, affordable and innovative medicines to help patients lead healthier lives.



— TPA website —

Taiwan Pharmaceutical Alliance

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