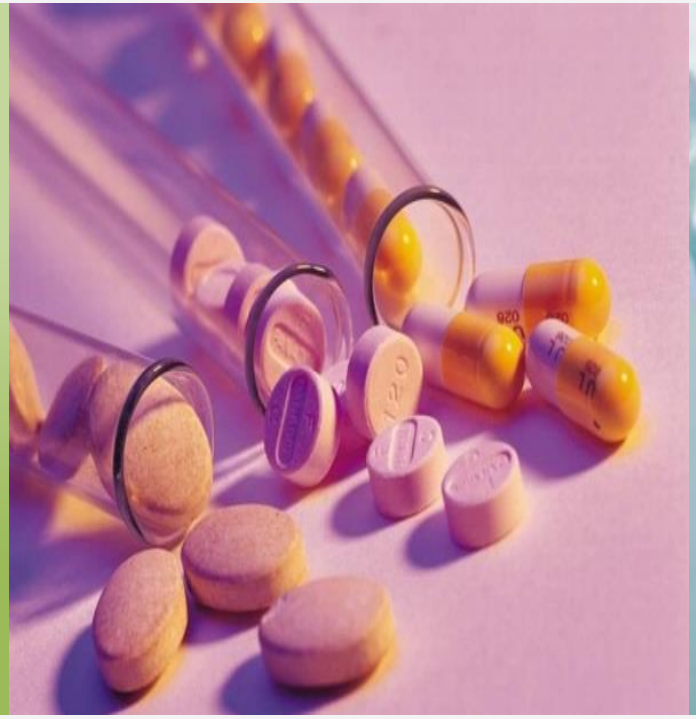
PHARMADAX INC

PHARMADAX

IMPROVING QUALITY OF LIFE



Content Overview

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- **Pharma**dax Advantage

Mission and Vision

Our Mission is to improve quality of life of all our stakeholders,

Of our Patients; by supplying high quality & cost effective generics

Of our Employees, by offering safe and healthy work environment

Of our Investors, by conducting business ethically and creating sustainable business value

Our Vision is to build a leading generic specialty pharmaceutical company with enhanced focus on specialty modified release and become preferred development partner for global organization.

PHARMADAX

IMPROVING QUALITY OF LIFE

Headquarter and R&D Facility

PHARMADAX focuses on developing and manufacturing specialty generic drugs and new drugs. PharmaDax commenced the business in 2008 and headquartered in Taipei, Taiwan.

Organization and Functions at headquarter:

- Research & Development
- GMP Management
- GMP Quality Center
- PK ,Biostatistics
- Clinical Management
- Quality Assurance
- Analytical & Validation Services
- Regulatory compliance
- Administration Management
- Finance& Accounting
- IT

5F-1, No. 236, Liancheng Rd.,
Zhonghe District, New Taipei City23553,
Taiwan(R.O.C.)

Total Headcount at HQ: 70 (28 in RnD)
Total Area: 1,600 square meters

Manufacturing Facilities

PHARMADAX has its manufacturing facilities located in Guangdong province, China, with combined capacity of 600 Mn tablets/year with expansion possibilities for other dosage forms.

- **Plant A**: Capacity of manufacturing 500 M tablets per year. It is inspected by FDA Twice in 2013 (No 483) and 2016 (One 483), which exhibits our commitment to compliance and quality

Land Scale: 6,666 square meters.

- **Plant B**: Construction completed in 2014 with potential manufacturing capacity of 100 M tablets per year, with large expansion capabilities for other dosage forms

Land Scale: 42,000 square meters.

Total Area: 48,666 square meters

Our Leaders

Yipin Huang
Chairman

- St. John's University MSc in Industrial Pharmacy
- Anchen Inc.(US)-Director of the Board & Senior VP
- Anchen Inc. Taiwan Branch-Director of the Board & President
- Empax Pharma Inc.-Founder& Director of the Board& President
- 25+ years of experience in pharmaceutical industry

Huiju Chan
President

- Soochow University EMBA management
- Anchen Inc. Taiwan Branch-Director of the Board& Vice President
- Empax Pharma Inc-Founder& Director of the Board& Vice President
- 28+ years of experience in pharmaceutical industry

Executive Team

Margaret Choy
EVP

- University of Southern California MSc in Regulatory
- Par Pharmaceuticals - Senior VP, Regulatory Affairs.
- Anchen Pharmaceuticals Inc. - Senior VP, Regulatory Affairs.
- Watson Pharmaceuticals – Director, Regulatory Affairs.
- Filed over 25 US "First to File" ANDA.
- 28+ years of experience in pharmaceutical industry

Name	Position	Education Background	Work Experience	Experience in industry
Susan Liao	Vice President	China Medical University, Taiwan; BA in Medicine	•Anchen Inc. Taiwan Branch-Senior Manager •Fujisawa Pharmaceutical Co. Ltd - QC/QA Manager (TW)	32+ years
Stephen Wong	Director	California State Polytechnic University, Pomona MBA	•Pharmadax Inc. the USBranch-Director •Anchen Inc.(US)-QC Manager •Associated Dir at Par and Anchen •Watson Pharma. (US)-R&D Lab. Manager	20+ years
Frank Lin PhD	Director of R&D	Tatung University; PhD in Chemical Eng.	•Anchen Inc. Taiwan Branch-Director of R&D department •Empax pharma Inc-Director of RD department of	25+ years
Jason Wu PhD	Director of Medical Affair	Taipei Medical University; PhD in Pharmaceutical Science	•China Chemical & Pharmaceutical Co., Ltd-Vice Chairman of R&D Dept. •Anchen Inc. Taiwan Branch - Vice Director	15+ years

Our journey so far…

FDA Submission: 3 approved, 2 under review and 7 in pipeline

2017 Quetiapine XR ANDA Approval and Launch in May '2017

2016 Plant A – USFDA GMP Renewal, One 483, EIR received Dec '16
FDA approval and launch of Glyburide Tablets in US

2015 Won the 12th annual Golden Torch Awards of Enterprise Excellence of R.O.C.

2014 ANDA Submission: Quetiapine Fumarate ER Tab (Para IV)

2013 Plant A passed USFDA inspection, Zero 483s, Listing on Taiwan Stock Exchange

2012 FDA approved Levetiracetam ER Tablet
The 2nd GMP plant (Plant B) was established in China
GMP QC Lab was set up in Taipei, Taiwan

2011 ANDA Submission : Glyburide Tab & Metoprolol Succinate ER Tab

2010 First ANDA Submission : Levetiracetam 750, 500mg ER Tablet (Keppra® XR)

2009 R&D center was set up in Taipei and China
Plant A got FDA production approval in China

2008 The GMP plant was set up in China (Plant A)
The headquarter of PharmaDax was established in Taipei

Business Reach

Pharmadax Group operates in the following geographies :

- ✓ Taiwan
- ✓ USA
- ✓ China
- ✓ Pharmerging

- **USA: Focused Market**

USA being the largest generic market and having management experience in the region, it was the first choice. Current pipeline is focused on blockbuster opportunities, riding the 2nd wave.

- **China: Next Frontier**

Generics had shown exponential growth and Pharmadax will leverage its manufacturing location and have initiated filings in the country, take for collaborative approach to push

Headquarter: New Taipei City, Taiwan
Manufacturing plant: Guangdong Province, China

US office: California, USA
European office: Slovenia, EU
India Office: Mumbai, India

Number of Total Employees : 210

Dosage Form Development

Pharmadax is committed to develop modified release generics in various therapeutic segments. The team brings a good mix of innovation and timely execution to ensure efficient dossier filing

We have a patented technology platform of high precision modified release, which can be used on large gamut molecules.

Our scientists ensure that this cutting edge technology continues to be flexible to bring more bioequivalent products to market.

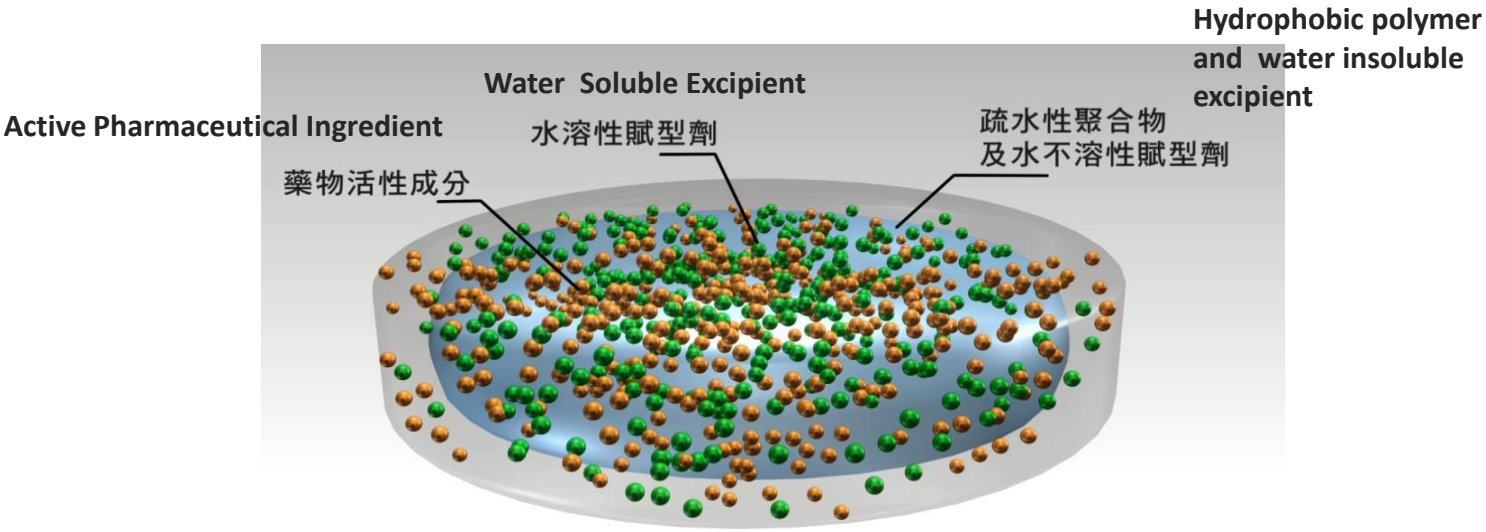
Number of employees of R&D in TW and CN: 35

Formulation Development

Manufacturing Process

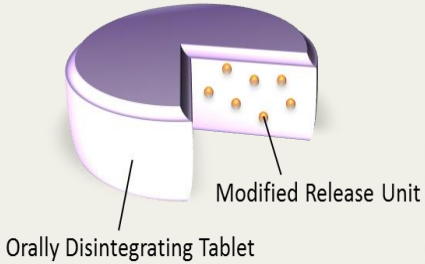
Core Technology

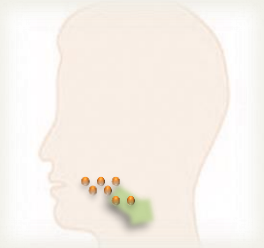
Pharmadax uses Non- Erosive Adjustable Drug Delivery System “NEADDS” technology to develop our products with better cost efficiency and enhance process stability.



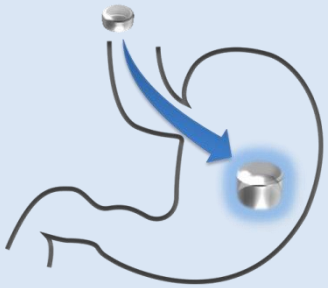
- US Patent : 62/066,917 Oral Controlled Release Matrix Tablet (NEADDS)
- Product Submitted to FDA Quetiapine Fumarate ER Tablets

Core Technology



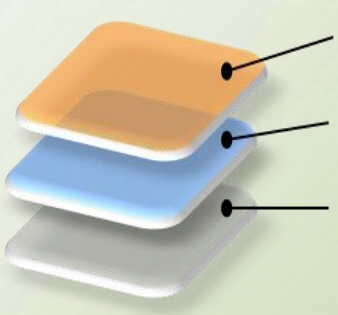


- **MR- ODT:**
  Modified- Release Orally Disintegrating Tablet



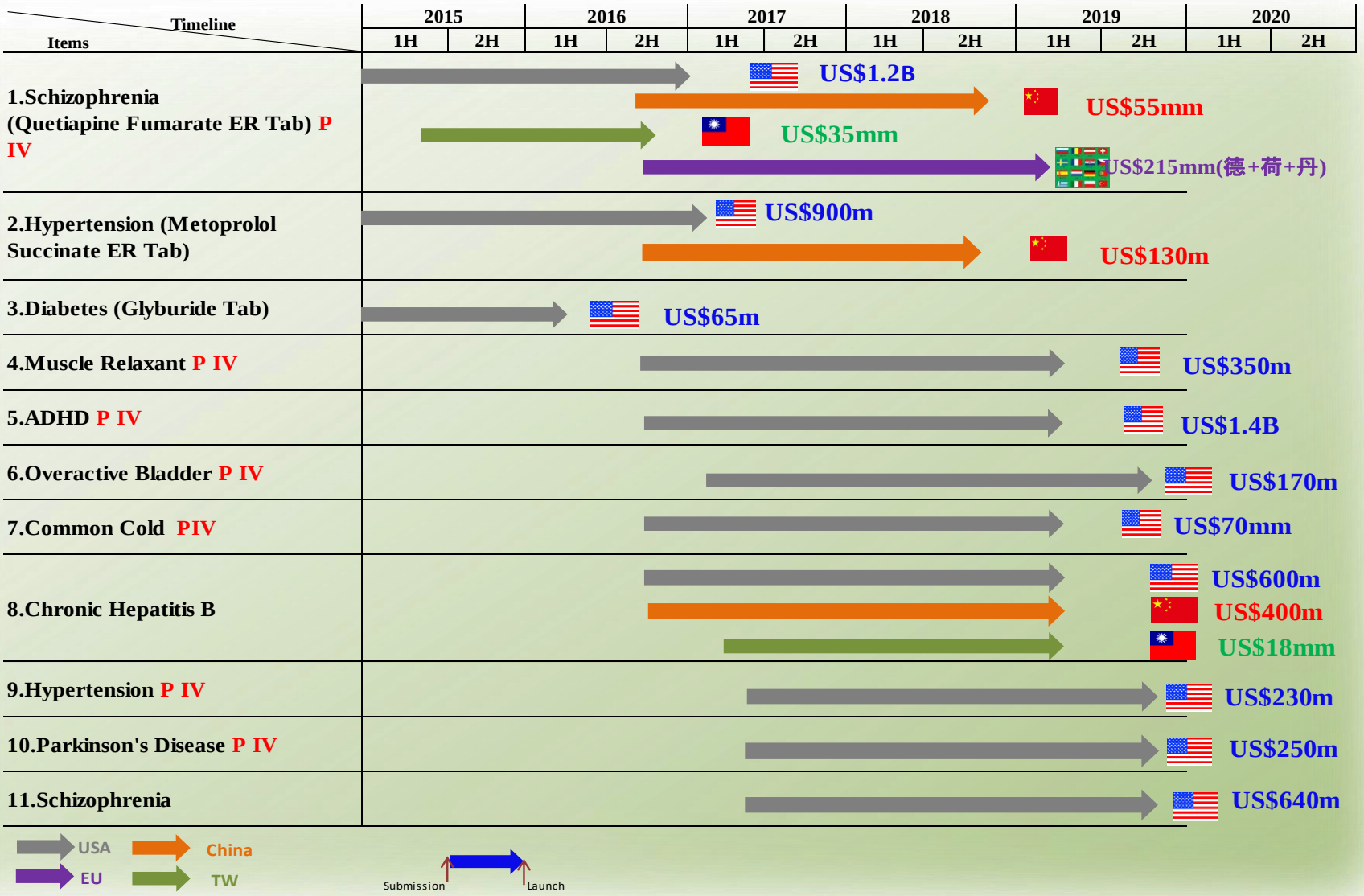
- **Sustained release gastroretentive dosage form**

Retention in stomach and sustained release



- **Ttransdermal patches:**

Portfolio


Timeline
Items
2015
2016
2017
2018
2019
2020
1H
2H
1.Schizophrenia (Quetiapine Fumarate ER Tab) P IV
US$1.2B
US$55mm
US$35mm
US$215mm(德+荷+丹)
2.Hypertension (Metoprolol Succinate ER Tab)
US$900m
US$130m
3.Diabetes (Glyburide Tab)
US$65m
4.Muscle Relaxant P IV
US$350m
5.ADHD P IV
US$1.4B
6.Overactive Bladder P IV
US$170m
7.Common Cold PIV
US$70mm
8.Chronic Hepatitis B
US$600m
US$400m
US$18mm
9.Hypertension P IV
US$230m
10.Parkinson's Disease P IV
US$250m
11.Schizophrenia
US$640m
USA
China
EU
TW
Submission
Launch

We Offer…..

Co-development and Marketing Partnerships

Regulated Market – CMO Opportunities – Tablets

Dossier development and Technology transfer services for Modified Release Products

Why Pharmadax

We believe in professionalism, we work with honesty and we perform decisively.

Quality and Compliance is our core ideology, which is exhibited in our products, people and partnerships.

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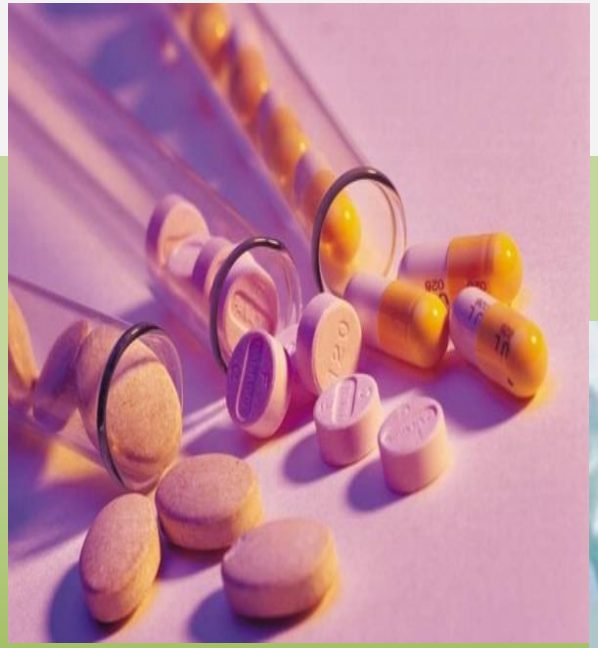

THANK YOU

For collaboration opportunities:
reachus@pharmadax.com OR
Call: +886-2- 2223-1552

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