



# A NEW DAWN IN HEALTHCARE FOR MYANMAR



pmipharma.com





## PARTNERING WITH PACIFIC MEDICAL INDUSTRIES (PMI)

 INTRODUCTION TO PMI, VISION & MISSION

 OUR MILESTONES

 OFFERED PRODUCTS & SERVICES

 COMPELLING REASONS TO WORK WITH PMI

 OUR QUALITY MANAGEMENT SYSTEM & ACCREDITATIONS



On July 23, 2017, we celebrated the official opening Ceremony of Pacific Medical Industries Ltd (PMI), a new US \$20 million manufacturing facility in Yangon, a symbol of progress for the entire nation. PMI is a sister company of A A Medical Products Ltd (Established in 1996), Myanmar's largest and most trusted pharmaceutical distribution company with branch offices throughout the country, as well as overseas subsidiaries in Singapore and the United States.



## VISION 2025

ISO compliance, WHO and USFDA approval of the multi-product generic facility with 3 ANDAs focused on Myanmar and targeting Global market with a value turnover of 30 Million USD by 2025.

## MISSION

1. PMI is a pioneer in the Myanmar pharmaceutical manufacturing environment of quality branded and generic drugs supported by innovations from a strong international R&D team.
2. We take pride in our cGMP compliant facility with necessary infrastructure and experienced technical personnel to manufacture products of reliable quality at affordable prices.
3. We strive to provide customer satisfaction to meet unmet market needs: making pharmaceuticals, nutraceuticals and OTC products accessible to the Myanmar and global market.



## PMI Milestones

AA Medical Products Ltd is founded by Mr. Zaw Moe Khine in 1996.

1996

Myanmar Investment Commission (MIC) Approval for construction of Pacific Medical Industries Ltd. was achieved on December 2011.

2011

GMP Conceptual and Basic Design of PMI was developed in accompany with Sure Assist (Korea) on January, 2013.

2013

Manufacturing Building was constructed by Min Dhama Company Ltd. (Myanmar), QC lab and offices Building was renovated by Sinma Construction Group Co., Ltd (Myanmar).

2014

Machines and Equipment Installation. Testing (FAT, SAT) & Commissioning Qualification (IQ, OQ, PQ) & Pilot Batch Manufacturing, etc. Recruitment Training (GMP, Technical & OJT).

2015-2016

2022

Achieved accreditation for 17025:2017 by ANAB

2020

Achieved ISO certification for 9001:2015 and Accreditation for 17025:2017 by SIS

2017

Manufacturing License Approval from FDA: Temp; (April 26, 2017), FDA Permanent License (Nov 20, 2017), FDA License for Powder/Granule Manufacturing (Feb 27, 2018), Opening Ceremony of PMI (July 23, 2017) Process Validation, Cleaning Validation and Commercial batch processing. Product Registration to FDA.

## OUR PRODUCTS

We are manufacturing the following wide range of Therapeutic Products (Oral solid dosage form: Tablet, Capsule, Powder) that are compliant against Safety, Efficacy, and International Quality Standards.

- Analgesic and Anti-Pyretic
- Anti-inflammatory
- Mucolytic
- Anti-infective
- Anti-Diabetic
- Antacid
- Anti-Hypertensive
- Cholesterol Lowering Agents
- Anti-Uric Acid/Anti-Gout
- Anti-Diarrhea
- Anti-Viral
- Anti-Helminthic
- Non Steroidal Anti-inflammatory
- Anti-Histamine
- Vitamin Supplement

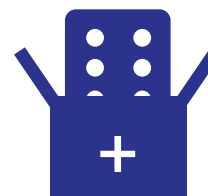
## DOSAGE FORM AND PACKAGING

PRODUCTS	DOSAGE FORM	PACKAGING
Oral Solid Products	Tablets (Plain/Coated)	Blister (Alu-Alu/Alu-PVC)
	Hard Capsule	Strip Bottle
	Powder	Sachet

## OUR SERVICES



Toll Manufacturing



Toll Packing



Toll Analytical Testing



Tablet Manufacturing  
(Wet and Dry Granulation) Method



Capsule Filling



Powder and Granules  
Manufacturing



Blister Packing



Sachet Packing



Bottle Packing

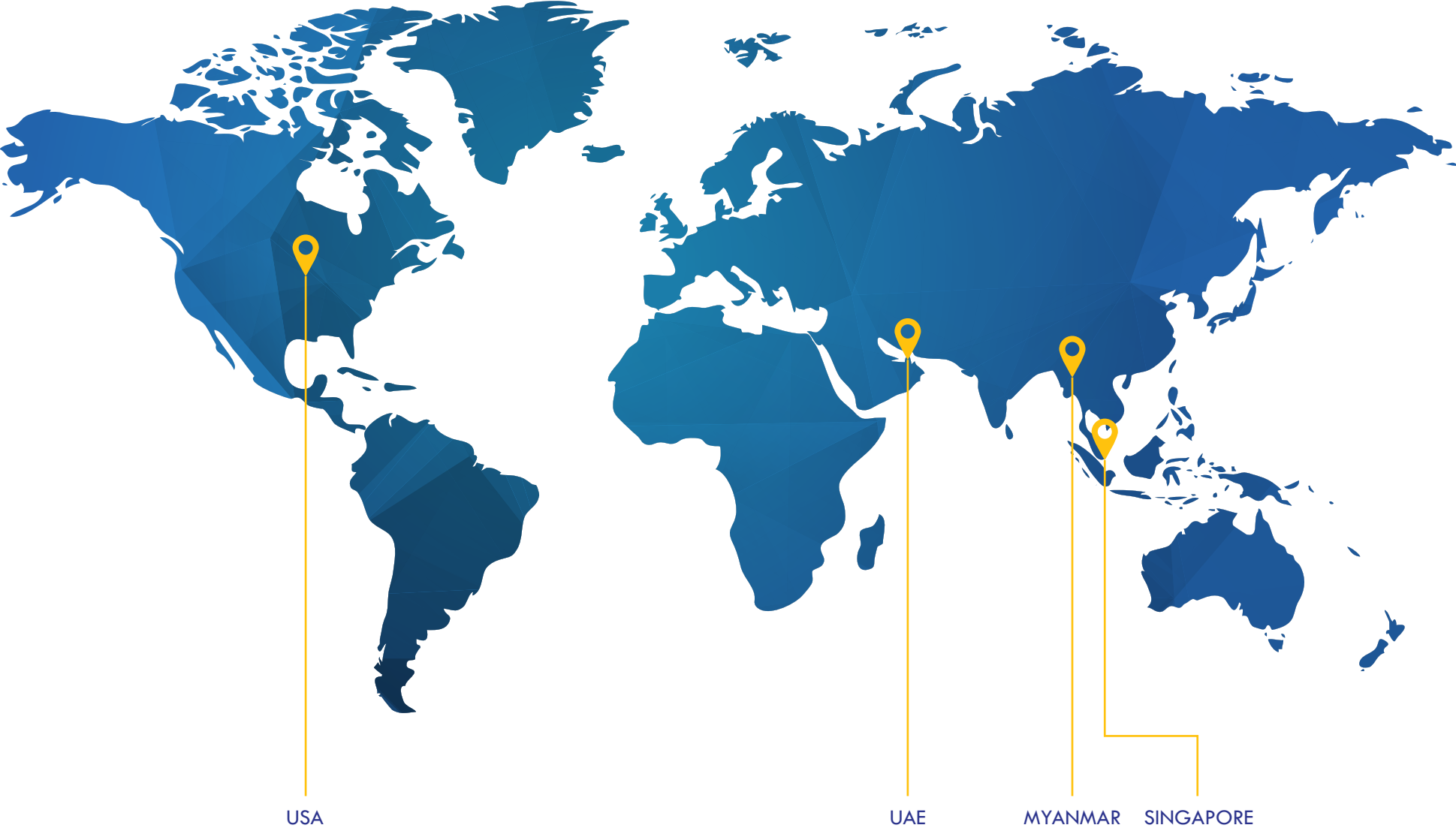


## COMPELLING REASONS TO WORK WITH PMI

1. Take advantage of local manufacturing by avoiding importation of products.
2. Ensure continuous and sustainable market supply of high quality products by manufacturing locally in a FDA GMP accredited facility with minimal risk of import restrictions.
3. Minimize currency, exchange rate & payment challenges by working with a local manufacturer like PMI.
4. Highly capable and competent expert local & expatriate staffs to seamlessly execute products transfer to the manufacturing site.
5. No more import challenges to keep product supply continuous.



# GLOBAL FOOTPRINTS



## QUALITY MANAGEMENT SYSTEM & ACCREDITATIONS

### OUR QUALITY SYSTEM

- Implementation of GMP
- R&D driven formulations
- Validated manufacturing Processes
- Qualified GMP Facility
- Dedicated Quality Control
- Qualified Competent Staffs
- Management Support
- Implement a quality management system
- Build a product strategy
- Consider competitors
- Customer Centric
- Always test your products
- Sustainable Supply
- Approachable Management Support



This certificate is valid only when accompanied by a current scope of accreditation document.  
The current scope of accreditation can be verified at [www.anab.org](http://www.anab.org).



This laboratory is accredited in accordance with the recognized International Standard ISO/IEC 17025:2017.  
This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory  
quality management system (refer to post BOMELAC-IAT Commencement dated April 2017).

### CERTIFICATE OF REGISTRATION



This Certificate has been awarded to

**PACIFIC MEDICAL INDUSTRIES LIMITED**  
No. 209/210/211, Corner of Khun Act Thar Minthar Street and Kanner Street,  
City of Industry, Dagon Seikkan Township, 11441 Yangon, Myanmar.

In recognition of the organization's Management System  
which complies with

**ISO 9001:2015 (QMS)**

The scope of activities covered by this certificate is defined below

**Development, Manufacturing, Packing, Holding, Testing, Distribution  
of Finished Pharmaceutical Products- Tablets, Capsules  
(Non-Betalactam), Powder, and Granules**

EA Code:- 13 & 35

SYNDICATE OF INTERNATIONAL SYSTEM CERTIFICATIONS

Certificate Number: **SISINDQ11202089688**  
Date of Issue of Original Certificate: **05.11.2020**  
Date of Issue of latest certificate: **23.09.2022**  
Expiry Date: **04.11.2023**  
Re-certification Due on: **05.10.2023**

*[Signature]*  
**Managing Director**



Note: This is an accredited certificate issued by SIS Certifications Pvt. Ltd.

Certified Organization is responsible for maintaining the compliance of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid

Corporate office- **SIS Certifications Pvt. Ltd.**  
Unit No. 514, 5th Floor, Vipul Business Park, Sector-48, Sohna Road, Gurgaon-122018, Haryana, India.  
International Key Locations: **Qatar, Egypt, Peru, Italy, KSA, Nigeria & Malaysia.**  
Email us- [support@siscertifications.com](mailto:support@siscertifications.com), Call/Whatsapp- +91-0643073391  
The status of this certificate can be verified on <https://siscertifications.com>  
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