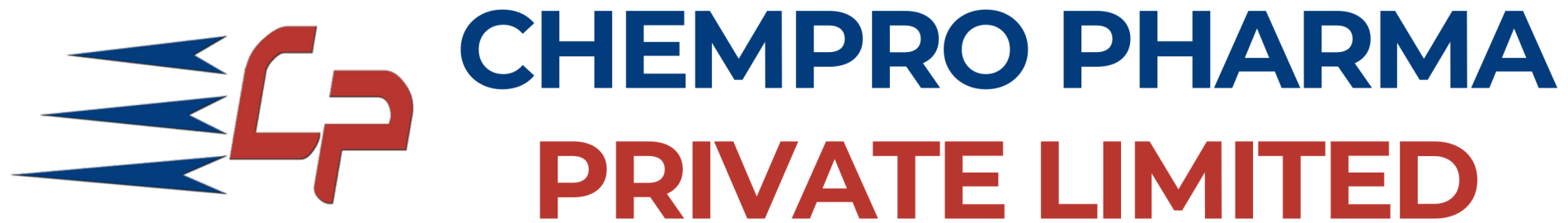


Serving The World Through Synergy



**A strategic tie-up that
leverages our strengths
to bring comprehensive
products, services and
tailored solutions for
developing the global
pharmaceutical industry**



- ❖ Focus on the development of pharmaceuticals and biosimilars
- ❖ Ensure regulatory compliance and commercial competitiveness
- ❖ Sustain profitable growth of business internationally while also supporting local manufacturing via technology transfers





18 Years of Excellence

100+ Successful Products & Technologies

1000+ Successful Solutions



CHEMPRO PHARMA PRIVATE LIMITED

Superior Quality

Higher Affordability

Reliable Service

50+ Manufacturing Partners (FDF)

200+ Manufacturing Partners (API & Intermediates)

15+ Manufacturing Partners (KSM)

10+ Impurity Manufacturing Partners (WS & RS)

5+ Regulatory Support Partners (BE Studies)

10+ Foreign Partners for Exclusive Raw Material Supplies



Mission & Vision

- ❖ Become one of the world's leading and most reliable organizations that supports all clients in ensuring global competitiveness
- ❖ Pursue excellence with commitment and support the global pharma industry through superior quality products and tailored solutions
- ❖ Sustain commercial viability on a global scale and empower our clients for import substitution
- ❖ Provide end-to-end solutions for all challenges related to pharmaceutical manufacturing

Global Presence



PharmSol
GmbH,
Germany



PharmSol
Europe Ltd.,
Malta



Eminus eGlobal
Institute Pty
Ltd., Australia



Pharmaceutical
Solutions FZCO,
Dubai



Pharmaceutical
Solutions
DMCC, Dubai



PharmSol
India Pvt Ltd,
India



PharmSol
(Nanjing)
Technology,
China



PharmSol
APAC Ltd,
Hong Kong



PharmSol -
Pharmaceutical
Products and
Trading,
Philippines



Agile FZCO,
Dubai



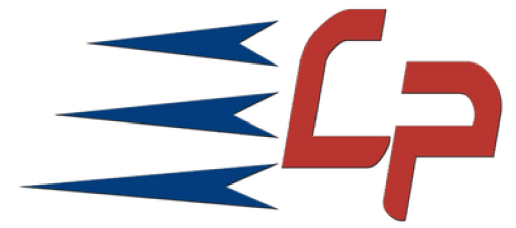
PharmSol,
Vietnam
(under
incorporation)



PharmSol,
Egypt (under
incorporation)



Global Presence



Products & Services



Licensing & Supply

**Contract
Development &
Manufacturing**

Co-developments

Tech Transfers

Product list available on request...

Custom Solutions





Products & Services



**Key Starting
Materials &
Intermediates**



**Active
Pharmaceutical
Ingredients**



**Finished
Dosage
Forms**



Products & Services



**Impurities
(Working & Reference
Standards)**



**Contract Research,
Development, &
Manufacturing**



**Custom Synthesis
(Fine Chemicals)**



Products & Services



**Regulatory Support
(Dossiers, Auditing,
BE & Clinical Studies)**



**Technology
Transfer**



**Cargo Consolidation
& Overseas Stock
Arrangements**

Track Record



25 (+) APIs
successfully developed
& commercialized



100 (+) FDF
successfully developed



300 (+) MAs



45 (+)
Tech Transfers



50 (+) BE
Studies



Strong bonds with 50 (+)
major manufacturer

Track Record



1200 (+)
GMP/QP Audits



40 (+) Facilities
certified by US / EU



12 (+) Facility
Compliance Remediations
(US/EU)



60 (+)
EU Registrations,
Ongoing **18 Procedures**



55 (+)
Analytical Methods



150 (+) Clients



CHEMPRO PHARMA PRIVATE LIMITED

Superior Quality

Higher Affordability

Reliable Service

3 Business Verticals - Pharmaceuticals, Chemicals & Cosmetics

40+ Years of Being a Preferred Partner in India

170+ International Clients

50+ Countries

35+ New Molecules Developed (CDMO)

25+ Warehouses for Overseas Stock Arrangements

Leadership



Sunil Doshi

Founder & Managing Director

36 Years of Global Techno-Commercial
Expertise In Pharma Industry



S. Lakshminarasimhan

Chief Operating Officer

25 Years of Pharma Technology,
Compliance & Regulatory

Leading a global team of 80 committed professionals.

Strategy



Christian Duchow
Director - EU Business

30 Years of Expertise in
EU Pharma Business



Arun Sehgal
Director - Strategy

40 Years of Expertise in
Global Pharma Business



Borut Strukelj
Senior Scientific Advisor

35 Years of Expertise in Biotech,
Biosimilars & Regulatory



Andrej Gasperlin
Senior Associate - US

35 Years of Expertise in
US Pharma Business

SMEs



Rajeev Patil

Director - Regulatory Affairs

40 Years of Expertise in USFDA
Regulatory Compliance



Ganesh Sangaiah

Director - NDDS

26 Years of Expertise in NDDS



Ashish Dasgupta

Senior Scientific Advisor

40 Years of Expertise in Clinical,
Quality & Regulatory

SMEs



Manish Rajeshirke
Vice President - BD

21 Years in Global
Pharma Business



Nancy Yao
Director - China,
Technical Operations
11 Years in China
Pharma Compliance



Felix Huang
Director - China, BD

10 Years in China
Pharma Business

Accreditations



BPI Bundesverband der
Pharmazeutischen Industrie e.V.

VERBAND DER
CHEMISCHEN INDUSTRIE e.V.
WIR GESTALTEN ZUKUNFT.



VCI



Clientele





Leadership



Arun Sehgal
Chairman & MD



Vijay Agarwal
Chief Executive Officer



Deepkiran Khanwalkar
Vice-President



Sunil Doshi
Director



Leadership



Shubhit Sehgal
Director



Vaandan Agarwal
Director

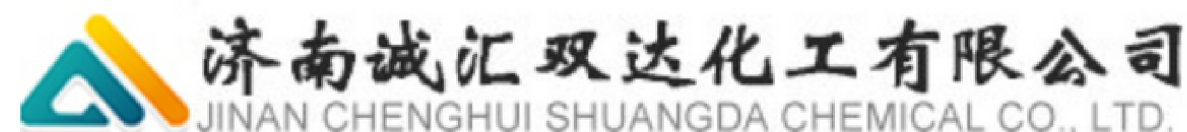
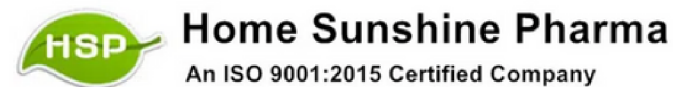


Abhishek Khanwalkar
Director



Vishal Agarwal
Director

Clientele

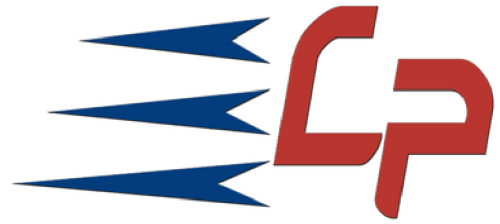




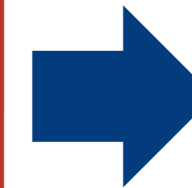
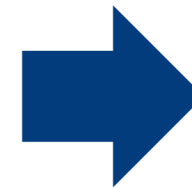
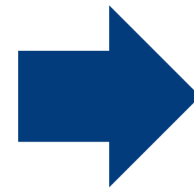
Building Global Competencies

- ❖ Facilitate technology transfers by bringing the best talent to clients
- ❖ Conduct regulatory compliance audits with our team of industry experts and consultants
- ❖ Perform regular cost competency audits with our team of global supply chain experts
- ❖ Develop efficient and competitive supply chains of manufacturers who are preferably already pre-approved by us
- ❖ Establish an infrastructure that ensures just-in-time inventory
- ❖ Support and expedite product registrations in the EU

Proposed Business Model



PERFORMING REGULAR COSTING AUDITS
FACILITATING SUPPLY CHAIN DEVELOPMENT
MAKING OVERSEAS STOCK ARRANGEMENTS



CONDUCTING REGULATORY COMPLIANCE AUDITS
PLANNING & EXECUTING TECHNOLOGY TRANSFERS

Note: We will also assist local manufacturers to get included in Group 1 for domestic tenders



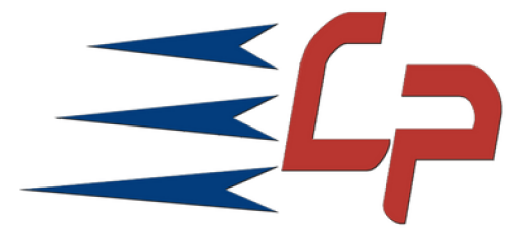
Combining Strengths

- ❖ 50+ years of cumulative experience
- ❖ Rich knowledge in developing, registering, and marketing generics in Europe (including 20+ First to File)
- ❖ Extensive experience in the domains of manufacturing, technology, compliance, regulatory, and marketing
- ❖ Offering innovative, comprehensive, and tailored solutions for all aspects of the Pharma Value Chain
- ❖ Cost-competitive and time-efficient turnkey projects
- ❖ Strong team of 75+ technocrats and growing...
- ❖ Associated with renowned industry experts specializing in Pharma, Biosimilars, Clinical, etc.



Combining Strengths

- ❖ 80+ years of cumulative experience
- ❖ Rich knowledge in contract development of new and innovative molecules with niche applications (35+ developed first in India)
- ❖ Extensive experience in the domains of manufacturing, technology, supply chain development, and marketing
- ❖ Offering innovative, comprehensive, and tailored solutions for value creation within the Pharma Supply Chain
- ❖ Designing cost competitive businesses to compete globally
- ❖ Strong local and international networks of industry experts



Empowering Industry... Globally

- ❖ Combining expertise, resources, and networks to assist manufacturers in achieving sustained profitable growth
- ❖ Supporting local production and providing global market access to our esteemed clientele for their products of interest
- ❖ Bringing the best of pharma competencies to YOU and introducing your newly enabled organization to the WORLD





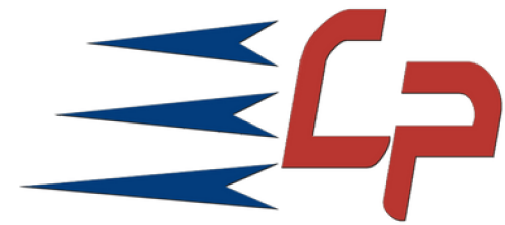
Our Approach

- ❖ **Transfer of efficient and cost competitive formulation technologies**
- ❖ **Build and develop a reliable supply chain for APIs**
- ❖ **Support clients in attaining EUGMP and other accreditations**
- ❖ **Register selected finished dosage forms in desired target markets**
- ❖ **Design and implement marketing strategies to generate a sustained stream of export revenue**
- ❖ **Helping manufacturers enter biopharmaceuticals via in-licensing and local development**



Our Approach

Cost Competitiveness	GMP Compliance	Market Access
<ul style="list-style-type: none">● Raw Material Pricing● API Inventory Management● Efficient Technology● Operational Efficiencies● Regular Audits of Supply Chain Partners● Regular Cost Competency Audits vis-à-vis Competition	<ul style="list-style-type: none">● GAP Audit of Facility● Compliance Guidance● Preparatory Audit Prior to● Authority Inspections● Training● Deputation of Experts at Site for Authority Inspection● Post Inspection CAPA Support● Continued Compliance	<ul style="list-style-type: none">● Product Registrations● Market Authorization● Batch Testing / Release● Finding, Liaising & Finalizing Marketing / Distribution Partnerships● Commercial Supply Chain Management● Post Marketing Follow Up



Conclusion

- ❖ Enabling manufacturers in gaining a competitive edge globally
- ❖ Enhancing GMP compliance provides new opportunities for entering regulated markets
- ❖ Getting an EU MA will help companies attain Group 1 status and increase the chances of commanding higher prices in local tenders
- ❖ Bringing top industry experts to assist entry into biopharmaceuticals
 - Developing capabilities for biosimilar production
 - Manufacturing with higher standards for quality & compliance
 - Swift implementation from conceptualization to commercialization

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THANK YOU!