SUANFARMA

INDUSTRIAL

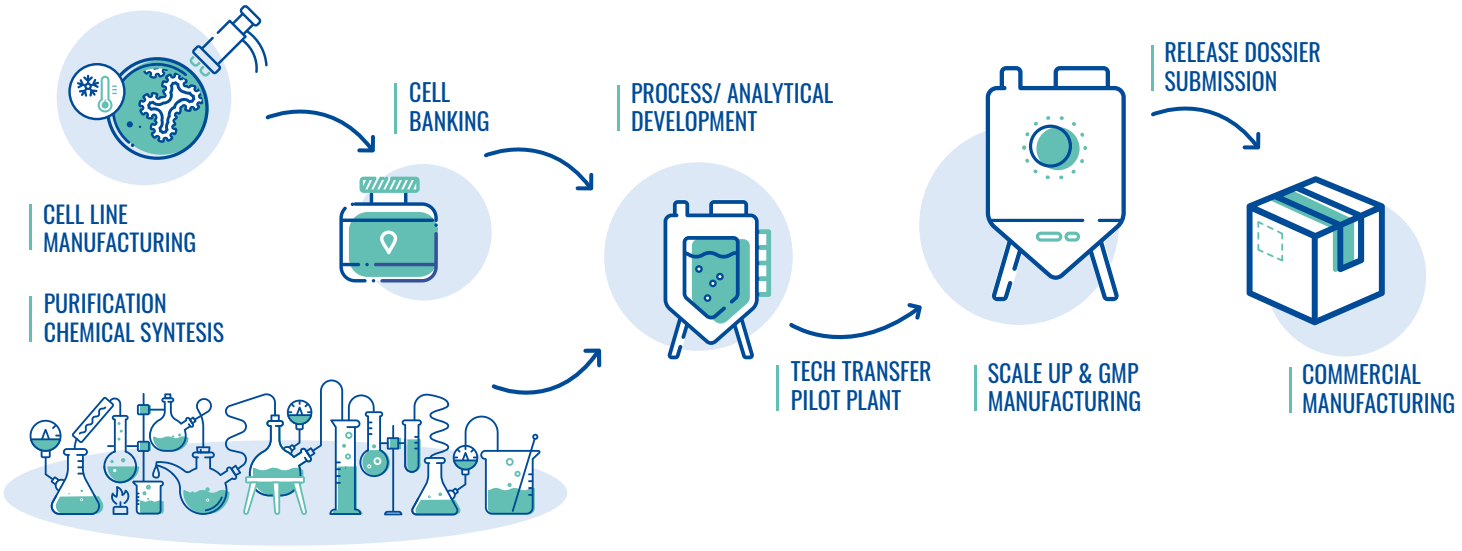
YOUR PARTNER TO SUCCESS

INNOVATIVE MOLECULES UNDER DEVELOPMENT OR CLINICAL TRIALS & FOR SUPPORT IN COMMERCIAL MANUFACTURING

At **SUANFARMA**, we are committed to meet our customers needs from a technical and commercial point of view. Our Industrial Division supports you in every phase of your project.

ONE STOP SHOP

DEVELOPMENT ➤ PRE-CLINICAL SUPPLY ➤ CLINICAL SUPPLY ➤ MARKET SUPPLY



We offer a tailored project which maximizes the chance of success in the achievement of our target.

Innovation race pursues to minimize the risks during early stages of development. For this reason, SUANFARMA has been working on strategies based on:

- Risks Based Management
- QbD (Quality by Design) - GAP & Risk Analysis
- A strong Tech Transfer Policy applied both to internal and external transfer processes

tt&GO
SPEED TECH TRANSFER 2 SUCCESS

PROVEN TRACK RECORD

CUSTOMER SUCCESS & SATISFACTION IS OUR VICTORY

WHY INDUSTRIAL SUANFARMA?

We have over 60 years of experience in the industry and we have been working in the development of our own molecules, ensuring the understanding against possible challenges and difficulties arising in these stages for the development of a molecule.

We provide a proven track record in development and commercial manufacturing. More than 28 molecules, 18 CDMO clients, 13 countries...

We ensure the sales capacity over the next 5 years and provide the client with all the investment needs to ensure a consensus on the growth management prior to the consumption of the available capacity.

We work under the strictest confidentiality policy. Our encoded processes ensure the non-disclosure of industrial secrets.

We work to guarantee a consensus in the decision-making process affecting any process performance.

We offer full transparency in communication. We build project teams adding optimal interaction and collaboration, understanding this as a key success factor.

TT&GO offers the highest degree of technological innovation throughout our acquired knowledge & experience.

We offer maximum flexibility to adapt ourselves to changes arising during initial stages of the project, providing the client with all resources required to avoid delays or idle times.

We continue working on energy saving strategies that allow a significant improvement in environmental sustainability with innovative measures which have an impact on lower cost in the energy bill. We are clearly committed to having efficient manufacturing plants, more competitive in our proposal to the market.

We are moving forward in our expansion plan to undertake the high demand for CDMO manufacturing capacity. This plan is a proof of confidence in our CDMO business and allows us to continue growing in number of projects, requests, and plant capabilities.

We have regulatory experience in the preparation of Registration Dossiers of pharmaceutical products around the world. This contributes to avoid delays in periods and opens up the possibility of accelerating the time to market.

We are a solid and reliable partner to your routine manufacturing.

FOR FURTHER INFORMATION PLEASE CONTACT:

David Blanco, PhD, PMP
Senior Industrial CDMO Manager
M + 34 610 665 080
davidblanco@suanfarma.com