



your
partner
that
cares

INTEGRATED CRDMO
ONE-STOP SHOP SOLUTION

Tailored solutions for product development

 **bluepharma**

Integrated CRDMO

END-TO-END SOLUTIONS

Unlocking Global Success:
Your European CRDMO
powering your product journey.

	Early Development	Clinical Phase	Validation	Regulatory Support	Commercial Stage
Pre-formulation Studies	●				
Formulation Development QbD Based	●				
Analytical Development	●				
Prototype Scale-up	●				
BioPharmaceutical Assessment	●				
Clinical Trial Materials (CTM)		●			
First in Human and PK Studies BA/BE Studies		●			
<i>In vitro/In vivo</i> Correlation		●			
Packaging of Investigational Medicinal Product		●			
QC and QP Release		●			
Clinical Supply Management (Phase I/II/III)		●			
Manufacturing Process Validation			●		
Analytical Validations			●		
ICH Stability Studies			●		
IMPD and CMC				●	
Dossier Preparation for Regulated Markets				●	
Support for the Post Regulatory Filing Queries				●	
Manufacturing and Supply Chain Management					●
Life Cycle Management Solutions					●

We offer **tailored solutions** and a **collaborative partnership** that truly values your vision.

Integrated CRDMO

TECHNOLOGIES AND PLATFORMS

Unlocking the full potential of your molecules with our comprehensive CRDMO-driven technical solutions.

Technologies & Platforms

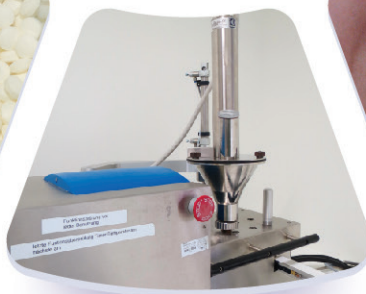
Oral Solids

- . High Potent Molecules
- . Controlled Substances



Modified Release

- . Matrix Release
- . Coating Agents
- . Pellets and Beads
- . Dual Release Layers



Hot Melt Extrusion

- . High-throughput Screening
- . Conventional and HPAPI Molecules
- . Lab and Commercial Scale

Complex Sterile

- . Liposomal Formulations
- . Lipid-based Nanoparticles
- . Polymer-based Nanoparticles
- . Solid Sterile Formulations



Oral Mucosal Delivery

- . Buccal Sprays **BlueEase™**
- . Oral Dispersible Tablets
- . Sublingual/Buccal Tablets
- . Oral Thin Films **BlueOS®**



Comprehensive Solutions backed by
End-to-End Analytical Expertise
and **Advanced Techniques.**

Integrated CRDMO

ASSETS AND INFRASTRUCTURE



3 R&D CENTERS

Dedicated to Innovation with expertise in Oral and Injectable Forms.



2 COMMERCIAL MANUFACTURING

Unlock Success with Bluepharma: Global Reach and a Proven Track Record of Success.



1 GMP PILOT SCALE AREA

- Tailored small GMP batches including highly potent products reducing the API needs.
- Ideal for clinical manufacturing from Phase I to III studies.



mRNA & ADVANCED THERAPIES HUB

A Technological Hub dedicated to the Innovation, Translation and Industrialization of complex injectable drugs.



scan me
State-of-the-art
industrial unit video



Integrated CRDMO

OUR UNIQUE VALUE PROPOSITION

One-stop Shop Solution

Tailored solutions for product development and clinical studies until market reach.



HARMONIZED PROCEDURES AND SWIFT COMMUNICATION

Integrated approach with harmonized procedures. A dedicated project manager to unify teams, capabilities, and ensure clear communication throughout your project.



AGILITY & MILESTONE SYNCHRONIZATION

Ensure that drug product development, clinical manufacturing, clinical supply forecasting, demand planning, and clinical trial supply services activities are synchronized.



INTEGRATED APPROACH: FROM IN VITRO TO IN VIVO

Bluepharma will be able to integrate all the pre-clinical, in vitro, and clinical data to define the best solution, avoiding common pitfalls and costly delays.



NO HANDOFFS AND REDUCED ADMINISTRATIVE BURDEN

We cover all stages from early development to commercialization and handle all clinical studies, eliminating the need for technology transfer to handover. Reducing time and costs!



STREAMLINED SOLUTIONS THAT ACCELERATE TIME TO MARKET

Knowing the development challenges, we can provide an end-to-end solution with scientific and technical insight to reduce redundancies and detours.

Speed to Market

Our agile processes accelerate your product's journey from development to market.



+750
Employees



PhDs
+40

Let's turn your ideas into reality

Take the next step and contact us to discuss how we can collaborate on your next pharmaceutical project.



Differentiation & focus on client needs



Integrated offer
(DEVELOPMENT, CLINICAL, MANUFACTURING)



Performance and track record



Focus on time to market



Reliability and sustainability

Bluepharma's facilities are **EU-GMP** certified and approved by:

- US FDA (2009, 2012, 2014, 2016, 2019)
- SFDA (Kingdom of Saudi Arabia)
- ANVISA (Brazil)
- MFDS (Republic of Korea)
- Minpromtorg (Russian Federation)
- MOH Libya
- TMMDA (Turkey)

We are registered in Iraq, Taiwan, Jordan, UAE, Kurdistan and Vietnam.



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