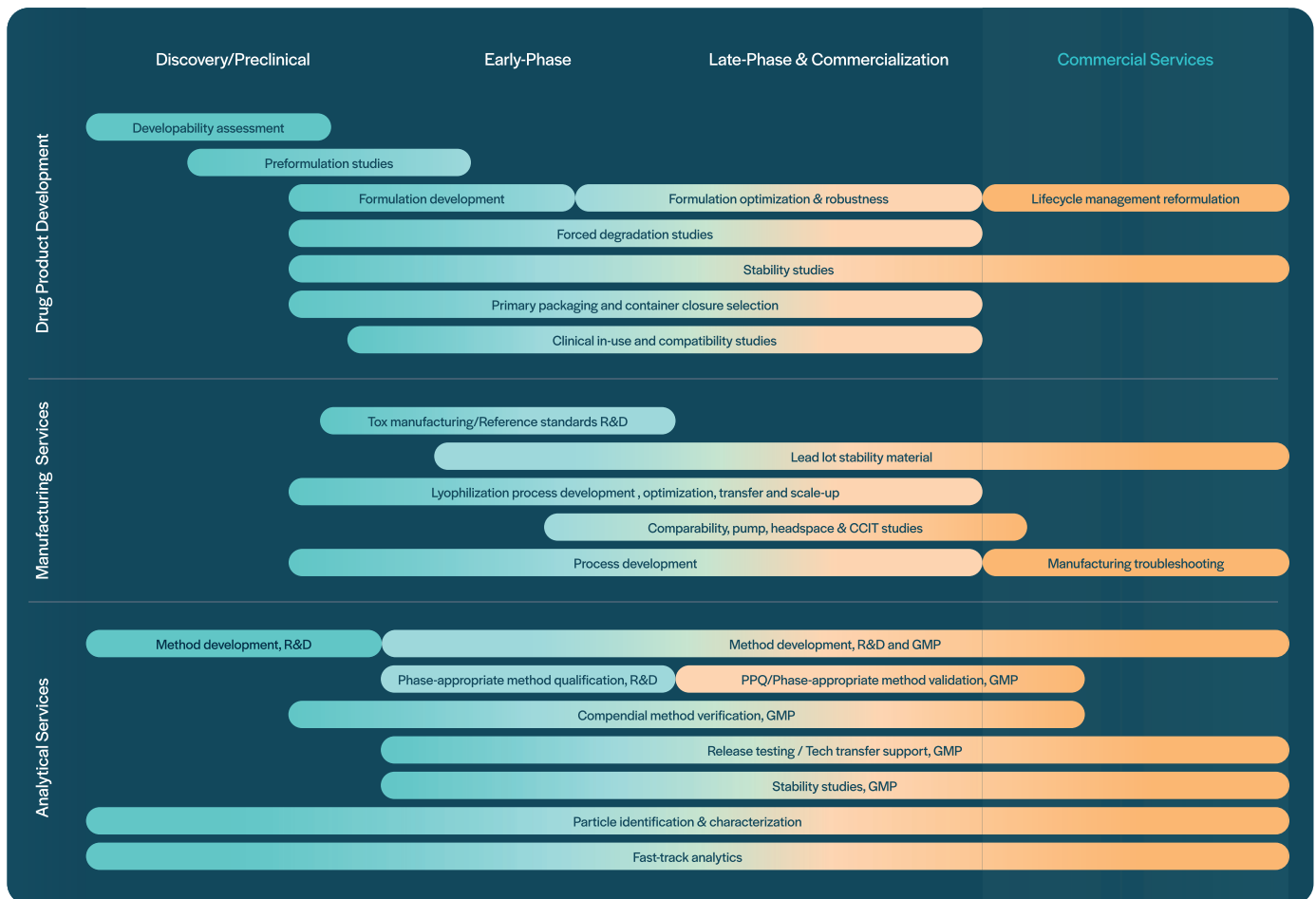


OUR SUPPORT FOR YOU  
**ACCELERATING AND DERISKING  
 YOUR PATH TO MARKET**

Coriolis Pharma is a global contract research and development organization (CRDO) and the premier partner for the development of liquid and lyophilized drug products, analytical services, and manufacturing services across a vast array of therapeutic modalities,

including biologics, gene and cell therapies and vaccines. Our expert scientists design and execute platform and custom services under R&D and GMP to accelerate and derisk your program right from the start and throughout the entire product lifecycle.

## DEVELOPMENT, ANALYTICAL AND MANUFACTURING SERVICES



## CORIOLIS BY THE NUMBERS



## OUR SUPPORT FOR YOUR MOLECULE

Coriolis provides expert support and high quality services across all biologics, cell and gene therapy classes.

- Antibody or Antibody-Related
- Peptide
- Therapeutic Protein
- Gene & Cell Therapy
- Viral Vector
- Nucleic Acid
- Vaccine
- Nanoparticulate Delivery Systems

## WHY CHOOSE CORIOLIS?

- Full lifecycle support with the end goal in mind from preclinical to commercial under R&D and GMP
- Development of drug products designed for global launches, specifically meeting the requirements of the FDA, EMA and Japanese regulatory standards.
- Client-focused collaboration with an independent service provider, offering dedicated attention to your specific project, molecule and development phase
- Support by Coriolis' scientific experts and its scientific advisors with multidisciplinary expertise
- Access to accelerator platform services and custom program capabilities
- High-quality, reliable and consistent release testing, including stability studies on clinical and commercial batches
- Short turn-around times and on-time delivery of results
- Dedicated ATMP facility up to biosafety level 2
- Seamless transition and continued collaboration with you or your DS and DP CMO
- Representatives and laboratories in close proximity to you and your team in the EU and USA

## DISCOVER THE CORIOLIS EFFECT NOW



Mutual Confidential Disclosure Agreement

Step 1



Request for Information/Proposal & Project Discussion

Step 2



Proposal Development  
(includes biosafety preassessment)

Step 3



Development Service Agreement/Master Service Agreement

Step 4



Project Start  
(includes completion of biosafety assessment)

Step 5

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