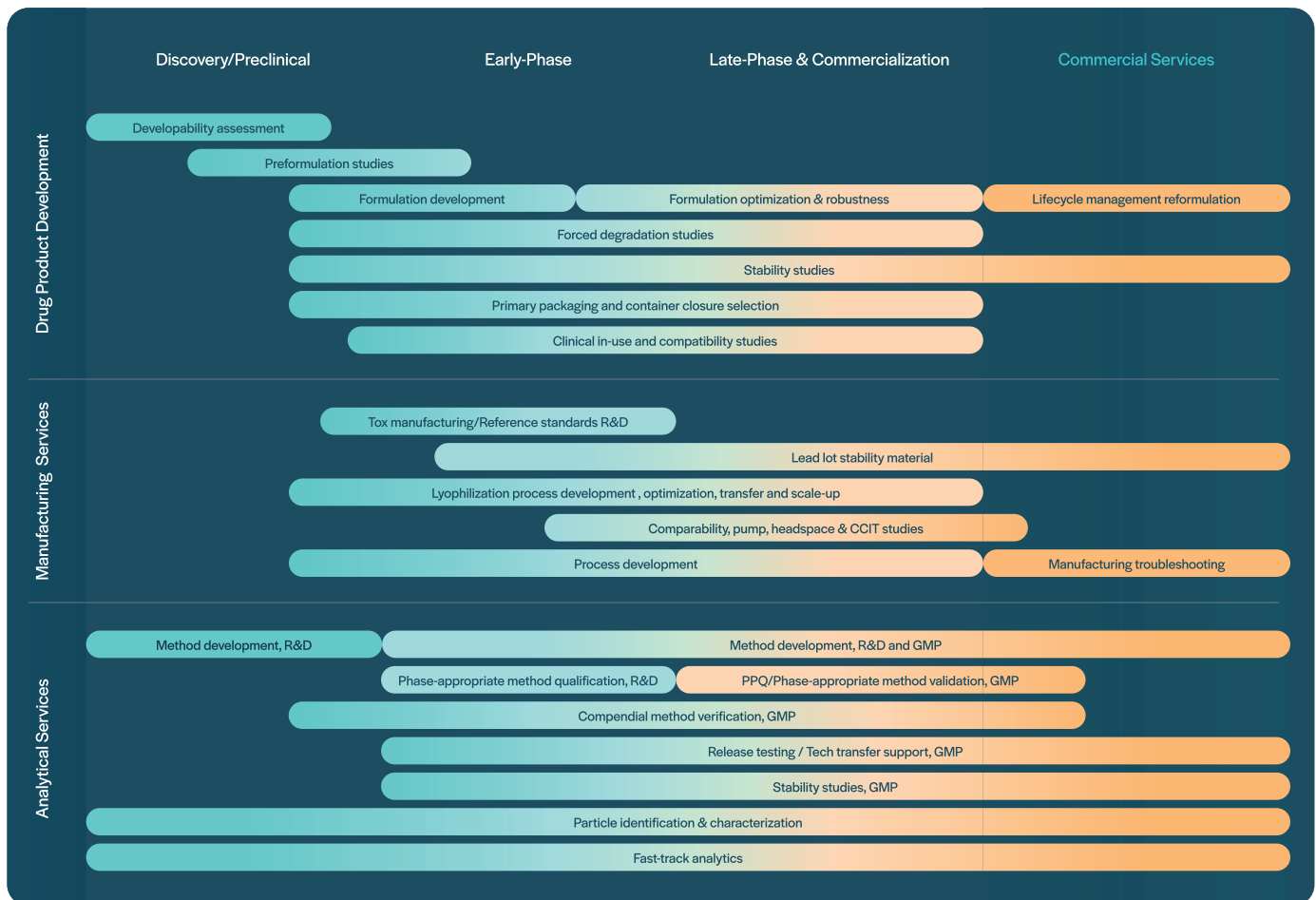


OUR SUPPORT FOR YOU
**ACCELERATING AND DERISKING
 YOUR PATH TO MARKET**

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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