

KYMOS
GROUP

Company Presentation

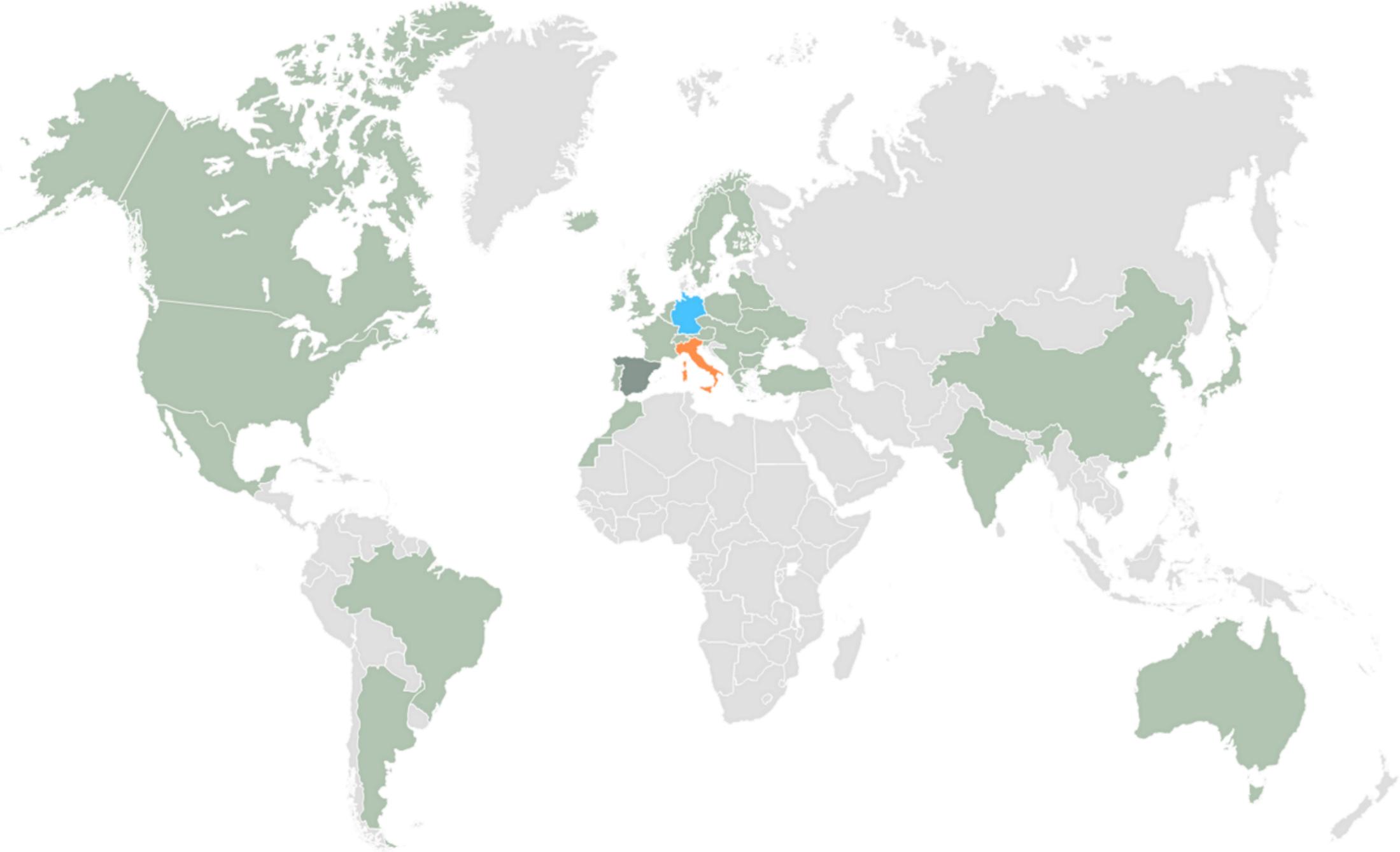
2026

European Laboratories

KYMOS GROUP is a European CRO/CLO Group with 3 sites in Barcelona (ES), Ancona (IT), and Frankfurt (DE) mainly devoted to biopharmaceutical industries. More than 5.700 sqm of laboratories with the latest technology.

GLP certificate, **GCP** compliance, and **GMP** certificate for importation, release and quality control of small molecules and biologics and advanced therapies. **FDA** and **EMA** inspected.

KYMOS GROUP has global presence, having clients from more than 60 countries, mainly in Europe and Asia, but many others in the USA, Australia, or South America.



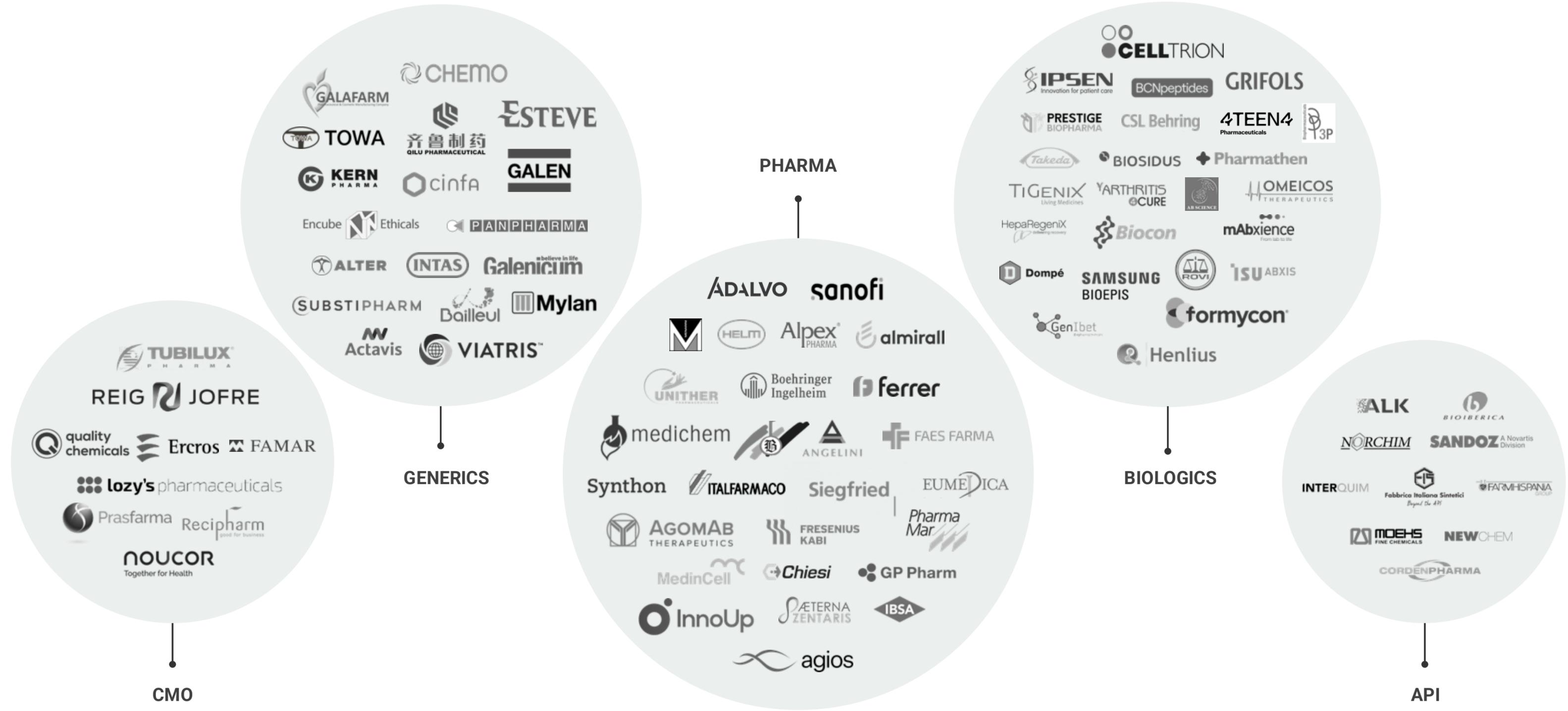


KYMOS SPAIN (Barcelona)
Kymos S.L.
Headquarters and Laboratories
Staff of 200 people

KYMOS HQ (Barcelona)
Kymos Group SL
Headquarters
Staff of 50 people

KYMOS GERMANY (Frankfurt)
Prolytic GmbH
Laboratories
Staff of 25 people

KYMOS ITALY (Ancona)
Pharmaprogress Srl
Laboratories NEW
Staff of 45 people



One-stop-shop

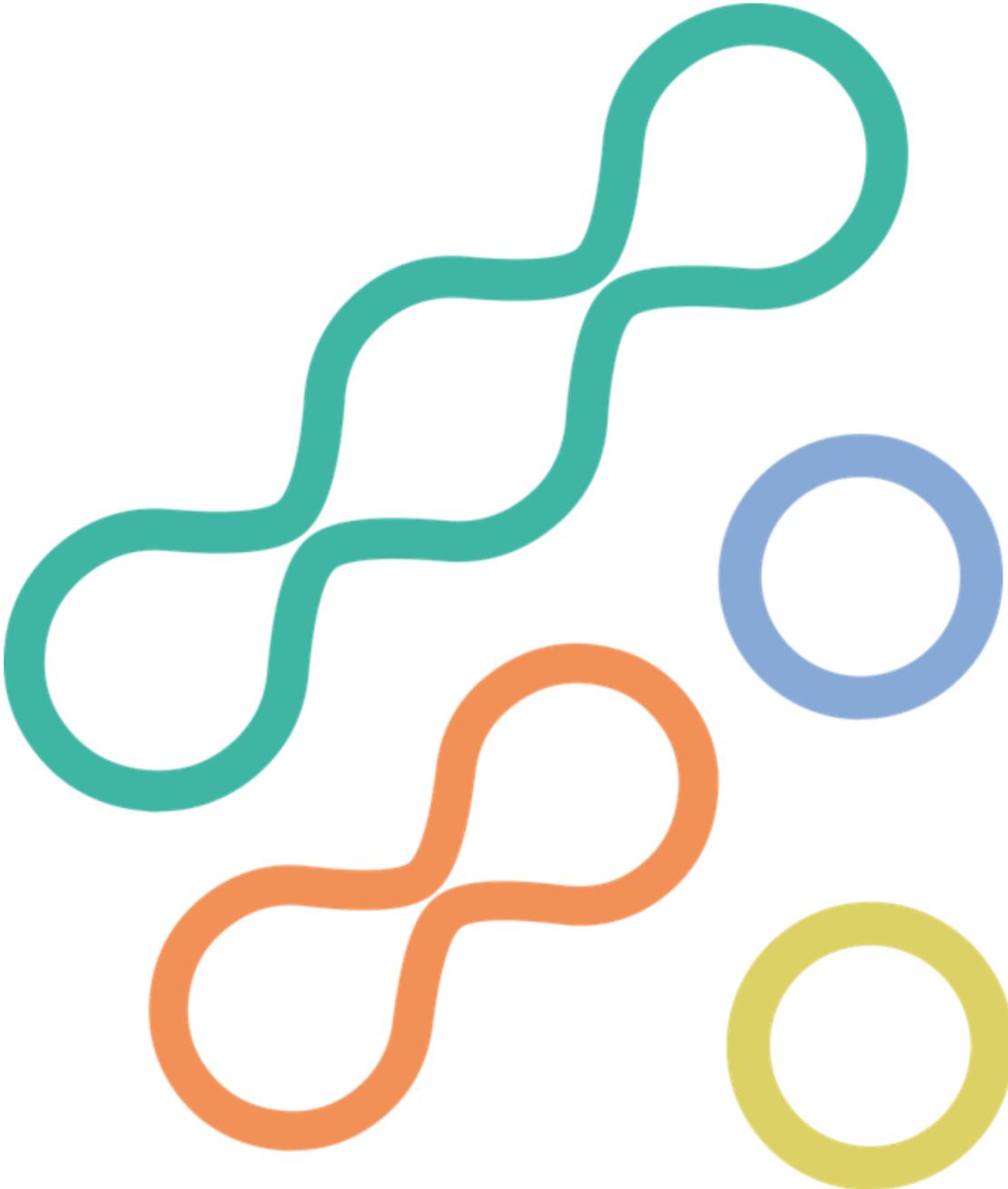
Provider with reliable and certified partners.



Our Value Proposition

KYMOS GROUP is a company devoted to providing analytical expertise and testing capabilities to third parties for research, development, and quality control of biopharmaceuticals.

Bioanalysis
(Preclinical & clinical studies)



CMC Analysis
(Chemistry, manufacturing & control)



—
**CRO Services based on
molecule specifications**

We are experts in the analysis of
small molecules, biologics,
TIDES and advanced therapies.



Small Molecules

GLP-certified bioanalysis
and GMP-certified quality
control of small molecules.



Biologics and Biosimilars

Bioanalytical and quality control services
for proteins, hormones, enzymes, vaccines,
biosimilars and ADCs



TIDES (Oligo & Peptides)

GMP and GLP-certified
bioanalysis and quality control
services for TIDES
(oligonucleotides and peptides)



Advanced Therapies

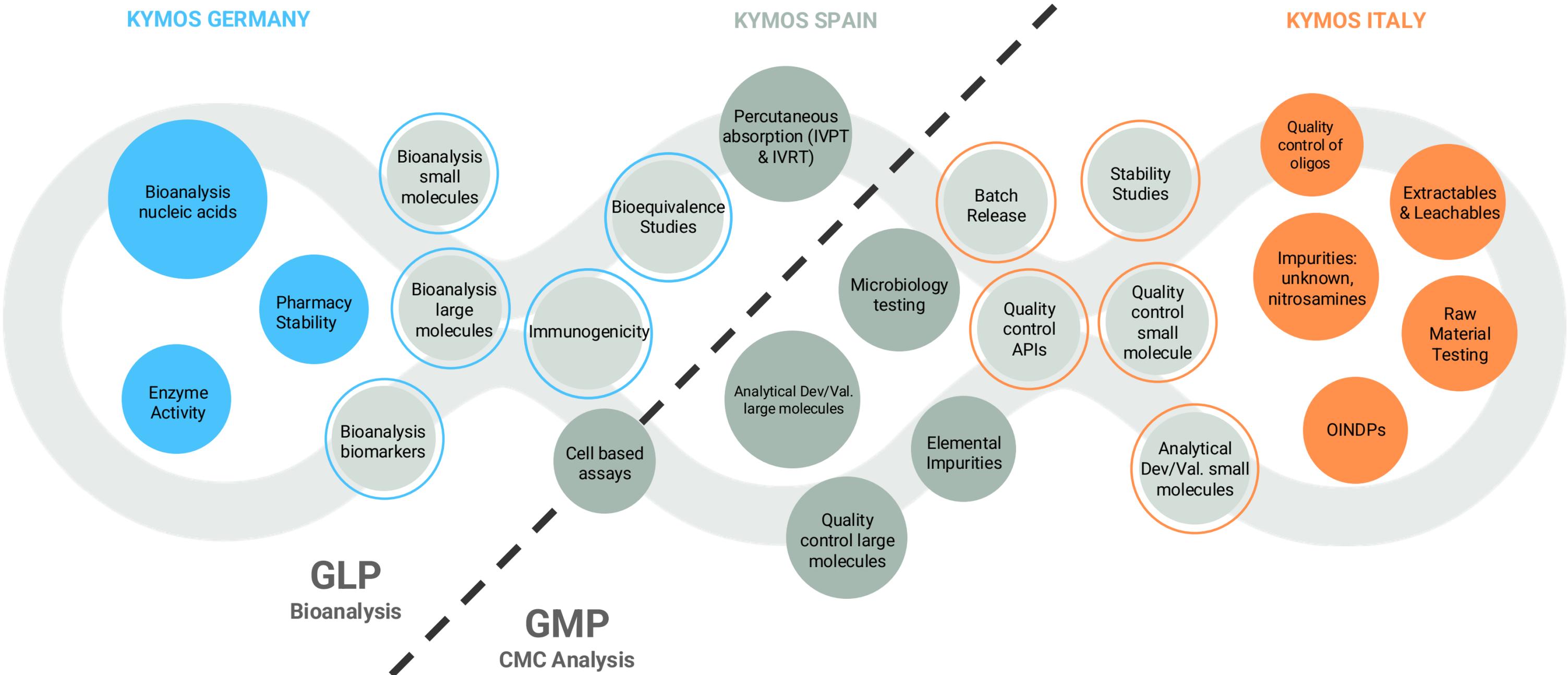
GLP and GMP-certified quality
control and bioanalysis of gene
therapy products.



KYMOS GERMANY

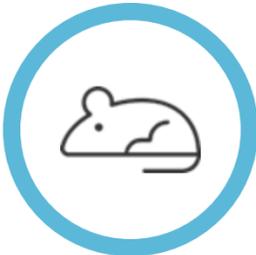
KYMOS SPAIN

KYMOS ITALY

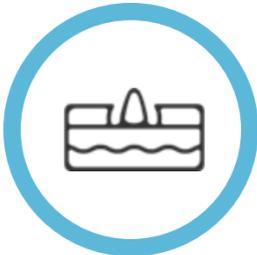


Bioanalysis

(Preclinical & Clinical Studies)



- Full preclinical study management



- In vitro release test (IVRT)
- In vitro permeation test (IVPT)



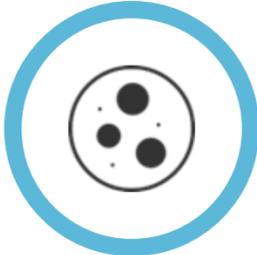
- Nucleic acids analysis by qPCR
- Nucleic acids bioanalysis by Hybridization ELISA & MS Spectrometry
- ASO bioanalysis by LC-MS



- Full bioequivalence studies



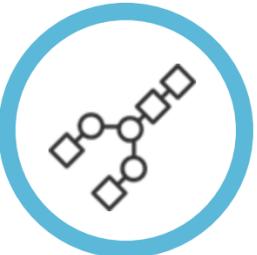
- PK studies of biologics by ELISA, ECLA, RIA
- ADA testing in immunogenicity studies
- Biomarkers determination by ECLA multiplex and LUMINEX



- Cell-based assays for neutralizing antibodies



- PK of small molecules by LC-MS/MS, HRMS, and ICP-MS
- Residue depletion studies (animal health)



- Binding assays by SPR
- Glycans by capillary electrophoresis



- Enzyme activity

CMC Analysis

(Chemistry,
manufacturing &
control)



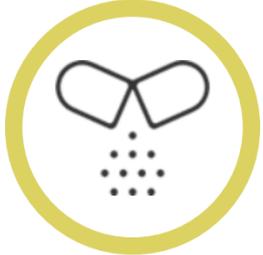
- Raw material testing (compendial and non-compendial methods)



- Stability studies and sample storage (ICH, on-going, photostability, in-use, holding time)



- Cell based potency assays for QC



- Method validation for assay, related substances, residual solvents by LC & GC
- Physical determinations (pH, melting point, density, osmolarity, particle, dissolution, etc.)



- Microbiology for sterile and non-sterile products and endotoxins



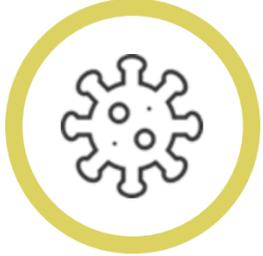
- Quality control of biologics and biosimilars
- Characterization and comparability of biologics and biosimilars



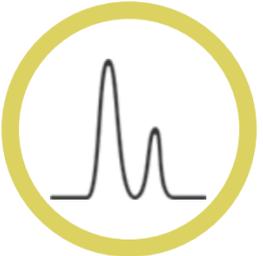
- Orally Inhaled & Nasal Drug Product testing



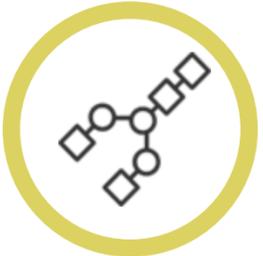
- TIDES (oligos & peptides) characterization and quality control



- Viral vector characterization and quality control



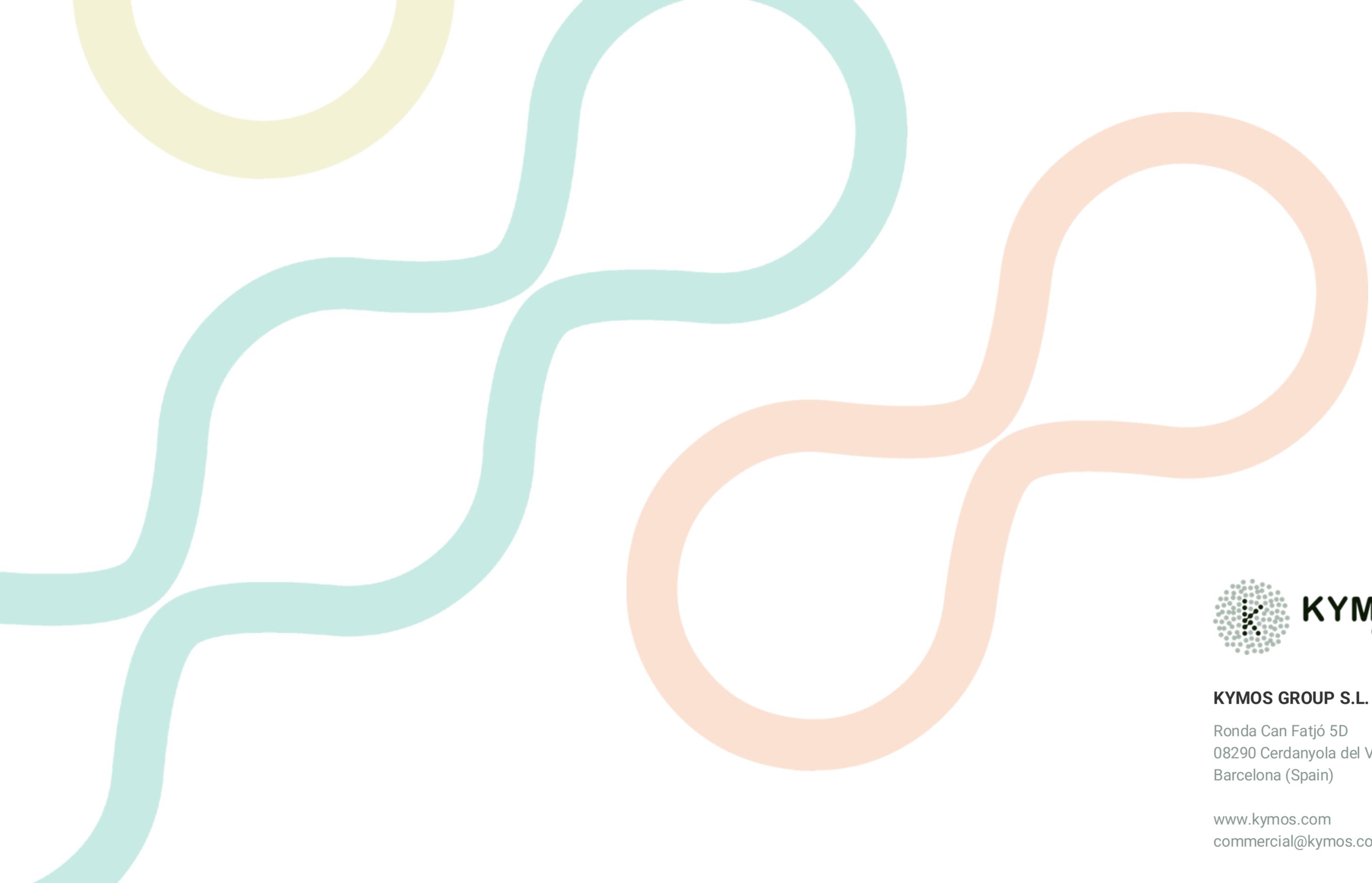
- Elemental impurities, Unknown Impurities, Nitrosamines and Extractables & Leachables



- Building blocks of glycans and antifactor testing in heparines



- Batch release into the European Union



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