



Drug development accelerated with compelling science

Open dialogue. Successful delivery.



Welcome to Pfizer CentreOne. We're the global CDMO backed by the scientific power of Pfizer. Working closely with you we combine our commercial drug development expertise with open dialogue, regulatory experience and global reach to help assure project success and your next patient breakthrough.



Take your project farther, faster with our scale up and technology transfer expertise

Our global network of sites have been developing and manufacturing Pfizer's broad array of innovative molecules for decades, and this legacy helps us in our mission to deliver on what we promise.

We offer an expert suite of development and optimization services

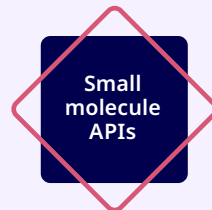


Our offerings include:

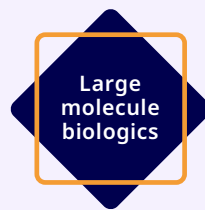
- Analytical methods development
- Pre and post-launch regulatory support
- Scale-up from lab to pilot to commercial
- Specification development
- Controlled substances (II-IV)
- Clinical manufacturing Ph I - III
- Full range of OEB 1-5

An altogether different approach to advanced therapeutics

Get promising candidates to commercial phases faster with an altogether different CDMO. Dedicated to your intellectual property's efficient advancement, we help you optimize your molecule's development and help it pass every milestone.



- API Characterization
- Synthetic route development
- Process safety & optimization
- Impurity assessment



- Vaccines & mAbs
- Cell culture medium optimization
- Upstream & downstream process optimization
- Biosafety levels 1-2



- Vials, Prefilled Syringes, Cartridges
- Formulation development
- Lyophilization optimization
- Aseptic process optimization



- Tablets, capsules, granules
- Early stage formulation development
- Process validation & optimization

Ready to shape your development masterpiece?

There is no limit to what we can achieve together. Get in touch with our experts to start your development journey today.

