

We navigate you through drug development

We as Ardena team-up with our global customer base to provide small molecule drug development, manufacturing, clinical logistics and (bio)analytical services. To streamline your drug's progress to the clinics and beyond, we compile your regulatory dossier in parallel. With our multidisciplinary approach we accelerate the drug development and approval process. Our services are seamlessly integrated to better mitigate development risks and ultimately reduce time-to-clinic.

Bringing your molecule into clinic, on time and within budget.

Ardena is your one source contractor for the chemical and pharmaceutical development of your valued compounds.

We provide full product analytical and bioanalytical support during development. We support early and late stage clinical development projects, all in one seamless offering.

Ardena provides full dossier development support. Our dossier-centric approach makes the difference to stay ahead of the game. We work with you to compile your regulatory dossier in parallel with drug product development, saving you time, identifying hurdles and fast tracking your product towards compliance. We offer expertise in CMC and regulatory support for both small and large molecules.







