PATHFINDER™

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Navigate Your Way to Clinical Success

Getting new molecules to the clinic is difficult and complex, with a host of technical hurdles to overcome.

Pathfinder™ is a flexible, end-to-end service that can be adapted to your specific requirements to ensure a clear path to success in first in human clinical trials. As a single provider we have the depth and breadth of scientific expertise - from discovery to clinical supply – to maximize scientific knowledge retention while significantly reducing time and cost for your program. Pathfinder will deliver your molecule to clinic faster, allowing you to obtain clinical data sooner.

Global, Integrated End-to-End CMC Services to Accelerate Time to Market

Cost-efficiency

Working with a single provider reduces the management and co-ordination of multiple partners, simplifying communication, while minimizing shipping requirements and logistics costs.

Time Savings

Streamline and expedite project timelines with seamless technology transfer to ensure consistency during scale up.

Scientific Continuity

Benefit from seamless knowledge transfer and retention, enabling scientific continuity with support from cross-functional expert teams.

Flexibility

Flexible solutions offer tailored approaches to meet the most complex requirements.

PATHFINDER™

Propel your candidate forward with PATHFINDER's streamlined API and drug product services:

API Process R&D, Analytical Development & Manufacturing

Route design, process design, analytical methods verification/validation, GMP and non-GMP manufacturing.

Materials Science & Preformulation Capabilities

Maximize chances of success with API characterization, pre-formulation, salt and polymorph screening, developability assessment, particle size assessment, and DMPK and bioanalysis studies.

Product Formulation, Analytical Development & Manufacturing

Ensure the best chance of clinical success by leveraging our formulation optimization technologies for oral solid dose development. Obtain the highest quality GMP-manufactured drug products for clinical trials, including packaging, labelling and QP services.

Regulatory Support & Consultation

Meet regulatory requirements with IND dossier preparation and submission, R&D consultation, and dedicated project management.

About Pharmaron

Pharmaron is a global R&D and manufacturing service provider supporting small molecule, biologics, and CGT product development — from drug discovery through clinical development and ultimately commercialization. Our globally integrated CMC platform — stretching across the UK, US, and China — offers flexible and bespoke end-to-end services. Our capabilities will streamline your route to first-in-human clinical trials and beyond.

Global CDMO Services

Contact Us

Ready to accelerate your drug product's path to the clinic? **Contact us at bd@pharmaron.com**