

## Helping you bring inhaled medicines to market



FORMULATION SCIENCE

DEVICE TECHNOLOGY DEVELOPMENT EXPERTISE

PRE-CLINICAL DEVELOPMENT

PHASE 1

PHASE 2

PHASE 3

SUBMISSION & APPROVAL COMMERCIALISATION





## Choose a partner to help you succeed

20 years' experience in inhaled product development

11 inhaled on-market DPI, pMDI and nebuliser medicines, launched by our partners and licensees

>\$10bn sales of products using our formulation and device technology since launch\*



## Fully-integrated services and expertise to progress your programme

- Formulation Development & Optimisation
- Small molecules and biologics
- Complex combinations
- Advanced powder blends
- Spray-dried, particle engineered powder formulations
- Aqueous & non-aqueous formulations

- Commercially-Validated Device platforms
- Device flexibility based on the needs of your programme
- Dry powder inhalers (unit-dose and multi-unit dose)
- Pressurised metered dose inhalers
- Smart vibrating mesh & jet nebulisers

- Product Manufacturing
- GMP drug product manufacturing (including biologics)
- GMP device manufacturing for clinical trials
- Equipment & facilities at relevant-to-commercial scale

- Comprehensive Analytical Services
- Physico-chemical (Q3 type) capability and "more realistic" test methods
- Dedicated physical properties capability
- Biologics evaluation
- Stability storage from small bespoke studies to pivotal prorammes

- Process Development& Scale-up
- Develop robust & reproducible manfacturing processes
- Seamless scale-up from laboratory to pilot to commercially-representative scale
- Installation engineering capability and technical support

- Medical & Regulatory Support
- Navigate the complex regulatory requirements for inhaled NCE & generic products
- All services are supported by our medical, regulatory, device vigilance & pharmacovigilance teams

See how our inhalation services can advance your programme through all stages of development