



Exova in Pharmaceuticals

Exova is one of the world's leading drug development, compliance and assurance service companies. We have been at the forefront of the Global pharmaceutical market for the last 25 years and have consistently helped leading pharmaceutical companies to safely bring their products to market faster and more cost effectively.

Regulatory Services at Exova

Please contact us for further information to help your needs in the following areas:

- Chemistry Manufacturing and Control (CMC) and Module 3 Support Services
- Common Technical Document (CTD) Formatting and Part II to CTD Conversion
- National Marketing Authorisations (Abridged Applications)
- License maintenance including National and European Variations
- Change of Ownership Submissions
- Certificate of Suitability and EDMF preparation and submission
- Response to Agency Questions
- GAP analyses and manufacturing compliance
- Quality Overall Summary (Expert Reports)
- Scientific Report Writing for dossier submission
- Veterinary applications

Development & Analytical Services at Exova

We also have a network internationally located laboratories providing our clients with easy access to specialised capabilities and services. These include:

- analytical testing
- method development and validation
- preformulation and formulation development for NCE's and existing molecules
- small scale GMP manufacturing of clinical trial materials
- stability storage studies (including all ICH conditions)
- bioanalytical and microbiological analyses
- QC testing for batch release

We possess a wide range of analytical and testing capabilities; are GMP MHRA approved; operate to stringent cGMP guidelines and have a UK Home Office licence for handling controlled drugs.

Please contact us on [+44 \(0\) 161 787 3291](tel:+44201617873291) or e-mail info@exova.com today.