

Regulatory consultancy available to cover the wide variety of areas encountered during the lifecycle of a medicinal product.

Particularly to include:

Electronic Common Technical Document (eCTD)
Traditional Herbal Medicines
Legal Category Switches
Regulatory Advice and expertise on Mergers and Acquisitions
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EU Regulation on Paediatric Medicines
Patient Information Leaflets for User Testing
Experience in Centralised and Mutual Recognition Filing
Clinical Trial Applications and Maintenance
Writing of Quality Overall Summary and Clinical Overviews
Food Supplements
Nutrition and Health Claims Legislation
Contract Regulatory Placements In-house
Provision of Regulatory Training

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