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Chris Holloway is Group Director of Regulatory Affairs at ERA Consulting, with overall responsibility for the company's operations in Germany, London, Washington, DC, and Australia.

He has worked in regulatory affairs, specializing in biotech products and other biologics, for more than



20 years, since leaving academia where he was Professor of Clinical Biochemistry at Hannover Medical School in Germany.

Chris graduated in 1971 and obtained his Ph.D. in 1975, after which he held a Royal Society European Research Fellowship at the Max-Planck-Institute in Germany. He then moved into university,

and was awarded the *venia legendi* in 1982.

Chris is particularly interested in regulatory strategies for novel therapeutics/platforms, interacting wherever possible with regulators to define expedient routes towards marketing authorizations.

He has worked on more than 350 different products, contributing particularly towards approaches to manufacturing and control as well as early non-clinical and clinical development. His team has compiled dossiers, and Chris authors expert reports for European submissions.