

Subin is a quality assessor specialising in the CMC assessment of small molecules as well as biological medicinal products, including monoclonal antibodies and vaccines. Before joining the authority in 2006, he worked in the generics industry for more than 8 years, gaining broad experience in regulatory affairs, quality assurance, product development and technology transfer.

He is a member of ICH Q12 IWG, Access Consortium- Generic Medicines Working Group and IPRP- Quality Working Group. Subin is a Pharmacist by training with a PhD in Pharmacy, a Masters in Public Health and certified in Regulatory Affairs.