Strenghts

Experience in the field of oncology
Professional knowledge in Regulatory Affairs
Used to work unsupervised

Professional Summary

Our consultant is a Regulatory Affairs consultant with more 5 year's regulatory affairs experience within a busy consultancy. Her main focus is in the areas of post-marketing pharmaceutical and drug regulatory affairs. She is specialist for redacting of spc, pal and labeling texts in connection with CP/DCP proceedings and adapting texts to QRD templates.

Prior to that she was more than 6 years an assistant of the medical scientific department of a CRO, which was specialized in oncology. There she was involved into various projects concerning pharmacovigilance, reviewing and adapting SOPs and controlling of informative texts regarding to content and form. She was monitoring e.g. approvals and variations. Additionally she gained full sales experience in being a Sales Representative and Area Sales Manager for original and generic products in different branches of medical science.

She graduated as state certified Biological Technical Assistant and IHK certified Area Sales Manager. 2013 she graduated from a German distance school with the course Psychotherapy HP with a testimony. 2014 -2015 she attended various courses of MEGRA, which have a main aspect of regulatory affairs.

Languages

German	mother tongue
English	proficient