Mrs. Hannie de Munnik

With over 20 years of experience in the pharmaceutical industry, Hannie de Munnik has made significant contributions in clinical research, medical affairs, regulatory affairs, and real-world evidence research. As the Executive Director of Regulatory Affairs for the EUCAN region at AstraZeneca, she plays a pivotal role in the implementation of electronic product information (ePI) initiatives across Europe and Canada. Her work involves close collaboration with key stakeholders to seamlessly integrate ePI into regulatory and medical frameworks, ensuring that product information is more accessible, up-to-date and compliant.