Karin Berndt
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Karin Berndt has been working with Medical Devices for 28 years.
She has worked in the industry in quality assurance, development and regulatory affairs departments. She also has experience with medical writing, having worked in a clinical research organization.
Currently she is team leader regulatory affairs at seleon GmbH, where she joined in 2019.
She has experience with national and international submissions, clinical evaluations and implementing quality management systems.