



Partner Profile – Generis and DocShifter

“With the integration of DocShifter as the default document conversion solution to the CARA foundational platform, we are able deliver our promise even better: drastically speeding up and simplifying the complex process of content and document conversion. DocShifter’s document conversion capabilities, combined with new CARA, are able to answer the complex requirements of highly regulated industries.”

Geert van Peteghem, CEO of DocShifter

DocShifter is an automated content and document conversion software. It is designed for high volume document conversion and enrichment, and allows maximum scalability and flexibility. Through automations and centralising processes companies can speed up their work, eliminate manual intervention and reduce risk.

Why DocShifter?

- Generate compliant PDFs for multiple Health Authorities automatically
- Reduce the dependency on manual PDF editing tools to create compliant PDFs
- Speed up time to market by moving technical content compliance earlier in your regulatory content creation process
- Convert +10.000 documents per hour with a 0.01% failure rate
- Proven technology since 2006.

Why Generis?

- Over 20 years' experience in Life sciences
- Industry leading foundational platform with a common UI and powerful process automation
- Out of the box solutions for modules including Quality, RIM, eTMF and Pharmacovigilance
- Flexibility to suit the clients' requirements
- Integrations with a wide range of systems

Previous Experience

Generis and DocShifter currently have multiple joint customers in the pharmaceutical industry with a large number of users. For several different use cases, CARA is the selected document management platform while DocShifter is used to allow large scale document conversion. The customers have highlighted the ease of use and the seamless integration of the two solutions which helps streamline their day-to-day work.

Future Plans

There is a growing need for a more interconnected approach to data and documents among clients within Life Sciences. Generis and DocShifter are making this transition easier through an integrated approach. Another key theme is a shorter development cycle for medicines, again this process is supported through automation, such as those found on the CARA Life Science Platform and in the DocShifter system. They allow users to reduce their manual work and scope for errors, making time to market ever faster. DocShifter and Generis will continue to work together to overcome these challenges for our clients.

About



DocShifter was created in 2007 to respond to the need for easy and fast content conversion in highly regulated environments. Customers can be found worldwide in Life Sciences, Public Sector, Engineering, Banking & Insurance. The company is headquartered in Ghent, Belgium.

At DocShifter, we believe that relevant digital content should never become obsolete. Unstructured information needs to be kept in a readable and searchable format, for generations to come. In our fast-paced digital world, standards appear and disappear. Transforming all this digital content, in whatever shape or form is our goal. To ensure your information continues to provide value.

To find out more visit: www.docshifter.com



Generis is a developer of world-class Content and Regulatory Information Management technology for regulated industries globally.

Eight of the top 10 global life sciences companies rely on Generis' flagship Intelligent Content Services platform, CARA™ for critical document and information management, including AbbVie, AstraZeneca, Biogen, Gilead, Bayer, Pfizer, and Merck KGaA.

Today Generis serves more than half a million users worldwide, across use cases ranging from RIM, Regulatory / R&D and Safety use cases to Clinical, Non-clinical, Quality GxP, CMC, Pharmacovigilance, Medical Information / Medical Affairs applications and more.

To find out more visit: www.generiscorp.com

