Automated Document Generation for Common Technical Documents and respective Modules and Chapters

LEXORO
The pulse of A.I.

lexoro is an expert for Intelligent Automation using cutting-edge Artificial Intelligence and Software Engineering technologies. With our technology we create and develop solutions for various well-known companies within the Pharmaceutical and Medical Device industries for already over 20 years.

With our lexoro Document Generation Suite we automate workflows for the creation of documents & tables including copying & populating of text & data. This includes the formatting and filling of regulatory documents such as Common Technical Documents with its Modules, or other dossiers for Registration. That way we can realize an automation degree of around 70%.

Copying, populating & formatting of data, text & information into the associated modules and chapters of the Common Technical Document:

lexoro Automation

| 1. Choose template or chapter | 2. Creation of data basis (upload or collect) | 3. Formatting and Aggregation | 4. Transfer, Filling and Population | 5. Preview and Export |
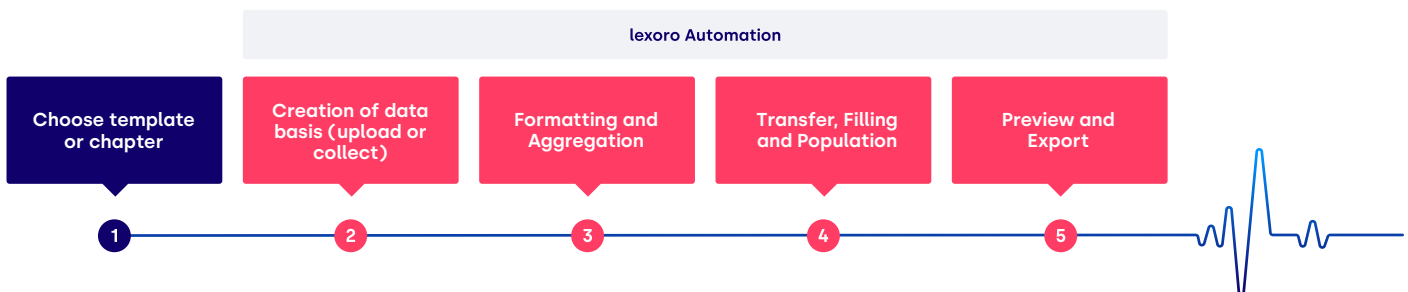
- Manual selection of the required template or chapter to be processed or generated.
- Automated collecting and copying of relevant raw data and text from in-house systems (e.g., LIMS, SAP, statistics systems, etc) to create a data basis.
- Extraction, preparation, editing, and formatting of data, text, and tables for the CTD modules - this includes adjusting tenses (present/future tense in past tense) or text generation for table descriptions.
- Fully automatic populating of dedicated chapters such as Quality & Clinical Summary, Non-clinical & Clinical Overview, etc.
- Preview of the document in a system with subsequent generation & export of the module or document for further manual processing.

Our offer & your benefits:

✓ Reduction of processing time: only approx. 25-30% have to be created and processed manually

✓ Individually tailored to your needs – Problem-free integration into your systems & workflows

✓ Standardization of workflow - EMA- and FDA-compliant creation of modules and chapters

✓ 20 years of experience in regulated industries with ISO certification and agile development

✓ Web-based Module & Document Generation. Secure data processing. Ready to use in 4-6 weeks.

Contact us now! We help you identify suitable routine tasks or standard processes quickly to be automated. info@lexoro.ai

Your first 4 steps for optimal success with Intelligent Automation:

| SCOPE First Digital Consultation | DISCOVER Definition Workflow | DEVELOP Development Pilot | IMPLEMENT Solution Introduction | www.lexoro.ai |