

PV SOFTWARE

Technology advancement and introduction of AI has the ability to improve PV application further. The application should be fully validated to ensure flawless execution, scalable to enhance functionalities as per future needs and cost effective for better adaptability. The software application must have ability to record, process, and store large volume of data in a way that allows its accurate assessment, interpretation, and verification with highest quality standard. Robust architecture is essential to automate and streamline the workflow. Selection of technology also important for better compatibility for database integration.

OUR PRODUCT

PVWRITER

PVWriter is a software application streamlining the entire process of aggregate report preparation, review and finalisation. The application is developed by technical experts, under supervision of domain experts with decades of experience. A fully validated application with inbuilt functionality to support data extraction, data analysis, natural language processing (NLP), and natural language generation (NLG), to produce aggregate report like PADER, PSUR, PBRER, DSUR etc. quickly as per regulatory requirements with minimal human intervention. The application has the ability to customise further to meet Client's requirements. PVWriter helps Medical Writer to improve performance, by assisting their day-to-day works.



OUR SPECIALISATION

Aggregate Report Writing
Periodic Adverse Drug Experience Report (PADER)
Periodic Safety Update Report (PSUR)
Periodic Benefit Risk Evaluation Report (PBRER)
Development Safety Update Report (DSUR)

GET IN TOUCH

Contact us to explore our products and solutions.



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www.pvwriter.com



ABOUT US

GVP Technology, a pioneering technology organisation dedicated to revolutionising Pharmacovigilance and contributing to the welfare of global healthcare.

Our technology-driven products and solutions are aligned with Clients' expectations and regulatory requirements through a blended approach for enrichment of analytics of safety information adopting innovative technologies.

GVP Technology, with extensive domain and technical expertise, can transform your Pharmacovigilance system to technology enabled futuristic process.



MANUAL EFFORT REDUCED
90%
TIME SAVING
90%

PVWRITER FEATURES

STREAMLINED PROCESS

Streamlined entire medical writing process, starting from data collection, report preparation, peer review, medical review, approval, and finalisation.

AUTOMATIC DATA ANALYSIS

Inbuilt data analysis and validation functionality to extract relevant information in structured format for report preparation.

DATABASE COMPATIBILITY

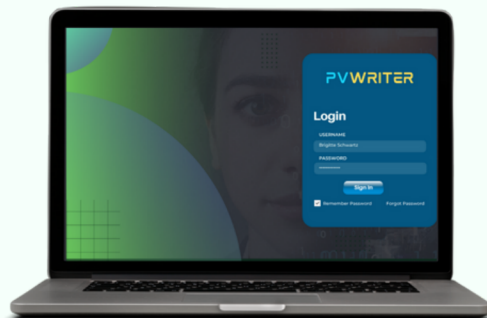
PVWriter is compatible with global Safety databases to sync or upload safety data.

AUTOMATIC REPORT PREPARATION

Report generation in Client's template with minimal human intervention.

ENHANCED COMPLIANCE

Fully validated, access control, with inbuilt audit trail functionality.



www.pvwriter.com



PVWRITER ADVANTAGE

TIME SAVING

Reducing manual effort through automation.

IMPROVED EFFICIENCY

Streamline and accelerate the medical writing process, leading to improved efficiency.

IMPROVE ACCURACY AND COMPLIANCE

Reducing the risk of human error thereby improving compliance.

IMPROVED DECISION MAKING

Enabling intelligent decision making to enhance benefit-risk assessment.

CONTACT US

Contact us to learn more about PVWriter and how PVWriter can support and simplify medical writing process.

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