Reduce Safety Case Processing Time by 50%

Case Study

Client Overview

The sponsor is a biopharmaceutical company, focused on developing therapies for high unmet medical needs. To enhance their pharmacovigilance (PV) operations, they sought a solution that would streamline case processing, improve data accuracy, and reduce manual data entry.

The Challenge

Sponsor faced several challenges in their PV processes:

- **Inefficient Data Entry:** Manual entry of safety data into multiple systems led to increased processing times and potential for errors.
- **Data Inconsistencies:** Disparate systems resulted in data discrepancies, affecting the quality and reliability of safety information.
- **Regulatory Compliance:** Ensuring adherence to global safety reporting requirements was resource-intensive and complex.

The Solution

Sponsor implemented Clinevo Safety, a comprehensive pharmacovigilance solution offering:

- **Integrated Data Management:** Seamless integration with existing Electronic Data Capture (EDC) systems to eliminate redundant data entry.
- **Automated Workflows:** Streamlined case processing with automated data transfer and validation checks.
- **Regulatory Compliance:** Built-in compliance with global safety reporting standards, ensuring timely and accurate submissions.

The Results

After adopting Clinevo Safety, sponsor achieved:

- **50% Reduction in Case Processing Time:** Automation and integration reduced manual efforts, accelerating case handling.
- **Enhanced Data Accuracy:** Elimination of duplicate data entry minimized errors, improving the quality of safety data.
- **Improved Compliance:** Streamlined processes ensured timely reporting, meeting global regulatory requirements.

Client Testimonial

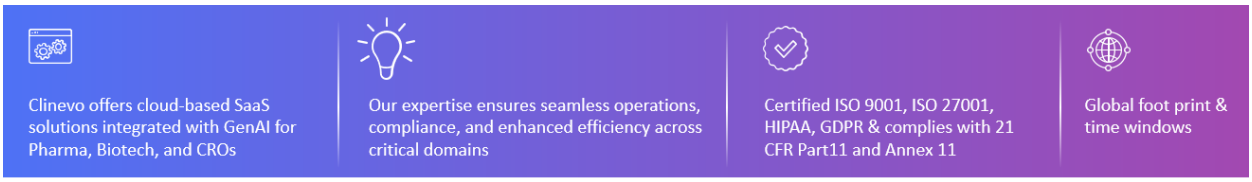
"Case processing time went down by 50% and data accuracy and quality went up because you're not doing double data entry from EDC to Safety."

Conclusion

Clinevo Safety provided sponsor with an integrated and automated pharmacovigilance solution, significantly enhancing efficiency, data accuracy, and regulatory compliance in their safety operations.

For further information contact us at info@clinevotech.com

Why Clinevo?



200+	Global clients support in the past 5 years	**50+**	Regulatory audits supported
100%	Customer retention rate YoY	**300+**	Gateway connections with authorities
99%	Customer satisfaction rate	**100+**	Data migrations from various systems
10Mn+	Cases processed till date	**100%**	Validated systems
1000+	Clinical trials supported till date	**24/7**	Unlimited global support