Data Migration from ArisG Using XML & Non-E2B Method

Case Study

Customer Overview

A US-based, fast-growing, research-oriented multinational pharmaceutical company with a portfolio of over 100 products distributed globally. The company regularly submits safety cases to global pharmacovigilance (PV) authorities, including the **EMA**, **MHRA**, **USFDA**, and **Health Canada**.

Business Challenge

The client was using ArisG for pharmacovigilance operations but faced challenges with scalability and data management due to their rapid growth and increasing case volumes. They needed a modern, cloud-based safety system capable of handling end-to-end case processing and regulatory submissions efficiently across spontaneous, clinical trial, and medical device reports.

Clinevo Solution

The client chose **Clinevo Safety** for its comprehensive capabilities in case processing, management, and regulatory submissions. Clinevo implemented a **non-E2B, XML-based data migration** approach to seamlessly migrate historical safety data from ArisG into the Clinevo Safety cloud platform.

Key highlights of the migration process:

- **XML-based methodology** ensured structured, accurate data transfer.
- Smooth migration of all legacy data without data loss or downtime.
- Clinevo handled migration, validation, and testing end-to-end.

Results & Impact

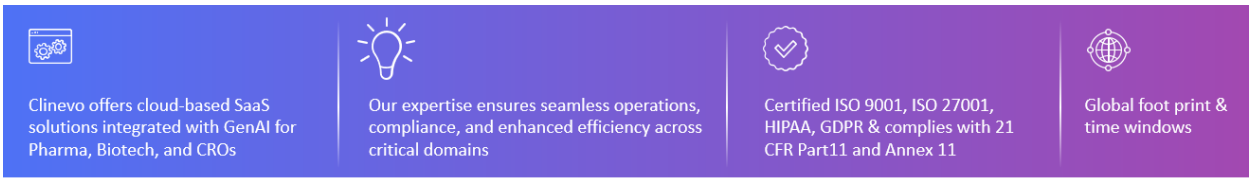
- **Seamless migration** of historical data from ArisG to Clinevo Safety.
- Enabled the client to perform **end-to-end safety case management and submissions** to global authorities (USFDA, EMA, MHRA, HC, RoW).
- **More than 3,000 cases processed annually** using the Clinevo Safety platform.
- Improved efficiency in managing **Spontaneous**, **Clinical Trial**, and **Medical Device** safety cases.

Conclusion

Clinevo's expertise in safety data migration and its robust Safety platform enabled the client to enhance pharmacovigilance operations, improve compliance, and scale their global submissions effectively.

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