TOOLS FOR ADVERSE REACTION ASSESMENT

Introducing our comprehensive, web-based pharmacovigilance safety database

TARA PV was designed by a team of pharmacovigilance professionals who saw the benefits in a user-driven approach to processing and storing drug, device and vaccine adverse events in a secure safety database.

WHO IS IT FOR?

– Pharmaceutical Companies
– CRO's
– Academic and Research Institutions
– Charitable Organisations

ISO 9001
Certificate No. GB2005816

ISO / IEC 27001
Certificate No. ISM7799365

CONNECT WITH US

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MedGenesis Ltd part of idash Group

With flexibility at its core, TARA PV is suitable for all pharmacovigilance requirements – with multiple product packages to match your needs and budget.

TARA PV

TOOLS FOR ADVERSE REACTION ASSESMENT

WHY TARA PV?

OUR PRICING
- Cost effective
- Flexible
- Competitive

OUR SUPPORT
- Full user training
- Responsive help
- Software upgrades
- Support portal access
- Project management

✔ QUALITY ASSURED
TARA PV is a fully tested and audited pre-validated software system from an ISO 9001 and ISO 27001 accredited vendor.

✔ FULLY COMPLIANT
TARA PV is 21 CFR Part 11 compliant, adheres to GxP and ICH standards and allows compliance with all European & Worldwide regulations.

✔ FAST AND INTEGRATED
As a hosted platform, TARA PV not only allows rapid implementation but also data migrations from other PV databases and third party integration (eg Medidata RAVE).

✔ LICENCING FLEXIBILITY
TARA PV offers a range of pricing models, dependent on your requirements.

✔ DATA ANALYSIS AND SIGNAL DETECTION
Microsoft Power BI is embedded in the TARA PV application providing a powerful data visualisation and interactive reporting tool.

✔ COMPLETE CONFIGURATION
With an independent Administration Module and test database, TARA PV offers an integrated workflow with individual case assignment, company product dictionary upload and flexible configuration options. Also included as standard are the ISO E2B vocabulary lists.

✔ CORE PV FUNCTIONALITY
Regulatory reporting (inc. E2B(R3), CIOMS, MedWatch, PSUR and DSUR) and full MedDRA and WHO Drug Dictionary integration. Adverse events from clinical trials and marketed products can be processed in the same database, multitenancy capability to separate data by study or sponsor, extensive case searching, ability to store external documents in individual case records, pre-submission case validation and action management.

✔ HOSTING PEACE OF MIND
Tier 4 ISO 27001 accredited Data Centres mean routine server hosting maintenance activities such as operating system critical security updates, triple layer backup, disaster recovery and GDPR/DPA compliance are all included as part of the service.

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