TOOLS FOR ADVERSE REACTION ASSESSMENT

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

Accreditations:

ISO 9001
Certificate No. GB2004632

ISO / IEC 27001
Certificate No. ISM7799158

With flexibility at its core, TARA PV is suitable for all pharmacovigilance requirements – with multiple product packages to match your needs and budget. To see TARA PV in action please contact us for a personalised demo.

CONTACT US

Thinki
NOIDA ONE
602, Tower B, Plot B8, Sector 62,
NOIDA 201309, UP

T +91 98100 68241 / 93547 92868
E bd@thinki.in
W www.thinki.in

TARA

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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 with ISO 9001 and ISO 27001 accreditation. Also available are user manuals and training webinars and validation documentation.

✓ 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 (e g 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 E2B vocabulary lists.

✓ CORE PV FUNCTIONALITY

Regulatory reporting (inc. E2B(R3), CIOMS, MedWatch 3500, PSUR and DSUR) and full MedDRA and WHO Drug Dictionary integration. Additional benefits include processing adverse events from clinical trials and marketed products in the same database (inc medical devices), extensive case searching, the capability to store external documents in individual case records and pre-submission case validation.

✓ 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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