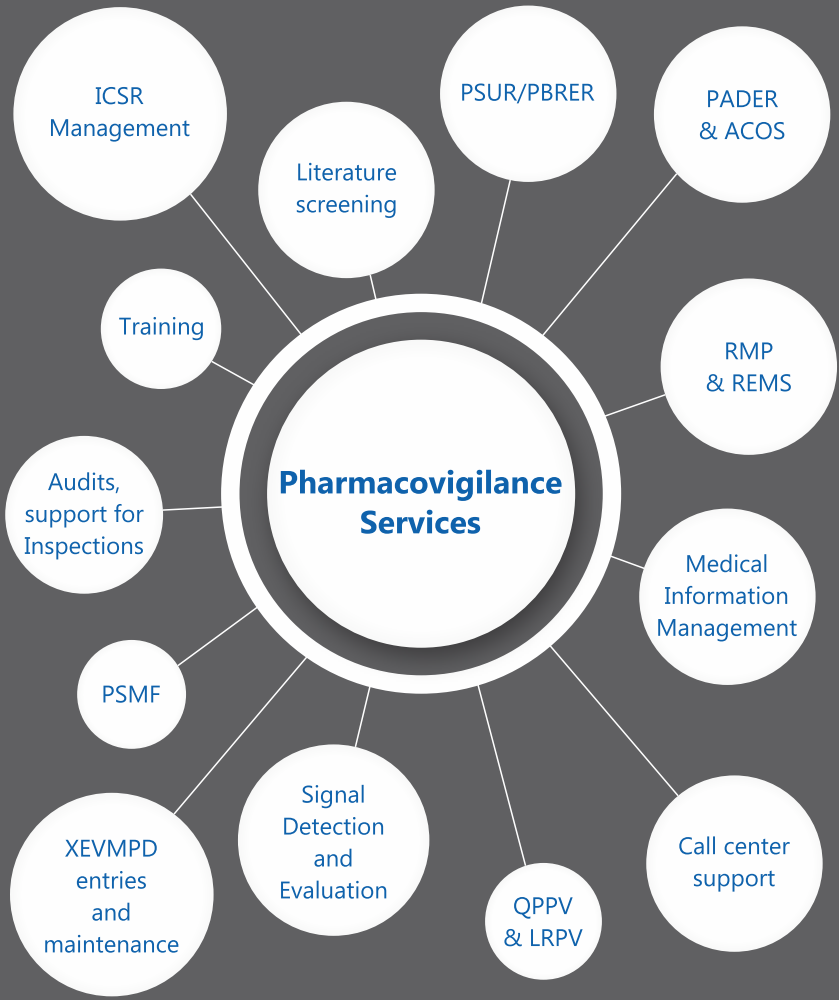
veeda clinical research®

To help our client meet ever increasing regulatory pharmacovigilance obligation, Veeda has created unique comprehensive offering of high quality, compliant service perform by highly trained professionals in cost effective manner

Our one stop Pharmacovigilance solutions can fit seamlessly into your process, meeting all regulatory compliance requirements.



Value Proposition of our Safety Monitoring Services :

- High Quality
- Flexible
- Cost Effective
- Scalable

Our Multi-Disciplinary Team Offers :

- Therapeutic, technical and operational excellence for pharmacovigilance and risk management during the complete lifecycle of a medicinal product
- Up-to-date information on country-specific regulatory requirements
- Robust quality system with a transparent and reliable approach to safety data

VINCURA
care for life

Our in-house safety database is an advanced software solution that fulfills all pharmacovigilance and risk management requirements while ensuing global regulatory compliance with bellow specifications:

- Compliant with ICH and 21 CFR Part 11 guidelines
- Integrates E2B (R3) requirement
- Easy Data Migration - XML file import and export
- GAMP 5 Validation

Features :

- Inbuilt global scientific literature management module, interface with any literature database
- Auto-narrative function limiting manual efforts
- Auto-Scheduling of Aggregate reports
- Global dictionary support (MedDRA, WHODD, etc)
- User productivity report
- Simple user friendly and cost effective

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