ClinFo 1T

We Deliver PV Consultancy, CTD Format Dossier Development & in the MEA, Gulf and Persian Countries

ClinFo1T is a leader in **Pharmacovigilance Consultancy and CTD Format Dossier Development,** offering comprehensive services with customers all over the Globe

Services

- **QPPV/Deputy** PV Services
- Literature Screening
- PV System Master File (**PSMF**)
- Risk Management Plan (**RMP**)
- Serious Adverse Event Case Processing and Reporting
- **SAE** Reconciliation
- Individual Case Safety Report
- PV **SOP**s writing
- PV Consultancy
- **CTD** (and **e-CTD**) Format Dossier Development

Why Us?

- **ISO 9001:2008 from Paris**
- **ICH GXP & US-FDA 21 CFR Compliance**
- **Over 70 International successful audits**
- **Over 90 International Pharmas and Biotechs customers**
- **Over 15 Successful PV MEA Projects**
- **Over 70 EMEA/FDA Submitted CTD**
- **30% More competitive for similar standards of deliverables**

16 YEARS *of*

Quality & Innovation in Healthcare Consultancy
&
Impactful Social Responsibility

Contact us

Website: www.clinfo1t.com **Email:** bd@clinfo1t.com