Pharmacovigilance

The safe way to monitor medicinal products

Diapharm ensures pharmacovigilance for your medicinal products - either as full service or as customized, modular service according to your company requirements. We fullfil all pharmacovigilance tasks according to current EU legislation and Good Pharmacovigilance Practices (GVP). In addition we take over responsibility, for example as Qualified Person for Pharmacovigilance (QPPV) or local contact person for pharmacovigilance for DACH and/or Benelux region.

Our range of services:

- Responsibility as EU-QPPV
- Responsibility as Graduated Plan Officer and/or local contact person for pharmacovigilance (for DACH and/or Benelux)
- Assessment, setup and and implementation of pharmacovigilance (PV) systems
- Preparation and maintenance of pharmacovigilance system master file (PSMF) and Standard Operating Procedures (SOPs)
- Internal and external PV audits
- Preparation and support for authority inspections
- Literature search (global, local, EudraVigilance-, and MLM-Sceening)
- Signal management and EVDAS
- Management of Individual Case Safety reports (ICSRs)
- Preparation of periodic safety update reports (PSURs)
- Preparation and updates of risk management plans (RMPs)
- Preparation of Educational Materials and related authority communication
- Authority and risk communication
- Eudravigilance and xEVMPD

Contact us!

Dr. Christoph Buss
Associate Director Regulatory Affairs & Pharmacovigilance

Further information:
Phone +49 (0)251-60935-0
info@diapharm.com

Your benefits:

- **Company- and product-specific pharmacovigilance systems**
- **Quality-assured processes according to current guidelines**
- **Qualified staff can assume responsibility (EU-QPPV, local QPPV etc.)**
- **Provision of resources according to clients' needs**

Diapharm