



Risk management in pharmacovigilance

[Risk management in pharmacovigilance](#) involves identifying, assessing, minimizing, and monitoring the risks associated with pharmaceutical products throughout their lifecycle. The goal is to ensure that the benefits of a drug outweigh its risks. Key components include risk assessment, the creation of Risk Management Plans (RMPs), and the implementation of risk minimization measures. Continuous monitoring and periodic evaluation help in updating safety measures and protecting public health. Risk management is crucial for regulatory compliance and maintaining patient safety in the use of medicines.