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Drug Safety and Risk Management in Clinical Practice

Guest Editor:

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Message from the Guest Editor

Dear Colleagues,

Safety monitoring of drugs throughout their whole life cycle is of paramount importance in order to protect public health and fulfill the whole scope of pharmacovigilance. Nevertheless, some adverse effects might not become apparent until after these products have been used for an extended period of time and by a large and diverse population, including those with concurrent illnesses and medications. Therefore, drug safety surveillance through spontaneous reporting systems (SRS) and post-authorization safety studies (PASS) using real-world data will provide information and new insights into the safety profiles of the drugs. This Special Issue welcomes reviews and original papers related to drug safety, such as, but not limited to, studies using active or passive methods for drug/vaccine monitoring (PASS, analysis of data from SRS, including signal detection/disproportionality analysis), studies on risks in special populations, and studies on risk minimization measures, including evaluations of the effectiveness of risk minimization measures.

Dr. Andreea Maria Farcas
Guest Editor

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