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Giri, Chandravanu Dash, 2020-08-07 Establishment of a normal phenotype involves dynamic epigenetic regulation of gene expression that when affected contributes to human diseases On a molecular level epigenetic regulation is marked by specific covalent modifications acetylation methylation phosphorylation sumoylation PARYlation and ubiquitylation of DNA and its associated histones Studies also suggest the influence of such epigenetic modifications on non coding RNA expression implicated in normal and diseased phenotypes Epigenetic control of genetic expression is a reversible process essential for normal development and function of an organism Alteration of epigenetic regulation leads to various disease forms such as

cancer diabetes inflammation and neuropsychiatric disorders Assessing these alterations provides a deeper insight into the changes induced in the genome which is often informative for identifying disease subtypes or developing suitable treatments Therefore epigenetics proves to be a key area of clinical investigation in diagnosis prognosis and treatment of complex diseases Genetic mutations environmental stress pathogens and drugs of abuse are some of the predominant factors that induce and impact changes on chromatin which directly dictate a diseased phenotype It is essential to consider the interaction between genetic and epigenetic factors to understand the molecular mechanisms of complex human diseases for safer and efficient drug development Furthermore genetic variation in absorption distribution metabolism and excretion ADME genes is insufficient to account for interindividual variability of drug response Therefore current efforts aim to identify epigenetic components of ADME gene regulation which include phase I and phase II enzymes uptake transporters efflux transporters and nuclear receptors involved in regulation of ADME genes Monitoring circulatory epigenetic biomarkers in liquid biopsies blood saliva urine cerebrospinal fluid of disease associated and drug associated epigenetic alterations may prove useful for decision support for routine clinical treatment and drug discovery Hence recent drug discovery efforts on targeting the epigenome has emerged an area of interest with several new drugs being developed tested and some already approved by the US Food and Drug Administration FDA These new insights into the complexities of epigenetic regulation are key contributors to our basic understanding of this process in human health and disease which will provide scope for innovative drug therapies It is of urgency to aid the present understanding of epigenomics driven diseased outcomes with the expectation that further studies will identify early markers of disease and targets for therapeutics

10 Years of Pharmacogenetics and Pharmacogenomics Jose Agundez, 2020-07-08 The Specialty Section Pharmacogenetics and Pharmacogenomics makes part of two different Journals Frontiers in Pharmacology and Frontiers in Genetics This Specialty Section focuses on the mechanisms by which genetic variations influence drug effects and adverse drug events and cover basic research clinical translation applications in drug development and regulatory issues related to this field Also studies addressing the role of other factors such as epigenetics phenotypic factors or drug drug interactions on drug pharmacokinetics or pharmacodynamics are welcome The editorial board is composed of 34 Associate Editors which together with the Guest Associate Editors and the Reviewer Editors constitute a team of nearly 340 leading experts in the field of Pharmacogenetics and Pharmacogenomics This guarantees high quality in the reviewing process as well as short review times A look back 10 years of Frontiers in Pharmacogenetics Pharmacogenomics Continued in eBook

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