



Editorial

Special Issue “Latest Review Papers in Molecular Genetics and Genomics 2023”

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In the rapidly evolving landscape of molecular genetics and genomics, this Special Issue brings together a collection of insightful review articles that delve into the forefront of scientific exploration.

The Special Issue titled “Latest Advances in Molecular Genetics and Genomics (2023)” of the International Journal of Molecular Sciences includes nine reviews underscoring the remarkable strides made in our understanding of fundamental biological processes and their implications for human health and disease. Spanning diverse topics, the nine articles presented herein illuminate various facets of molecular genetics and genomics, shedding light on the intricate interplay of genetic information, cellular pathways, and disease mechanisms.

A pivotal theme that emerges from these articles is the burgeoning role of non-coding RNAs in shaping gene expression and cellular function. The intricate regulatory networks orchestrated by microRNAs (miRNAs) and circular RNAs (circRNAs) have garnered significant attention as they control gene expression at the transcriptional and post-transcriptional levels and, thus, play a critical role in tumorigenesis and cancer progression. The work by Kim et al. [1] reveals the intricate circular RNA–microRNA axis in cancer, where these non-coding RNAs engage in a complex interplay, influencing tumorigenesis and progression across multiple cancer types. The authors focus on the biological regulation of the correlative axis among circular RNAs, miRNAs, and their target genes in various cancer types and suggest the biological importance of miRNAs derived from transposable elements (MDTEs) interacting with oncogenic or tumor-suppressive circRNAs in tumor progression [1].

Similarly, the study by Pierzynowska et al. [2] underscores the multifaceted involvement of the oxytocin receptor (OXTR) and its ligand in various diseases, highlighting the delicate balance between behavior, molecular signaling, and pathogenesis. The authors summarize the involvement of OXTR dysfunctions and polymorphisms in the development of different diseases, including cancer, cardiac disorders, osteoporosis, and obesity. Their analysis suggests that changes in OXTR expression and OXTR abundance and activity are not specific to individual diseases but rather influence processes that might modulate the course of various disorders. The authors also propose a possible explanation of the discrepancies in the published results of the effects of the OXTR gene polymorphisms and methylation on different diseases [2].

Advances in technology have also enabled comprehensive exploration of genetic alterations and their clinical implications. The investigation by Aveta et al. [3] comprehensively surveys urinary microRNAs (umiRNAs) as potential biomarkers for urological cancers. This systematic review offers insights into the diagnostic and prognostic potential of umiRNAs, illuminating new avenues for non-invasive cancer detection and monitoring. Their systematic review suggests that umiRNAs could play an important role in the diagnosis, prognosis, and therapy of urological cancers [3].

Additionally, the study by Moeckel et al. [4] delves into the promise of tumor mutation burden (TMB) as a predictive biomarker for immune checkpoint inhibitor response. The



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