



## Molecular and Genetic Studies on Tourette Syndrome and Comorbid Disorders

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submissions:

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### Message from the Guest Editors

Gilles de la Tourette syndrome (GTS) is a developmental neuropsychiatric disorder characterized by multiple involuntary motor and vocal tics. The clinical phenotype of GTS belongs to the spectrum of tic disorders (TDs). In general, GTS and TDs have a significant genetic component, with the heritability estimated at 60–80%. However, the clinical phenotype may be influenced by environmental, prenatal, and perinatal factors, hormonal disturbances, and interactions with multiple proteins.

Authors are invited to submit research articles and reviews on related topics, which will improve our understanding of the etiology and pathomechanism of neuropsychiatric disorders. The topics may include, but are not limited to, genetics and genomics, molecular neurobiology, neuromolecular imaging, molecular experimental models, and treatment. Our Special Issue is mainly focused on molecular research. Therefore, we wish to avoid attracting pure clinical manuscripts. Please notice this point when you prepare your manuscript for submission to this Special Issue.





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## Message from the Editor-in-Chief

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