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# **Advances of Brain Transcriptomics**

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Deadline for manuscript submissions: closed (20 July 2022)



#### Message from the Guest Editors

Advancements in RNA-Seq technology have underlined its power for elucidating the brain gene networks responsible for various stressful factors, as well as pathologies like Alzheimer and Parkinson, and other neurological diseases including schizophrenia and depressive disorders. Single cell RNA-seq allows for ascertaining various neurons and glia cell identities by elucidating the specific marker genes within them. Multiple databases like ASCOT and Genome (for elucidating the tissue/cell specific AS profiles for each gene) are arising, underlining the research of brain transcriptome structural and expression variability as a top priority.

Additionally, there is a genetic based vs. acquired trait paradigm, which would be represented by both genetic studies based on animal model strains/breeds (e.g., tame foxes, aggressive strains of rats, etc.), and those that acquired the trait within single generation upon administering certain stress-related protocols.

In this Issue, we hope to address the spectra of physiological studies, including, but not limited to, animal models of social stress response and various brain disease related data/models using the abovementioned approaches.







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### **Editor-in-Chief**

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### Message from the Editor-in-Chief

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