

## Special Issue

# Glioblastoma: State of the Art and Future Perspectives

### Message from the Guest Editors

Glioblastoma is one of the most common types of primary cancer in the central nervous system. Many features contribute to this therapeutic challenge including high intratumoral and intertumoral heterogeneity, resistance to therapy, migration and invasion, immunosuppression. With the entry of novel highthroughput technologies, significant progress has been made to understand molecular and immunological signatures underlying the pathology of glioblastoma. Here, we aim at updating researchers and clinicians to provide a systemic view on glioblastoma by highlighting the progress made in basic, preclinical and clinical research and outlining future challenges. To this end, we welcome articles on basic, preclinical and clinical research in glioblastoma. In addition, we will present invited review articles on the topics of neurosurgical approaches, neuropathology and genetics, preclinical research and mouse models of glioblastoma, imaging, radiation therapy and tumor-treating fields, alkylating chemotherapy, immunotherapy, investigational therapies and precision oncology trials in glioblastoma.

### Guest Editors

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### Deadline for manuscript submissions

closed (31 December 2018)



## Cancers

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## About the Journal

### Message from the Editor-in-Chief

*Cancers* (ISSN 2072-6694) is an international, online journal addressing both clinical and basic science issues related to cancer research. The journal will continue its open access format, which will certainly evolve to ensure that the journal takes full advantage of the rapidly changing world of information and knowledge dissemination. It publishes high-quality clinical, translational, and basic science research on cancer prevention, initiation, progression, and treatment, as well as other related topics, particularly to capture the most seminal studies in the rapidly growing area of immunology, immunotherapy, and tumor microenvironment.

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### Editor-in-Chief

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