

Special Issue

RAS Proteins and Their Regulators in Human Cancer

Message from the Guest Editor

RAS proteins (KRAS, NRAS, and HRAS) are small GTPases that function as binary molecular switches dependent on the loading of GTP (active state) or GDP (inactive state) and which govern critical cellular processes such as cell proliferation, differentiation, and survival. State transitions of RAS proteins are regulated either by intrinsic or GTPase-activating protein (GAP)-stimulated GTP hydrolysis or by guanine nucleotide exchange factor (GEF)-facilitated nucleotide exchange. RAS mutations are found in a large proportion of deadly human tumors, occurring in ~20–30% of malignant tumors overall. Furthermore, RAS mutations are often essential for tumor initiation and maintenance, making RAS a high-priority target in cancer therapy. This Special Issue will cover subjects related to recent progress in RAS proteins, RAS regulators, and their directed therapeutic approaches in cancer therapy. It aims to increase our understanding of RAS regulation and challenges in therapy targeting RAS in human cancers.

Guest Editor

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About the Journal

Message from the Editor-in-Chief

Cancers is an international online journal addressing both clinical and basic science issues related to cancer research. The journal is publishing in 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.

Editor-in-Chief

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