

Special Issue

Molecular Diagnosis and Treatment of Colorectal Cancer

Message from the Guest Editors

This Special Issue aims to focus on the complex genetic and epigenetic alterations that drive malignant transformations in colorectal cancer (CRC).

Understanding these molecular pathways is crucial for diagnosis, treatment, and prognosis. CIN is the most common genetic mechanism in CRC, accounting for approximately 85% of cases. It results from mutations that lead to large-scale chromosomal abnormalities, including gains, losses, and aneuploidy. The key genes involved are APC, KRAS, TP53, BRAF V600E, and SMAD4. MSI occurs due to defects in DNA mismatch repair (MMR) genes, including MLH1, MSH2, MSH6, and PMS2. In this Special Issue, the current understanding of the role of these genes in CRC will be discussed.

Approximately 5–10% of CRC cases arise from inherited genetic mutations, mostly presenting as Lynch Syndrome (Hereditary Non-Polyposis Colorectal Cancer) and Familial Adenomatous Polyposis. Understanding these molecular pathways is crucial for diagnosis, treatment, and prognosis. The emergence of targeted therapies for CRC will also be presented in this Special Issue.

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Deadline for manuscript submissions

31 August 2025



International Journal of Molecular Sciences

an Open Access Journal
by MDPI

Impact Factor 4.9
CiteScore 9.0
Indexed in PubMed



mdpi.com/si/233920

*International Journal of
Molecular Sciences*
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The International Journal of Molecular Sciences (*IJMS*, ISSN 1422-0067) is an open access journal, which was established in 2000. The journal aims to provide a forum for scholarly research on a range of topics, including biochemistry, molecular and cell biology, molecular biophysics, molecular medicine, and all aspects of molecular research in chemistry. *IJMS* publishes both original research and review articles, and regularly publishes special issues to highlight advances at the cutting edge of research. We invite you to read recent articles published in *IJMS* and consider publishing your next paper with us.

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