Special Issue

Innovative Treatments for Anemias

Message from the Guest Editor

Sickle cell anemia (SCA) is a hereditary blood disorder characterized by the presence of abnormal hemoglobin, known as hemoglobin S (HbS), which causes red blood cells to become rigid and assume a sickle shape. This condition can lead to a range of complications, including chronic pain, organ damage, and increased susceptibility to infections. Over the years, there have been significant advancements in understanding the genetic epidemiology of sickle cell anemia and the development of new therapeutic strategie.

Guest Editor

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Deadline for manuscript submissions

closed (24 September 2024)

Journal of Clinical Medicine

an Open Access Journal by MDPI

Impact Factor 2.9
CiteScore 5.2
Indexed in PubMed

mdpi.com/si/175536

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There has been an explosion of gene and target based research and therapeutics in the multitude of fields that compose clinical medicine. The *Journal of Clinical Medicine*'s (*JCM*) staff and editorial board are dedicated to providing cutting edge, timely, and peer-reviewed articles covering the diverse subspecialties of clinical medicine. The journal publishes concise, innovative, and exciting research articles as well as clinically significant articles and reviews that are pertinent to the myriad of disciplines within medicine. The articles published are relevant to both primary care physicians and specialists. The journal's full-texts are archived in PubMed Central and indexed in PubMed. Please consider submitting your manuscripts for publication to our journal and check us out on-line!

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