

## Special Issue

# New Frontiers in Molecular Hematology

### Message from the Guest Editor

The severe complications that still exist on hematopoietic stem cell transplants in the absence of a compatible donor led us, 20 years ago, to explore another therapeutic strategy for genetic diseases of the hematopoietic system. The key idea behind the treatment of these inherited diseases is simple: The diseased bone marrow is replaced with autologous gene-modified cells, in order to establish a stable expression of the missing protein and thus restore its function. Several inherited immune deficiencies and blood disorders can be cured with the use of a lentiviral vector for people affected by immune deficiencies worldwide. Our aim is to bring this powerful innovation to all affected patients.

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### Guest Editor

Prof. Dr. Marina Cavazzana

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2. Head of the Biotherapy Department, Hôpital Necker-Enfants Malades, Assistance Publique-Hôpitaux de Paris, Paris, France
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### Deadline for manuscript submissions

closed (15 December 2020)



## Journal of Clinical Medicine

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