



Pathogenesis and Novel Diagnostic in Juvenile Idiopathic Arthritis

Guest Editor:

Prof. Dr. Katarzyna Winsz-Szczotka

Department of Clinical Chemistry
and Laboratory Diagnostics,
Faculty of Pharmaceutical
Sciences in Sosnowiec, Medical
University of Silesia, Sosnowiec,
Poland

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Message from the Guest Editor

The term ‘juvenile idiopathic arthritis’ (JIA) defines a heterogeneous collection of autoimmune or autoinflammatory forms of chronic arthritis with onset in childhood, with origins that are not yet entirely understood.

There are no pathognomonic symptom or examination findings for JIA, and the diagnosis is made by exclusion and differentiation. JIA that is diagnosed too late or treated poorly treated may contribute to the disability of an afflicted child, due to disturbances in the structure and function of the osteoarticular system. Hence, prompt detection of structural disorders of articular cartilage would allow clinicians to initiate appropriate therapies, which is essential for the course of the arthropathy in question. Too late application of appropriate treatment, resulting from the lack of specific diagnostic biomarkers, may result in the perpetuation of pathological changes in the motor system in patients, or the development of systemic disorders, especially in those with high disease activity who require an aggressive therapy.





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Editor-in-Chief

Prof. Dr. Felipe Fregni

1. Neuromodulation Center and
Center for Clinical Research

Learning, Spaulding
Rehabilitation Hospital and
Massachusetts General Hospital,
Harvard Medical School, Boston,
MA 02114, USA

2. Department of Epidemiology,
Harvard T.H. Chan School of
Public Health, Boston, MA 02115,
USA

Message from the Editor-in-Chief

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MDPI, St. Alban-Anlage 66
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