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Nephrotic Syndrome: Pathomechanism, Diagnostics and Novel Treatment Options

Guest Editor:

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Deadline for manuscript submissions:
closed (31 December 2024)

Message from the Guest Editor

Dear Colleagues,

Nephrotic syndrome (NS) is characterized by massive proteinuria, hypoproteinemia, and edema, and it is divided into steroid-sensitive NS (SSNS) and steroid-resistant NS (SRNS). While SSNS typically presents with minimal change NS, SRNS typically presents with focal segmental glomerulosclerosis (FSGS), which often leads to end-stage kidney failure. About 30% of cases of childhood-onset FSGS have been found to be hereditary FSGS caused by dozens of podocyte-related genes, and about 5-10% of cases of adult-onset FSGS have also been found to be hereditary FSGS.

The aim of this Special Issue is to gather original research articles and review articles that focus on NS. Articles focused on genetic forms of kidney disease are especially encouraged.

Dr. Kan Katayama
Guest Editor

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Message from the Editor-in-Chief

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