Diagnosis and management of inherited kidney diseases: What’s new?

Chairs: Nine V.A.M. Knoers, Utrecht, The Netherlands
Franz Schaefer, Heidelberg, Germany

Part I

Collagen IV glomerulopathies: an underdiagnosed phenotypic chameleon?
Constantinos Deltas, Nicosia, Cyprus

Alport Syndrome: a treatable hereditary kidney disease
Oliver Gross, Göttingen, Germany

NGS podocytopathy panel screening in adults with CKD of unknown origin: findings from the GCKD Study
Anna Koettgen, Freiburg, Germany

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C3 glomerulopathy: role of complement for pathogenesis and treatment
Marina Vivarelli, Rome, Italy

Advances in molecular understanding of cystinosis: implications for therapy
Corinne Antignac, Paris, France

Preimplantation genetic diagnosis for inherited renal diseases
Nine V.A.M. Knoers, Utrecht, The Netherlands

The European Reference Network for Rare Kidney Diseases
Franz Schaefer, Heidelberg, Germany