

Pitch Inherited Kidney Disease I	
Chair: Rina Rus	
<u>PI-86</u>	Novel cellular model of inhibition of glycolate oxidase dependent oxalate production by BBP-711 THULASHITHA RAJASINGHAM 1 ANDY WHITNEY 2 JUSTIN LAFOUNTAINNE 1 UMA SINHA 1
<u>PI-87</u>	Correlation between the metabolic profile of urolithiasis in children with idiopathic hypercalciuria and the composition of the deposit assessed by infrared spectroscopy MALGORZATA PLACZYNSKA 1 JOANNA MILART 1 KATARZYNA JOBS 1 ARKADIUSZ LUBAS 1
<u>PI-88</u>	RESPONSE TO LUMASIRAN IN PRIMARY HYPEROXALURIA TYPE 1 (PH1): FIRST CLINICAL EXPERIENCES SINA SAFFE 1 MARKUS J KEMPER 1 MATTHIAS HANSEN 2 NELE KANZELMEYER 3 ANJA BÜSCHER 4 JUN OH 5 KATJA DOERRY 5
<u>PI-89</u>	Genotype-phenotype correlation in a cohort of individuals with disease-causing variants in COL4A3/COL4A4 associated with type-IV-collagen-related nephropathy (Alport syndrome and thin basement membrane nephropathy) SIMMENDINGER HANNES 1 RIEDHAMMER KORBINIAN MARIA 1 TASIC VELIBOR 3 PUTNIK JOVANA 4 ABAZI-EMINI NORA 3 STAJIC NATASA 4 WEIDENBUSCH MARC 5 PATZER LUDWIG 6 LUNGU ADRIAN 7 MILOSEVSKI-LOMIC GORDANA 4 BRAUNISCH MATTHIAS 2 GÜNTNER ROMAN 2 COMIC JASMINA 1 HOEFELE JULIA 1
<u>PI-90</u>	A rare cause of combined hepatic and renal failure: NPHP19 due to a novel DCDC2 variant in two siblings GIZEM YILDIZ 1 MERAL TORUN BAYRAM 1 AHMET OKAY ÇAĞLAYAN 2 AYFER ÜLGENALP 2 ALPER SOYLU 1 SALIH KAVUKCU 1
<u>PI-91</u>	EARLY PRO-TECT Alport XXL: A Worldwide Observational Study to Improve Evidence for Preemptive Start of ACE Inhibitor Therapy in Children With Alport Syndrome OLIVER GROSS 1 JAN BOECKHAUS 1
<u>PI-92</u>	Autosomal dominant hypocalcemia. When Bartter syndrome "pushes boundaries" MARTA JIMÉNEZ MORENO 1 PEDRO ARANGO SANCHO 1 ANA CRISTINA AGUILAR RODRÍGUEZ 1 RAQUEL JIMÉNEZ GARCÍA 1 ELENA CODINA SAMPERA 1 YOLANDA CALZADA BAÑOS 1 ÁLVARO MADRID ARIS 1