Symposium 11	
Inherited renal disorders	
OP-30	**Novel syndrome: Craniofacial and central nervous system malformations, and atypical hemolytic uremic syndrome associated with TSEN2 mutation** NUR CANPOLAT 1 DINGXIAO LIU 2 EMINE ATAYAR 3 SEHA SAYGILI 1 NAZLI SILA KARA 4 TRUDI A. WESTFALL 5 QIONG DING 2 BARTLEY J. BROWN 6 TERRY A. BRAUN 6 DIANE SLUSARSKI 6 KADER KARLI OGUZ 7 YASEMIN OZLUK 8 BEYHAN TUYSUZ 9 TUGBA TASTEMEL OZTURK 10 LALE SEVER 1 OSMAN UGUR SEZERMAN 4 REZAN TOPALOGLU 10 SALIM CALISKAN 1 MASSIMO ATTANASIO 2 FATIH OZALTIN 10
OP-31	**Effects of burosumab treatment on mineral homeostasis in children and adolescents with X-linked hypophosphatemia: lessons from the German XLH Registry** EWERT ANNIKA REHBERG MIRKO 2 SCHLINGMANN KARL PETER 3 KEMPER MARKUS 4 DERICHS UTE 5 PATZER LUDWIG 6 STAUDE HAGEN 8 JOHN ULRIKE 7 METZING OLIVER 7 WEITZ MARCUS 9 FREIBERG CLEMENS 10 WÜHL ELKE 11 SCHAEFER FRANZ 11 HIORT OLAF 12 SCHNABEL DIRK 13 HAFFNER DIETER 1
OP-32	**A Randomized, Placebo -Controlled, Phase 2/3 study of glycolate oxidase (GO) inhibitor BBP-711 in Children and Adults with Primary Hyperoxaluria Type 1** SCOTT ADLER 1 JIA MA 1 RAMEI SANI-GROSSO 1 LILLIAN LEE 1 GUSTAVO LORENTE 1 JUSTIN LAFOUNTAINE 1 JONATHAN FOX 1