



**2018**

1. Gonçalves S, Patat J, Guida MC, (...) Boyer O, (...) Antignac C, Simons M. **Correction: A homozygous KAT2B variant modulates the clinical phenotype of ADD3 deficiency in humans and flies.**  
PLoS Genet. 2018 Oct 26;14(10):e1007748. doi: 10.1371/journal.pgen.1007748. eCollection 2018 Oct. PubMed PMID: 30365502.
2. Gribouval O, Boyer O, Hummel A, (...) Tête MJ, Heidet L, Antignac C, Servais A. **Identification of genetic causes for sporadic steroid-resistant nephrotic syndrome in adults.**  
Kidney Int. 2018 Nov;94(5):1013-1022. doi: 10.1016/j.kint.2018.07.024. PubMed PMID: 30348286.
3. Gribouval O, Boyer O, Knebelmann B, Karras A, Dantal J, Fourrage C, Alibeu O, Hogan J, Dossier C, Tête MJ, Antignac C, Servais A. **APOL1 risk genotype in European steroid-resistant nephrotic syndrome and/or focal segmental glomerulosclerosis patients of different African ancestries.**  
Nephrol Dial Transplant. 2018 Jul 9. doi: 10.1093/ndt/gfy176. [Epub ahead of print] PubMed PMID: 29992269.
4. Gonçalves S, Patat J, Guida MC, Lachaussée N, Arrondel C, Helmstädter M, Boyer O, Gribouval O, Gubler MC, Mollet G, Rio M, Charbit M, Bole-Feysot C, Nitschke P, Huber TB, Wheeler PG, Haynes D, Juusola J, Billette de Villemeur T, Nava C, Afenjar A, Keren B, Bodmer R, Antignac C, Simons M. **A homozygous KAT2B variant modulates the clinical phenotype of ADD3 deficiency in humans and flies.**  
PLoS Genet. 2018 May 16;14(5):e1007386. doi: 10.1371/journal.pgen.1007386. eCollection 2018 May. Erratum in: PLoS Genet. 2018 Oct 26;14(10):e1007748. PubMed PMID: 29768408; PubMed Central PMCID: PMC5973622.
5. Bérody S, Heidet L, Gribouval O, Harambat J, Niaudet P, Baudouin V, Bacchetta J, Boudailliez B, Dehennault M, de Parscau L, Dunand O, Flodrops H, Fila M, Garnier A, Louillet F, Macher MA, May A, Merieau E, Monceaux F, Pietrement C, Rousset-Rouvière C, Roussey G, Taque S, Tenenbaum J, Ulinski T, Vieux R, Zaloszc A, Morinière V, Salomon R, Boyer O. **Treatment and outcome of congenital nephrotic syndrome.**  
Nephrol Dial Transplant. 2018 Feb 20. doi: 10.1093/ndt/gfy015. [Epub ahead of print] PubMed PMID: 29474669.

6. Dorval G, Gribouval O, Martinez-Barquero V, Machuca E, Tête MJ, Baudouin V, Benoit S, Chabchoub I, Champion G, Chauveau D, Chehade H, Chouchane C, Cloarec S, Cochat P, Dahan K, Dantal J, Delmas Y, Deschênes G, Dolhem P, Durand D, Ekinci Z, El Karoui K, Fischbach M, Grunfeld JP, Guignon V, Hachicha M, Hogan J, Hourmant M, Hummel A, Kamar N, Krummel T, Lacombe D, Llanas B, Mesnard L, Mohsin N, Niaudet P, Nivet H, Parvex P, Pietrement C, de Pontual L, Noble CP, Ribes D, Ronco P, Rondeau E, Sallee M, Tsimaratos M, Ulinski T, Salomon R, Antignac C, Boyer O. **Clinical and genetic heterogeneity in familial steroid-sensitive nephrotic syndrome.**  
Pediatr Nephrol. 2018 Mar;33(3):473-483. doi: 10.1007/s00467-017-3819-9. Epub 2017 Oct 23. PubMed PMID: 29058154.
7. Braun DA, Rao J, Mollet G, (...) Boyer O, R(...) Antignac C, Hildebrandt F. **Mutations in KEOPS-complex genes cause nephrotic syndrome with primary microcephaly.**  
Nat Genet. 2017 Oct;49(10):1529-1538. doi: 10.1038/ng.3933. Epub 2017 Aug 14. PubMed PMID: 28805828; PubMed Central PMCID: PMC5819591.
8. Lipska-Ziętkiewicz BS, Gellermann J, Boyer O, Gribouval O, (...) Antignac C, Schaefer F; PodoNet Consortium.  
**Low renal but high extrarenal phenotype variability in Schimke immuno-osseous dysplasia.**  
PLoS One. 2017 Aug 10;12(8):e0180926. doi:10.1371/journal.pone.0180926. eCollection 2017. PubMed PMID: 28796785; PubMed Central PMCID: PMC5552097.
9. Gruppen MP, Bouts AH, Jansen-van der Weide MC, (...) Niaudet P, Cornelissen EAM, Schurmans T, Raes A, van de Walle J, van Dyck M, Gulati A, Bagga A, Davin JC; all members of the Levamisole Study Group.  
**A randomized clinical trial indicates that levamisole increases the time to relapse in children with steroid-sensitive idiopathic nephrotic syndrome.**  
Kidney Int. 2017 Oct 17.
10. Boyer O, Dorval G, Servais A. **Hereditary Podocytopathies in Adults: The Next Generation.**  
Kidney Dis (Basel). 2017 Jul;3(2):50-56. doi: 10.1159/000477243. Epub 2017 May 31. Review. PubMed PMID: 28868292; PubMed Central PMCID: PMC5566765.
11. Dossier C, Lapidus N, Bayer F, Sellier-Leclerc AL, Boyer O, de Pontual L, May A, Nathanson S, Orzechowski C, Simon T, Carrat F, Deschênes G. **Epidemiology of idiopathic nephrotic syndrome in children: endemic or epidemic?**  
Pediatr Nephrol. 2016 Dec;31(12):2299-2308.
12. Ding WY, Koziell A, McCarthy HJ, Bierzynska A, (...) Antignac C, Boyer O, Saleem MA. **Initial steroid sensitivity in children with steroid-resistant nephrotic syndrome predicts post-transplant recurrence.**  
J Am Soc Nephrol. 2014 Jun;25(6):1342-8.
13. Greff B, Faivre J, Carli PA, Niaudet P, Orliaguet GA. **Intra- and postoperative adverse events in children with nephrotic syndrome requiring surgery under general anesthesia.**  
Paediatr Anaesth. 2012 Mar;22(3):244-9.

14. Dossier C, Sellier-Leclerc AL, Rousseau A, Michel Y, Gautheret-Dejean A, Englender M, Madhi F, Charbit M, Ulinski T, Simon T, Jacqz-Aigrain E, Deschênes G. **Prevalence of herpesviruses at onset of idiopathic nephrotic syndrome.**  
Pediatr Nephrol. 2014 Dec;29(12):2325-31.
15. Bouchireb K, Boyer O, Gribouval O, Dantal J, Antignac C. **NPHS2 mutations in steroid-resistant nephrotic syndrome: a mutation update and the associated phenotypic spectrum.**  
Hum Mutat. 2014 Feb;35(2):178-86.
16. Boyer O, Niaudet P. **Nephrotic syndrome: Rituximab in childhood steroid-dependent nephrotic syndrome.**  
Nat Rev Nephrol. 2013 Oct;9(10):562-3.
17. Krug P, Boyer O, Balzamo E, Sidi D, Lehnert A, Niaudet P. **Nephrotic syndrome in Kawasaki disease.**  
Pediatr Nephrol. 2012 Sep;27(9):1547-50.
18. Baudouin V, Alberti C, Lapeyraque AL, Bensman A, André JL, Broux F, Cailliez M, Decramer S, Niaudet P, Deschênes G, Jacqz-Aigrain E, Loirat C. **Mycophenolate mofetil for steroid-dependent nephrotic syndrome: a phase II Bayesian trial.**  
Pediatr Nephrol. 2012 Mar;27(3):389-96.