

Publications for Hugh McCarthy

2019

Jayasinghe, K., Stark, Z., Patel, C., Mallawaarachchi, A., McCarthy, H., Faull, R., Chakera, A., Sundaram, M., Jose, M., Kerr, P., et al (2019). Comprehensive evaluation of a prospective Australian patient cohort with suspected genetic kidney disease undergoing clinical genomic testing: A study protocol. *BMJ Open*, 9(8). [More Information]

2018

Francis, A., Didsbury, M., McCarthy, H., Kara, T. (2018). Treatment of recurrent focal segmental glomerulosclerosis post-kidney transplantation in Australian and New Zealand children: A retrospective cohort study. *Pediatric Transplantation*, 22(5), 1-7. [More Information]

2017

Bierzynska, A., Soderquest, K., Dean, P., Colby, E., Rollason, R., Jones, C., Inward, C., McCarthy, H., Simpson, M., et al (2017). MAGI2 Mutations Cause Congenital Nephrotic Syndrome. *Journal of the American Society of Nephrology*, 28(5), 1614-1621. [More Information]

Mallett, A., McCarthy, H., Ho, G., Holman, K., Farnsworth, E., Patel, C., Fletcher, J., Mallawaarachchi, A., Quinlan, C., Bennetts, B., Alexander, S. (2017). Massively parallel sequencing and targeted exomes in familial kidney disease can diagnose underlying genetic disorders. *Kidney International*, 92(6), 1493-1506. [More Information]

2015

Ebarasi, L., Ashraf, S., Bierzynska, A., Gee, H., McCarthy, H., Lovric, S., Sadowski, C., Pabst, W., Vega-Warner, V., et al (2015). Defects of CRB2 Cause Steroid-Resistant Nephrotic Syndrome. *American Journal of Human Genetics*, 96(1), 153-161. [More Information]

Mallett, A., Corney, C., McCarthy, H., Alexander, S., Healy, H. (2015). Genomics in the renal clinic-translating nephrogenetics for clinical practice. *Human Genomics (Online)*, 9(1), 1-4. [More Information]

2014

Ding, W., Koziell, A., McCarthy, H., Bierzynska, A., Bhagavatula, M., Dudley, J., Inward, C., Coward, R., Tizard, E., et al (2014). Initial steroid sensitivity in children with steroid-resistant nephrotic syndrome predicts post-transplant recurrence. *Journal of the American Society of Nephrology*, 25(6), 1342-1348. [More Information]

2013

Harris, J., McCarthy, H., Ni, L., Wherlock, M., Kang, H., Wetzels, J., Welsh, G., Saleem, M. (2013). Active proteases in nephrotic plasma lead to a podocin-dependent phosphorylation of VASP in podocytes via protease activated receptor-1. *Journal of Pathology*, 229(5), 660-671. [More Information]

McCarthy, H., Bierzynska, A., Wherlock, M., Ognjanovic, M., Kerecuk, L., Hegde, S., Feather, S., Gilbert, R., Krischock, L., Jones, C., et al (2013). Simultaneous Sequencing of 24 Genes Associated with Steroid-Resistant Nephrotic Syndrome. *Clinical Journal of the American Society of Nephrology*, 8(4), 637-648. [More Information]

2011

McCarthy, H., Saleem, M. (2011). Genetics in clinical practice: nephrotic and proteinuric syndromes. *Nephron Experimental Nephrology*, 118(1), e1-e8. [More Information]

McCarthy, H., Inward, C., Marriage, S., Astley, P., Tizard, E. (2011). Red cell exchange transfusion as a rescue therapy for tacrolimus toxicity in a paediatric renal transplant. *Pediatric Nephrology*, 26(12), 2245-2248. [More Information]

2010

McCarthy, H., Tizard, E. (2010). Clinical practice: diagnosis and management of Henoch-Schonlein purpura. *European Journal of Pediatrics*, 169(6), 643-650. [More Information]

Edwards, A., McCarthy, H., Morgan, J., Saleem, M. (2010). Paediatric non-heart-beating renal transplantation. *Archives of Disease in Childhood*, 95(10), 843-844. [More Information]