5. Anlagen

5.1 Der Habilitationsschrift zugrunde liegende Originalpublikationen

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B. Köhler*, **V. Schumacher****,* D. l'Allemand, B. Royer-Pokora, A. Grüters. Germline Wilms tumor suppressor gene (*WT1*) mutation leading to isolated genital malformation without Wilms tumor or nephropathy. *J Pediatr,* 138: 421-424, 2001. * = Gleichwertige Erstautorenschaft

P. Kaltenis*, **V. Schumacher****, A. Jankauskienė, A. Laurinavičius, B. Royer-Pokora. Slow progressive FSGS associated with a F392L *WT1*-mutation. *Pediatr Nephrol,* 19(3): 353-6, 2004. * = Gleichwertige Erstautorenschaft

V. Schumacher, S. Jeruschke, F. Eitner, Jan Becker, G. Pitschke, Y. Ince, J. Miner, I. Leuschner, R. Engers, A. Schulze Everding, M. Bulla, B. Royer-Pokora. Impaired glomerular maturation and lack of VEGF165b in Denys-Drash Syndrome. *J Am Soc Nephrol,* 18: 719-29, 2007

V. Schumacher, J. Thumfart, M. Drechsler, M. Essayie, B. Royer-Pokora, U. Querfeld und D Müller. A novel *WT1* missense mutation presenting with Denys-Drash Syndrome and cortical atrophy. *Nephrol Dial Transplant,* 21: 518-521, 2005

V. Schumacher, B. Gueler, L. HJ Looijenga, J. Ulrich Becker, K. Amann, R. Engers, J. Doetsch, H. Stoop, W. Schulz, B. Royer-Pokora. Characteristics of Testicular Dysgenesis Syndrome and Decreased Expression of *SRY* and *SOX9* in Frasier Syndrome. *Mol Reprod Dev*, zum Druck angenommen.