Steroid-resistant nephrotic syndrome with mutations in NPHS2 (podocin): report from a three-generation family

Vani Jain1, John Feehally2, Gabriela Jones1, Lisa Robertson1, Dheepa Nair1 and Pradeep Vasudevan1

1Department of Clinical Genetics, Leicester Royal Infirmary, University Hospitals of Leicester, Leicester, UK and 2Department of Nephrology, Leicester General Hospital, University Hospitals of Leicester, Leicester, UK

Correspondence and offprint requests to: Pradeep Vasudevan; E-mail: pradeep.vasudevan@uhl-tr.nhs.uk

Abstract
Genetic causes of steroid-resistant nephrotic syndrome are being increasingly recognized. Mutations in NPHS2, which encodes the glomerular protein podocin, account for up to 17% of sporadic and 40% of familial cases, where they display an autosomal-recessive pattern of inheritance. This report describes a non-consanguineous family with three generations of individuals who are either compound heterozygotes for mutations in NPHS2 or who have inherited a mutation and a non-neutral polymorphism (R229Q). As well as providing an aetiological explanation, identifying pathogenic mutations and considering genotype-phenotype correlations can provide prognostic information and lead to changes in genetic counselling and management.

Keywords: focal segmental glomerulosclerosis; NPHS2; podocin; steroid-resistant nephrotic syndrome

Background
Nephrotic syndrome (NS) is defined by the presence of proteinuria, hypoalbuminaemia and oedema. The peak age of presentation is in pre-school children with the majority of cases responding to steroids [1]. However, around 10% of children and 40% of adults with NS fall into the ‘steroid-resistant’ category (steroid-resistant nephrotic syndrome: SRNS) [2]. The progression to end-stage renal disease (ESRD) occurs in 50–70% of patients with SRNS [3]. Inherited structural defects of the glomerular filtration barrier have been identified in isolated as well as familial cases of SRNS [2, 4, 5]. The genes involved in non-syndromic SRNS include NPHS1, NPHS2, CD2AP, PLCE1, ACTN4, TRPC6, INF2, MOY1E, PTPRO, and ARHGDI1. Syndromic SRNS is less common but the known genes are WT1, LMX1B, LAMB2, ITGB4, SCARB2, COQ2, PDSS2, MTL1, SMARCAL1, MYH9, and NXF5 [6, 7].

The protein podocin, encoded by the NPHS2 gene, is expressed exclusively in glomerular podocytes at the insertion site of the slit diaphragm [8, 9]. Mutations in NPHS2 have been found in 40% of familial SRNS cases [where it follows an autosomal-recessive (AR) pattern of inheritance] as well as in 6–17% of sporadic SRNS cases [10]. NPHS2 mutations lead to dysfunction of the glomerular filtration barrier with the prevailing renal pathology being focal segmental glomerulosclerosis (FSGS) [3, 9, 11]. The age of onset is variable but typically before 6 years of age [3, 12]. Late-onset (>18 years) cases of SRNS associated with NPHS2 are also well recognized. These patients are frequently found to be compound heterozygotes for a mutation and the non-neutral polymorphism, R229Q [3, 12–14].

This report describes a non-consanguineous Caucasian family in which two generations are affected with SRNS due to compound heterozygous mutations in NPHS2 and with an older unaffected family member in an earlier generation who has inherited a mutation and a non-neutral polymorphism (R229Q) in the same gene. We illustrate how genetic testing in SRNS can lead to modified management for patients and define the risks for the wider family.

Case report
The proband (III.1 in Figure 1) was 13 months old when she presented with ‘puffy eyes’, proteinuria and generally feeling unwell. She was diagnosed with NS and a renal biopsy showed minimal-change disease. Her family history was significant in that her father (II.1) had a renal transplant at the age of 10. He had been admitted at 11 months of age with respiratory symptoms, constant wet nappies and excessive thirst. Urinalysis revealed significant proteinuria and he was diagnosed with NS. The initial renal biopsy was suggestive of minimal-change disease but a repeat biopsy demonstrated FSGS. Both father and daughter had trials of various immunosuppressants that failed to improve their condition. The proband is now 9 years old and on furosemide and enalapril. Recent blood results showed sodium 140 mmol/L, potassium 5.0 mmol/L, urea 8.5 mmol/L, creatinine 27 μmol/L, calcium...
2.32 mmol/L, phosphate 1.61 mmol/L and albumin 21 g/L. Her father remains well since his renal transplant.

After referral to a regional clinical genetics centre, the family was counselled about SRNS following what appeared to be an autosomal-dominant (AD) pattern of inheritance but was also told that AR inheritance was a possibility. Genetic testing was undertaken and the father and daughter were both found to have two pathogenic NPHS2 mutations (R138Q and Q215X). This particular gene and the presence of two mutations suggested an AR form of SRNS. In order to confirm this and to provide accurate recurrence risks, NPHS2 genetic testing was offered to other family members. The mother (II.2) of the proband was a carrier for mutation Q215X, and II.1 and II.2 were told that they have a 50% chance of having another child affected with SRNS and that they may want to consider pre-natal testing in a future pregnancy if they feel that they would terminate an affected fetus. The genotype of an affected fetus may be either R138Q/Q215X or R138Q/R138Q. The latter is one of the genotypes associated with the earliest onset of NPHS2-associated SRNS, although there is no significant difference in progression to ESRD [12]. SRNS is not currently on the Human Fertilization and Embryology Authorities (HFEA) list of conditions licensed for pre-implantation genetic diagnosis (PGD) [17] but in theory would be possible, as causative mutations have been found. Although a recurrence risk of 50% is the same as the risk that this couple would have been given if the condition had been confirmed as following an AD pattern of inheritance, the risk to any future children of III.1 is no longer 50%. She now has a 25% risk of having a child with SRNS, but only if her partner also happens to be a carrier of a mutation in NPHS2.

Cascade testing in the proband’s paternal grandfather (I.1) revealed an additional complexity in this case. He carried a pathogenic mutation (R138Q) along with a variant (R229Q). Variants (also called ‘polymorphisms’) are found in all genes. Many are not pathogenic, but some may have an as yet undefined disease-modifying role (a ‘variant of unknown significance’). R229Q is the commonest NPHS2 variant in people of European descent, with a frequency of between 2 and 3% in this population [3, 4]. However, it has a higher frequency in SRNS cases (5.3% in a study by Santin et al [15]). When R229Q is inherited with a pathogenic NPHS2 mutation (in trans), some individuals have developed SRNS with a significantly later onset compared to those with two NPHS2 mutations [12–14]. Studies by Santin et al. and Machuca et al. reported that adult-onset SRNS cases with one NPHS2 mutation all had the R229Q variant in trans, rather than a second NPHS2 mutation [3, 14].

Fig. 1. Family pedigree showing the genotypes of those individuals who had NPHS2 testing (written below the symbol) and their phenotype.
Therefore, the evidence suggests that R229Q had some role to play in the development of SRNS (i.e. a non-neutral polymorphism). Because of this, some advocated that the spouses of individuals with NPHS2-associated SRNS be offered genetic testing for the R229Q variant as there is up to a 50% risk of their children developing later onset SRNS [10]. However, R229Q inherited with a pathogenic mutation is not disease causing in all cases and at the age of 68, individual I.1 remains well. A recently published study by Tory et al. [18] shows that R229Q is only pathogenic when in trans association with specific NPHS2 mutations. Those mutations located in exons 1–6 (which includes R138Q) are unlikely to be pathogenic in association with R229Q, which is supported by our case. A greater understanding of the contribution of R229Q to the development of SRNS will enable more accurate counselling for families in the future.

This case, no doubt, illustrates that rare recessive conditions can manifest in more than one individual in a non-consanguineous family. The two carriers with R138Q (I.1 and II.2) are to the best of their knowledge, unrelated, as are II.1 and II.2. The R138Q mutation is the most frequently found pathogenic NPHS2 mutation and is believed to be a founder mutation in Europe [13, 19], therefore possibly explaining its presence in multiple individuals in the family.

We have shown that genetic testing in SRNS can be useful for aetiological confirmation and for providing accurate risk assessment and counselling in families. Significant progress has been made in understanding the genetics of SRNS. Further research is needed to understand the role of genetic variants and identify modifiers of disease that may explain why, for example, siblings with the same NPHS2 mutations have variability in the onset and presentation of SRNS [12, 20]. The ‘mainstreaming’ of genetic testing, and increasing use of next-generation sequencing means that clinicians will be faced more frequently with questions of how to interpret and use genetic information in the management and counselling of patients and their families.

Acknowledgements. The authors would like to thank the family for agreeing to publication.

Conflict of interest statement. The results presented in this paper have not been published previously in whole or part, except in abstract format.

References


Received for publication: 24.8.13; Accepted in revised form: 28.2.14