Characterisation of a novel TSC2 missense mutation in the GAP related domain associated with minimal clinical manifestations of tuberous sclerosis

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BACKGROUND

Tuberous sclerosis (TSC) is an autosomal dominant disorder caused by mutations in either the TSC1 or TSC2 gene. Both genes encode tumour suppressor genes and encode the proteins hamartin and tuberin, respectively. Recent work has established that tuberin and hamartin interact to form a complex, and that this tuberin-hamartin complex antagonises signal transduction by preventing the activation of p70 S6 kinase, either directly or through mTOR. The inhibition of p70 S6 kinase activity results in the dephosphorylation of ribosomal protein S6 and consequently leads to the down-regulation of cell growth. 4

The clinical manifestations of TSC vary considerably between and within families. The phenotypic spectrum ranges from minor, asymptomatic skin affections, such as hypomelanotic macules, to drug resistant epilepsy, mental retardation, and increased morbidity due to cardiac tumours, cerebral astrocytomas, renal insufficiency, and lung affections. Although some studies suggest that TSC1 mutations are more often associated with less severe disease, considerable overlap exists in the phenotypic spectrum associated with TSC1 and TSC2 mutations.

Approximately 50% of cases result from new mutations which arise during meiosis or maternal oogenesis. The remainder are due to de novo mutations, and most of these are maternally inherited. Investigations into the effect of amino acid substitutions on tuberin can help identify important functional domains and residues, and indicate whether putative mutations are likely to be pathogenic, either by disrupting the tuberin-hamartin complex, or by preventing the inhibition of S6 phosphorylation.

METHODS

Mutation analysis and genotyping

Genomic DNA was extracted from peripheral lymphocytes by standard methods. All 21 coding exons of the TSC1 gene and all 41 coding exons of the TSC2 gene, including flanking intronic sequences, were amplified by PCR. Both strands were sequenced using a direct dideoxy method (ABI Prism). All 41 coding exons of the TSC2 gene, including flanking intronic sequences, were amplified by PCR. Both strands were sequenced using a direct dideoxy method (ABI Prism).

Confirmation of sequence variants as well as haplotyping of unrelated controls was performed by RFLP analysis and PyrosequencingTM (www.pyrosequencing.com). All the constructs used were sequenced using the Stratagene QuickChange kit (La Jolla, CA, USA) and analysed on an ABI3100 sequencer. Functional analysis

To obtain additional evidence that the TSC2 4684G>A mutation was the pathogenic mutation in this family, the effect of the G1556S substitution on tuberin function was investigated. Compared to wild type tuberin, the G1556S variant was hypophosphorylated, showed a reduced ability to form a complex with hamartin, and was unable to inhibit the phosphorylation of ribosomal protein S6, supporting the conclusion that the TSC2 4684G>A substitution was the pathogenic mutation in this family. The implications of these results for the diagnosis of TSC patients are discussed.

Key points

- Tuberous sclerosis (TSC) is an autosomal dominant disorder with a broad phenotypic spectrum. The clinical manifestations of the disease can vary considerably, both between and within families.
- Here we describe a family segregating a novel missense mutation, 4684G>A (G1556S), in exon 36 of the TSC2 gene. Only the index patient fulfilled the present diagnostic criteria for TSC. However, clinical signs associated with TSC but insufficient for a definitive diagnosis were identified in additional family members carrying the same substitution.
- The mutation leads to a change of a highly conserved glycine residue in the GAP related domain of tuberin. To obtain additional evidence that the TSC2 4684G>A mutation was the pathogenic mutation in this family, the effect of the G1556S substitution on tuberin function was investigated.
- Compared to wild type tuberin, the G1556S variant was hypophosphorylated, showed a reduced ability to form a complex with hamartin, and was unable to inhibit the phosphorylation of ribosomal protein S6, supporting the conclusion that the TSC2 4684G>A substitution was the pathogenic mutation in this family. The implications of these results for the diagnosis of TSC patients are discussed.
antibodies used in this study have been described previously. Phosphospecific antibodies against S6 (S235/236) and the PKB target sequence ([(R/K)(X/R/K)(X/T-P/S-P)]) were purchased from Cell Signaling Technology (www.cellsignal.com); antibodies against the polyhistidine tag were purchased from Qiagen (www.qiagen.com). Coimmunoprecipitation experiments were performed on transfected COS-1 cells as described previously. The effect of tuberin expression on S6 phosphorylation was assayed in homozygote TSC2 knockout mouse embryonic fibroblasts kindly provided by H Onda, Boston, USA. The cells were transfected with the wild type or G1556S TSC2 expression construct using lipofectAMINE PLUS reagent, according to the manufacturer’s guidelines (Invitrogen, Carlsbad, CA, USA). Twenty four hours after transfection the medium was replaced with serum free medium and the cells incubated for an additional 24 hours before being processed for double label fluorescent microscopy, according to the recommended protocol (Cell Signaling Technology). Transfected cells expressing tuberin and showing inhibition of S6 phosphorylation were counted in three separate experiments, according to established methods. At least 50 cells were counted per experiment.

RESULTS

Clinical findings

A pedigree of the family is given in fig 1, and the clinical symptoms of the mutation carriers are listed in table 1. In the index patient (IV-3), fetal sonography revealed a cardiac “white spot” in the 22nd week of gestation, possibly due to the presence of a cardiac rhabdomyoma. Fourteen weeks later, three intracardial tumours of up to 5 mm in size were recorded. After birth these tumours were confirmed as 9×6 mm and 8×6 mm in size. Cerebral structures were normal and postnatal development was uncomplicated. No other cardiac abnormalities were reported. At the age of 2 months, clinical examination including Woods light revealed three typical hypomelanotic macules, sufficient for the diagnosis of TSC. Ophthalmological examination and CT were normal. The child is now 3 years of age and has attained normal developmental landmarks. Seizures have never been observed.

The sibling of the index patient (IV-2) was delivered by vacuum extraction after an uncomplicated pregnancy. At the age of 2 years he had one single febrile convulsion. He never received anticonvulsive medical treatment. Mental and speech development were slow. Clinical examination including Woods light at the age of 4 years revealed no cutaneous TSC symptoms. Cerebral MRT and CT as well as echocardiography were normal. At the age of 7 years an attention deficit hyperactivity disorder (ADHD) was diagnosed and treatment with Ritalin (methylphenidat) was initiated. At that time he had remained free of seizures and displayed no cutaneous signs of TSC. This individual does not meet the diagnostic criteria for TSC.

Individual III-3, the father of IV-2 and IV-3, is 35 years of age and healthy. Clinical examination including Woods light revealed a single hypomelanotic macule of coin size on the left buttock. Renal ultrasound revealed bilateral local increases in echogeneity of the upper poles and a single echogenic structure of 1 cm in size, compatible with an angiomyolipoma. Liver sonography and cerebral CT were normal. Individual III-2, the paternal aunt of the index patient, had been examined at the age of 20 years because of multiple depigmented lesions and suspected TSC. CT, eye fundus and nails were found to be normal. Examination at the age of 36 years revealed multiple characteristic hypomelanotic macules, but no complaints. She had a minor shagreen patch in the lumbar region of 2 mm in size, consistent with TSC.

Her son, individual IV-1, was admitted to hospital at the age of 11 years because of epileptic attacks which ceased after anticonvulsive treatment. EEG showed a focal activity of both temporoparietal regions and MRT revealed an increase in signal intensity in the mesial temporal lobes, which were interpreted as beginning hippocampal sclerosis, although inflammation or an astrocytoma could not be excluded at that time. A causal relationship of this finding to TSC is neither proven nor excluded. There were no cortical tubera and no calcified subependymal nodules. Sonography revealed one single kidney on the right. Eye fundi were normal. Dermatological examination at the age of 12 years revealed three hypomelanotic macules of up to 1.5 cm × 2.5 cm in size.

Individual II-2, the grandfather of the index patient, was not available for examination. However, there were no reports of epileptic seizures or facial angiofibromas or any other TSC symptoms. Individual II-1 died at the age of 30 years from renal bleeding. The possibility that the bleeding was due to a renal angiomyolipoma was not excluded.

Genetic analysis

Complete sequencing of the coding regions of both TSC genes from DNA of peripheral leukocytes in the index patient (IV-3) identified a germline mutation 4684G>A in the TSC2 gene (nucleotide numbering is as originally described and therefore does not include the 69 nucleotides of the alternatively spliced exon 31). No other sequence changes were identified at either the TSC1 or TSC2 gene loci in the index patient.

Subsequent analysis of exon 36 of the TSC2 gene in the rest of the family identified the 4684G>A substitution in individuals III-3, III-2, and IV-1. The TSC2 4684G>A mutation was also suspected in individual II-2 who was not available for testing, and possibly individual II-1 who died from renal bleeding, consistent with the presence of a renal angiomyolipoma. The TSC2 4684G>A substitution was excluded in individuals IV-2, III-4, and II-3.

To determine whether the TSC2 4684G>A substitution represented a simple polymorphism, we screened exon 36 in an additional 50 control DNAs isolated from individuals without TSC. None of the individuals tested was positive for the 4684G>A substitution. In addition, the 4684G>A substitution was not detected in >300 TSC patients tested in Rotterdam, Erlangen, and Martinsried and, as far as we are aware, has not been described elsewhere in the literature.

To exclude the possibility of an undetected TSC1 mutation in this family, linkage analysis was performed using four microsatellite markers, D9S1216, D9S1830, D9S1199, and D9S1198, flanking the TSC1 gene. Individuals III-2, III-3, IV-1, and IV-3 did not share a common haplotype (data not shown). In contrast, linkage analysis using the markers KG8, 16AC2.5, and SM7, flanking the TSC2 gene, revealed a common haplotype in all mutation carriers, but not in the unaffected individual IV-2 (fig 1). For individual II-2, the same haplotype as in the mutation carriers could be reconstructed, supporting the hypothesis that the G1556S mutation was inherited from the grandfather.

Functional analysis

Coimmunoprecipitation and Western blot experiments were performed on COS-1 cells transfected with a TSC2 expression construct containing the 4684G>A substitution. The cDNA used in these experiments lacks the 23 amino acids encoded by the alternatively spliced exon 31, and this tuberin variant is therefore referred to as G1556S. The functional significance of the amino acids encoded by exon 31 appears minimal. Previous work had indicated that the majority of TSC2 mRNA transcripts do not contain the sequence corresponding to exon 31 and that the function of tuberin is not affected by the presence or absence of this sequence in the assays used.
Expression of the G1556S tuberin variant

As shown in fig 2A, wild type tuberin migrates as a broad band on SDS-PAGE gels. This is due to the presence of differentially phosphorylated tuberin isoforms. The tuberin G1556S variant, however, migrated as a single high mobility band, consistent with a reduction in tuberin phosphorylation. Similar effects have been observed for other tuberin missense variants. Immunofluorescence microscopy indicated that there were no differences in the transfection efficiencies of the wild type compared with the G1556S tuberin constructs. Identical results were obtained using different preparations of both the wild type and G1556S variant expression constructs (data not shown).

As shown in fig 2B, coexpression of an active PKB isoform with the G1556S variant resulted in a shift to the low mobility, phosphorylated tuberin isoforms, similar to the wild type protein but distinct from another tuberin variant (R611Q) that is completely resistant to PKB dependent phosphorylation. Compared with either wild type tuberin or the R611Q variant, the changes in the mobility of the G1556S variant upon coexpression of PKB indicates that, although the G1556S variant is less susceptible to phosphorylation by PKB than wild type tuberin, it is not completely resistant to PKB, in contrast to the R611Q variant.

Phosphorylation of wild type tuberin and the G1556S variant by PKB was confirmed using an antibody specific for the phosphorylated PKB target sequence. The phosphorylation resistant, high mobility R611Q tuberin variant was not detected with the PKB target sequence antibody.
Interaction between hamartin and the tuberin G1556S variant

Hamartin and tuberin are products of the TSC1 and TSC2 genes, respectively. They form a complex that is involved in the regulation of cell growth. The G1556S variant of tuberin was found to interact with hamartin, and this complex formation was studied using coimmunoprecipitation experiments. The results showed that the G1556S variant reduced the stability of the tuberin-hamartin complex, as evidenced by a decrease in the amount of coimmunoprecipitated hamartin.

**Figure 2** Functional comparison of wild type tuberin with the G1556S variant. (A) Interaction with hamartin. Cotransfected COS-1 cells expressing hamartin and either wild type tuberin (wt) or the G1556S variant (G1556S) were lysed and the tuberin-hamartin complex was immunoprecipitated using antibodies specific for tuberin (Tuberin IP) and antibodies specific for hamartin (Hamartin IP). The amounts of (co)immunoprecipitated tuberin and hamartin were estimated by Western blotting. The examples shown are representative of at least three separate experiments. (B) PKB dependent phosphorylation. Cotransfected COS-1 cells expressing an activated isoform of PKB and either wild type tuberin (wt), the tuberin R611Q variant (R611Q), or the G1556S variant (G1556S) were lysed and analysed by Western blotting with an antibody specific for tuberin (lysate, above). Tuberin phosphorylated by PKB was detected by immunoprecipitation of tuberin, followed by incubation with the PKB target sequence antibody (Cell Signaling Technology) (Tuberin IP, below). The examples shown are representative of at least three separate experiments. (C) Inhibition of phosphorylation. TSC2 -/- MEFs were transfected with either the wild type TSC2 expression construct or the G1556S variant expression construct. After 24 hours the cells were serum starved for 24 hours prior to analysis by double label fluorescent microscopy. Transfected cells expressing wild type tuberin (or the G1556S variant) and either positive or negative for phosphorylated S6 were counted. The proportion of transfected cells without S6 phosphorylation is indicated.

Interaction between hamartin and the tuberin G1556S variant

Tuberin forms a complex with hamartin, the product of the TSC1 gene, and the formation of the complex stabilises both proteins. To investigate whether the tuberin G1556S variant was able to interact with hamartin, coimmunoprecipitation experiments were performed. Hamartin and the tuberin G1556S variant could be coimmunoprecipitated. However, as shown in fig 2A, using antibodies specific for tuberin less hamartin was coimmunoprecipitated with the G1556S variant than with wild type tuberin and, using antibodies specific for hamartin, compared with wild type tuberin less of the G1556S variant was coimmunoprecipitated. Therefore, although complex formation was not abrogated completely, the G1556S substitution reduced the stability of the tuberin-hamartin interaction.

Tuberin is necessary to maintain hamartin in a soluble form in the cytosol. The reduction in the amount of hamartin in the lysate of the cells expressing the G1556S tuberin variant is therefore consistent with the reduction in stability of the tuberin-hamartin complex caused by the G1556S substitution.
Inhibition of S6 phosphorylation by the tuberin G1556S variant

The tuberin-hamartin complex inhibits the activation of S6 kinase, thereby preventing phosphorylation of the ribosomal protein S6, and tuberin missense mutations have been shown to prevent this inhibition. The effect of the tuberin G1556S variant on S6 phosphorylation was investigated in mouse embryo fibroblasts (MEFs) lacking endogenous tuberin using an established assay. As shown in fig 2C, no S6 phosphorylation was visible in 70% of the cells expressing wild type tuberin. Co-expression of hamartin resulted in inhibition of S6 phosphorylation in almost 90% of the transfected cells. In contrast, the inhibition of S6 phosphorylation by the G1556S variant, either expressed alone or co-expressed with hamartin, was reduced approximately fourfold. Less than 20% of the cells expressing the G1556S variant showed inhibition of S6 phosphorylation.

DISCUSSION

We report on the identification of a novel TSC2 missense mutation, 4684G>A (G1556S), in a patient with the clinical diagnosis of TSC due to prenatal cardiac rhabdomyomas and the development of hypomelanotic macules in the first months of life. Subsequent analysis of the parents of the index patient and further family members to distinguish between a disease causing mutation and a non-pathogenic polymorphism identified three additional individuals with the 4684G>A substitution. Although only the index patient fulfilled the established criteria for the diagnosis of TSC, the other family members with the 4684G>A substitution all displayed minor signs of TSC, preferentially hypomelanotic macules (table 1). None of them had cortical tubera or calcifications and only one (IV-1) suffered from epileptic seizures suspected to result from a hippocampus sclerosis. These findings are so far consistent with the hypothesis that the 4684G>A substitution is indeed a pathogenic mutation. No other changes were identified in either the TSC2 gene or the TSC1 gene of the index patient. Analysis of microsatellite markers across both loci excluded linkage to TSC1, but confirmed identical haplotypes for TSC2 in all mutation carriers. The possibility of a second undetected mutation on the same allele as the 4684G>A substitution, although not excluded, seems unlikely.

The 4684G>A mutation leads to a change of glycine, a non-polar residue which is completely conserved between human, mouse, rat, Fugu and Drosophila (fig 3) for serine, a polar amino acid in the C-terminal domain of tuberin. Several regions with possible functional importance have been reported for tuberin: the hamartin binding region at the N-terminus, 21 the putative rap1GAP homology region with GAP activity for Rap1 22 and Rab5, 23 two transcriptional activating domains, 24 as well as binding regions for Rabaptin5, 24 steroid hormone receptors, 25 and Calmodulin. 26 These findings confirm the pathogenicity of TSC2 4684G>A and are consistent with the mild clinical phenotype in the mutation carriers in this family. The approach described here will help to define which domains and amino acid residues are critical for tuberin function and provide insight into the possibility that additional phenotype-genotype correlations exist in TSC. Moreover, the family presented here highlights the necessity for careful and complete clinical evaluation of both apparently affected and unaffected members in families with TSC.

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Figure 3 Evolutionary conservation of TSC2 exon 36. Identical amino acids are shaded in grey. The G1556 residue affected by the mutation is additionally highlighted in bold.
REFERENCES


