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# Germline *SMARCB1* mutation and somatic *NF2* mutations in familial multiple meningiomas

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## ABSTRACT

**Background** Multiple meningiomas occur in <10% of meningioma patients. Their development may be caused by the presence of a predisposing germline mutation in the neurofibromatosis type 2 (*NF2*) gene. The predisposing gene in patients with non-*NF2* associated multiple meningiomas remains to be identified. Recently, *SMARCB1* was reported to be a potential predisposing gene for multiple meningiomas in a family with schwannomatosis and multiple meningiomas. However, involvement of this gene in the development of the meningiomas was not demonstrated.

**Results** Five affected members of a large family with multiple meningiomas were investigated for the presence of mutations in *SMARCB1* and *NF2*. A missense mutation was identified in exon 2 of *SMARCB1* as the causative germline mutation predisposing to multiple meningiomas; furthermore, it was demonstrated that, in accordance with the two-hit hypothesis for tumourigenesis, the mutant allele was retained and the wild-type allele lost in all four investigated meningiomas. In addition, independent somatically acquired *NF2* mutations were identified in two meningiomas of one patient with concomitant losses of the wild-type *NF2* allele.

**Conclusion** It is concluded that, analogous to the genetic events in a subset of schwannomatosis associated schwannomas, a four-hit mechanism of tumour suppressor gene inactivation, involving *SMARCB1* and *NF2*, might be operative in familial multiple meningiomas associated meningiomas.

## INTRODUCTION

Meningiomas are among the most common intracranial primary tumours (20% of all brain tumours), usually occurring as slowly growing sporadic solitary lesions.<sup>1</sup> They also develop in about 50% of patients with neurofibromatosis type 2 (*NF2*). The hallmark of *NF2* is the presence of bilateral vestibular schwannomas. Schwannomas at other locations are found in more than 50% of patients.<sup>2–4</sup> *NF2* patients carry a heterozygous germline mutation in the *NF2* gene and the wild-type copy of this gene is often found to be lost in the *NF2* associated tumours.<sup>5–7</sup>

Multiple meningiomas occur in <10% of patients with meningioma.<sup>8</sup> These may be *NF2* associated and in those cases develop as the consequence of the presence of a predisposing *NF2* germline mutation. Non-*NF2* associated multiple meningiomas may occur as sporadic or familial cases. The limited number of meningiomas of familial

multiple meningioma patients investigated thus far did not display somatic or constitutional *NF2* mutations.<sup>9–10</sup> The multiple tumours of sporadic patients were found to carry an identical somatic *NF2* mutation, which was not present in their constitutional DNAs.<sup>9–13</sup> This might be explained by spread via the cerebrospinal fluid or distant metastasis of a single tumour or as a consequence of mosaicism with undetectable contribution of the *NF2* mutation to the constitutional DNA. Alternatively, the non-*NF2* sporadic and familial cases suggest that a gene different from *NF2* is implicated in the development of non-*NF2* associated multiple meningiomas. This contention is supported by an earlier linkage study in a family with meningiomas in which a 15 cM region around and including *NF2* was excluded from harbouring the meningioma predisposing gene.<sup>14</sup>

*SMARCB1* is a predisposing gene in schwannomatosis, which is characterised by the presence of multiple schwannomas in the absence of vestibular schwannomas. The gene is involved in about 30–40% of the familial cases, but in no more than 10% of the sporadic cases. In the schwannomas of these patients, the wild-type copy of *SMARCB1* is found to be inactivated, usually by deletion. In addition, somatically acquired *NF2* mutations have been found in schwannomas of patients with a germline *SMARCB1* mutation.<sup>15–18</sup>

We wondered whether *SMARCB1* might be a predisposing gene in multiple meningiomas as well. Support for this candidacy is given by the location of *SMARCB1*, which is proximal to marker D22S1, that is outside the region excluded by linkage analysis.<sup>14</sup> Furthermore, somatic *SMARCB1* mutations have been shown to occur in sporadic meningiomas, albeit at low frequency, that is <3% of tumours.<sup>19–21</sup> Evidence for the involvement of *SMARCB1* in multiple meningiomas was recently reported by Bacci *et al*,<sup>22</sup> who showed the inheritance of a germline *SMARCB1* mutation in a family with schwannomatosis and multiple meningiomas. However, the mutational status of *SMARCB1* (and of *NF2*) in the meningiomas of the patients was not clarified. In another recent study, no germline *SMARCB1* mutation could be detected in 47 patients with multiple meningiomas, including eight familial cases.<sup>23</sup>

In this study, we investigated a large family with three members affected by multiple meningiomas and two with a single meningioma. We identified the causative germline *SMARCB1* mutation segregating with (multiple) meningiomas in this family and demonstrated inactivation by deletion of the wild-type copy of *SMARCB1* and additional bi-allelic

## Original article

and independent inactivation of *NF2* in meningiomas of the patients.

### PATIENTS AND METHODS

#### Patients

Informed consent was obtained from all family members for using their body materials.

The proband (III-6 in figure 1) was referred to our outpatient clinic with a suspected diagnosis of NF2 because of multiple meningiomas. At the age of 34 years she presented with epilepsy. CT of the brain showed three extra-axial lesions along the falx cerebri, which were suspected of being meningiomas. Two lesions were resected and pathological examination confirmed the diagnosis of meningioma. At the age of 51 years, our proband developed sensory problems and paraplegia of the legs. MRI of the brain and spinal cord showed an intradural extramedullary lesion at thoracic level (T) 7, suspected of being a meningioma. The lesion was removed and turned out to be a psammomatous meningioma, WHO grade I. Her family history revealed a twin sister (III-7 in figure 1) with a chest wall schwannoma at age 37. At age 49, she developed epilepsy and brain MRI showed multiple extra-axial lesions, which were on histology transitional/mixed meningiomas grade I. She underwent additional surgery for a spinal fibroblastic meningioma at T4, multiple schwannomas of the chest wall and elbow, and was diagnosed with multiple spinal lesions suspected of being schwannomas. The father of these twin sisters died at age 63 from coronary artery disease. Two of his 10 siblings had been undergone surgery for meningioma. An uncle of our proband (II-6 in figure 1) developed left-sided deafness caused by a cerebellopontine angle mass, suspected of being a vestibular schwannoma, at age 56. At age 64, a spinal fibroblastic meningioma at T1 was removed; more recently, at age 77, he developed gait difficulties and MRI showed multiple spinal intradural extramedullary lesions, suspected of being meningiomas. An aunt of our proband (II-9 in

figure 1) underwent surgery on a spinal psammomatous meningioma at T10 at age 53. A fibrous meningioma at T6 was removed from the spinal cord of her daughter (III-1 in figure 1) at the age of 40 years.

None of the affected family members had neurofibromas, café-au-lait spots or freckling. Their clinical characteristics are summarised in table 1.

#### DNA and RNA samples

A frozen tissue sample of a brain meningioma M1 of patient III-6 and formalin fixed, paraffin embedded (FFPE) tissue samples of one meningioma each of patients II-6, II-9, and III-1, and the brain and spinal cord meningiomas (M1, M2) of patient III-6, were available for molecular analysis. DNA was extracted and purified from blood samples and the tumour tissue samples according to standard methods. DNA extracted from the meningioma of patient II-9 was too heavily degraded for reliable molecular analyses.

Total RNA was extracted and purified from another part of the frozen tissue sample of meningioma M1 of patient III-6 using TRIzol LS Reagent according to the protocol of the manufacturer (Invitrogen, Leek, The Netherlands). One microgram of total RNA was used to synthesise cDNA with random priming according to standard methods.

#### Microsatellite analysis

Genotyping of family members and loss of heterozygosity (LOH) analysis of tumours were performed by using the chromosome 22-specific microsatellite markers of the ABI Prism Linkage Mapping Set version 2.5-HD5 and additional marker D22S929. Primer sequences for the latter marker were taken from Legoux *et al.*<sup>24</sup> Data were analysed with Genemapper software version 4.0. (Applied Biosystems, Foster City, California, USA).

#### Mutation analysis

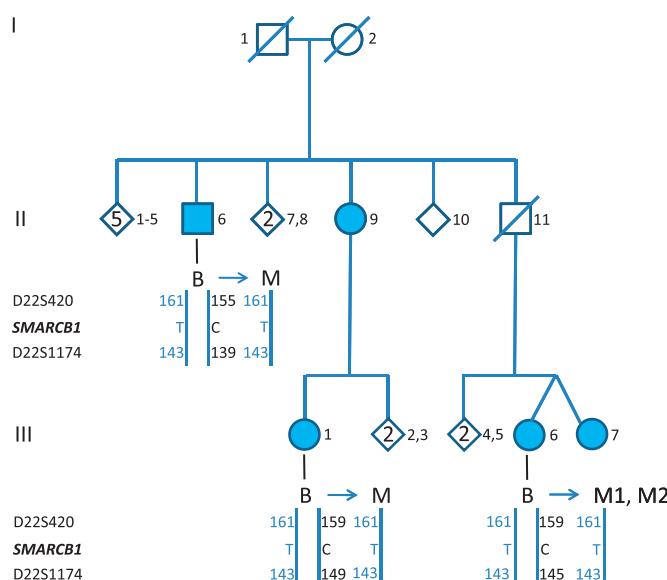
*SMARCB1* and *NF2* were sequenced by using genomic or cDNA as substrate for amplification by PCR. Primer sequences for mutation analysis of the *SMARCB1* and *NF2* exons have been published previously.<sup>15</sup> For improved performance, some *SMARCB1* primers were newly designed (sequences available on request). In degraded DNA extracted from FFPE material, the *SMARCB1* exon 2 segment harbouring the missense mutation was sequenced by using forward primer 5'-CCCTTATAATGAGCCTCTTGCT-3' and reverse primer 5'-TCTTCCACAGTGGCTAGTCG-3', generating a PCR product of 116 bp.

The cDNA product from meningioma M1 of patient III-6 was sequenced by using forward primer 5'-GACGGCGAGTTC-TACAT-3' in exon 1 and reverse primer 5'-TTTACCATGT-GACGATGCAA-3' at the 3'-end of exon 2, generating a PCR product of 148 bp. All sequence reactions were performed using ABI Big Dye v3.1 chemistry, and the products were sequenced with an ABI 3730 capillary system (Applied Biosystems). Sequences were analysed with CodonCode Aligner (CodonCode Corp, Dedham, Massachusetts, USA).

The HomoloGene program in PubMed was used to perform amino acid sequence alignments (<http://www.ncbi.nlm.nih.gov/pubmed>). The PolyPhen and SIFT programs were used to predict the impact of amino acid substitutions on protein function and structure (<http://genetics.bwh.harvard.edu/pph> and <http://sift.jcvi.org>, respectively).

## RESULTS

Sequencing of the nine exons of *SMARCB1* in constitutional DNA of proband III-6 revealed a missense mutation in exon 2,



**Figure 1** Genotyping of *SMARCB1* and flanking microsatellite markers in constitutional blood (B) and corresponding meningioma (M) DNA of affected members in a multiple meningiomas family. Patients with (multiple) meningiomas are marked by blue symbols. Diamond symbols indicate family members with insufficient clinical details (under clinical and molecular investigation). The wild-type and mutant allele of *SMARCB1* are indicated by C and T, respectively. Marker alleles are given by the length of their PCR products (bp).

**Table 1** Clinical characteristics of affected family members

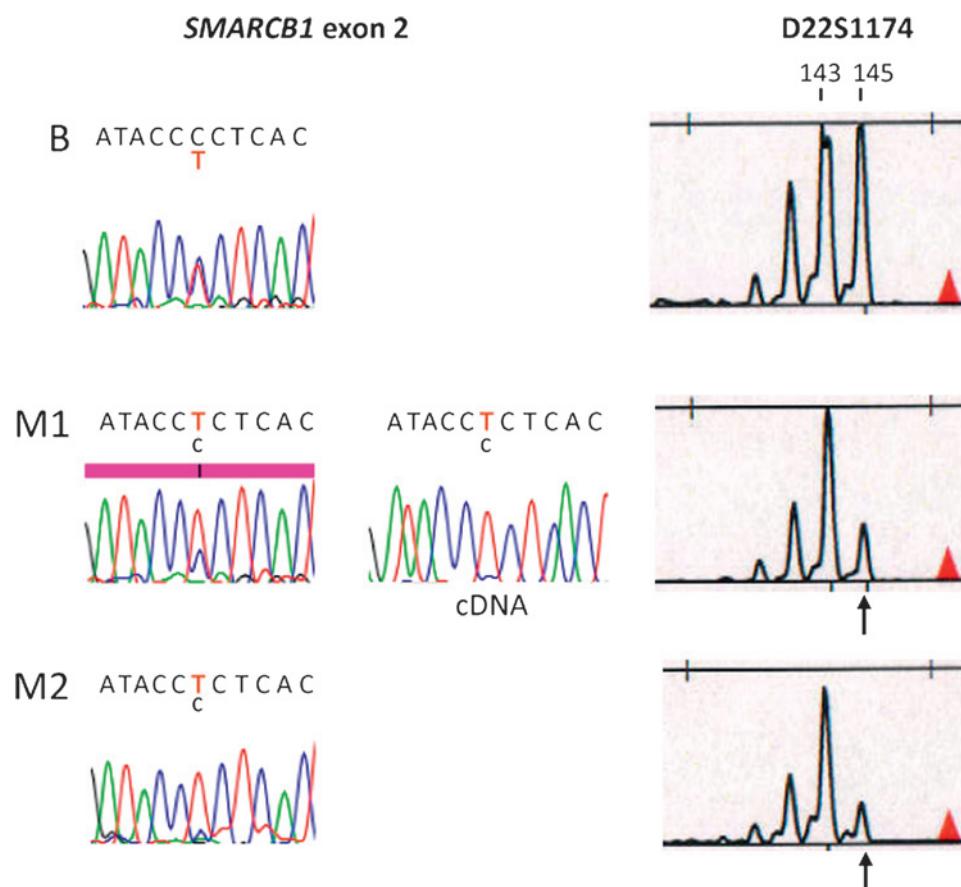
Patient	Sex (M/F)	Age (years) at diagnosis	Presenting symptom	Meningiomas (n, location)	Schwannomas (n, location)
III-6	F	34	Epilepsy	3 cranial, 1 spinal	—
III-7	F	37	Chest wall mass	Multiple cranial, 1 spinal	Multiple chest wall, multiple spinal, 1 elbow
II-6	M	56	Deafness	Multiple spinal	1 vestibular
II-9	F	53	Paresis legs	1 spinal	—
III-1	F	40	Sensory problems legs	1 spinal	—

c.143C→T, P48L (figure 2), which also proved to be present in constitutional DNA of the other affected family members (III-7, II-6, II-9, and III-1). The mutation was not found in constitutional DNA of more than 100 control individuals. The proline residue that is replaced by leucine in the resulting SMARCB1 protein is evolutionarily conserved across all species for which sequence information is available (data not shown). In silico analyses with the Polyphen and SIFT programs predicted that the amino acid substitution affects SMARCB1 protein function and is probably damaging.

The four meningiomas that were available for molecular analysis all showed loss of the wild-type C allele, as demonstrated in figure 2 for the brain (M1) and spinal cord (M2) meningioma of the proband (III-6). In accordance with this, genotyping with flanking markers revealed retention of the chromosome 22 segment containing the mutant T-allele in the investigated tumours, as exemplified in figure 2 for marker D22S1174. The data are summarised in figure 1.

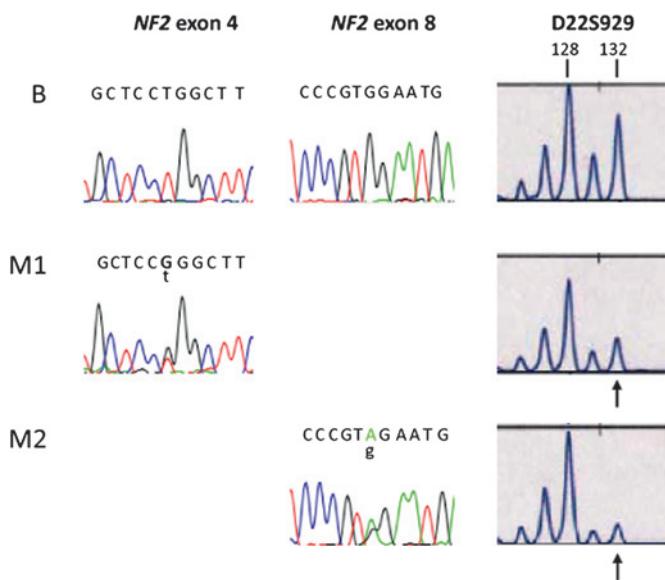
To investigate whether the retained SMARCB1 allele was transcribed, we extracted total RNA from frozen tissue of meningioma M1 and reverse transcribed it into cDNA. The cDNA sequence demonstrates that a stable mRNA is transcribed from the retained mutant allele (figure 2).

**Figure 2** Sequence analysis of part of exon 2 of SMARCB1 and loss of heterozygosity (LOH) analysis with marker D22S1174 in blood and meningioma DNAs of patient III-6. Left: presence of the mutant T allele in constitutional blood (B) DNA and considerable loss of the wild-type C allele in genomic and cDNA of meningioma M1 and in genomic DNA of meningioma M2. Right: presence of two alleles of marker D22S1174 in blood DNA and considerable loss of the 145 bp allele, marked by arrow, in meningioma M1 and M2 DNA.



Schwannomatosis associated schwannomas harbour somatically acquired *NF2* mutations. Considering the apparent analogy with schwannomatosis, we wondered whether these mutations might also be present in the meningiomas of the family under study. Although attempts were made to sequence *NF2* in all available meningiomas, reliable sequences could only be derived from the frozen tumour M1 and the freshly fixed tumour M2, both of proband III-6. We found a missense mutation in exon 4 of meningioma M1 (c.422T→G, p.L141R) and a nonsense mutation in exon 8 (c.773G→A, p.W258X) of meningioma M2, both being absent in the corresponding constitutional DNA of the patient (figure 3). The leucine residue at position 141 is evolutionarily conserved (data not shown). The Polyphen and SIFT programs predicted that the amino acid substitution caused by the missense mutation affects *NF2* protein function and is probably damaging. The sequence recordings indicate loss of the wild-type allele in both tumours. This is substantiated by LOH analysis with the intragenic *NF2* marker D22S929, which shows considerable loss of the 132 bp allele in tumour M1 as well as tumour M2 (figure 3). Finally, we sequenced all exons of the *NF2* gene in constitutional DNA of the affected family members (III-6, III-7, II-6, II-9, III-1), but did not detect any deviation from the wild-type sequence.

## Original article



**Figure 3** Sequence analysis of part of exons 4 and 8 of *NF2* and loss of heterozygosity (LOH) analysis with marker D22S929 in blood and meningioma of patient III-6. Left: presence of the mutant G allele in exon 4 in meningioma M1 DNA and the mutant A allele in exon 8 in meningioma M2 DNA, but absence of these mutant alleles in constitutional blood (B) DNA. Right: Presence of two alleles of marker D22S929 in blood DNA and considerable loss of the 132 bp allele, marked by arrow, in meningioma M1 and M2 DNA.

In addition, we performed multiple ligation dependent probe amplification (MLPA) analysis of the *NF2* gene in constitutional DNA of patient III-7. However, this analysis did not reveal any deletions or duplications.

## DISCUSSION

We identified the germline mutation c.143C→T in *SMARCB1* that most probably predisposes to the development of multiple meningiomas. The mutation segregates with the presence of (multiple) meningiomas in this family and, in accordance with Knudsons' two-hit hypothesis for tumourigenesis,<sup>25</sup> remains in the four investigated meningiomas, each with loss of the wild-type *SMARCB1* allele by LOH (figure 2). The causative nature of this missense mutation may be inferred from our in silico analyses, which demonstrated that the proline residue at position 48 is evolutionary conserved and that the change to leucine affects *SMARCB1* protein function and is probably damaging. The importance of proline at that position is furthermore supported by the finding of another missense mutation causing its replacement (c.142C→T, p.P48S) in a malignant rhabdoid tumour of the kidney.<sup>26</sup> Another missense mutation in the *SMARCB1* gene, c.92A→T, p.E31V, was recently identified as the predisposing genetic event in a family with schwannomatosis and multiple meningioma.<sup>22</sup> RNA studies in whole blood of two patients in this family demonstrated absence of transcripts from the mutant *SMARCB1* allele. We, however, detected a stable transcript derived from the mutant *SMARCB1* allele that was retained in meningioma M1 of patient III-6 (figure 2). Accordingly, we found uniform staining of nuclei with a *SMARCB1* antibody (anti-BAF47) in all four meningiomas, suggesting that an intact *SMARCB1* protein was formed (data not shown).

The presence of a left sided cerebellopontine angle mass, probably a vestibular schwannoma, in patient II-6 and of multiple schwannomas in patient III-7 suggest that the *SMARCB1* mutation in this family may predispose to the

development of schwannomas as well. However, the vestibular schwannoma was diagnosed at age 56 and such a sporadic unilateral tumour is not uncommon at that age in the general population.<sup>27</sup> Unfortunately, the schwannomas were not available to further investigate the possible involvement of *SMARCB1*. A schwannoma analysed in the family with schwannomatosis and multiple meningiomas indeed displayed bi-allelic inactivation of *SMARCB1*.<sup>22</sup> Two other schwannomatosis families, each with one patient developing one or two meningiomas, have been reported. However, the status of *SMARCB1* in these tumours was not investigated.<sup>16 17 28 29</sup> Considering the frequent occurrence of meningiomas in the general population,<sup>1</sup> it cannot be excluded that their association with schwannomatosis in the latter families was coincidental. On the other hand, the missense mutation in *SMARCB1* in our family, affecting amino acid residue 48, and most probably also in the family presented by Bacci *et al*,<sup>22</sup> affecting nearby amino acid residue 31, clearly predispose to the development of multiple meningiomas. Therefore, it is possible that the preference for meningioma development is, at least in part, determined by type and location of the *SMARCB1* mutation. Alternatively, the co-occurrence of schwannomas and meningiomas in some *SMARCB1* families and the sole occurrence of schwannomas in most other *SMARCB1* families may point to the existence of modifying genes segregating in these families and affecting the phenotypic expression of the *SMARCB1* mutation. Our finding, and that of Bacci *et al*,<sup>22</sup> that meningiomas are part of the schwannomatosis tumour spectrum may require the diagnostic criteria for this disease be reconsidered. Further investigation of the possible association of type and location of the *SMARCB1* mutation and the development of multiple meningiomas and of possible modifying factors will require the identification and clinical evaluation of additional carriers of the mutated gene in our family and in other multiple meningiomas patients and families.

We identified additional independent somatic mutations in the *NF2* gene in the brain (M1) and spinal cord (M2) meningiomas of patient III-6 (figure 3). The missense mutation in exon 4 in meningioma M1 caused replacement of the evolutionary conserved leucine residue at position 141 in the *NF2* protein and its change into arginine affects, according to the in silico analyses, protein function and is probably damaging. The importance of leucine at position 141 is furthermore supported by the finding of another missense mutation causing its replacement (c.422T→C, p.L141P), which was reported as a constitutional *NF2* mutation in a patient with bilateral schwannomas.<sup>30</sup> The exon 8 mutation in meningioma M2 is a typical *NF2* nonsense mutation, causing truncation of the protein.<sup>30</sup> In accordance with Knudsons' two-hit model, loss of the wild-type copy of *NF2* by LOH occurred in both tumours.<sup>25</sup> The finding of two causative hits (*NF2* mutation and deletion) in these meningiomas and our failure to detect a constitutional *NF2* mutation or deletion in the affected family members strongly argue against this gene as being the multiple meningioma predisposing gene in this family.

Summarising our data, we conclude that, analogous to the genetic events in a subset of schwannomatosis associated schwannomas,<sup>16–18</sup> a four-hit mechanism of tumour suppressor gene inactivation, involving *SMARCB1* and *NF2*, might also be operative in familial multiple meningiomas associated meningiomas. In a recent study, no germline *SMARCB1* mutations were detected in 47 patients with multiple meningiomas, including eight familial cases.<sup>23</sup> Germline *SMARCB1* mutations are infrequently found in sporadic schwannomatosis patients

(<10% of cases). It is possible that, as in familial schwannomatosis (30–40% of cases), *SMARCB1* involvement is more frequent in familial multiple meningiomas. Alternatively, other genes might be implicated as predisposing genetic factors in both diseases.

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**Competing interests** None.

**Patient consent** Obtained.

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## Germline *SMARCB1* mutation and somatic *NF2* mutations in familial multiple meningiomas

I Christiaans, S B Kenter, H C Brink, et al.

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