Germline SMARCB1 mutation and somatic NF2 mutations in familial multiple meningiomas
Christiaans, I.; Kenter, S.B.; Brink, H.C.; van Os, T.A.M.; Baas, F.; van den Munckhof, P.; Kidd, A.M.J.; Hulsebos, T.J.M.

Published in:
Journal of Medical Genetics

DOI:
10.1136/jmg.2010.082420

Link to publication

Citation for published version (APA):

General rights
It is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), other than for strictly personal, individual use, unless the work is under an open content license (like Creative Commons).

Disclaimer/Complaints regulations
If you believe that digital publication of certain material infringes any of your rights or (privacy) interests, please let the Library know, stating your reasons. In case of a legitimate complaint, the Library will make the material inaccessible and/or remove it from the website. Please Ask the Library: http://uba.uva.nl/en/contact, or a letter to: Library of the University of Amsterdam, Secretariat, Singel 425, 1012 WP Amsterdam, The Netherlands. You will be contacted as soon as possible.
Germline SMARCB1 mutation and somatic NF2 mutations in familial multiple meningiomas

I Christiaans,1 S B Kenter,2 H C Brink,1 T A M van Os,1 F Baas,2 P van den Munckhof,3 A M J Kidd,4 T J M Hulsebos2

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.1 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.2–4 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.5–7

Multiple meningiomas occur in <10% of patients with meningioma.5 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.9-10 The multiple tumours of sporadic patients were found to carry an identical somatic NF2 mutation, which was not present in their constitutional DNAs.9-10 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.11

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.15-18

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.14 Furthermore, somatic SMARCB1 mutations have been shown to occur in sporadic meningiomas, albeit at low frequency, that is <5% of tumours.19–21 Evidence for the involvement of SMARCB1 in multiple meningiomas was recently reported by Bacci et al.,22 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.23

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
and independent inactivation of \textit{NF2} in meningiomas of the patients.

\textbf{PATIENTS AND METHODS}

\textbf{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 54 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 57. 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 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.

\textbf{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.

\textbf{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-HDS and additional marker D22S292. Primer sequences for the latter marker were taken from Legoix \textit{et al.} \cite{24}. Data were analysed with Genemapper software version 4.0. (Applied Biosystems, Foster City, California, USA).

\textbf{Mutation analysis}

\textit{SMARCB1} and \textit{NF2} were sequenced by using genomic or cDNA as substrate for amplification by PCR. Primer sequences for mutation analysis of the \textit{SMARCB1} and \textit{NF2} exons have been published previously. \cite{15} For improved performance, some \textit{SMARCB1} primers were newly designed (sequences available on request). In degraded DNA extracted from FFPE material, the \textit{SMARCB1} exon 2 segment harbouring the missense mutation was sequenced by using forward primer 5'-CCCTTATAATGAGCCTTCTTGCT-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'-TTTACCATGTT-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).

\textbf{RESULTS}

Sequencing of the nine exons of \textit{SMARCB1} in constitutional DNA of proband III-6 revealed a missense mutation in exon 2,
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).

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.

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

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.
meningiomas. The mutation segregates with the presence of an intact SMARCB1 protein was formed (data not shown).

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.145C→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 Knudson’s two-hit hypothesis for tumourigenesis, 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.E31V), which was reported as a constitutional NF2 mutation (c.422T→C, p.L141P), which was reported as a constitutional NF2 mutation in a patient with bilateral schwannomas. The exon 8 mutation in meningioma M2 is a typical NF2 nonsense mutation, causing truncation of the protein. In accordance with Knudson’s two-hit model, loss of the wild-type copy of NF2 by LOH occurred in both tumours. 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, 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. 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.

Acknowledgements We thank the family members for their participation in this study. We are indebted to Dr D Troost (Neuropathology Department, Academic Medical Center, Amsterdam), Dr M G Havenith (Pathology Department, Isala Clinics, Zwolle), D Stomphorst (Pathology Department, Leiden University Medical Center, Leiden), and M Waajar (Pathology Department, Medical Center Alkmaar, Alkmaar) for providing tumour materials.

Competing interests None.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES

Germline SMARCB1 mutation and somatic NF2 mutations in familial multiple meningiomas

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

J Med Genet 2011 48: 93-97 originally published online October 7, 2010
doi: 10.1136/jmg.2010.082420

Updated information and services can be found at:
http://jmg.bmj.com/content/48/2/93.full.html

References

This article cites 30 articles, 12 of which can be accessed free at:
http://jmg.bmj.com/content/48/2/93.full.html#ref-list-1

Article cited in:
http://jmg.bmj.com/content/48/2/93.full.html#related-urls

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections

Articles on similar topics can be found in the following collections

CNS cancer (37 articles)
Neurooncology (69 articles)
Molecular genetics (1086 articles)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/