CDKN2A mutations in Spanish cutaneous malignant melanoma families and patients with multiple melanomas and other neoplasia

Anna Ruiz, Susana Puig, Josep Malvehy, Conxi Lázaro, Michael Lynch, Anna M Gimenez-Arnau, Lluís Puig, Julian Sánchez-Conejo, Xavier Estivill, Teresa Castel

Abstract

The CDKN2A gene has been implicated in cutaneous malignant melanoma (CMM) in about 40% of families with linkage to chromosome 9p21, while a small proportion of families have mutations in the CDK4 gene. In order to estimate the importance of these genes in the predisposition to CMM in Spanish families and patients we have analysed, by SSCP analysis, a total of 56 subjects belonging to 34 CMM families, and nine patients with multiple CMM and other neoplasia. We have detected germline CDKN2A mutations in six out of the 34 families (17%). A frameshift mutation (358delG) and four missense mutations (G59V, G101W (two cases), D84Y, and R87W) were identified. Five CMM patients from different families (14%) carried the A148T variant, which is known not to affect p16 activity. No mutations were detected in the patients with multiple CMM or other neoplasms. We have not found mutations either in exon 1 of the CDKN2A gene or in exon 2A of CDK4. Linkage analysis of the 9p21 region showed exclusion for one of the families for CMM and for four families for CMM/dysplastic naevi. This study indicates a small role for CDKN2A in Spanish CMM families and suggests that other genes are also responsible for CMM predisposition.

Keywords: cutaneous malignant melanoma; CDKN2A; p16; CDK4

Cutaneous malignant melanoma (CMM) is one of the most serious forms of skin cancer and has a heterogeneous and complex aetiology. Sun exposure and genetic susceptibility contribute to melanoma risk. The CDKN2A gene has been implicated in CMM in 15% of such patients.

The CDKN2A gene also encodes an alternative transcript known as p19ARF which is derived from a different promoter and has a unique exon 1 (exon1b), but the same exon 2 as CDKN2A in a different codon reading frame. p19ARF has been shown to inhibit the cell cycle in a p53 dependent manner not involving CDK4/6 binding. It binds to MDM2 and promotes its degradation, leading to p53 stabilisation, accumulation, and cell cycle arrest.

The CDKN2A gene has been implicated as the melanoma susceptibility gene in some families with linkage to 9p21. CDKN2A mutations detected in families impair p16 function by decreasing its ability to bind to CDK4. In spite of this, no mutations have been found in other families which also show linkage to 9p21, suggesting that either these families have mutations that lie outside the coding region of the CDKN2A gene or that there is another susceptibility gene on 9p. The CDK4 gene has been found to be mutated in two out of 31 melanoma families indicating that CDK4 is also a melanoma predisposing gene. Both families carried the same missense mutation, R24C, which makes CDK4 resistant to the inhibition of p16. Another missense mutation located at the same position has also been detected in one French melanoma family. Patients with multiple primary melanomas, but without family history of melanoma, may also have a genetic predisposition for the disease. In fact, Monzon et al found germline mutations in the CDKN2A gene in 15% of such patients.

A total of 56 subjects belonging to 34 CMM families and nine patients with multiple CMM and other neoplasias were screened for mutations in the CDKN2A and the CDK4 genes by SSCP analysis and subsequent sequencing of abnormal bands. Patients with multiple melanomas and cases with CMM and other neoplasia could have susceptibility mutations in the CDKN2A or CDK4 genes. In table 1 we present a summary of families and patients analysed for mutations. The number of affected family members and association with...
CDKN2A mutations in malignant melanoma

†Pancreas adenocarcinoma, bladder carcinoma, mesothelioma and retinoblastoma.
*Rates correspond to families or to sporadic cases.

AFM, a\textsuperscript{d}encc of other malignancies.\textsuperscript{16} A missense mutation (176T\textrightarrow\textit{G}) was detected in a patient with CMM associated with DN and a high incidence of other malignancies.\textsuperscript{16} A missense mutation (R87P) has been described previously in familial melanoma.\textsuperscript{17} Although no functional studies have been performed, this substitution is located at the second ankyrin repeat. Ankyrin repeats are protein binding motifs which are conserved among the human INK4 family of proteins and are predicted to be important for their function.\textsuperscript{17} Thus, V59G is likely to be important in this family's susceptibility to CMM.

In two other unrelated families, the previously described G101W (301G\textrightarrow\textit{T}) mutation was detected. This mutation has been shown to have a very severe effect, the mutant protein having no capacity to inhibit the cyclin D1/CDK4 complex,\textsuperscript{10} and has been reported several times, suggesting a mutational hot spot or a common founder effect. In Italian families with the G101W mutation, a common founder effect has been shown for eight of 16 families with this mutation.\textsuperscript{12} Haplotype analysis in our two families has shown that they do not share the same haplotypes at the three markers used in this study, but they share a common allele for D9S162. All the melanoma patients in these two families carry the mutation.

We detected two missense mutations that have not been described previously. Both are located at the third ankyrin repeat. One mutation changes aspartic acid to tyrosine (D84Y) at nucleotide 250 (250\textrightarrow\textit{T}) and the other arginine to tryptophan (R87W) at nucleotide 259 (259\textrightarrow\textit{C}). At position 87 another missense mutation (R87P) has been described in familial melanoma and proven to fail to inhibit the cyclin complex.\textsuperscript{9}

<table>
<thead>
<tr>
<th>Family</th>
<th>Mutation</th>
<th>Patients with CMM</th>
<th>Patients with CMM and DN</th>
<th>Patients with DN</th>
<th>Mean age at first diagnosis</th>
<th>CDKN2A mutations (%)</th>
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<tbody>
<tr>
<td>29</td>
<td>358delG</td>
<td>4</td>
<td>4</td>
<td>1</td>
<td>35.7</td>
<td>6/34 (17)</td>
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<td>32</td>
<td>V59G</td>
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<td>1</td>
<td>1</td>
<td>1</td>
<td>Yes</td>
</tr>
<tr>
<td>40</td>
<td>G101W</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>1</td>
<td>Yes</td>
</tr>
<tr>
<td>42</td>
<td>R87W</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>No</td>
</tr>
<tr>
<td>43</td>
<td>D84Y</td>
<td>3</td>
<td>1</td>
<td>2</td>
<td>1</td>
<td>Unknown</td>
</tr>
<tr>
<td>47</td>
<td>G101W</td>
<td>3</td>
<td>3</td>
<td>0</td>
<td>0</td>
<td>No</td>
</tr>
</tbody>
</table>
Five melanoma patients belonging to five different families carried the A148T variant, which is a well-defined polymorphism that does not affect p16 activity. No other CDKN2A variants were detected in the other 40 subjects. We have not found mutations either in exon 1β of the CDKN2A gene or in exon 2A of the CDK4 gene.

It has been estimated that about 50% of the melanoma families linked to human chromosome 9p have mutations at the CDKN2A gene. We have studied 34 families with suspected susceptibility to melanoma and nine subjects affected with multiple melanosomes or melanoma associated with another type of cancer. Six families (17%) carried mutations in the CDKN2A gene. This low prevalence of CDKN2A mutations in our families has been observed previously. It is possible that some patients had mutations outside the coding regions or have inactivation by methylation of the CDKN2A gene. In contrast, a higher prevalence of mutations has been found in families where linkage to 9p21 has been obtained or where stringent criteria in defining families (based on the number of affected subjects) has been used. Most of the families studied here were small and some could represent chance clustering of sporadic cases. Only nine families had at least three affected family members and three of these carry mutations in CDKN2A (33.3%). In two of these families (20, 47), segregation of melanoma with the mutation has been shown, while in the third family (43) DNA from the other two members affected with CMM was not available (table 2). Soufir et al. reported 87% of families with mutations in the CDKN2A gene when one member of the family has multiple melanomas. In contrast, Monzon et al. reported that 15% of patients with multiple melanomas, but without a family history of the disease, have mutations in CDKN2A, although in some cases a family history of melanoma was shown afterwards. In agreement with this, we found two mutations in two patients with multiple melanomas and discovered later that other members of the family were also affected with melanoma. In our study, 43% of the families, where one of the patients had more than one primary melanoma, harboured CDKN2A mutations, whereas mutations were not detected in patients with multiple melanomas and no family history of the disease. This would indicate that multiple primary melanomas with a family history of at least three affected family members are suggestive of mutations in the CDKN2A gene. The CDKN2A gene is probably a high penetrance melanoma susceptibility gene, but other genes at 9p21 or at other locations should account for the rest of the families.

While the CDKN2A mutations described here segregate with the CMM phenotype, DN segregates independently of these mutations, indicating that the DN syndrome may segregate with other genes in these families. Linkage analysis with 9p21 markers in nine families showed exclusion for one of them, when CMM was considered the affected phenotype (table 3), while the other eight families were not informative, giving a total lod score of 0.722. When DN/CMM was considered as the affected phenotype four families showed exclusion for the D9S162 marker with a total lod score of −22.696 (data not shown), indicating that either CMM/DN is not located here or that DN and CMM are genetically distinct entities. The other families were uninformative for CMM/DN owing to the low number of affected patients from whom samples were obtained.

The alternative transcript from the CDKN2A gene, termed p19ARF, has been shown to inhibit the cell cycle progression, the amino-terminal domain of the p19ARF protein which is encoded by exon 1β being sufficient for its function. Interestingly, mutations affecting exon 2 of CDKN2A, which also affect the p19ARF protein, have been shown not to affect its ability to inhibit the cell cycle progression.

We have not found mutations in exon 1β of the CDKN2A gene, concluding that this gene is not responsible for melanoma susceptibility in our families. Supporting this finding, other groups have not found mutations in the p19ARF coding region in 10 melanoma families with linkage at 9p21 or in 64 other melanoma kindreds. No mutations were detected in exon 2A of CDK4, which has been implicated in three melanoma families, highlighting it as a melanoma susceptibility gene. Other groups have also not found the CDK4 R24C mutation in their families. This gene is probably implicated in few melanoma families, not being a major contributor to the melanoma susceptibility in our families.

In summary, we have studied a total of 56 subjects belonging to 34 CMM families with or without DN, or patients with multiple CMM with or without other neoplasms and DN. Mutations in the CDKN2A gene were detected in six of 34 families (17%). No mutations were detected in the other patients or families and no mutations were found in the p19ARF and CDK4 genes. This study indicates the small role of CDKN2A in Spanish families and that other genes are also responsible for CMM predisposition.

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