IMPORTANCE The protection of telomeres 1 protein (POT1) is a critical component of the shelterin complex, a multiple-protein machine that regulates telomere length and protects telomere ends. Germline variants in \textit{POT1} have been linked to familial melanoma, and somatic mutations are associated with a range of cancers including cutaneous T-cell lymphoma (CTCL).

OBJECTIVE To characterize pathogenic variation in \textit{POT1} in families with melanoma to inform clinical management.

DESIGN, SETTING, AND PARTICIPANTS In this case study and pedigree evaluation, analysis of the pedigree of 1 patient with melanoma revealed a novel germline \textit{POT1} variant (p.I78T, c.233T>C, chromosome 7, g.124870933A>G, GRCh38) that was subsequently found in 2 other pedigrees obtained from the GenoMEL Consortium.

MAIN OUTCOMES AND MEASURES (1) Identification of the \textit{POT1} p.I78T variant; (2) evaluation of the clinical features and characteristics of patients with this variant; (3) analysis of 3 pedigrees; (4) genomewide single-nucleotide polymorphism genotyping of germline DNA; and (5) a somatic genetic analysis of available nevi and 1 melanoma lesion.

RESULTS The \textit{POT1} p.I78T variant was found in 3 melanoma pedigrees, all of persons who self-reported as being of Jewish descent, and was shown to disrupt POT1-telomere binding. A UV mutation signature was associated with nevus and melanoma formation in \textit{POT1} variant carriers, and somatic mutations in driver genes such as \textit{BRAF}, \textit{NRAS}, and \textit{KIT} were associated with lesion development in these patients.

CONCLUSIONS AND RELEVANCE \textit{POT1} p.I78T is a newly identified, likely pathogenic, variant meritng screening for in families with melanoma after more common predisposition genes such as \textit{CDKN2A} have been excluded. It could also be included as part of gene panel testing.
About 10% of patients with melanoma have a family history of the disease. The major melanoma predisposition genes are CDKN2A, CDK4, and BAP1. Extremely rare germline variants in the TERT promoter have also been reported. More recently, components of the shelterin complex, including in the Protection of Telomeres 1 (POT1) gene, ACD, and TERF2IP, have been implicated. Pathogenic POT1 variants promote telomere lengthening and disrupt telomere stability. Importantly, these variants are rare and potentially contribute to 1% to 5% of familial melanoma cases. Pathogenic POT1 variants have also been associated with predisposition to chronic lymphocytic leukemia (CLL), cardiac angiosarcoma, glioma, and, more recently, colon cancer.

Methods

Institutional review board approval for this study was obtained from the University of Leeds, University of Michigan, Hospital Clinic de Barcelona, and Gustave Roussy. All patients included in this study signed informed consent. Exome capture was performed using the SureSelect XT Human All-Exon V5 platform (Agilent Technologies Inc), and sequencing was performed using the Illumina HiSeq 2000 platform (Illumina Inc). Telomere binding and length assays were performed as previously described. For analysis methods, see the Supplement.

Results

Our analysis began with the identification of a North American proband who had developed primary melanoma at age 31 years. The primary melanoma on the neck was from the proband’s father, not the proband, but is added here to illustrate its position. This melanoma progressed to metastatic disease in the father. All numbered lesions were sequenced. Samples numbered 550/551, 554/555, and 609/610 indicate 2 samples taken from the same lesion in these 3 cases.

Key Points

Question What is the association of the p.I78T variant in the Protection of Telomeres 1 gene (POT1) with familial melanoma?

Findings We identified the POT1 p.I78T variant in 3 families of Jewish heritage with melanoma in the family and provide evidence that p.I78T interferes with POT1-telomere binding, a critical disruption associated with tumor predisposition. These findings suggest that the POT1 p.I78T variant may represent a founder allele.

Meaning The identification of high-penetrance germline variants, such as POT1 p.I78T, facilitates screening and counseling of at-risk patients.

In this proband and all 3 family pedigrees, the POT1 p.I78T variant was identified as the likely pathogenic variant; all 3 pedigrees show the same base change. A, Locations of lesions identified and excised from the proband, a POT1 variant carrier (p.I78T, c.233T>C, chromosome 7, g.124870933A>G, GRCH38) at age 31 years. The primary melanoma on the neck was from the proband’s father, not the proband, but is added here to illustrate its position. This melanoma progressed to metastatic disease in the father. All numbered lesions were sequenced. Samples numbered 550/551, 554/555, and 609/610 indicate 2 samples taken from the same lesion in these 3 cases. B, Pedigree structures for all families identified to carry the p.I78T variant. Pedigree 1 corresponds to that of the proband, and pedigrees 2 and 3 were identified from the GenoMEL Consortium. Individuals represented by black shapes are variant carriers who presented with melanoma. Individuals with no genotype were not tested. Ages, reported in years, at the times of cancer diagnoses are shown in parentheses when available.
31 years along with multiple dysplastic and compound and dermal nevi (Figure 1A). He had also developed cutaneous T-cell lymphoma (CTCL) and CLL and at ages 29 and 40 years, respectively. His father had developed CTCL, CLL, melanoma in situ, and melanoma at ages 58, early 60s, 72, and 74 years, respectively (Figure 1B, pedigree 1). The proband had sequenced his own genome on a next-generation platform and had identified a POT1 variant (p.I78T, c.237T>C, chromosome 7, g.124870933A>G, GRch38), the pathogenicity of which was unknown, but it was predicted to be damaging by SIFT (score 0)9 and Polyphen-2 (score 1.0)10 analysis. To determine the frequency of this variant in melanoma families, we ascertained cases by consulting The Melanoma Genetics Consortium (GenoMEL, http://www.genomel.org) and found 2 more carrier families; one of Spanish ancestry (1 case among 171 families) (Figure 1B, pedigree 1; more detail on the Spanish pedigree will be published elsewhere) and the other of French descent (1 case among 157 families) (Figure 1B, pedigree 3), both of whom had tested negative for other high-penetrance genes (eMethods in the Supplement).

We next investigated whether the p.I78T variant had a functional effect. An analysis of orthologous protein sequences spanning 450 million years of evolutionary distance showed complete conservation of the isoleucine at position 78 (Figure 2A), and the protein model showed that this residue is in close proximity to arginine 137, another position found to be mutated in melanoma pedigrees (Figure 2B).6 This suggested that alteration of this residue might impair POT1 single-stranded DNA binding. To test this, we generated a POT1 complementary DNA (cDNA) construct encoding the POT1 p.I78T protein and found that it was severely impaired for its ability to bind to a polynucleotide telomere-like probe (IR700-GGTTAGGGTTAGGGTTAGGG) (Figure 2C), as previously seen with established melanoma-predisposing alleles.5 To provide further functional proof, we transduced...
telomerase-positive HT1080 cells with either wild-type or p.178T POT1 cDNA constructs and cultured them through 45 population doublings. Telomeres in cells expressing the p.178T POT1 cDNA were elongated compared with cells expressing wild-type POT1 (eMethods and eFigure 1 in the Supplement).

After identifying 3 apparently unrelated families living in different countries carrying the POT1 p.178T variant, we genotyped individuals using CoreExome single-nucleotide polymorphism arrays (Illumina Inc) to estimate the coefficient of relatedness between carriers of the variant from different families and performed an analysis that suggested no recent co-ancestry between the families. We were also able to confirm the reported genetic relationships within each family. Although the families were not closely related, the sharing of genotypes around the POT1 gene in p.178T variant carriers was consistent with a common haplotype across some 4 megabase pairs (Mb) upstream and downstream of the variant, suggesting that the p.178T variant is a founder event.

Interestingly, all 3 families self-reported being Jewish, and we noted that the chromosome 7 g.124870933A>G change has been reported in 3 individuals of Ashkenazi Jewish descent in the Genome Aggregation Database (gnomAD); chromosome 7 g.124510987A>G in GRCh37; allele frequency in the Ashkenazi Jewish population, 0.0003105; in all other populations, 0.000004305. Since patients with germline POT1 variants are highly predisposed to melanoma development, we wondered if there would be any defining features of the somatic landscape of melanocytic lesions in these patients. Thus, we whole-exome sequenced 30 previously excised, formalin-fixed, paraffin-embedded nevi from the proband in pedigree 1 (Figure 1B) revealed that there were no defining features of the somatic landscape of melanocytic lesions in these patients. Thus, we whole-exome sequenced 30 previously excised, formalin-fixed, paraffin-embedded nevi from the proband in pedigree 1 (Figure 1) and a large malignant melanoma from the proband’s father (anatomic location illustrated in Figure 1A; eMethods in the Supplement). Hotspot NRAS (Q61L, Q61K and Q61R) and BRAF (V600E, V600D) mutations were identified in nevi, and the melanoma was found to be KIT L576P positive and to carry truncating NF1 and RASA2 mutations (eTable 1 in the Supplement), suggesting a prominent role for the MAPK pathway. No TERT promoter mutations were found.

Since a major driver of melanoma formation is exposure to UV light, we next attempted to identify the imprint of this mutagen on the melanoma genome. This analysis revealed an average of 40.67 somatically acquired mutations per Mb of exome, of which 87% were C>T/G>A mutations. Extraction of mutation signatures from mutations per Mb of exome, of which 87% were found an average of 40.67 somatically acquired mutations. The number of mutations we identified in the father’s melanoma was found to be enriched for C>T mutations (Figure 3; eTable 2 in the Supplement). All of the carriers in pedigrees 2 and 3 (Figure 1B) who had presented with melanoma had been diagnosed with either superficial spreading or nodular cutaneous melanoma. A histopathological analysis of the melanoma from the p.178T variant carrier (father, pedigree 1, Figure 1B) revealed that there were no defining features distinguishable from sporadic melanoma (eFigure 2 in the Supplement).

Discussion

The identification of pathogenic germline variants is of critical importance because it informs genetic counselling and allows individuals at high risk to be prioritized for surveillance programs. In this study, we identified the POT1 p.178T variant in 3 unrelated melanoma pedigrees, with evidence for the presence of this allele at a higher frequency among Ashkenazi Jews. Interestingly, only 1 founder allele (p.V59G; CDKN2A) has previously been associated with melanoma risk in people of Jewish heritage. Notably, as illustrated in Figure 1B, in addition to melanoma, members of pedigree 1 have been diagnosed with both CTCL and CLL, while pedigree 2 contains a variant carrier who developed melanoma and thyroid cancer. Collectively, this clinical picture may suggest that this variant contributes to a range of cancers. Indeed, somatic POT1 p.178T mutations have been found in a case of ovarian cancer (COSMIC ID: COSM16403) and in a case of hepatosplenic T-cell lymphoma.
Limitations

Our study ascertained cases from GenoMEL, meaning that we may not have a full picture of the spectrum of cancers acquired by variant carriers. Additionally, while 4 of the 8 variant carriers illustrated in Figure 1B have developed melanoma, 4 variant carriers in pedigree 2 have not yet presented with melanoma or any other malignancy, and these individuals were 19, 25, 50, and 55 years old, respectively, at last follow-up. Thus, the penetrance of melanoma and other cancers in p.I78T variant carriers is yet to be defined. Ultimately the identification of further pedigrees with the p.I78T variant will help define its role in tumorigenesis.

ARTICLE INFORMATION

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