Germline microsatellite genotypes differentiate children with medulloblastoma

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Abstract

Background. The germline genetic events underpinning medulloblastoma (MB) initiation, and therefore the ability to determine who is at risk, are still unknown for the majority of cases. Microsatellites are short repeated sequences that make up ~3% of the genome. Repeat lengths vary among individuals and are often nonrandomly associated with disease, including several cancers such as breast, glioma, lung, and ovarian. Due to their effects on gene function, they have been called the “tuning knobs of the genome.”

Methods. We have developed a novel approach for identifying a microsatellite-based signature to differentiate MB patients from controls using germline DNA.

Results. Analyzing germline whole exome sequencing data from a training set of 120 MB subjects and 425 controls, we identified 139 individual microsatellite loci whose genotypes differ significantly between the groups. Using a genetic algorithm, we identified a subset of 43 microsatellites that distinguish MB subjects from controls with a sensitivity and specificity of 92% and 88%, respectively. This microsatellite signature was validated in an independent dataset consisting of 102 subjects and 428 controls, with comparable sensitivity and specificity of 95% and 90%, respectively. Analysis of the allele genotypes of those 139 informative loci demonstrates that their association with MB is a consequence of individual microsatellites’ genotypes rather than their hypermutability. Finally, an analysis of the genes harboring these microsatellite loci reveals cellular functions important for tumorigenesis.

Conclusion. This study demonstrates that MB-specific germline microsatellite variations mark those at risk for MB development and suggests mechanisms of predisposition.

Key Points

1. Microsatellites in germline DNA differentiate children with medulloblastoma.
2. Children with medulloblastoma do not have constitutional microsatellite instability.
3. Genes harboring the tumor associated microsatellites are cancer associated.

Medulloblastoma (MB) is the most common malignant childhood brain tumor. Extensive genomic characterization has divided MB tumors into 4 molecular subgroups: wingless (WNT), sonic hedgehog (SHH), Group 3, and Group 4, with distinct transcriptional profiles, copy number alterations, somatic mutations, and clinical outcomes. Overall, 5–6% of MB patients harbor germline mutations that are considered to be pathogenic, including in adenomatous polyposis...
this is the first study establishing a link between germline microsatellite genotypes and the presence of a pediatric cancer. Given the dearth of environmental influences and the relative paucity of germline mutations associated with pediatric cancers, microsatellites constitute an important novel and understudied source of genetic predisposition. Future screening and prevention strategies applied to rare diseases will require the means to identify populations at risk, a purpose that microsatellite signatures could fulfill. In addition to this potential, the establishment of the biological influence of specific microsatellites upon their companion genes could delineate avenues for preventative intervention.

Methods

Patient Samples

The majority of sequencing data used in this work has been previously published and can be found in Supplementary Table 1 and the Supplementary Methods. Additionally, whole-exome sequencing (WES) from 6 MB patients’ blood DNA was newly generated using the TruSeq exome target enrichment kit and Illumina Sequencer HiSeq 2500 (data available upon request). All patient material was collected after informed consent under a CNMC institutional review board approved protocol. Power calculations were performed to determine the sample size of training and validation cohorts (see Supplementary Methods).

Microsatellite Genotyping

A list of 625 195 unique MS in version GRCh38/hg38 of the human reference genome was generated with a custom Perl script ‘searchTandemRepeats.pl’ using default parameters (Supplementary Methods).

Reads from WES and whole-genome sequencing (WGS) were mapped to the human GRCh38/hg38 reference genome using Bowtie2 and reads mapping to MS locations were locally realigned using the Genome Analysis
Toolkit IndelRealigner tool after duplicated reads were marked. We used the program Repeatseq to determine the genotype of MS in WES or WGS (Supplementary Methods).

For each microsatellite, we calculated whether the distribution of genotypes differs in the germline DNA from 2 groups of samples in the training dataset. For each microsatellite, a contingency table is populated with genotype counts for the 2 groups: MB and normal (Supplementary Fig. 1). Then, statistical differences were quantified using a generalized Fisher’s exact test. The Benjamini–Hochberg multiple testing correction ($n = 43457$ tested MS) was applied to control the false discovery rate. The numbers used to describe each genotype denote the length of that microsatellite in each allele.

Microsatellite Filtering to Control for Age, Ethnicity, and Sequencing Protocol

This study was designed to identify germline variations of MS specific to MB. The publicly available sequence data used contain limited metadata (sex, age, and ethnicity), variation in the sequencing protocol used, and relatively few children. Thus it was not possible to confine our analysis to age and ethnically matched controls, and therefore our analysis ran the risk of identifying MS with age, sequencing, and ethnic bias rather than disease status alone. In an attempt to eliminate this risk, we identified MS with potential bias—for age, sequencing protocol, or ethnicity—and excluded them from subsequent analysis. Details about how this filtering was done can be found in the Supplementary Methods and Supplementary Figures 2–4.

Metric to Score Samples and ROC Analysis

Metric to score samples

We designed a metric to score samples based on their unique distribution of MS genotypes. Essentially, the metric is a weighted sum of the genotypes belonging to each sample: weights stem from the frequency in each genotype in the MB and healthy groups. A visual summary of the metric is provided in Supplementary Fig. 5. Scores calculated for every sample in our cohorts can be found in Supplementary Table 1.

ROC analysis

We used receiver operating characteristic (ROC) analysis to design a classification scheme capable of differentiating samples with MB from healthy controls. The area under the ROC curve (AUC) was used as a measure of how well scores differentiate the 2 groups. Then, a cutoff was selected for all future classification (see the “Validation” section below). Here, the cutoff is a single score that minimizes sensitivity and simultaneously maximizes specificity; it was identified using the Youden index (ie, sensitivity + specificity – 1)$^{32}$ (Supplementary Methods).

Subset of Microsatellites (Genetic Algorithm)

Genetic algorithms are a class of biologically inspired algorithms and have been described extensively elsewhere.$^{33}$ In this work we use a genetic algorithm to identify the most predictive subset of markers—from the set of 139 found to be correlated with MB—using a 2-step process. In initialization, an initial population is generated with random subsets of the 139 MS markers. In optimization, subsets are iteratively ranked, recombined, and replaced (see Supplementary Fig. 6). The algorithm has 4 adjustable hyperparameters: (i) the maximum population size, (ii) the size of each subset, (iii) the metric used to rank each subset, and (iv) the number of subsets replaced in each iteration. Details of each step and hyperparameters are provided in the Supplementary Methods.

Validation

Samples used for validation can be found in Supplementary Table 1 and the Supplementary Methods. Each validation sample was scored with the same metric used for the training samples (see above). The cutoff (identified in training) was used to predict which of the 530 validation samples had MB and which were healthy controls. MB was predicted for validation samples above the cutoff and compared with the known identity of the validation samples.

Additionally, an extended validation cohort using a recently published MB germline WGS dataset contained 290 samples that did not overlap with the samples we previously used. These samples were scored with the procedure described above.

Microsatellite Mutability

In order to test whether individuals with MB are more prone to MS variation, we used the total number of alleles genotyped for each microsatellite as a measure of its mutability and compared this metric across disease and control cohorts. Details regarding the identification of minor alleles can be found in the Supplementary Methods.

Downstream Analysis

For functional analysis, we used genes associated with the 139 loci of MS whose genotypes are significantly different between MB subjects and controls. Pathway analysis was performed using Ingenuity Pathway Analysis (Qiagen, https://www.qiagenbioinformatics.com/products/ingenuity-pathway-analysis). Mutations and co-occurrence were analyzed using PedcBioPortal (www.ncbiportal.org). In the mutation analysis, the numbers indicate the percent of tumors in each dataset with mutations in at least 1 of the 124 genes associated with the identified MS. Construction of a protein-protein interaction (PPI) network was conducted with STRING (Search Tool for the Retrieval
RNA Sequencing

In order to investigate the effect of the MS genotypes on their associated genes, we downloaded 170 files from RNA sequencing (RNA-seq) (European Genome-phenome Archive dataset EGAD00001003279; Supplementary Table 1) partially overlapping with our validation and extended validation cohorts. Details of the expression analysis can be found in the Supplementary Methods.

Stability Prediction of mRNA

To analyze the effect of the untranslated region (UTR)/exonic MS on the stability of the RNA, we predicted the minimum free energy (MFE) secondary structures for each of the MS variants using the RNAfold tool from the ViennaRNA package. Because the MFE depends on the length of the RNA, we normalized the MFE by the size of the RNA and then calculated the difference in MFE between the different RNA variants (ΔMFE).

Results

Identification of Medulloblastoma Microsatellite Informative Loci

To explore the impact of variations of MS in MB predisposition, we have developed a new computational workflow to identify germline MS whose genotypes differ between children with MB and control subjects (Fig. 1). We have applied this approach to germline DNA sequencing data divided into training and validation groups. The training set contained 120 MB and 425 control individuals, and the validation set contained 102 MB and 428 control individuals (Supplementary Table 1).

Using the training set, we first genotyped 43,457 different MS present in both the 120 MB samples and 425 healthy controls. For each of these microsatellites, a generalized Fisher's exact test was used to assess the statistical difference in genotype distribution between the 2 groups, revealing 2094 MS with P < 0.05. After Benjamini–Hochberg multiple testing correction (α = 0.05), 422 passed false discovery. We then performed 3 additional steps to remove MS that nonrandomly vary with age, ethnicity, and DNA sequencing protocol (Fig. 1, Supplementary Figures 2–4 and Supplementary Methods).

Fig. 1 Schematic representation of the approach used for the identification and validation of medulloblastoma associated microsatellites. The approach used in this work is divided into 3 stages: computational identification of informative microsatellite loci using the training set, validation of the microsatellite markers in an independent validation cohort, and finally downstream analysis of the genes associated with those microsatellites. The first stage includes a filter to eliminate microsatellites that vary with age, ethnicity, and sequencing technology.
In total, 283 MS were removed from the list of 422, resulting in a reduced list of 139 markers (Supplementary Table 2). In summary, our approach identified 139 MS from germline DNA whose genotypes were significantly different between MB subjects and healthy controls.

**Medulloblastoma Microsatellite Classifier Set**

In order to identify a subset of MS with the best performance in distinguishing MB samples and healthy controls, we used the set of 139 MS to train an MB classifier. First, we designed a metric to score each sample based on the genotypes of the 139 MS (Supplementary Methods, Supplementary Fig. 5). We used a genetic algorithm\(^{33}\) to identify a subset of 43 MS that best distinguish MB samples from healthy controls (Supplementary Methods, Supplementary Fig. 6). Briefly, subsets of MS (drawn from the set of 139) are ranked by their precision and recall. Subsets with the lowest ROC results are discarded; those that remain are continuously mixed and re-ranked. The algorithm converged in 87 cycles to reveal a subset of 43 MS with an \(F\)-measure of 0.90 and an AUC of 0.962 (Fig. 2, Table 1, Supplementary Table 3, Supplementary Fig. 6). Using Youden’s index,\(^{32}\) we determined that the optimal cutoff score for differentiating MB samples from healthy controls is 0.155 (Supplementary Fig. 7). Thus, we have identified a set of 43 MS whose genotype distributions are able to distinguish MB patients from healthy controls with 88% sensitivity and 92% specificity (Fig. 2A, C and Supplementary Fig. 8).

In order to validate this result, we used an independent cohort of germline DNA including 102 MB and 428 control subjects. This number of sample was selected to ensure that the study would be robustly powered (Supplementary Methods). We applied the optimal cutoff (0.155), using Youden’s index, to the independent validation sample set and found that the classifier could distinguish cases from controls with a sensitivity of 95% and specificity of 90% (Fig. 2B, D). We further tested our signature in a larger, recently published independent dataset of 290 germline MB WGS with deeper coverage\(^{35}\) (Supplementary Table 1) and found it to perform similarly well (Supplementary Fig. 9).

In summary, we have identified and validated a set of 43 MS whose genotype distributions are able to distinguish MB patients from healthy controls using germline DNA with high sensitivity and specificity.

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Fig. 2  Distribution of metric scores in the training (A) and validation (B) cohorts. ROC analysis performed on training (C) and validation (D) cohorts. The \(y\)-axis indicates the normalized proportion of individuals in each cohort per score value bin on the \(x\)-axis.
To determine whether the performance of the signature set varies across the 4 consensus molecular subgroups of MB, we tested the 43 MS signature across MB subgroups combining the training and the validation cohorts to achieve adequate sample sizes. It performed equally well for all MB subgroups (Kolmogorov–Smirnov test) (Supplementary Fig. 10).

Table 1  List of 43 MS loci in the MB germline DNA classifier

| Position       | Fisher’s P-value | Benjamini Adj. P-value | Repeat Unit | Gene | Region     |
|----------------|------------------|------------------------|-------------|------|------------|
| chr20:30215031-30215040 | 0                | 0                      | A           | –    | Intergenic |
| chr4:71756149-71755177  | 0                | 3.60E-07               | TATT        | GC   | Intron     |
| chr2:216610248-216610286 | 0                | 3.05E-06               | TTTTC       | –    | Intergenic |
| chr4:102949102-102949116 | 1.00E-08         | 5.17E-06               | AC          | SLC9B1 | Intron     |
| chr13:44943352-44943377  | 2.00E-08         | 1.21E-05               | AC          | NUFP1 | Intron     |
| chr22:50215636-50215648  | 1.60E-07         | 7.59E-05               | G           | SELENOO | Intron    |
| chr17:17793870-17793820  | 3.54E-06         | 1.07E-03               | CAG          | RAI1 | Exon       |
| chr1:153770135-153770152 | 3.76E-06         | 1.11E-03               | GT           | INTS3 | Intron     |
| chr6:106518976-106519008 | 4.94E-06         | 1.36E-03               | CA           | CRYBG1 | Intron     |
| chr2:42923771-42923785  | 1.16E-05         | 2.69E-03               | A            | RIT2  | Intron     |
| chr17:7024701-7024730    | 3.54E-06         | 1.07E-03               | CAG          | BCL6B | Exon       |
| chr16:4407278-4407290    | 1.10E-05         | 2.58E-03               | AT           | ESCO1 | Intron     |
| chr8:1278442-1278456     | 1.10E-05         | 2.59E-03               | CA           | TERT  | Intron     |
| chr18:12653323-12653334  | 2.12E-05         | 4.39E-03               | T            | LARP7 | Intron     |
| chr4:7773174-7773189     | 2.42E-05         | 4.88E-03               | A            | CNOT8L | Intron    |
| chr5:64507682-64507694   | 2.69E-05         | 5.31E-03               | A            | RGS7BP | Intron     |
| chr1:3836468-3836492     | 3.11E-05         | 5.99E-03               | T            | CEP104 | Intron     |
| chr20:33015600-33015613  | 3.24E-05         | 6.11E-03               | A            | BPIFB2 | Intron     |
| chr10:26223687-26223728  | 6.49E-05         | 1.06E-02               | GT           | GAD2  | Intron     |
| chr15:91749825-91749843  | 6.80E-05         | 1.10E-02               | GGTGGA       | –     | Intergenic |
| chr15:64395361-64395385  | 7.25E-05         | 1.16E-02               | TG           | TRIP4  | Intron     |
| chr10:36522760-36522794  | 8.32E-05         | 1.29E-02               | GT           | –     | Intergenic |
| chr3:189907237-189907252 | 9.32E-05         | 1.40E-02               | T            | –     | Intergenic |
| chr5:91149985-91150025   | 1.04E-04         | 1.54E-02               | TTCTTT       | ADGRV1 | Intron     |
| chr20:33623955-33623983  | 1.07E-04         | 1.58E-02               | A            | CBFA2T2 | Intron    |
| chr16:75234097-75234113  | 1.47E-04         | 2.07E-02               | AC           | BCAR1  | Intron     |
| chr4:151662756-151662771 | 2.06E-04         | 2.71E-02               | T            | FAM160A1 | 3’ UTR     |
| chr7:37033493-37033506   | 2.43E-04         | 3.03E-02               | A            | ELMO1  | Intron     |
| chr10:27145290-27145311  | 2.51E-04         | 3.11E-02               | A            | YME1L1  | Intron     |
| chr5:79088359-79088398   | 2.81E-04         | 3.37E-02               | TG           | BHMT2  | Intron     |
| chr4:52062187-52062204   | 2.99E-04         | 3.51E-02               | T            | SPATA18 | Intron     |
| chr1:153645035-153645049 | 3.45E-04         | 3.92E-02               | T            | CH10P  | Intron     |
| chr15:85123338-85123386  | 3.44E-04         | 3.92E-02               | AC           | PDE8A  | Intron     |
| chr17:80822939-80822950  | 3.56E-04         | 3.99E-02               | GT           | RPTOR  | Intron     |
| chr20:5106079-5106120    | 3.76E-04         | 4.17E-02               | AC           | TMEM230 | Intron     |
| chr10:104037288-104037301 | 3.86E-04       | 4.25E-02               | T            | COL17A1 | Intron     |
| chr4:6084968-6084994     | 4.24E-04         | 4.55E-02               | A            | JAKMIP1 | Intron     |
| chr14:80905756-80905768  | 4.63E-04         | 4.84E-02               | A            | CEP128  | Intron     |
| chr6:49470141-49470153   | 4.84E-04         | 4.99E-02               | T            | CENPQ   | Intron     |
Medulloblastoma Informative Microsatellite Loci Mutability

In the germline, indels of MS are significantly more common than elsewhere in the genome, $10^{-4}$ to $10^{-3}$ compared with $10^{-5}$ per locus per generation respectively. However, mutation rates also vary for different MS based on the length of the repeat, their repetitive motif, and influence on DNA folding. We hypothesized that the differences found for the 139 MB-associated MS (Supplementary Table 2) could be the result of increased MS genotype variation inherent in the individual with MB. In order to test whether individuals with MB are more prone to MS variation, we used the total number of alleles genotyped for each microsatellite as a measure of its mutability and compared this metric across disease and control cohorts. We didn’t find significant differences in the number of genotyped alleles between healthy and MB individuals, supporting the conclusion that there is not a general instability of MS in MB patients. This is consistent with a previous report that found only 1 case out of 36 with increased MS instability. We then investigated whether the predictive capability could be related to a characteristic of the informative MS themselves by ranking all identified MS by allelic load. We found that while the 139 were among the more mutable MS (higher number of alleles), they did not include the most mutable sites. Additionally, we compared the number of homozygote and heterozygote genotypes and the MS array lengths as potential sources of variability and found no statistically significant differences between MB and control germline DNA. From these data, we conclude that the association of the 139 MS with MB is a consequence of those individual microsatellite genotypes rather than simply being a result of constitutional hypermutability.

Specificity of Microsatellite Informative Loci for Medulloblastoma

Interestingly, the group of 139 MS significantly associated with MB contained some intronic MS previously implicated in other cancer types, nuclear fragile X mental retardation protein interacting protein 1 (NUFIP1) and kinesin family member 1B (KIF1B), raising the possibility that the effects of germline MS genotypes are not histologically restricted. To analyze the specificity of the 43 signature MS markers for MB, we tested their performance on a cohort of 153 adult pancreatic ductal adenocarcinoma germline samples from The Cancer Genome Atlas. The signature does differentiate pancreatic cancer samples from the MB validation cohort; however, the pancreatic cancer sample scores are clearly elevated compared with healthy controls (Fig. 3). From this, one may expect other pediatric brain tumors to also have elevated prediction scores. To test this, we performed MS genotyping on 72 ependymomas, 65 high-grade gliomas, and 230 low-grade gliomas from the Children’s Brain Tumor Tissue Consortium and observed elevated MS genotype scores compared with normal controls and pancreatic cancer samples (Fig. 3). Finally, we cross-referenced our set of 139 markers with 55 previously published informative MS markers for breast cancer and 105 for lung cancer. Two markers are in common with breast cancer, NUFIP1 and KIF1B, and 3 markers are shared with lung cancer: NUFIP1, KIF1B, and UBXN7. Interestingly, NUFIP1 and KIF1B had been identified as pan-cancer markers before. In summary, these findings support the overarching hypothesis that variations of MS play a role in cancer susceptibility at an organismal level and, like predisposing gene mutations, do not respect histologic boundaries.

Role of the Informative MS Associated Genes

Of the 139 MS whose genotypes differed between MB and control samples, 114 were located in intronic regions, 15 in intergenic regions, 6 in 3’ UTRs, 3 in exonic regions, and 1 in a 5’ UTR (Fig. 4A). To understand the potential mechanistic roles of these genes, we conducted an Ingenuity Pathway Analysis of the 124 genes associated with non-intergenic informative MS loci. The analysis revealed statistically significant associations with cancer and molecular cellular functions such as cell cycle, DNA replication, recombination and repair, and cellular growth and proliferation (Fig. 4B, Supplementary Table 4). The occurrence of mutations in these 124 genes was examined in 4 MB

Fig. 3 Distribution of metric scores for the signature set of 43 MS in the normal, adult pancreatic ductal adenocarcinoma, ependymoma, high-grade glioma, low-grade glioma, and MB cohorts. The y-axis indicates the normalized proportion of individuals in each cohort per score value bin on the x-axis.
Fig. 4  (A) Pie chart displaying the genomic locations of the 139 informative MS loci. (B) Gene Ontology analysis of the informative MB MS. Y-axis indicates the $-\log_{10}(p\text{-value})$. (C) PPI network of the 124 genes associated with the informative MS loci.
in this study, we have identified a set of 139 MS whose genotypes differ between MB patients and healthy controls. We have shown that a subset of 43 of these MS is able to differentiate MB individuals from controls based upon their germline DNA with a sensitivity and specificity of 95% and 90%, respectively. Although the prospective validation of a diagnostic test for MB predisposition will need to wait for a time when germline sequencing early in life is more ubiquitous, these findings represent the first evidence of germline predisposition to a pediatric cancer that is not attributable to a germline coding mutation.

Discussion

This study identifies a subset of MS that has genotypes that statistically differ between MB samples and healthy controls. Beyond marking individuals with MB, the role these MS may play in medulloblastoma etiology is not yet defined. MS array length variations are known to affect nucleosome positioning, the spacing of DNA binding sites, DNA secondary structure alteration, mRNA and protein stability, gene expression, alternative splicing, and protein sequence. MSs located in intronic regions can alter mRNA splicing by altering immature RNA secondary structure. It has been described that MS proximate to splice sites can alter mRNA splicing by altering immature RNA secondary structure. Out of 139 informative MS, 114 are located in intronic regions. As shown in Supplementary Fig. 11, all the intronic MS are within 1 to 260 bp of an exon and thus a splice site. To detect a possible effect on splicing, we determined the genotypes of the 114 intronic MS in RNAseq data from 170 MB tumors and then correlated them with the abundance of each splice isoform (Supplementary Table 1); 49 of the 837 isoforms were significantly correlated (ANOVA P < 0.05; Supplementary Table 7). Exonic or UTR MS can affect the stability of the mature mRNA. We found 6 in 3' UTRs, 3 in exonic regions, and 1 in a 5' UTR (Fig. 4A). To analyze the effect of the exonic/UTR MS on the stability of the RNA, we predicted the MFE secondary structures for each of the MS variants using the RNAfold tool from the ViennaRNA package (see Supplementary Methods). We calculated the difference in MFE between each RNA variant and the shortest (Supplementary Fig. 12a). We found that differing MS lengths affect the predicted stability of the RNA. For example, we found that the retinoic acid induced 1 (RAI1) mRNA with a microsatellite of 38 nucleotides is the most stable, as opposed to that with a 41 nucleotide microsatellite, which is the least stable. The insertion of an extra CAG triplet increases the positional entropy of the loop formed by the CAG MS in RAI1 mRNA, resulting in the lower predicted stability (Supplementary Fig. 12b). We found that the genotypes associated with MB have the least stable mRNAs for the exonic MS in the RAI1 and B-cell chronic lymphocytic leukemia/lymphoma 6 member B (BCL6B) genes, as well as for the 3' UTR MS in the ZBTB3 and MIDN genes. On the other hand, the genotype associated with MB on the 3' UTR of the FAM160A1 gene has the most stable mRNA. To extend this analysis to mRNA abundance, we then analyzed RNA-seq data from 170 MB tumors (Supplementary Table 1) and correlated them with the MS genotypes in tumor DNA but did not find any significant correlation with gene expression (ANOVA; Supplementary Fig. 12c).
Supplementary Material

Supplementary data are available at Neuro-Oncology online.

Keywords

germline predisposition | medulloblastoma | microsatellites

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