Neurofibromatosis 2 (NF2) is a rare autosomal dominant disease that is caused by inactivating mutations of the NF2 tumour suppressor gene. The NF2 gene has 17 exons and codes for a protein (termed merlin or schwannomin) that links the cytoskeleton to the cell membrane. The protein has 595 amino acids with three functional domains: an amino terminal domain, an α helical domain, and a carboxy terminal domain. NF2 codons 1–302 (exons 1–9) correspond to the predicted amino terminal domain, codons 303–478 (exons 10–13) to the α helical domain, and codons 479–595 (exons 14–17) to the carboxy terminal domain. Pathogenic NF2 mutations have not been found in exons 1 or 17.

Clinically, NF2 is characterised by nervous system tumours and ocular abnormalities. Vestibular schwannomas, usually bilateral, occur in more than 90% of adult patients and intracranial meningiomas occur in about 50% of patients. Mild and severe disease often can be defined by the age at onset of symptoms of NF2 and the number of cerebral tumours other than vestibular schwannomas. In general, constitutional nonsense and frameshift NF2 mutations are associated with severe disease, and missense mutations and somatic mosaicism are associated with mild disease. In contrast, patients with constitutional splice site NF2 mutations have highly variable disease severity. The cause of the high phenotypic variability in NF2 patients with splice site mutations is not known. One hypothesis concerns the location of constitutional NF2 mutations. The clinical characteristics examined were age at onset of symptoms of NF2 and number of intracranial meningiomas, which are the primary indices of the severity of NF2. Two regression models were used to analyse genotype-phenotype correlations. People with splice site mutations in exons 1–5 had more severe disease than those with splice site mutations in exons 11–15. This result is compatible with studies showing that exons 2 and 3 are required for self-association of the amino terminal of the NF2 protein in vitro, and that deletions of exons 2 and 3 in transgenic and knockout mouse models of NF2 cause a high prevalence of Schwann cell derived tumours.

Neurofibromatosis 2 (NF2) patients with constitutional splice site NF2 mutations have greater variability in disease severity than NF2 patients with other types of mutations; the cause of this variability is unknown. We evaluated genotype-phenotype correlations, with particular focus on the location of splice site mutations, using mutation and clinical information on 831 patients from 528 NF2 families with identified constitutional NF2 mutations. The clinical characteristics examined were age at onset of symptoms of NF2 and number of intracranial meningiomas, which are the primary indices of the severity of NF2. Two regression models were used to analyse genotype-phenotype correlations. People with splice site mutations in exons 1–5 had more severe disease than those with splice site mutations in exons 11–15. This result is compatible with studies showing that exons 2 and 3 are required for self-association of the amino terminal of the NF2 protein in vitro, and that deletions of exons 2 and 3 in transgenic and knockout mouse models of NF2 cause a high prevalence of Schwann cell derived tumours.

METHODS
Patient data
We compiled an international database from all published studies of NF2 mutations and related clinical data, the United Kingdom NF2 registry, and unpublished data from investigators. The database has the advantage of a very large sample size, although clinical data vary in completeness. Currently, the database has information on 1112 patients from 738 NF2 families with identified constitutional NF2 mutations. The database is described in the Human Gene Mutation Database and additional information is available from MEB.

People were eligible for this study if they had an identified constitutional NF2 mutation and information on the age at onset of symptoms of...
NF2, or the number of intracranial meningiomas, or both (subsequently, age at onset of symptoms refers to symptoms of NF2 and age at diagnosis refers to diagnosis of NF2). Age at onset of symptoms and number of intracranial meningiomas are the primary clinical indices of the severity of NF2. The analyses excluded people with rare types of NF2 mutations (in-frame deletions or insertions, large insertions, or chromosomal translocations) and somatic mosaics with mutations other than nonsense or frameshift mutations. A total of 831 patients from 528 NF2 families (409 people with new mutations and 422 inherited cases) met these criteria and were included in the study.

Statistical methods
Cross sectional associations of the type of constitutional NF2 mutation with the age at onset of symptoms and the number of meningiomas were assessed by regression models. The models were chosen on the basis of the nature of the dependent variable. A multivariate normal model with right censoring was used to model the association of the type of constitutional NF2 mutation with the age at onset of symptoms. For asymptomatic patients, the age at onset of symptoms was right censored at the older of two other age variables (age at diagnosis and age at last examination or death). A gamma mixture of negative binomials model was used to evaluate the association of the type of constitutional NF2 mutation with the number of intracranial meningiomas because the count distribution of meningiomas was heavy right skewed. Models were fit by maximum likelihood estimated using the quasi-Newton method. Computations were done using C programs developed at the Department of Statistics, University of British Columbia (available from HJ).

Each model had an exchangeable correlation structure within families that permitted assessment of intrafamilial effects beyond those due to the type of constitutional NF2 mutation. The other covariates in each model were gender and inheritance (new mutation or inherited case). In the meningioma model, the age at last examination or at death (or the age at diagnosis if this information was missing) was a covariate because the penetrance of NF2 associated meningiomas increases with age.

The type of mutation variable was classified categorically as seven binary variables. These variables were indicators of somatic mosaicism (defined at the molecular level), splice site mutations (subdivided by location), missense mutations, large deletions, and patients with full constitutional nonsense or frameshift mutations. The latter group of patients, who typically have severe disease, was the reference group for statistical comparisons.

For the purpose of the analysis, we divided the location of splice site mutations into three groups using two different methods. There were 128 families with splice site mutations, but these mutations were distributed unevenly among the 15 NF2 exons in which pathogenic mutations have been reported. There was only one mutation in exon 1 and only two mutations each in exons 9 and 10. There are several possible ways to group the exons, and two different approaches were used in the analysis. In the first analysis, splice site mutations were divided according to the predicted merlin functional domains (exons 1–9, 10–13, and 14–15). In the second analysis, splice site mutations were divided in three equal groups by exon number without a priori reference to functional domains (exons 1–5, 6–10, and 11–15).

Genotype-phenotype correlations were evaluated using the model based dependent variables and their 95% confidence intervals (CI). CIs that excluded zero were considered to be statistically significant. CIs were calculated in two ways: for different types of mutations relative to the reference group (that is, with splice site mutations analysed as a single class), and between the three NF2 regions of splice site mutations. To evaluate differences between people with splice site mutations in the three NF2 regions, the 95% CI of the difference between the model coefficients for the exon categories was calculated (for example, the difference in the age at onset of symptoms between people with splice site mutations in exons 1–5 or 6–10, exons 1–5 or 11–15, and exons 6–10 or 11–15).

Kaplan-Meier analysis was used to compare survival in people with splice site mutations in different regions of the NF2 gene. Only patients from the United Kingdom NF2 registry were used in the survival analysis because the registry has complete ascertainment of survival. The United Kingdom NF2 registry has 94 NF2 patients from 51 families with splice site mutations. To account for the positive association within families, the p value was based on the null permutation distribution of the log rank statistic, with permutation of families. Under the null hypothesis of equal survival distribution in the two groups, families can be permuted to each of the two groups (while keeping the same number of families in each group as in the original data), and the log rank statistic can be computed for each permutation.

RESULTS
The characteristics of the study population are presented in table 1. As in previously published studies that were included in the international NF2 mutation database, somatic mosaics and people with non-truncating mutations had milder disease than people with full constitutional nonsense or frameshift mutations (the reference group). Somatic mosaics and people with splice site mutations, missense mutations, or large deletions were older at onset of symptoms and at diagnosis than the reference group. The prevalence of meningiomas was lower in people with splice site mutations, missense mutations, or large deletions than in the reference group.

The genotype-phenotype correlations are presented in tables 2 and 3. These tables have the observed and model based maximum likelihood estimates for average age at onset of symptoms (table 2) and average number of meningiomas (table 3). The model based estimate of the average age at onset of symptoms was significantly higher in somatic mosaics (29 years) and in people with missense mutations (31 years) or large deletions (23 years) than in the reference group (16 years) (table 2). The model based estimate of the average number of meningiomas was significantly lower in people with missense mutations (0.4) or large deletions (0.8) than in the reference group (1.7) (table 3). When splice site mutations were analysed as a single class, people with splice site mutations were significantly older at onset of symptoms and had significantly fewer meningiomas than the reference group.

The genotype-phenotype correlations for the location of splice site mutations are of particular interest. The results for the two groupings were similar for age at onset of symptoms, although the model based on grouping of exons by functional domains produced mean ages of onset that were slightly closer to the observed averages (table 2). People with splice site mutations in exons 1–9 were significantly younger at onset of symptoms than people with mutations in exons 14–15. Similarly, people with NF2 splice site mutations in exons 1–5 or 6–10 were significantly younger at onset of symptoms than people with splice site mutations in exons 11–15.

For number of meningiomas, the grouping based on equal number of exons provided more precise genotype-phenotype correlations within the predicted amino terminal domain. People with splice site mutations in exons 1–9 had significantly more meningiomas than people with mutations in exons 14–15, but people with mutations in exons 1–5 had
significantly more meningiomas than people with mutations in exons 6–10 or 11–15. Overall, when the results for age at onset of symptoms and number of meningiomas are taken together, people with splice site mutations in exons 1–5 have more severe disease than people with mutations in exons 11–15.

Other covariates contributed significantly to the models after adjustment for the type of constitutional NF2 mutation. There was a weak but significant intrafamilial correlation for age at onset of symptoms (0.21, 95% CI 0.13 to 0.30), but neither age, inheritance, nor gender were significant explanatory variables in these models. Similarly, there was a weak but significant intrafamilial correlation for number of meningiomas (0.18, 95% CI 0.06 to 0.30). Inherited cases had significantly fewer meningiomas than people with new mutations (0.5-fold, 95% CI 0.2– to 0.8-fold). The number of meningiomas increased slightly but not significantly with increasing age, and gender did not significantly influence the number of meningiomas.

For the Kaplan-Meier survival analysis, the location of splice site mutations was dichotomised into exons 1–5 or 6–15 because people with splice site mutations in exons 1–5 had the most severe disease. There were too few people with splice site mutations in exons 14–15 to support a mortality

<p>| Table 1 | Characteristics of 831 people with NF2 by type of NF2 mutation |
|-----------------|-------------------------------------------------|-----------------|-----------------|-----------------|-----------------|-----------------|</p>
<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Type of NF2 mutation</th>
<th>Nonsense or frameshift</th>
<th>Splice site</th>
<th>Missense</th>
<th>Large deletion</th>
<th>Splice site</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of people/families</td>
<td>Classical*</td>
<td>340/272</td>
<td>40</td>
<td>263/128</td>
<td>73/25</td>
<td>115/63</td>
</tr>
<tr>
<td>Inheritance (%)</td>
<td>New mutations</td>
<td>60</td>
<td>100</td>
<td>32</td>
<td>15</td>
<td>40</td>
</tr>
<tr>
<td>Age, mean (SD) (years)</td>
<td>Exons 6–10</td>
<td>32</td>
<td>15</td>
<td>29</td>
<td>12</td>
<td>22</td>
</tr>
<tr>
<td>Intracranial meningioma (%)</td>
<td>Splice site (total)</td>
<td>7</td>
<td>11</td>
<td>13</td>
<td>7</td>
<td>5</td>
</tr>
<tr>
<td>Vestibular schwannoma (%)</td>
<td>Splice site (divided by functional domains)</td>
<td>None</td>
<td>Exons 1–9</td>
<td>23</td>
<td>12</td>
<td>29</td>
</tr>
<tr>
<td>Male</td>
<td>Exons 10–13</td>
<td>26</td>
<td>29</td>
<td>14</td>
<td>28</td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>Exons 14–15</td>
<td>25</td>
<td>23</td>
<td>31</td>
<td>29</td>
<td></td>
</tr>
<tr>
<td>Infiltrated cases</td>
<td>Absent</td>
<td>55</td>
<td>50</td>
<td>44</td>
<td>49</td>
<td>31</td>
</tr>
<tr>
<td>New mutations</td>
<td>Present</td>
<td>31</td>
<td>41</td>
<td>71</td>
<td>69</td>
<td>66</td>
</tr>
<tr>
<td>Male</td>
<td>Diagnosis of NF2</td>
<td>69</td>
<td>59</td>
<td>40</td>
<td>29</td>
<td>31</td>
</tr>
<tr>
<td>Female</td>
<td>Non</td>
<td>69</td>
<td>59</td>
<td>40</td>
<td>29</td>
<td>31</td>
</tr>
</tbody>
</table>

**Note:**
*Classical NF2* refers to people with full constitutional NF2 mutations, as opposed to somatic mosaics defined at the molecular level. †Excludes 67 inherited cases who were asymptomatic at the time of diagnosis of NF2.

<p>| Table 2 | Genotype-phenotype correlations for the age at onset of symptoms of NF2 |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Covariate | Sample average age at onset of symptoms (years) | Model based estimated average age at onset of symptoms (years) | 95% CI |</p>
<table>
<thead>
<tr>
<th>Type of NF2 mutation</th>
<th>Nonsense or frameshift</th>
<th>Classical NF2†</th>
<th>Somatic mosaic</th>
<th>Splice site</th>
<th>Missense</th>
<th>Large deletion</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exons 1–9</td>
<td>17</td>
<td>29*</td>
<td>25*</td>
<td>21*</td>
<td>26*</td>
<td>32*</td>
</tr>
<tr>
<td>Exons 10–13</td>
<td>23</td>
<td>25*</td>
<td>31*</td>
<td>21*</td>
<td>26*</td>
<td>32*</td>
</tr>
<tr>
<td>Exons 14–15</td>
<td>21</td>
<td>20</td>
<td>20</td>
<td>20</td>
<td>20</td>
<td>20</td>
</tr>
<tr>
<td>Exons 1–5</td>
<td>25</td>
<td>23</td>
<td>23</td>
<td>23</td>
<td>23</td>
<td>23</td>
</tr>
</tbody>
</table>

Statistically significant.
†The NF2 mutations in the body of the table are comparisons to the reference group. The 95% CI of differences in age at onset of symptoms (in years) between people with splice site mutations in different regions of the NF2 gene are as follows:

- Exons 1–9: 10–13: −11 to +1
- Exons 1–9: 14–15: −15 to −5
- Exons 10–13: 11–15: −12 to −2
- Exons 1–5: 6–10: −5 to +3
- Exons 1–5: 11–15: −13 to −4
- Exons 6–10: 11–15: −12 to −3

**CI:** confidence interval.
Splice site mutations in NF2

Table 3  Genotype-phenotype correlations for the number of intracranial meningiomas

<table>
<thead>
<tr>
<th>Covariate</th>
<th>Sample average number of intracranial meningiomas</th>
<th>Model based estimated average number of intracranial meningiomas</th>
<th>95% CI for count ratios (compared to people with classical NF2 and nonsense or frameshift mutations)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type of NF2 mutation</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nonsense or frameshift</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Classical NF2†</td>
<td>1.7</td>
<td>1.7</td>
<td>1.00 (reference group)</td>
</tr>
<tr>
<td>Somatic mosaic</td>
<td>2.1</td>
<td>1.4</td>
<td>(0.5 to 1.2)</td>
</tr>
<tr>
<td>Splice site (total)‡</td>
<td>0.9*</td>
<td>0.9*</td>
<td>(0.4 to 0.7)*</td>
</tr>
<tr>
<td>Splice site (divided by functional domains)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exons 1–9</td>
<td>1.1*</td>
<td>1.1*</td>
<td>(0.5 to 0.9)*</td>
</tr>
<tr>
<td>Exons 10–13</td>
<td>0.8</td>
<td>0.7</td>
<td>(0.2 to 1.1)</td>
</tr>
<tr>
<td>Exons 14–15</td>
<td>0.2*</td>
<td>0.3*</td>
<td>(0.1 to 0.4)*</td>
</tr>
<tr>
<td>Splice site (divided by exon number)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exons 1–5</td>
<td>1.7</td>
<td>1.7</td>
<td>(0.6 to 1.6)</td>
</tr>
<tr>
<td>Exons 6–10</td>
<td>0.7*</td>
<td>0.7*</td>
<td>(0.2 to 0.7)*</td>
</tr>
<tr>
<td>Exons 11–15</td>
<td>0.4*</td>
<td>0.5*</td>
<td>(0.1 to 0.5)*</td>
</tr>
<tr>
<td>Missense</td>
<td>0.4*</td>
<td>0.4*</td>
<td>(0.1 to 0.5)*</td>
</tr>
<tr>
<td>Large deletion</td>
<td>0.7*</td>
<td>0.8*</td>
<td>(0.3 to 0.7)*</td>
</tr>
</tbody>
</table>

*Statistically significant.
‡There are three separate models. In the first model, splice site mutations are analysed as a single class ("total"). In the second model, splice site mutations are assigned to the nearest exon, and the exons are divided by functional domains within the merlin protein (exons 1–9, 10–13, or 14–15). In the third model, splice site mutations are assigned to the nearest exon, and exons are divided into three equal groups by exon number (exons 1–5, 6–10, or 11–15). In each instance, the coefficients for the other covariates (for example, the intrafamilial correlation coefficient) are those from the model in which splice site mutations are analysed by equal groups by exon number. The values for these other covariates are almost identical in each of the three models.
§Statistical tests in the body of the table are comparisons to the reference group. The 95% CIs of differences in number of meningiomas between people with splice site mutations in different regions of the NF2 gene are as follows:
- Exons 1–9 v 10–13: −0.5 to +1.4
- Exons 1–9 v 14–15: 0.6 to 2.2
- Exons 10–13 v 11–15: −0.2 to +2.2
- Exons 1–5 v 6–10: 0.2 to +1.3
- Exons 1–5 v 11–15: 0.6 to 2.0
- Exons 6–10 v 11–15: −0.2 to +1.2

CI, confidence interval.

**DISCUSSION**

Almost all previous studies of genotype-phenotype correlations in NF2 have been based on relatively few patients due to the rarity of the disease.19 Larger studies with detailed clinical data are needed to evaluate specific genotype-phenotype correlations, such as correlations for individual types of NF2 associated abnormalities and correlations for the location of a single type of mutation within the NF2 gene. Recently, we evaluated genotype-phenotype correlations for individual types of NF2 associated abnormalities using the population based United Kingdom NF2 registry, which has information on 560 people with NF2. We found that there were genotype-phenotype correlations for intracranial meningiomas, spinal tumours, peripheral nerve tumours, and cataracts.27 Genotype-phenotype correlations for specific types of abnormalities have been found in other diseases that are caused by mutations in other tumour suppressor genes, notably APC, VHL, and NF1.

Yet even the United Kingdom NF2 registry is not large enough to evaluate genotype-phenotype correlations for the location of a single type of mutation within the NF2 gene. In this study, we used an international NF2 mutation database to extend NF2 genotype-phenotype correlations to the location of splice site mutations. Our findings are analogous to those that have been observed for the location of APC mutations and the number of colorectal adenomas,20,21 the occurrence of congenital hypertrophy of the retinal pigment epithelium,22–24 the occurrence of desmoid tumours,25,26 and the age at onset of colon cancer.27 However, our findings differ from the genotype-phenotype correlations in APC because there is not a genotype-phenotype correlation for the location of all types of mutations within the NF2 gene. The location of nonsense and frameshift NF2 mutations, which occur in 56% of NF2 families,28 is not significantly associated with the age at onset of symptoms of NF2 or the number of meningiomas (data not shown).

Genotype-phenotype correlations in NF2 have been studied experimentally. Naturally occurring single-base pair substitutions (missense mutations and nonsense mutations) have
been used to identify the location of amino acid residues that are critical for the tumour suppressor function of merlin. Missense mutations produce merlin that is defective in negative growth regulation, while nonsense mutations do not produce stable merlin. Missense mutations also produce merlin with reduced but not absent binding to β1 integrin. In this study, we confirmed that missense mutations are usually associated with mild NF2.

Missense mutations occur in only 5% of NF2 families, and the number of currently available patients is insufficient to test the association of the location of missense mutations with phenotype. Splice site mutations in exons 1–5. Each had significantly fewer meningiomas than people with splice site mutations in 3 exons also had older onset of symptoms. In addition, the effect on number of meningiomas does not appear to be limited to exons 14–15. People with splice site mutations in exon 6–10 and 11–15 each had significantly fewer meningiomas than people with splice site mutations in exons 1–5.

Age at onset of symptoms and number of meningiomas provide complementary information for genotype-phenotype correlations in NF2 because presenting symptoms in NF2 usually are not related to meningiomas. The most common presenting symptoms of NF2 are hearing loss, tinnitus, vertigo, or imbalance, which are usually caused by vestibular schwannomas. Seizures or headaches, the usual symptoms of meningiomas, are the presenting symptoms in only 6–8% of symptomatic NF2 patients. Age at onset of symptoms of NF2 and age at diagnosis of NF2 are strong predictors of the risk of mortality in NF2. The association between the location of splice site mutations and survival in the present study probably reflects the differences in age at onset of symptoms in people with splice site mutations in different regions of the NF2 gene. When the location of splice site mutations and age at onset of symptoms are both included as covariates in a Cox regression model, mutation location is not an independent predictor of mortality (data not shown).

We found that, in general, NF2 large deletions were associated with mild clinical manifestations of NF2 (in the sub-group analysis, the position of intragenic large deletions with the NF2 gene was associated with the age at onset of symptoms). The cause of the association of NF2 large deletions with mild disease is unknown. Constitutional NF2 large deletions are common, occurring in 21–32% of NF2 families, but conventional mutation screening techniques such as single strand conformation polymorphism analysis do not detect heterozygous large deletions. There have been few published data on genotype-phenotype correlations for NF2 large deletions, but our findings are consistent with these earlier observations. Bruder et al hypothesised that there was a modifier gene for NF2 located telomeric to NF2 on chromosome 22. These investigators then used microarray-comparative genomic hybridisation in a series of NF2 patients with large deletions and demonstrated that the deletions in people with moderate or severe NF2 often extended beyond the NF2 locus, while the deletions in people with mild NF2 only affected the NF2 locus itself. When NF2 large deletions in the international database are classified as either whole gene deletions or deletions that involve part of the NF2 gene, people with these two types of large deletions do not have significantly different mean ages at onset of symptoms or numbers of intracranial meningiomas (data not shown). However, this broad categorisation of the size of NF2 large deletions may not be sufficiently precise to identify the phenotypic effects of putative modifier genes near the NF2 locus.
Splice site mutations in NF2

Somatic mosaicism occurs in an estimated 25–30% of NF2 patients with new mutations. Our data suggest that there is ascertainment bias in the identification of somatic mosaics. In the international database, 65 (79%) of 82 known mosaics have nonsense mutations, frameshift mutations, or indels, while only 241 (37%) of 640 people with new mutations and full constitutional mutations have these types of mutations (Fisher’s exact test, p = 0.001). Full constitutional nonsense mutations and frameshift mutations usually cause severe NF2. Somatic mosaics with nonsense or frameshift mutations may be more likely to be diagnosed because their phenotype, although attenuated by mosaicism, is still severe enough to be classified as NF2, while mosaicism in people with other types of mutations may produce a phenotype that is so mild that it escapes medical attention or fails to meet the standard clinical diagnostic criteria for NF2.

After accounting for genotype-phenotype correlations in regression models, there were weak but significant intrafamilial correlations for the age at onset of symptoms and the number of meningiomas. We previously demonstrated these correlations in a subset of patients from the present study. Intratumoral correlations may reflect differences in the phenotype of individual NF2 alleles that comprise a mutation type, or these correlations might be evidence of modifying genes.

The finding that inherited cases had fewer meningiomas than people with new mutations may be due to reproductive selection. As seen in table 1, the proportion of inherited cases is higher in people with splice site mutations, missense mutations, or large deletions than in people with full constitutional nonsense or frameshift mutations. Full constitutional nonsense or frameshift mutations usually cause severe NF2, and these patients are less likely to reproduce because they often die in the third or fourth decade of life and are typically debilitated for a lengthy time beforehand.


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15. Evans DGR, Ruttledge MH, Mautner VF. Splice site mutations in NF2 54554
germline mutant APC associated with APC mutations beyond codon 1444. polyposis: desmoid tumours and lack of ophthalmic lesions (CHPRE).


The location of constitutional neurofibromatosis 2 (NF2) splice site mutations is associated with the severity of NF2


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