Risk of Colorectal Cancer in Monoallelic and Biallelic Carriers of MYH Mutations: A Population-Based Case-Family Study

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Abstract

Previous case-control studies have suggested that carriers of monoallelic germine mutations in the MYH gene may be at increased risk of colorectal cancer. We applied a kin-cohort design, using a modified segregation analysis, to estimate the colorectal cancer risk using 300 first-degree relatives of 35 colorectal cancer cases who were monoallelic or biallelic carriers of MYH mutations. We found that monoallelic carriers had a 3-fold increased risk of colorectal cancer (hazard ratio, 2.9; 95% confidence interval, 1.2-7.0; P = 0.02) and biallelic carriers a 50-fold increased risk (hazard ratio, 53; 95% confidence interval, 14-200; P < 0.0001). This analysis illustrates the potential of family analysis to estimate cancer risk for low-frequency mutations and, based on the proportion of relatives predicted to be carriers, we believe that this constitutes the largest study of monoallelic carriers to date. (Cancer Epidemiol Biomarkers Prev 2006;15(2):312–4)

Introduction

A recessive colorectal cancer risk associated with mutations in the base excision repair gene MYH was first shown by Al-Tassan et al. (1) and Sieber et al. (2). Since then, we (3) and several other groups (4-8) have observed a higher prevalence of monoallelic mutation carriers in colorectal cases compared with controls. Although no individual study alone found statistically significant evidence, when combined, these studies suggest an additional dominant effect (3). Most recently, Farrington et al. (9) found 45 monoallelic MYH mutation carriers in 2,239 colorectal cases and 24 monoallelic MYH mutation carriers in 1,845 controls, and concluded that monoallelic MYH mutation carriers may be at increased risk of colorectal cancer. However, whereas the increased risk was observed only in carriers over the age of 55 years [risk ratio, 1.68; 95% confidence interval (95% CI), 1.07-2.95], there was no difference between the increased risk in this age group and the risk in carriers 55 years old or younger.

We have applied a kin-cohort design to assess colorectal cancer risk in monoallelic and biallelic MYH mutation carriers, all of whom are a first-degree relative of a colorectal cancer case identified in a population-based study. We analyzed the cancer histories of ungenotyped first-degree relatives of the colorectal cancer cases we previously found to be carriers of MYH mutations (3).

Materials and Methods

As explained in detail by Cotterchio et al. (10), the Ontario Familial Colorectal Cancer Registry identified individuals diagnosed with colorectal cancer (excluding familial adenomatous polyposis) of ages 20 to 74 years at diagnosis (probands). Attempts were made to confirm all reports of colorectal cancer in relatives using clinical records and/or death certificates. All probands defined by strong family histories of colorectal cancer and age of diagnosis as being either at “high risk” (Amsterdam Criteria ref. 11) or “intermediate risk” of carrying a genetic mutation and 25% of the remaining “low-risk” cases were invited to provide a blood sample. As described by Croitoru et al. (3), MYH Y165C and G382D mutations were identified in the probands by denaturing high-performance liquid chromatography and confirmed by sequencing. In subjects heterozygous for the MYH Y165C and G382D mutations, the entire coding region of the MYH gene was analyzed by denaturing high-performance liquid chromatography to identify other pathogenic MYH mutations. This identified 41 mutation carrying probands. To exclude the possibility that a proportion of colorectal cancer in the probands, and hence in their relatives, was due to germline mutations in the mismatch repair genes, available tumors of probands were tested for microsatellite instability and loss of mismatch repair protein expression. If tumors showed high-frequency microsatellite instability and/or were immunodeficient for a mismatch repair protein, the germline was tested for mutations in mismatch repair genes hMLH1, hMSH2, and hMSH6 by sequencing and multiplex ligation-dependent probe amplification. This led to the identification of two germline mismatch repair gene mutation carriers and these probands and their relatives were excluded from analysis. Of the remaining 39 probands, 27 were monoallelic carriers for Y165C (n = 8) or G382D (n = 19), 6 were homozygous carriers for Y165C (n = 2) or G382D (n = 4), and 6 were compound heterozygous carriers for G382D/Y165C (n = 3), Y165C/Y90X (n = 1), Y165C/891+3A>C (n = 1), and G382D/891+3A>C (n = 1). Of the 39 probands, 25 were from the 100% sampled high-risk or intermediate-risk group and 14 were from the 25% sampled low-risk group.

The observed incidence of colorectal cancer in relatives (number of relatives with colorectal cancer divided by the total person years of relatives) was compared with the expected incidence calculated by applying age- and sex-specific cancer incidence rates obtained for the Ontario population 1993-2002. Standardized incidence ratios and 95% CIs were calculated using STATA version 8.0 (12).
Table 1. Frequency of monoallelic MYH carriers colorectal cancer cases and controls from previous case-control studies

<table>
<thead>
<tr>
<th>Study</th>
<th>Frequency in cases* (%)</th>
<th>Frequency in controls* (%)</th>
</tr>
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<tbody>
<tr>
<td>Farrington et al. (9)</td>
<td>45 of 2,205 (2.0%)</td>
<td>28 of 1,822 (1.5%)</td>
</tr>
<tr>
<td>Croitoru et al. (3)</td>
<td>29 of 1,226 (2.4%)</td>
<td>21 of 1,255 (1.7%)</td>
</tr>
<tr>
<td>Fleischmann et al. (5)</td>
<td>6 of 356 (1.7%)</td>
<td>2 of 354 (0.6%)</td>
</tr>
<tr>
<td>Peterlongo et al. (8)</td>
<td>4 of 553 (0.7%)</td>
<td>7 of 918 (0.8%)</td>
</tr>
<tr>
<td>Enholm et al. (4)</td>
<td>5 of 1,038 (0.5%)</td>
<td>0 of 424 (0.0%)</td>
</tr>
<tr>
<td>Kambara et al. (6)</td>
<td>2 of 92 (2.2%)</td>
<td>1 of 53 (1.9%)</td>
</tr>
<tr>
<td>Wang et al. (7)</td>
<td>10 of 442 (2.3%)</td>
<td>4 of 313 (1.3%)</td>
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</table>

*Excluding biallelic carriers.

We estimated the average hazard ratio (the ratio of the age-specific incidence in carriers of a germline MYH mutation to that in noncarriers) by a modified segregation analysis fitted under maximum likelihood theory using the statistical package MENDEL (13-15). Biallelic carriers were assumed to have both alleles mutated and treated as homozygous carriers. The colorectal cancer incidence for each family member was assumed to follow a Cox proportional hazards model where the incidence depends on the genotype (carrier or noncarrier), colorectal cancer diagnosed in at least one parent, and colorectal cancer diagnosed in at least one sibling. The latter two covariates account for familial aggregation unrelated to the mutations. Specifically, based on unpublished control data from the same population, the Ontario Familial Colorectal Cancer Registry, the probability of having at least one parent with colorectal cancer was set at 0.08 (8% of population controls reported having at least one parent with colorectal cancer) and of having at least one sibling with colorectal cancer was set at 0.04 (4% of population controls reported having at least one sibling with colorectal cancer). We assumed that in the population, having at least one first-degree relative with colorectal cancer doubled colorectal cancer risk (16, 17). The population allele frequency of germline mutations was taken to be 0.0085 based on a 1.7% monoallelic carrier frequency in the control population (3). The likelihood of each family was expressed as a function of the colorectal cancer status and the ages at last contact, death, or diagnosis of family members. Recessive, dominant, and codominant modes of inheritance were compared using the likelihood ratio test.

To adjust for ascertainment, the likelihood for each family was conditioned on the colorectal cancer status and MYH genotype status of the proband, thus ensuring unbiased risk estimates. The incidence in noncarriers was approximated by the population incidence of colorectal cancer for Ontario for 1993-2002. The cumulative risk in carriers was calculated by integrating over the age-specific incidence in carriers (equal to the population age-specific incidence multiplied by the estimated average hazard ratio). To account for stratified sampling based on family history, the low-risk families were given a weight of 4.0 compared with a weight of 1.0 for the high-risk and intermediate-risk families. Standard errors for calculation of confidence intervals were estimated using the robust Huber-White sandwich estimator (18).

The overall evidence from previous studies for an association between monoallelic MYH mutation carriers and colorectal cancer was estimated by a meta-analysis on the data from seven case-control studies (3-9). These constituted all identified studies that tested for monoallelic and biallelic carriers of MYH mutations in colorectal cancer cases and in controls. Exposed cases were defined as colorectal cancer cases who were monoallelic carriers of MYH mutations and nonexposed cases were colorectal cancer cases with no germline mutations. Controls were subjects without colorectal cancer (see Table 1). Random and fixed effects models were fitted; heterogeneity was tested using Cochran’s Q; and a Forrest plot of the odds ratios was generated. Odds ratios, their 95% CIs, and P values were estimated using the software Comprehensive Meta-analysis (http://www.meta-analysis.com).

Results

A total of 300 first-degree relatives of 39 mutation-carrying probands were identified (204 of the 27 monoallelic probands and 96 of the 12 biallelic probands). Of the 123 first-degree relatives of the high-risk or intermediate-risk monoallelic probands, 10 (8%) were diagnosed with colorectal cancer, an incidence which is 5.0 (95% CI, 2.6-9.3) times that expected based on their age distribution. Of the 66 first-degree relatives of the biallelic probands, 5 (8%) were diagnosed with colorectal cancer, which is 5.2 (95% CI, 2.1-12.5) times more than expected. Of the 81 first-degree relatives of the low-risk monoallelic probands, 5 (6%) were diagnosed with colorectal cancer, which is 3.0 (95% CI, 1.2-7.3) times that expected, and of the 30 first-degree relatives of the low-risk biallelic probands, 2 (7%) were diagnosed with colorectal cancer, which is 4.5 (95% CI, 1.1-18.1) times that expected.

A codominant model (homozygote risk is greater than heterozygote risk which is greater than wild-type risk) gave a better fit than a recessive model (P = 0.002). Table 2 shows that monoallelic carriers were, on average, 2.9 (95% CI, 1.2-7.0) times more likely to be diagnosed with colorectal cancer compared with the general population (P = 0.02). This hazard ratio translates to an 8% cumulative risk to age 70 years. Biallelic carriers were, on average, 53 (95% CI, 14-200) times more likely to be diagnosed with colorectal cancer compared with the general population (P < 0.0001) and were almost certain to be diagnosed with colorectal cancer by age 70 years.

There was no difference in risk of colorectal cancer for monoallelic carriers between those over the age of 55 years (hazard ratio, 2.9; 95% CI, 1.0-8.2; P = 0.04) and those 55 years old or younger (hazard ratio, 2.8; 95% CI, 0.5-17; P = 0.3 and P for difference = 0.9).

Figure 1 shows that whereas no single previous study showed conclusively that monoallelic carriers were at increased risk, the combined estimate from all studies was significant. The widths of the confidence intervals in Fig. 1 illustrate the limited statistical power for most of the individual studies. The pooled odds ratio estimate from the meta-analysis was 1.4 (95% CI, 1.0-2.0; P = 0.04) with no evidence to reject homogeneity (Cochran’s Q = 2.076 with 6 degrees of freedom; P = 0.9).

Discussion

We have applied a kin-cohort design to assess colorectal cancer risk in monoallelic and biallelic MYH mutation carriers, all of whom are a first-degree relative of a colorectal cancer case. We analyzed the cancer histories of ungenotyped first-degree relatives of the 39 colorectal cancer cases we previously found to be carriers of MYH mutations from a population-based study (3). We found an ~3-fold increased risk of colorectal cancer in monoallelic carriers and a 50-fold increased risk in biallelic carriers of colorectal cancer.

Table 2. Hazard ratios of risk of colorectal cancer in MYH carriers compared with that in the general population, and associated cumulative risk to age 70 years, estimated using modified segregation analysis

<table>
<thead>
<tr>
<th>Carriers</th>
<th>Hazard ratio (compared with population risk)</th>
<th>P</th>
<th>Cumulative risk to age 70 y</th>
</tr>
</thead>
<tbody>
<tr>
<td>Monoallelic</td>
<td>2.9 (1.2-7.0)</td>
<td>0.02</td>
<td>8% (4-19%)</td>
</tr>
<tr>
<td>Biallelic</td>
<td>53 (14-200)</td>
<td>&lt;0.0001</td>
<td>80% (35-100%)</td>
</tr>
</tbody>
</table>

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increased risk in biallelic carriers. This analysis illustrates the power of family-based data for the study of rare variants. It is more efficient than a case-control analysis because for every monoallelic carrier, on average, half their first-degree relatives will also be monoallelic carriers, and for every biallelic carrier (assuming mutations are on different alleles), both parents, all children, and, on average, half their siblings will be monoallelic carriers. Based on Mendelian inheritance, the carrier (assuming mutations are on different alleles), both monoallelic carrier, on average, half their first-degree relatives.

We found that, on average, such carriers were approximately three times the population risk for colorectal cancer. Further studies using genotyped relatives will be necessary to refine these estimates for the MYH165C and G382D mutations, as well as for the less common pathogenic MYH mutations.

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References

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