SHORT COMMUNICATION

Deletion analysis of chromosome 8p in sporadic colorectal adenomas

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Summary In order to assess the stage of colorectal tumorigenesis at which chromosome 8p loss of heterozygosity (LOH) occurs, 56 sporadic adenomas were examined for LOH at four polymorphic loci which show frequent LOH in carcinomas. LOH was found in only 5 out of 51 (9.8%) informative adenomas, whereas studies with the same markers in 85 informative carcinomas showed a LOH of 45%. The adenomas showing LOH were all in the 'high-risk' clinicopathological category, being 10 mm or more in diameter and showing tubulovillous architecture. It is concluded that the chromosome 8p locus is involved preferentially in the development of carcinomas rather than adenomas.

The development of colorectal cancer incorporates several discrete genetic events. Some are largely restricted to carcinomas, whereas others are found in both adenomas and carcinomas. Inactivation of tumour-suppressor genes APC and DCC by mutation or loss occurs in the majority of both colorectal adenomas and carcinomas (Fearon & Vogelstein, 1990; Powell et al., 1992). Similarly, activating mutations in K-ras oncogene occur with nearly identical frequency in carcinomas and larger adenomas (Fearon & Vogelstein, 1990). Presumably such lesions affect the expansion of neoplastic populations without directly conferring a malignant phenotype. In contrast, the p53 gene is inactivated in 75% of colorectal cancers but is not abnormal in adenomas except those displaying features of severe dysplasia, and this gene would appear to have a major role initiating malignant behaviour (Baker et al., 1990; Kikuchi-Yanoshita et al., 1992; Carder et al., 1993). Such genetic lesions are of interest because they suggest the existence of genes involved in the critical transition from benign to malignant growth. We (Cunningham et al., 1993) and others (Fujiiwara et al., 1993) have recently identified a region on chromosome 8p that exhibits frequent LOH in colorectal cancer, indicating a further putative oncosuppressor locus or loci. In this paper we assess the frequency of LOH at this region in sporadic colorectal adenomas to determine at what stage in evolution of colorectal tumours this molecular lesion exerts preferential selective advantage.

Loss of heterozygosity affecting chromosome 8p is found in over 50% of colorectal cancers (Cunningham et al., 1993). A similar frequency of chromosome 8p LOH has been determined in bladder, prostate, lung and hepatocellular cancer (Bergenheim et al., 1991; Emi et al., 1992; Knowles et al., 1993). In bladder and hepatocellular cancer, there is a correlation between chromosome 8p LOH and advanced tumour stage and grade (Emi et al., 1993; Knowles et al., 1993). In colorectal cancer we were unable to identify a correlation between chromosome 8p LOH and tumour site or Duke’s stage (Cunningham et al., 1993), although such an association with clinicopathological stage has been suggested by others (Fujiiwara et al., 1993). In addition, the possibility of two separate chromosome 8p colorectal oncosuppressor loci has been proposed (Fujiiwara et al., 1993). This work addresses the question of the role of the chromosome 8p locus or loci in tumorigenesis by determining the frequency of loss of heterozygosity in 56 sporadic colorectal adenomas at four chromosome 8p loci which have shown a high frequency of LOH in colorectal cancers. The adenomas were gathered from both cancer-bearing and cancer-free bowel. The results suggest preferential involvement of the chromosome 8p locus in the later stages of colorectal tumorigenesis.

Materials and methods

DNA was purified from 56 sporadic colorectal adenomas and matched normal tissue, either blood or histologically normal colonic mucosa, obtained from 49 individuals. Thirty-three adenomas were collected from fresh colorectal specimens resected for malignant disease. Of the remainder, 16 were removed endoscopically from cancer-free bowel as part of a presymptomatic screening programme.

Analysis of LOH using (CA)n repeat polymorphisms

Three microsatellite (CA)n repeat markers were employed which had shown a high frequency of LOH in colorectal cancers. ANK1 (Polymeropoulos et al., 1991) maps to 8p21–p11.2, LPLS GT (Tomforde et al., 1992) maps to 8p22 and D8S137 (Tomforde et al., 1992) to 8p21–p12. Polymerase chain reactions comprised 25 μl volumes with 200 μM of each nucleotide, 0.625 units of Taq polymerase enzyme (Promega, UK) 2.5 μl of buffer (Promega, UK) 2.5 mm magnesium sulphate, 50 ng of each primer and 50 ng of template DNA. In each reaction 10 pg of one primer was end labelled with [32P]dATP, using T4 polymerase kinase. PCR reactions were carried out in microwell plates in an Omnitegen thermal cycler (Hybaid, Middlesex, UK) and consisted of 29 cycles of 95°C for 1 min, 55°C for 1 min and 72°C for 2 min. A 10 μl aliquot of loading buffer (95% deionised formamide, 10 mm disodium EDTA, 0.1% xylene cyanol and 0.1% bromophenol blue) was added to each reaction and 4 μl aliquots were run on 6% denaturing polyacrylamide gels. Gels were dried and exposed to Kodak X-OMAT AR film for 24 h. Autoradiographs were assessed visually by two observers.

HindIII polymorphism detected by PCR

HindIII polymorphism in intron 8 of the lipoprotein lipase gene (8p.22) was demonstrated by PCR under conditions described by Bruin et al. (1991). PCR products were digested with HindIII, separated by electrophoresis on 2% agarose gels, stained with ethidium bromide and visualised under ultraviolet light.
Results

We have previously reported LOH analysis of three of the markers used in this analysis (ANK1, LPL3GT and LPLHdIII) in 120 colorectal cancers (Cunningham et al., 1993). In this study we further analysed the same colorectal cancers at the D8S137 locus and found a frequency of LOH of 50% (27 out of 54 informative cases). The percentage LOH for each of these markers in colorectal cancers and adenomas is presented in Figure 1. Of the 56 adenomas, 51 were informative with at least one marker, and of these only five (9.8%) showed loss of heterozygosity. In contrast, when used in the study of 120 colorectal cancers, these four markers detected a LOH in 38 out of 85 informative cases (45%), a difference that is highly significant ($\chi^2 = 16.38; P < 0.00005$). The adenomas consisted of 35 tubulovillous lesions (mean size 27 mm, range 6–80 mm), 20 tubular lesions (mean size 14 mm, range 5–20 mm) and one villous lesion (15 mm). The five shown to have LOH at chromosome 8p were tubulovillous adenomas of 10 mm or more in diameter. Chromosome 8p LOH was not detected in any of the 10 adenomas less than 10 mm in diameter, seven of which were tubular adenomas and three tubulovillous lesions.

Discussion

We have detected loss of heterozygosity in less than 10% (5/51) of adenomas examined in this series, which is significantly less than the frequency of LOH in adenomas of colorectal cancers (45%) in a similar analysis of 85 informative malignant tumours. As detailed above, this was a mixed group of adenomas in terms of size and histological types. The five lesions showing LOH were all tubulovillous adenomas, 10 mm or more in diameter. Although the numbers are small, chromosome 8p LOH was only detected in this study in the subgroup of adenomas which are known to carry a greater malignant potential. We are unaware of any study of chromosome 8p LOH in sporadic adenomas. However, a recent report of adenomas arising in familial adenomatous polyposis (FAP) describes no chromosome 8p LOH in 37 informative adenomas from two individuals (Ichii et al., 1993). Our series is likely to have included a higher proportion of tubulovillous lesions than this group of FAP adenomas, and this may account for the absence, albeit rare, of chromosome 8p LOH in the sporadic lesions reported here. Overall, our data suggest that the putative chromosome 8p tumour-suppressor gene is important in the later stages of tumorigenesis in the colon and rectum. This pattern is similar to that noted for the p53 gene (Baker et al., 1990; Kikuchi-Yanoshita et al., 1992) and strikingly different to that found in APC, DCC and K-ras (Fearon & Vogelstein, 1990; Powell et al., 1992).

Thus, LOH at the 8p locus appears to be one of a select group of acquired genetic lesions preferentially associated with malignant change in colorectal epithelium. Others include abnormalities of p53 and aneuploidy. In several cell lineages, including colorectal mucosa, abnormalities of p53 are known to induce instability of the genome, of which aneuploidy is an example (Livingstone et al., 1992; Carder et al., 1993). A further lesion, at the MSH2 gene on chromosome 2p, is associated with hereditary non-polyposis colorectal cancer (HNPPC), a familial disorder characterised by the development of carcinoma without prior proliferation of benign lesions (Fishel et al., 1993; Leach et al., 1993). This also appears to involve infidelity in DNA replication, characterised by variability in the length of microsatellite repeats between normal and tumour DNA (Aaltonen et al., 1993; Ionov et al., 1993; Thibodeau et al., 1993). Such instability has been recorded in up to 28% (Thibodeau et al., 1993) of apparently sporadic colorectal cancers but is rare in adenomas (Young et al., 1993). At the three microsatellite loci examined in this paper we detected instability in only one adenoma, an 18-mm-diameter tubulovillous lesion removed from non-cancer-bearing bowel. Published data indicate that genomic instability manifest as either aneuploidy or microsatellite instability is commonly acquired in malignant colorectal lesions. It is interesting to speculate that the defects in the putative 8p oncogene may also relax the fidelity of DNA or chromosomal replication or impair DNA repair mechanisms.

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