Peutz-Jeghers polyps, dysplasia, and K-ras codon 12 mutations


Abstract

Background—Peutz-Jeghers syndrome (PJS) is a rare, autosomal dominant, polyposis syndrome, associated with an increased risk of gastrointestinal and extragastrointestinal malignancy. Occasionally dysplasia occurs in PJS polyps. Aims—In colorectal carcinomas, mutations in codon 12 of the K-ras oncogene are common and are found at similar frequency in precursor adenomas. Therefore, K-ras codon 12 point mutations in PJS polyps were evaluated.

Materials and methods—Fifty two PJS polyps, including four with dysplasia, collected from 19 patients with PJS, were analysed for mutations in the K-ras codon 12 by a mutant enriched polymerase chain reaction procedure, followed by allele specific oligodeoxynucleotide hybridisation.

Results—A K-ras codon 12 mutation was identified in one colonic polyp with dysplasia. The mutation was found in the non-neoplastic epithelial cells and not in the dysplastic component of the polyp.

Conclusions—K-ras codon 12 point mutations are very rare in PJS polyps, by contrast with colorectal adenomas. The findings support previous evidence that there seems to be no intrinsic relation between K-ras codon 12 mutation and dysplasia.

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Keywords: Peutz-Jeghers syndrome; polyps; dysplasia; K-ras codon 12 mutations

Peutz-Jeghers syndrome (PJS) is a rare autosomal dominant disorder defined by hamartomatous polyposis of the gastrointestinal tract and the occurrence of melanin spots on the lips and buccal mucosa. Polyps primarily occur in the small bowel, but are also found in the stomach and colon, and they range in size from a few millimeters to several centimeters. PJS polyps are classified as hamartomas because they are composed of epithelium consisting of the cellular elements normally present at that particular site of the gut and an excessive proliferation of smooth muscle extending into the lamina propria in an arborisation-like fashion. Patients with PJS can present with intussusception, anaemia, gastrointestinal bleeding, and intestinal obstruction. The germline mutation(s) responsible for PJS are as yet unknown, although linkage to chromosome 19p is reported.

The hamartomatous polyps are not generally regarded as premalignant lesions. However, patients with PJS have a high relative risk of both intestinal and extraintestinal malignancies. Although no apparent direct relation exists between polyp formation and the risk of carcinoma formation, dysplasia is occasionally found in PJS polyps. By analogy to the adenoma—carcinoma sequence in colorectal cancer, carcinomas in the gastrointestinal tract may form due to progression of hamartoma to dysplasia to carcinoma. Molecular genetic alterations identifiable in hamartomatous PJS polyps might give an insight into the nature of the neoplastic risk. Tumorigenesis in the gastrointestinal tract results from an interaction between activation of dominantly acting oncogenes and inactivation of recessive tumour suppressor pathways. In the large bowel, the ras oncogene family is known to be involved in the adenoma—carcinoma sequence. The ras oncogenes encode for closely related 21 kDa proteins that function in cellular signal transduction, providing growth advantage. These proteins acquire transforming potential when altered at certain critical positions, in codons 12, 13, and 61. Activation of the ras oncogenes (H-ras, K-ras and N-ras) is seen in about 50% of colorectal cancers and their precursor adenomas. The most often encountered mutations are in codon 12 of K-ras and account for about 75% of ras mutations in tumours of the gastrointestinal tract. In this report we describe the analysis of 52 hamartomatous PJS polyps for K-ras mutations.

Methods

Tissue specimens
Fifty two PJS polyps were collected from the pathology archives of the Academic Medical Centre of the University of Amsterdam, the University Hospital Rotterdam, and the Bowel Tumor Working Group Registry at The Johns Hopkins University School of Medicine. These were all hamartomatous polyps from the gastrointestinal tract (stomach n = 4, duodenum n = 14, small intestine n = 18, large bowel...
n=16), obtained from 19 patients with PJS (table 1). The diagnosis of dysplasia was derived from the pathology reports of the original institutions and confirmed by review of the slides. Specimens of normal gastroenterocolonic mucosa were also analysed in all the patients.

TISSUE PREPARATION AND K-ras CODON 12 ANALYSIS
Archival specimens were cut into 5 µm sections, mounted on to glass slides, and non-dysplastic and dysplastic areas of polyp tissue were microdissected. The tissue was collected in microcentrifuge tubes containing 50–200 µl DNA isolation buffer (50 mM TRIS-HCl, pH 8.0, 0.2% Tween-20, and 100 mg/ml proteinase K) depending on the size of the tissue fragments and incubated at 56°C for 18–24 hours. Samples were heated to 95°C for 10 minutes to inactivate the proteinase K and to denature the DNA. After this treatment, the samples were analysed for K-ras codon 12 using a modified version of a previously described protocol.13 K-ras point mutations were evaluated by amplification of the sample DNAs, restriction enzyme digestion using MvaI (an isoschizomer of BstNI; Boehringer Mannheim, Mannheim, Germany), reamplification of the digested and undigested polymerase chain reaction (PCR) products, and allele specific oligodeoxynucleotide hybridisation. For the first round of PCR sample DNA was amplified for 15 cycles (94°C 30 seconds; 62°C 30 seconds; 72°C 60 seconds) with primers A (5’-ACT GAA TAT AAA CTT GTG GTA GTT GGA CCT-3’) and D (5’-TCA TGA AAA TGG TCA GAG AAACC-3’). To enrich for the K-ras codon 12 mutants, the PCR products were then digested with MvaI to concentrate the mutant K-ras DNA fragments. Subsequently, a second round PCR was performed on both the digested and undigested first round PCR products. In the second round PCR, DNA was amplified for 35 cycles (94°C 30 seconds; 55°C 30 seconds; 72°C 60 seconds) using primers A and B (5’-TCAAAGAATGGTCCTGGACC-3’). After denaturation the undigested and digested PCR products were spotted on to seven different nylon membranes (Genescreen Plus, NEN Research Products, Boston, MA, USA) and hybridised to each of the K-ras codon 12 specific oligodeoxynucleotides, as described previously.14 Positive controls comprised plasmid clones harbouring each of the possible sequence variations in codon 12. Subsequent washes were carried out at 63°C, followed by autoradiography. All analyses for K-ras codon 12 mutations were performed in duplicate in two independently isolated DNA specimens.
Results
A K-ras codon 12 mutation—with the wild type codon GGT (glycine) mutated to GTT (valine)—was identified in the DNA of one colonic polyp (figs 1 and 2). After separating the dysplastic epithelium from the remaining polyp tissue by microdissection, the mutation was solely found in the non-dysplastic area (fig 1), whereas the dysplastic epithelium lacked the mutation. No additional K-ras mutations were identified in the remaining 51 hamartomatous gastrointestinal polyps, of which three contained dysplasia (table 1). Also, in the 52 samples of the normal adjacent mucosa, no K-ras codon 12 mutations were found.

Discussion
Hamartomatous PJS polyps have a neoplastic potential,13 but our study shows that activated K-ras plays no obvious part in the process. In a previous cytogenetic study a normal karyotype was found in PJS polyps.16 To our knowledge, these are the first findings of a difference in genetic make up between hamartomatous PJS polyps and adenomatous polyps of the colorectum.

Of particular note, the DNA from the dysplastic epithelium of the PJS polyp with the mutation showed wild type K-ras at codon 12 and the mutation was found in the non-dysplastic epithelium. Our finding raises questions about the role of ras mutations in colorectal carcinogenesis. Aberrant crypt foci are microscopic lesions speculated to precede adenomatous poly development. Ras mutations are often found in aberrant crypt foci in the mucosa of patients, but these lesions usually lack dysplasia.17,18 By contrast, another event—for example, inactivation of the APC gene—seems to be needed to provide the dysplastic phenotype.17 Apparently, ras activation provides a growth advantage but does not contribute to neoplastic transformation.

In summary, K-ras codon 12 mutations are very rare events in PJS polyps, by contrast with the high frequency of ras activation in adenomas and carcinomas of the large bowel. Furthermore, an intrinsic relation between K-ras codon 12 mutation and dysplasia was not noted. Further investigation is needed to elucidate the role and timing of K-ras oncogene mutation in various forms of colorectal tumorigenesis.

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