Elevated expression of the human ras oncogene family in premalignant and malignant tumours of the colorectum
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Summary  Study of expression of ras-related oncogenes in human premalignant polyps and malignant tumours of the colorectum, as well as in normal colorectal mucosa, shows a significant elevation in the premalignant and malignant tissues as compared to their respective colorectal mucosa. These results suggest that activation of the ras oncogene family occurs in the development of colorectal tumours and that elevated expression at a premalignant stage may well be critical in the process of carcinogenesis but not in itself sufficient.

Materials and methods

Tissue specimens were collected and stored in liquid nitrogen. These were subsequently pulverized under liquid nitrogen and RNA and DNA was extracted as previously described (Spandidos & Paul, 1982). Briefly, the tissue or cells were homogenized in guanidine-HCl buffer (8.0 M guanidine HCl, 20 mM sodium acetate, 50 mM EDTA, 5% β-mercaptoethanol, pH 7.0). Cell lysates were made 2% with SDS and heated at 65°C for 2 min. After vortexing, 5 ml of cell lysate were placed on a 3 ml cushion of CsCl solution (5.7 M CsCl, 50 mM EDTA pH 8.0) and centrifuged for 48 h at 40 K rpm at 15°C in a 10 x 10 Ti rotor. The RNA pellet was resuspended in 2.0 M LiCl, 4.0 M urea and left at 4°C overnight. RNA was pelleted at 10 K rpm for 15 min in a Sorvall centrifuge, resuspended in 0.1 x MOPS buffer (1 x MOPS = 20 M NaMOPS, 5 mM sodium acetate, 1 mM EDTA, pH 7.0) and dialyzed in the same buffer for 2 h before lyophilization. Before each experiment the quality of RNA preparations was examined by formaldehyde-agarose gel electrophoresis, followed by ethidiun bromide staining, transfer to nitrocellulose and hybridization to DNA probes (see below). Ten μg of total cell RNA was spotted per dot as described (Spandidos et al., 1981). Hybridizations were performed in 5 x SSC, 50% formamide for 24 h at 42°C with 10 ng ml⁻¹ probe as described (Wahl et al., 1979) using 2 x Denhardt's solution (Denhardt, 1966). ³²P-labelled DNA probes with specific activities of 2-3 x 10⁶ cpm μg⁻¹ DNA were made by nick-translation (Rigby et al., 1977). The nitrocellulose sheets were washed in 0.5 x SSC at 60°C and exposed to hypersensitized X-ray films at -70°C (Lasky & Mills, 1977). The filters were hybridized sequentially with ³²P-labelled nick-translated HiHi3 (Ellis et al., 1981), BS9 (Ellis et al., 1980), pL335 (Dalla Favera et al., 1981), pHR28 (Sproul & Birnie, unpublished results) or pAM91 (Minty et al., 1982) recombinant probes carrying the viral Kirsten ras (v-Ki-ras), viral Harvey ras (v-
Ha-ras), human cellular sis (H-c-sis) and human 28S ribosomal and mouse actin DNA sequences respectively. Probes were removed by washing the nitrocellulose at 65°C with double distilled H2O for 2h. Approximately 100 pg of each insert oncogene DNA were spotted as a positive control. Fractionation of RNA in formaldehyde containing agarose gels and blotting on to nitrocellulose have been described elsewhere (Spandidos & Paul, 1982).

Results

RNA spot hybridization analysis

The relative levels of human Ki-ras, Ha-ras and sis transcripts in total cell RNA made from premalignant and malignant tissue, normal colonic mucosa and cell lines were determined using an RNA spot hybridization assay (Spandidos et al., 1981). Quantification of the intensities of the autoradiographic spots was carried out using densitometric scanning as previously described (Spandidos et al., 1981). Probe excess was confirmed by obtaining a linear autoradiographic response to serial dilutions of the various RNAs (data not shown). Results of the RNA spot hybridization analysis are shown in Figure 1 and Table I. These show firstly that transcripts from the human Ki-ras and Ha-ras related oncogenes could easily be detected in most premalignant, malignant tissues and cell line RNAs but are barely detectable in normal tissue. The human sis oncogene is expressed at very low levels in all types of tissue examined. Secondly, the amount of human Ki-ras and Ha-ras specific RNAs varied in different cells and tissues whereas little variation was observed in sis RNA levels. In particular, in the first three patients where samples were available from all three types of tissue, Ki-ras RNA levels in premalignant and malignant tissues varied between 9.5-20 and 3.5-19× higher respectively than the levels seen in normal colorectal mucosa. A slightly different picture was seen when expression of the Ha-ras oncogene family

| Tissue histology | Normal* | Premalignant* | Malignant* |
|------------------|---------|---------------|------------|
| Patient no. and cell line⁹ | Ki-ras Ha-ras Ki-ras Ha-ras Ki-ras Ha-ras |
| 1                | 1.0     | 0.3           | 19         | 31         | 6.3       | 4.5       |
| 2                | 2.0     | 3.5           | 20         | 8.0        | 3.5       | 14        |
| 3                | 2.0     | 2.3           | 9.5        | 9.1        | 19        | 9.0       |
| 4                | 1.7     | 1.0           | 7.0        | 7.6        |           |           |
| 5                | 2.3     | 1.0           | 3.5        | 1.5        |           |           |
| 6                | 1.0     | 1.0           | 6.5        | 1.5        |           |           |
| 7                | 1.0     | 1.3           | 7.0        | 1.5        |           |           |
| 8                | 4.1     | 1.5           | 22         | 8.0        |           |           |
| 9                |         |               |            |            | 5.0       | 1.5       |
| 10               |         |               |            |            | 4.4       | 1.5       |
| 11               |         |               |            |            | 6.5       | 11        |
| 12               |         |               |            |            | 1.9       | 14        |
| 13               |         |               |            |            | 3.0       | 6.3       |
| CHB              |         |               |            |            | 6.5       | 2.0       |
| HL60             |         |               |            |            | 13        | 1.5       |
| K562             |         |               |            |            | 8.0       | 1.5       |

*The autoradiographs (Figure 1) were scanned and the concentrations of HiHi3 (v-Ki-ras) or (v-Ha-ras) specific RNAs are given at arbitrary units for each probe.

¹Histological examination was carried out in part of the specimen and the remaining tissue was stored in liquid nitrogen until RNA and DNA were isolated. No. 1-12, colorectal carcinoma; No. 13, a breast carcinoma; CHB, an established cell line from an adenocarcinoma of the colon; HL60, a promyelocytic and K562 an erythroleukaemic cell line.

²Colorectal mucosa.

³1-2 Colorectal polyps (predominantly tubular), 3, Colorectal polyp (metaplastic) and 8, colorectal polyp (tubulovillous).

⁴Adenocarcinoma of the colorectum.
**Figure 1** RNA spot hybridization analysis of (a) Ki-ras; (b) Ha-ras; (c) sis; (d) rRNA and (e) actin gene expression in human cells. Extraction of RNA from cells and spotting on to nitrocellulose is described in Materials and methods. Malignant: 1–12 = colorectal adenocarcinoma; 13 = breast adenocarcinoma; CHB = an established cell line from an adenocarcinoma of the colon; HL60, a promyelocytic and K562 an erythroleukemic cell line. Premalignant: 1–2 = predominantly tubular adenomatous polyps; 3 = metaplastic polyp, and 8 = tubulo villous polyp. Normal: histologically normal colorectal mucosa removed from colectomy specimens several centimetres distant from tumour site.
was examined. The relative levels of Ha-ras transcripts varied between 8.0–31 and 4.5–14× higher in premalignant and malignant tissues respectively as compared to normal colorectal mucosa for the same first three patients as described above (Figure 1 and Table 1). In patient No. 1 RNA from two separate polyps (P1 and P2) was examined gave similar results. In other malignant tumours there was some variation in the degree of elevation of Ki and Ha-ras expression (Table 1). Oncogene expression in some but not all of the premalignant lesions was in fact significantly higher in comparison to the corresponding malignant tumours. Among the latter a more marked variation in expression was observed. Finally in the four other samples, a breast carcinoma RNA (No. 13), an established cell line from adenocarcinoma of the colon (CHB), a promyelocytic (HL60) and an erythroleukaemic (K562) cell line, expression of Ki-ras was elevated, but Ha-ras RNA levels were increased only in patient No. 13. As an additional control to check the relative amount of RNA from each sample spotted on to nitrocellulose, the filter was hybridized with pHK28, (a human ribosomal), or pAM91 (a mouse actin) DNA probe. As shown in the autoradiographs (Figure 1d and 1e) and confirmed by scanning the dots, there is no substantial difference in the amount of ribosomal or actin RNA present in these samples.

Northern blot hybridization analysis
Northern blot hybridization analysis was carried out to measure the sizes of c-onc related transcripts. A v-Ki-ras probe, HiHi3 recombinant (Ellis et al. 1981), revealed the presence of one main band of ~5.8 kb in total cellular RNA (Figure 2a and c). In several cases a much less intense band of ~2.2 kb was seen as well as some other nondiscrete hybridization probably due to degradation or possibly to cross hybridization with other ras gene family transcripts. As shown in Figure 2b,d after scanning across this 5.8 kb band, the ki-ras related transcript was found in higher amounts in premalignant and malignant tissues of the colorectum as compared to normal mucosa. A similar sized major transcript was also found in RNA from HL60 and K562 cells. As shown in Figure 2e when a v-Ha-ras probe, BS9 recombinant (Ellis et al., 1980) was used, the same size band of 5.8 kb was also present, again more intensely in some premalignant and malignant tissues as compared to normal colorectal mucosa (Figure 2f). These results confirm and extend the spot hybridization analyses. The nature of these transcripts was further investigated by isolating polyA+ RNA and Northern blot hybridization analysis. As shown in Figure 2g,i, Ki-ras and Ha-ras related transcripts of 5.8 kb in size were again found to be elevated in premalignant and malignant tissues as compared to normal colorectal mucosa.

We also detected the 5.8 kb sized transcripts using pT24-C3 (Santos et al., 1982), a recombinant carrying a 6.6 kb BamHI human DNA fragment containing the whole bladder carcinoma oncogene c-Ha-ras 1 (data not shown). The exact nature of these transcripts is still incompletely understood. However mechanisms accounting for varied Ki-ras transcripts in terms of alternative splicing patterns have been recently described which include the generation of such a 5.8 kb species (Shimizu et al., 1983; McGrath et al. 1983). It seems most likely that the same sized transcript seen with the Ha-ras probe represents in fact hybridisation with products of another member of the ras gene family. Northern blot hybridization analysis of total of polyA+ RNA using a sis probe failed, however, to show any discrete RNA transcripts.

Discussion
Other studies with RNA from human cells have demonstrated the presence of 1.2 kb Ha-ras related transcripts in the T24 human bladder carcinoma cell line (Goldfarb et al., 1982) and two ~6.0 kb Ha-ras related transcripts in human haematopoietic cell lines (Westin et al., 1982b). More recently, using an N-ras probe 3 different sized transcripts of 5.8, 2.2 and 1.5 kb have been found in normal human fibroblasts and established human cell lines and it has been claimed that the 2.2 kb transcript is related to the N-ras oncogene (Hall et al., 1983).

The human genome contains at least four genes homologous to the transforming genes of Kirsten

Figure 2 Northern blot hybridization analysis of transcripts related to human Ki-ras and Ha-ras oncogenes in RNAs from samples of normal premalignant and malignant tissues of the colorectum. Total RNAs were isolated as described in Materials and methods. Poly A+ RNA was isolated using an oligo(dT)-cellulose Type 3 from Collaborative Research Inc. (Spandios & Paul, 1982). In (a), (c) and (e), 20 μg of total cell RNA and in (g) and (i) poly A+ RNA isolate from 100 μg total cell RNA were analyzed in 1% agarose-formaldehyde-containing gels, blotted on to nitrocellulose and hybridized with one probes. The HiHi3 (Ellis et al., 1981) recombinant containing the v-Ki-ras sequences was used as a probe in panels (a), (c) and (g). The BS9 (Ellis et al., 1980) recombinant containing the v-Ha-ras sequences was used in (e) and (i). N=normal colorectal mucosa, P=premalignant polyps, M=malignant adenocarcinoma. The autoradiographs are shown in (a), (c), (e), (g) and (i) and the scans across the 5.8 kb bands in (b), (d), (f), (h) and (j).
and Harvey murine sarcoma viruses (Chang et al., 1982) which are dispersed in different chromosomes (O'Brien et al., 1983). Moreover, recent transfection studies have revealed the presence of a distantly related N-ras oncogene (Hall et al., 1983). Our results demonstrate that cellular sequences related to the transforming genes of Kirsten and Harvey murine retroviruses are actively transcribed in human tissues. The demonstration that Ki-ras and Ha-ras related transcripts are elevated in premalignant and malignant tissues as compared to normal colorectal mucosa shows that the expression of these onc genes is associated with the transformed state of the cells and suggests that elevated expression of these genes in premalignant state(s) may be critical in the process of carcinogenesis. The fact, however, that only a relatively small proportion of these premalignant polyps progress to frank malignancy although elevated oncogene expression was observed in all, suggests that, consistent with the concept of carcinogenesis being a multi-step process (Spandidos, 1983) the elevation observed here is not in itself sufficient to produce malignant change. Parallel studies in this laboratory, of carcinogen induced mouse skin papillomata, which have a similar potential for malignancy, have also demonstrated elevated Ha-ras oncogene expression and in addition, the DNA from these tumours has acquired transforming activity in transfection assays (Balmain et al., 1984).

The cellular homologues of several retroviral oncgenes have been shown to exhibit tissuespecific patterns of transcriptional activity (Westin et al., 1982a,b, Gonda et al., 1982). Expression of c-onc genes during mouse development (Muller et al., 1982; 1983) and liver regeneration (Goyette et al., 1983) has lent further support to the hypothesis that cellular oncogenes play a role in normal developmental processes. Abnormal expression of these genes could be directly involved in the development of the transformed phenotype of tumour cells. Although meaningful in vivo studies of proliferation rates in these tumours involving repeated sampling and labelling would present obvious ethical problems, the most recent studies on human material in vitro comparing malignant tumours to normal mucosa using thymomikine (Pritchett et al., 1982) and 3H-Thymidine labelling (Bleiburg et al., 1976) techniques suggest that cell birth rate and turnover time respectively, were only very slightly increased in tumours or not significantly different. Similar findings have been reported for premalignant polyps in human (Weisburger et al., 1975), and in experimentally induced rodent tumours, where similar proliferation rates for benign and malignant tumours were reported (Sunter et al., 1980). In the tumours, furthermore, the growth fraction is probably actually lower than in the normal mucosa. Thus the marked elevation of ras related transcripts we observe would not appear to be comparable to the two–three fold increase seen in regenerating liver (Goyette et al., 1983).

Gene amplification seems not to be involved in the generation of elevated onc transcript levels found in our present study since Ha-ras and Ki-ras related DNA sequences in the various tissues were at approximately the same level when examined by DNA spot hybridization analysis (data not shown). However, oncogene amplification remains a possibility particularly since we have observed such a phenomenon in DNA from a different adenocarcinoma of the colon (our unpublished results). Such a phenomenon has recently been described for c-myc (Collins & Groudine, 1982; Dala-Favera et al., 1982) and Ki-ras (Schwab et al., 1983).

Since our results demonstrate that both the premalignant and malignant tumours examined here are characterized by elevated levels of ras family transcripts and if the gene product is unaltered, the question obviously arises as to the nature of the further event(s) involved in the acquisition of the malignant phenotype and their relationship with the changes in gene expression observed here. To address this question may require the use of further assay systems, although as an initial step it will clearly be of interest to clone the ras genes involved directly from the tumours since they may not readily be detected in transfection experiments, and such studies, as well as transfection studies using DNA from these tumours, are currently in progress in our laboratory.

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