Loss of heterozygosity occurs at the D11S29 locus on chromosome 11q23 in invasive cervical carcinoma

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Summary
Allelotypic detection of loss of heterozygosity (LOH) has been used to identify putative tumour-suppressor genes. Loci on human chromosome 11q23 are frequently altered in malignant disease, and LOH has been reported at an anonymous D11S29 locus at 11q23 in a proportion of breast and ovarian cancers and malignant melanomas. Previous studies have reported a high frequency of LOH in cervical carcinoma mapping to 11q23. Using polymerase chain reaction techniques employing probes for a recently described polymorphic dinucleotide microsatellite within this locus, we have searched for LOH in 69 cases of invasive cervical carcinoma. Genomic material was microdissected from sections cut from archival paraffin-embedded material, using the patients' constitutional genotype as a control. Sixty-two (90%) of the cases were informative, and LOH occurred in 25/62 (40%) of tumours. Loss of an arm or single chromosome 11 is a well-recognised event in cervical carcinoma, and by employing other microsatellite polymorphisms mapping to 11q13 and 11p11–p12 we excluded those cases with widespread allelic loss. By doing so, LOH at D11S29 was found in 16/53 (30%) of tumours. The findings suggest a putative tumour-suppressor gene on 11q involved in cervical carcinogenesis.

Keywords: cervix neoplasms; human chromosome 11; loss of heterozygosity; tumour-suppressor genes; microsatellites; polymerase chain reaction

Cytogenetic and molecular biology studies of human tumours have identified chromosomal abnormalities, including deletions of specific regions, that are associated with malignancy. Many deletions are believed to involve loss of tumour-suppressor genes, only a few of which have been cloned and sequenced. In the tumour cell, mutation of a tumour-suppressor gene is frequently accompanied by loss of the other non-mutated allele together with portions of adjacent DNA (Solomon et al., 1991). This DNA loss may be detected using polymorphic markers for the affected locus and, where a marker is informative, deletion of the locus on one chromosome leads to loss of heterozygosity (LOH) for the polymorphism. Analysis of LOH in tumours may be used to indicate the chromosomal location of putative tumour-suppressor genes.

Cytogenetic studies of cervical carcinoma show a high degree of karyotypic variability, although the breakpoints involved in the structural rearrangements are non-randomly distributed (Atkin et al., 1990). Numerical changes together with non-random deletions, inversions, isochromosome formation and translocations are found to frequently involve chromosome 11. The breakpoints here tended to cluster around the locus 11q23–q25 (Sreekantaiah et al., 1991).

LOH has been identified at a number of loci in cervical cancer. Deletion mapping has shown loss at chromosome 3p13–p14.3 in 75% (Jones and Nakamura, 1992) and 3p14–p21 in 100% (Yokota et al., 1989) of small series of cervical carcinomas, with a suggestion of LOH at a locus mapping to chromosome 17p (Poulsen et al., 1992). Restriction fragment length polymorphism (RFLP) studies using probes that detect different polymorphisms have shown a somatic loss of chromosome 11 heterozygosity in 30% of primary cervical tumours (Srivatsan et al., 1991). Preliminary data from this laboratory (Bethwaite et al., 1994) and others (Hampton et al., 1994) have demonstrated a high frequency of LOH (43–52%) at loci mapping to 11q23 in cervical tumours by RFLP and microsatellite polymerase chain reaction (PCR) analysis.

LOH has been identified at an anonymous D11S29 locus on chromosome 11q23 in a proportion of breast cancers (60%) (Stickland et al., 1992) and cutaneous malignant melanomas (67%) (Tomlinson et al., 1993). The 11q23 region is of special interest in human oncogenesis, being frequently altered in haematological malignancies, colorectal tumours and ovarian cancer. Previously undertaken RFLP analysis using probes for D11S29 was hampered by low rate of informativeness (Stickland et al., 1992; Tomlinson et al., 1993). However this situation is greatly improved by the identification of a highly polymorphic (GT) microsatellite variable number of tandem repeats (VNTR) within the locus (Warmich et al., 1992). Using probes for this microsatellite polymorphism, we have searched for LOH at locus D11S29 on chromosome 11q23 in 72 cases of invasive carcinoma of the uterine cervix, using the patients' constitutional genotype as a control.

Materials and methods

Materials
Eighty-one cases of invasive cervical carcinoma were retrieved randomly from the archives of the Department of Histopathology, John Radcliffe Hospital, UK (64 cases), and the Department of Histopathology, Johannesburg General Hospital, South Africa (17 cases). All material consisted of formalin-fixed, routinely processed and paraffin wax-embedded surgical specimens. Cases were selected to reflect a balance of histological tumour type only. Nine cases (11%) were excluded from the study as adequate constitutional tissue was unavailable or the tumour tissue showed significant necrosis which often inhibits the PCR.

DNA extraction
Six-micron-thick sections were cut from the tissue blocks selected, fixed to glass slides, deixed, stained with haematoxylin–eosin and dehydrated without a final xylene step. Using a dissecting microscope, approximately 2 mm2 of invasive tumour and constitutional tissue was microdissected from the slide, using a modification of the technique of Pan et al. (1994). The areas of interest were lifted from the slide using a 22 gauge syringe needle fixed to a 1 ml insulin syringe, under a drop of 50% ethanol. In the tumour tissue, three sites were randomly selected from each case, comprising...
approximately 150–200 cells, and digested together. A similar quantity of constitutional tissue was selected usually from normal cervical epithelium, endometrium, pelvic lymph nodes or ovarian tissue. Material from a wax block was included every 20 cases to exclude cross-contamination.

DNA was extracted in sterile microcentrifuge tubes in 25 μl of 100 mM Tris–HCl, 1 mM EDTA and proteinase K (Boehringer Mannheim) at a final concentration of 400 μg ml⁻¹ at 55°C for 2 h, and heated to 95°C for 40 min. Combination of the two steps was found to improve the yield of DNA compared with either proteinase K digestion or heating alone.

**PCR technique**

PCR was performed using oligonucleotide primers flanking an informative (GT), microsatellite polymorphism at D11S29 (Warnich et al., 1992). Reaction volumes were 50 μl and the reaction constituents are given in Table I. Taq polymerase was added to each tube after heating samples in a Perkin-Elmer Cetus thermal cycler to 94°C for 5 min. The samples were cycled using a ‘touchdown’ technique, in which the primer annealing temperature is reduced by 1°C every second cycle until a final temperature is reached (Don et al., 1991).

The amplified product was visualised using a 16 × 16 cm × 0.75 mm, 12% non-denaturing polyacrylamide gel (Sigma Chemicals, UK) run, with cooling, at 8 W for 5 h (Protein II xi cell, Bio-Rad Laboratories, UK). The alleles were revealed after staining with 0.5 μg ml⁻¹ ethidium bromide and UV transillumination. In cases where a partial loss was detected, the gel photograph was scanned into a computerised image analysis system and the relative densities of the allelic bands compared using appropriate software (Quantiscan, Biosoft UK). Because of differential loading or PCR amplification between tumour and normal samples, relative allelic ratios were calculated using the technique outlined by Cawkwell et al. (1993). For a heterozygous sample, the peak areas of the two dominant bands were calculated for each constitutional (C) and tumour (T) sample, and the relative allelic ratio calculated as [T1:T2]/[C1:C2], where T1 and C1 are the values derived from the shorter length allele peak and T2 and C2 from the longer length peak. A priori, a relative ratio of less than 0.50 was chosen as indicative of LOH, based on previous studies (Cawkwell et al., 1993; Orphanos et al., 1993; Hampton et al., 1994).

Informative cases identified as having LOH at D11S29 were subject to further analysis using other informative microsatellite dinucleotide polymorphisms at loci on 11q13 (D11S554) (Hauge et al., 1991) and 11p12–p11.2 (D11S554) (Phromchotikul et al., 1992). The PCR approach was similar; the reaction conditions are given in Table I.

**Results**

DNA amplification of the microsatellite polymorphism at D11S29 from tumour and constitutional tissue was successful in 69 of 72 cases. Despite repeated microdissections, amplification failed from two tumours and one constitutional case, presumably because of variations in fixation techniques. Sixty-two (90%) of the cases were informative for the D11S29 polymorphism. Results of the hybridisation on 20 cases using the (CA), probe confirmed that the bands scored corresponded to the dinucleotide repeat sequences of interest (Figure 1).

Two patterns of loss were detected. The most common pattern showed complete loss of one allele in the tumour sample (16 cases) (Figure 2a). The other pattern showed reduced intensity of one allele without complete loss owing to contamination of the sample by tumour-infiltrating leukocytes (14 cases) (Figure 2b). These 14 cases were subject to

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**Table 1** Parameters for polymerase chain reaction in the amplification of three microsatellite polymorphisms on human chromosome 11

| Parameters                  | D11S29 | D11S554 | D11S554 |
|-----------------------------|--------|---------|---------|
| Extracted DNA               | 5 μl   | 5 μl    | 5 μl    |
| 10 × reaction buffer*       | 5 μl   | 5 μl    | 5 μl    |
| Primer concentration*       | 12.5 pmol | 12.5 pmol | 12.5 pmol |
| dNTP concentration          | 25 μM  | 60 μM   | 60 μM   |
| Taq polymerase              | 0.25 U | 0.5 U   | 0.5 U   |
| PCR cycle number/type       | 35/touchdown | 40/touchdown | 40/touchdown |
| Denaturing                  | 94°C × 1 min | 94°C × 1 min | 94°C × 1 min |
| Annealing – initial         | 72°C × 1 min | 65°C × 1 min | 65°C × 1 min |
| Annealing – final           | 61°C × 1 min | 55°C × 1 min | 55°C × 1 min |
| Extension                   | 72°C × 1 min | 72°C × 1 min | 72°C × 1 min |
| Final elongation            | 72°C × 10 min | 72°C × 10 min | 72°C × 10 min |

*100 mM Tris–HCl, 15 mM magnesium chloride, 500 mM potassium chloride, 0.1% (w/v) gelatin, 1% Triton X-100. The primer sequences (from published data) are as follows: D11S29 primer 1, 5'-TCTAAGCTTCCACCATCGTGT-3'; primer 2, 5'-ACA-ACACACTGCGCACAAAGC-3' (Warnich et al., 1992); D11S554 primer 1, 5'-ATA-TGAGAACCTCTCCGTACT-3'; primer 2, 5'-GAAACATGGAGAAGTCTGGA-3' (Hauge et al., 1991); D11S554 primer 1, 5'-GGTACGCAGGCAAGACTGTC-3'; primer 2, 5'-CACACCTCTCATCTTTAGGCA-3' (Phromchotikul et al., 1992).
Discussion

Allelic loss had initially been detected in tumours using RFLP techniques, which are limited by their expense, time requirements, safety concerns, interpretative difficulties and the high level of non-informative homozygous cases. Many of these drawbacks have been overcome by the use of microsatellites (Cawkwell et al., 1993). These are short tandem repeat sequences exhibiting length polymorphisms which occur throughout the genome and which are highly informative. This study demonstrates that PCR analysis of microsatellite polymorphisms is easily applied to routine paraffin-embedded archival material to provide rapid and sensitive detection of allelic imbalance. Allelic imbalance in tumour tissue is a useful marker for putative allelic loss, although as extracted DNA is not quantified the method does not distinguish allelic deletions from mitotic recombination and allele amplification (Nordenskjöld, 1990).

Tradtitionally, molecular analysis on archival material has involved chemical genomic DNA extractions from whole tissue sections, which often included invasive and in situ tumour tissue together with a variable proportion of constitutional tissue. In addition to obtaining widely varying amounts of DNA from case to case, histological correlations were limited and interpretation of allelic imbalance was subjective. In order to circumvent these problems a range of microdissection techniques have been developed, which allow the selective identification and molecular analysis of cells of interest (Whetsell et al., 1992; Lakhani et al., 1994; Pan et al., 1994).

The modified microdissection technique used in this study was easy and cheap to apply after a short learning period. A 22 gauge syringe needle was found to be as useful and as easy to manipulate as the micropipettes described by Pan et al. (1994). Dissection under a drop of 50% alcohol prevented any flaking and dispersion of tissue and at the same time allowed the dissected tissue to clump and be easily lifted from the slide. The major advantage of the technique was the reduction in 'contamination' of tumour tissue by constitutional cells, hence giving a largely clean pattern of allelic loss in most instances.

Scoring of allelic loss using microsatellites is often undertaken using radiographic techniques. However, a problem with employing dinucleotide repeat polymorphisms is the production of so-called 'stutter bands' caused by slippage of the Taq polymerase in the PCR resulting in production of smaller fragments and making autoradiographs difficult to interpret. However, in the current study, by using a combination of careful optimisation of PCR reaction conditions, very small quantities of PCR primers in each reaction and employing a 'touchdown' PCR technique, we found that 'stutter bands' were minimised and that scoring was possible following ethidium bromide staining alone.

This study adds to our preliminary report (Bethwaite et al., 1994), the first to report allelic loss at locus D11S29 in human cervical carcinoma, 40% of the cases studied showing loss at this locus. We have made the conservative assumption that additional LOH demonstrated using markers for other loci on 11p and/or 11q indicate more general loss of a whole or hemichromosome 11, and therefore these cases have been excluded from further analysis. There is support for this assumption using data from RFLP deletion mapping studies (Srivatsan et al., 1991) and from cytogenetic studies of cervical carcinoma which find such patterns of chromosomal 11 loss in between 10% and 30% of tumours (Atkin et al., 1990; Sreekantaiyah et al., 1991). After exclusion of these cases, allelic loss at locus D11S29 occurred in 30% of cervical tumours in our hands.

Despite cytogenetic data showing chromosome 11 to be over-represented in structural aberrations in cervical malignancy, there has been little previous work looking at the molecular changes on chromosome 11. Srivatsan et al. (1991) using RFLP analysis on 15 cases of cervical carcinoma found a diversity of loci deleted in six tumour samples, the majority at sites on 11q, leading the authors to implicate the long arm as a putative location for a cervical cancer tumour-
suppressor gene. Hampton et al. (1994) reported on an allelic mapping study of 32 cervical carcinomas using polymorphic markers, the majority using PCR-detected microsatellite polymorphisms on chromosome 11. This study found evidence of clonal LOH for one or more markers on 11q in 14 cases; the highest frequencies were observed with polymorphic loci at the D11S144 locus (52%) and the APOC3 gene (43%), both of which map to 11q23.

Foulkes et al. (1993), using microsatellite polymorphisms, produced a deletion map of chromosome 11q in 28 epithelial ovarian tumours. At the D11S29 locus they found LOH in 7 of 15 cases (47%). Interestingly, this study found a higher rate of LOH at adjacent more distal loci, implying that the area of maximum interest for a putative tumour-suppressor gene is distal to the CD3D locus on 11q23.3, which is some 1.8 cm distal from the D11S29 locus. 11q23 has also been of importance in haematological malignancies, with rearrangements of this band being very common in acute leukaemias. Here a number of translocations of the MLL gene (also known as the ALLI or HRX gene) from 11q23 lead to the creation of a fusion gene which encodes chimaeric oncogenic proteins (Rowley, 1993). The role of 11q23 in proto-oncogene activation in cervical malignancy remains to be investigated.

Here the existence of a tumour-suppressor gene on human chromosome 11q fits into our current understanding of the aetiology of cervical carcinoma is uncertain. It is known that transfer of a single human chromosome 11 to either the HeLa (Saxon et al., 1986) or SiHa (Koi et al., 1989) cervical carcinoma cell lines suppresses the tumorigenic phenotype. It is now appreciated that several subtypes of human papillomavirus (HPV) occur frequently in cervical carcinoma and its precursor lesions, and that viral integration into host DNA affecting viral early (E) or late protein-encoding regions appears important in the oncogenic process (Lane and Wells, 1994). Compared with many other tumour types, inactivation of the p53 gene by allelic loss or mutation is an uncommon event in cervical carcinoma, however HPV E6 protein is known to bind to and neutralise normal p53 protein function. HPV integration sites have been shown to be located on chromosomes close to oncogenes, fragile sites and cancer chromosome breakpoints (Cannizzaro et al., 1988), although chromosome 11 has not been implicated (De Braekeleer et al., 1992). Interestingly, in experiments on human fibroblast cells co-expressed with an HPV 16 DNA containing plasmid with an activated EJ-ras oncogene, a translocation, t(1;11), was found to be a frequent event; the breakpoint was invariably at 11q23 (Matlashewski et al., 1988). However, in the present series, LOH at D11S29 was correlated with the presence of HPV as detected by PCR techniques. It is likely that HPV integration is a necessary but not sufficient precondition for cervical cancer, and there is a synergistic role for other oncogenes including c-myc and H-ras (Chen and Defendi, 1992). Loss or inactivation of putative tumour-suppressor genes on chromosomes 3p (Yokota et al., 1989; Jones and Nakamura, 1992) and 17p (Russell et al., 1992) has been reported in a proportion of cervical tumours, and the data from this and other studies suggest that allelic loss in the region 11q22–q24 may be implicated in the chain of malignant transformation in the cervix.

In summary, we have shown that loss of heterozygosity at the D11S29 locus on human chromosome 11q23 occurs in 40% of invasive cervical carcinomas. By excluding those cases where LOH is occurring secondarily to loss of a whole chromosome 11 or its long arm, then we find LOH at D11S29 in 30% of cervical carcinomas. Previous reports have shown LOH at this site in other tumour types (Stickland et al., 1992; Foulkes et al., 1993; Tomlinson et al., 1993) with a high frequency of LOH at 11q22–q24 in small series of ovarian and cervical carcinomas (Foulkes et al., 1993; Hampton et al., 1994), implicating a candidate tumour-suppressor gene distally on chromosome 11q. The findings in the current study, while not providing a detailed deletion map of the 11q region, are consistent with the involvement of chromosome 11q in cervical carcinogenesis. Further work is now under way to produce a more detailed deletion map using other loci on chromosome 11q23 and distal sites in an attempt to pinpoint more accurately the region of interest. In addition, it is important to examine the pattern of loss in high- and low-grade cervical intraepithelial lesions to assess where events on chromosome 11 are acting in the chain of cervical malignant transformation.

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