Genomic homogeneity in fibrolamellar carcinomas

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Abstract

Background—Fibrolamellar carcinoma (FLC) is a variant of hepatocellular carcinoma (HCC) with distinctive clinical and histological features. To date there have been few studies on the genotypic aspects of FLC and no previous attempts have been made to use the arbitrarily primed-polymerase chain reaction (AP-PCR) technique to detect genetic alterations in this disease.

Aim—The aim of this study was to assess the degree of genomic heterogeneity of FLC using the AP-PCR technique.

Methods—A total of 50 tissue samples of primary and metastatic FLCs from seven patients were microdissected. AP-PCR amplification of each genomic DNA sample was carried out using two arbitrary primers.

Results—DNA fingerprints of the primary FLCs and all their metastatic lesions (both synchronous and metachronous disease) were identical in an individual patient. The fingerprints were different between tumours of different patients. No evidence of intratumour heterogeneity was observed.

Conclusions—Such genomic homogeneity in FLCs may explain their indolent growth. The absence of clonal evolution, which is present in other tumours (particularly HCCs), may explain the distinct behaviour in this tumour. The tumorigenic pathway and degree of somatic genomic changes in this disease may be less complex than in HCC.

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Keywords: fibrolamellar carcinoma; hepatocellular carcinoma; DNA fingerprint; arbitrarily primed-polymerase chain reaction; laser capture microdissection

Fibrolamellar carcinoma (FLC) is an uncommon variant of hepatocellular carcinoma (HCC) which has a particular type of histological appearance and clinical behaviour. It constitutes approximately 7% of primary liver cancers and is relatively common among Whites, occurring with equal frequency in men and women. The clinical symptoms are usually non-specific and often comprise discomfort related to the presence of a large hepatic mass. The tumour typically arises in non-cirrhotic liver and has serum biochemical markers that differ from the usual HCC. Alpha fetoprotein is usually normal in FLC but levels of serum vitamin B12 binding globulin and neurotensin are often increased. Histologically, FLC is characterised by large polygonal shaped cells with eosinophilic granular cytoplasm separated into nests and sheets by lamellar bands of collagen. The nuclei are vesicular and contain prominent eosinophilic nucleoli. Pleomorphism and mitoses are minimal.

Surgical resection is the treatment of choice for this tumour. Complete surgical resection results in a cure in 50–75% of patients. Extensive surgical resection is warranted in FLC even with very large tumours. In a large study of 41 patients with FLC, long term survival was frequently achieved with radical surgical treatment; cumulative survival rates at 1, 3, 5, and 10 years were 98%, 72%, 66%, and 47%, respectively. Surgery may even be beneficial with recurrent and metastatic tumour either with or without chemotherapy.

Recently, HCC has been shown to contain subpopulations of heterogeneous cellular differentiation within each tumour which can be demonstrated using the arbitrarily primed-polymerase chain reaction (AP-PCR) technique. This is one of the novel fingerprinting techniques described by Welsh and McClelland (1990) and is widely used to amplify sections of DNA/RNA for identifying gene structure or matching tissue specimens. AP-PCR allows the detection of polymorphisms without prior knowledge of nucleotide sequence and is based on the selective amplification of genomic sequences that, by chance, are flanked by adequate matches to an arbitrarily chosen primer. Despite the obvious clinical differences in the natural history of FLC in comparison with HCC, to date there have been few studies on the genotypic aspects of FLC and no previous attempts have been made to use AP-PCR to investigate evidence of genetic alterations in this disease. In this study, the degree of genomic heterogeneity of FLC was assessed using the AP-PCR technique.

Materials and methods

TISSUE SAMPLES

Tissue samples used in this study were formalin fixed, paraffin embedded tissues of primary and metastatic FLC lesions from seven patients undergoing elective surgical resection. Details of tissue samples are summarised in table 1.

NEEDLE MICRODISSECTION AND DNA PREPARATION (PATIENTS NOS 1–3)

One 5 µm section and 10 µm sections were serially cut from each archival block onto glass slides by microtome. The first 5 µm section was

Abbreviations used in this paper: FLC, fibrolamellar carcinoma; HCC, hepatocellular carcinoma; AP-PCR, arbitrarily primed-polymerase chain reaction; LCM, laser capture microdissection.
stained with haematoxylin and eosin for histological confirmation. The remaining 10 µm sections were stained with 0.1% toluidine blue solution for 10 minutes. Tissue samples were separately prepared by microdissection using sterile 27 gauge needles with the assistance of a dissection microscope and referenced to a section stained with haematoxylin and eosin. Deparaffinisation was achieved by incubating in xylene for 20 minutes at 55°C and then washing twice with absolute ethanol. DNA was prepared by incubating tissue in lysis buffer (10 mM Tris Cl, pH 8.0, 50 mM KCl, 2.5 mM MgCl₂, 0.1 mg/ml gelatin, 0.45% NP-40, 0.45% Tween 20, proteinase K 500 µg/ml) overnight at 55°C. The lysate was subjected to three phenol/chloroform/isoamyl alcohol (25:24:1, by volume) extractions. After ethanol precipitating, washing, and drying the DNA, it was resuspended in sterile water.

LASER CAPTURE MICRODISSECTION (LCM) AND DNA PREPARATION (PATIENTS NOS 4–7)

Two 10 µm sections were serially cut from each archival block onto glass slides by microtome and stained with haematoxylin and eosin. The first section was for histological confirmation. The tissues were separately microdissected by laser capture microdissection (LCM) (Arcturus Engineering Inc., California, USA). The system employs an ethylene vinyl acetate polymer layer coated onto the underside of a rigid flat vial cap. Under the microscope, the cells of choice can be viewed through the film, and the pulsed laser beam activates a precise spot on the transfer film. At this precise location, the film immediately above the cells of interest melts and fuses with the underlying cells. The cap is lifted off the tissue and placed directly onto a 0.5 ml microfuge tube containing 50 µl of proteinase K buffer. The tube was inverted and incubated overnight at 37°C. After the incubation period, the tube was centrifuged at 13 000 rpm for five minutes and the cap was removed. Then the buffer was inactivated at 95°C for 10 minutes and the solution was ready to use as a template for PCR. Two sectors of each section were microdissected separately.

The DNA samples obtained by the microdissection techniques were qualitatively assessed; 4 µl of DNA solution were used for amplification with GAPDH primers in a total volume of 50 µl containing 50 mM KCl, 10 mM Tris Cl, pH 8.3, 1.5 mM MgCl₂, 100 pM primers, and 5 units of Taq polymerase (Bioline, London, UK). Templates were denatured for five minutes at 95°C and subjected to 35 cycles at 94°C for one minute, 55°C for one minute, and 72°C for two minutes. The PCR products were run on 1.5% agarose gel staining with ethidium bromide (data not shown). Fingerprinting of DNA samples was performed using the AP-PCR technique.

AP-PCR AMPLIFICATION

Two different arbitrary primers (AR3: 5'-GGCGATTTCTGCTAGTGCG-3' and ZF3: 5'-GCCGCACGGAGAAGACC-3') were used to amplify the genomic DNA of each sample. These primers have previously been shown to provide useful information on the genotypic profile of HCC. An arbitrary primer (5 µM) was labelled with γ-[32P] (American Pharmacia Biotech, Buckinghamshire, UK) using T4 polynucleotide kinase (Bioline, London, UK) at 37°C for one hour. AP-PCR amplifications were carried out using 200 ng of template DNA in a reaction mixture containing 5 µM 32P labelled arbitrary primer, 0.2 mM each dNTP (Bioline, London, UK), 10 mM Tris Cl, pH 9.2, 3.5 mM MgCl₂, 75 mM KCl, and 0.5 units of Taq DNA polymerase (Bioline, London, UK) in a final volume of 25 µl. The reactions were overlaid with mineral oil and cycled through the following temperature profile: 94°C for one minute to denature, 45°C for...
five minutes for low stringency annealing of primer, and 72°C for five minutes for extension for four cycles. This temperature profile was followed by high stringency cycles: 94°C for one minute, 60°C for one minute, and 72°C for two minutes for 35 cycles, with a final extension at 72°C for 10 minutes. To ensure reproducibility, all amplifications were performed in duplicate on each sample with a Hybaid Thermoreactor (Hybaid, Middlesex, UK). The PCR product was added to 5 µl of dye mixture and was subsequently run on 8% polyacrylamide gel electrophoresis in a Model S2 sequencing gel electrophoresis apparatus (Life Technologies, Paisley, UK). After electrophoresis, the gel was transferred to a piece of Whatman 3MM paper, dried under vacuum, and exposed to x-ray film (BioMax MR, Kodak, USA) for 24–48 hours at room temperature.

**Results**

Comparison between the needle based and LCM techniques have been previously described. Although manual microdissection (patients Nos 1–3) was time consuming, labour intensive, and required a high degree of manual skill, it was successfully used to isolate tumour cells from sheets of collagen. The LCM technique (patients Nos 4–7) was simple, fast, required no moving parts, involved no manual microdissection or manipulations, and enabled one step transfers. Moreover, the manually microdissected cells can detach from the tip of the needle during transfer whereas cells obtained by LCM adhere to the film where they retain their morphological features and the operator can verify that the correct cells have been procured (fig 1). Tumour cells are large and polygonal. Islands of tumour are separated by dense collagenous fibrous tissue.

A total of 50 samples of primary and metastatic FLCS from seven patients were analysed (table 1). Each tumour was microdissected and tissue was taken from two sectors of each tumour (for assessment of intratumoral
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were the same when compared between di
from different patients. The AP-PCR results
were the same when compared between differ-
ent sectors of the same tumour mass, inde-
pendent of whether these were primary or
metastatic lesions. No evidence of intratumour
heterogeneity was observed.

Discussion

Laser assisted microdissection assists in the
molecular study of FLCs because these cancers
contain dense fibrous stromal components
which may complicate analysis of a particular
tumour cell population. The identity of the
transferred cells attached to the film can be
viewed and recorded by image capture. It is
obviously an advantage to use microdissected
cell samples in molecular analysis because the
confounding effect of contaminating cells is
eliminated. An increase in sensitivity of more
than 50% in allelic imbalance analysis has been
reported using microdissected cell populations
compared with crushed tumour samples. Precise microdissection of phenotypically simi-
lar prostatic cancer tissue samples and analysis
of the DNA for loss of heterozygosity revealed
intratumoral genetic heterogeneity and pro-
vided evidence for the multifocality of tumour
development in the prostate. AP-PCR is a
highly specific technique that can generate spe-
cies specific and individual diagnostic DNA
fingerprints. This method has been applied to
detect and isolate novel DNA fragments
associated with somatic genomic changes in
colorectal cancer, lung cancer, and astrocy-
toma. Moreover, it has been shown to be
an uncomplicated and effective approach for
scanning the genomes of tumour samples to
show the evolution of differences.

FLC has distinctive histological and clinical
features. It occurs predominantly in children
and young adults without a predilection for
either sex, and it carries a better prognosis than
the more common HCC. The indolent growth
of FLC is especially obvious when recurrences
develop. Even after the diagnosis of tumour
recurrence, survival at one, three, and five years
has been reported to be 75%, 48%, and 28%
respectively. Several studies have attempted to
associate the differences in natural history of
this cancer from those of HCC. Flow cytometa-
ic analysis suggested that the DNA content in
FLC is not directly related to clinical behav-
ior. In contrast, the overall survival rate of
HCC patients with DNA aneuploid tumours
was significantly poorer than that of patients
with diploid tumours. Positive staining for
neurone specific enolase, the finding of neuro-
secretory type granules at the ultrastructural
level, and raised serum levels of neurotensin
suggest neuroendocrine differentiation of this
disease. Neurotensin, an important regulato-
ry peptide in the gut that facilitates transloca-
tion of fatty acids from the intestinal lumen, is
a useful marker to differentiate FLC from other
tumours. Neurotensin is expressed in fetal
human liver and in FLC, but not in the adult
liver, focal nodular hyperplasia, or in HCC. However, the absence of neurotensin receptor
expression emphasises the fact that neuro-
tensin probably does not play a primary role in
liver growth.

This is the first report of genomic homoge-
inity in FLC and metastatic lesions. Absence
of the clonal evolution that is found in other
tumours, particularly HCC, may explain the
distinct behaviour of FLC. Patients with this
tumour differ from those with HCC in terms of
age at presentation, absence of underlying liver
disease, and tumour markers. These character-
istics may reflect different aetiological factors
because there is no recognised association of
FLC with viral hepatitis, alcohol, cirrhosis, oral
contraceptives, or genetic disorders such as
Wilson’s disease. Tumorigenesis and degree of
somatic genomic changes (rearrangements,
losses, or gains) in this disease may be less
complex than in HCC. None of nine patients

Figure 3 Arbitrarily primed-polymerase chain reaction (AP-PCR) analysis of primary fibrolamellar carcinomas (FLC) and their metastatic lesions from patient No 6. Autoradiograms of polyacrylamide gel electrophoresis of 32P labelled DNA fragments amplified by AP-PCR. Genomic DNA from each sample was amplified with the AR3 (A) and ZF3 (B) primers. P, primary FLC; LN, lymph nodes metastases; (number of sectors is represented at the top).
with FLCs had p53 mutations (exons 5–8) analysed by denaturing gradient gel electrophoresis, single strand conformation polymorphism, and gene sequencing. Moreover, in a loss of heterozygosity study using restriction fragment length polymorphism probes for various chromosomes, the rate of allelic loss in FLC was infrequent (3.6%) compared with that in HCC (16.1%). The cell of origin of FLC remains uncertain. Whether FLC and HCC represent extremes of the spectrum of one disease entity remains uncertain. Interestingly, there are few reports of the synchronous development of FLC and ordinary HCC.

There is no defined at risk group to screen other than the young, and thus there is little prospect of improving the prognosis in this way.

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