Production and characterisation of a monoclonal antibody to human papillomavirus type 16 using recombinant vaccinia virus

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Abstract
A monoclonal antibody was raised against the major capsid protein L1 of human papillomavirus type 16, using a recombinant vaccinia virus that expresses the L1 protein, as a target for screening. This antibody, designated CAMVIR-1, reacted with a 56 kilodalton protein in cells infected with L1-vaccinia virus, and the protein was present in a predominantly nuclear location. The antibody also detects the HPV-16 L1 antigen in formalin fixed, paraffin wax embedded biopsy specimens and on routine cervical smears. The antibody reacts strongly and consistently with biopsy specimens containing HPV-16 or HPV-33, but very weak reactions were occasionally observed with biopsy specimens or smears containing HPV-6 or HPV-11.

The potential advantages of using a vaccinia recombinant are (i) the target protein is synthesised in a eukaryotic cell so that its “processing” and location are normal; (ii) cells infected with vaccinia recombinants can be subjected to various fixing procedures similar to those used for routine clinical material. This greatly increases the probability that an identified antibody will be useful in a clinical setting.

Human papillomaviruses (HPV) are small DNA viruses which cause a variety of proliferative lesions on epithelial or mucosal surfaces. More than 50 HPV types have now been identified, and several of these are associated with premalignant and malignant disease of the anogenital region. In particular, HPV 6 and 11 have been identified in benign lesions; HPV 16, 18, 31 and 33 are highly significantly associated with all grades of cervical intraepithelial neoplasia (CIN) and cervical cancer.

There is no suitable tissue culture system for the propagation of these viruses and the identification of viral DNA and classification of viruses is based on DNA homology using hybridisation techniques. Viral DNA sequences have been identified and typed in abnormal cervical lesions and in cervical smears using blot hybridisation techniques, and more recently by the polymerase chain reaction. The exquisite sensitivity of the latter has resulted in the detection of low copy viral DNA in histologically and cytologically normal subjects. The ubiquity of viral DNA highlights the need for reagents that will detect viral proteins. Several groups have recently tried to overcome the problems associated with the lack of a permissive in vitro culture system by cloning and expressing viral genes in prokaryotic and eukaryotic expression systems.

In an earlier paper we reported the construction of a vaccinia virus recombinant containing the HPV 16 L1 gene (encoding the major capsid antigen), and the preparation of polyclonal antisera raised against a bacterial fusion protein containing HPV 16 L1 sequences.

Polyclonal antisera, however, are of limited use in that they cannot give information about the presence of type specific epitopes on the L1 protein and tend to result in high background in immunohistochemical and immunoprecipitation studies. There is a need for type specific reagents for diagnostic as well as for research purposes for viruses such as HPV 16, which has been identified in 50–60%, of cases of cervical carcinoma subjected to viral typing.

In this report we describe the production of a monoclonal antibody to HPV 16 L1 using a novel protocol in which viral antigen is generated for immunisation using a bacterial expression system and hybridoma supernatants are screened for specific antibody using a vaccinia recombinant expressing the target protein.

Methods
The myeloma cell line NSO and established hybridoma cell lines were maintained in Glasgow modified Eagle’s medium (GMEM) containing 5% fetal calf serum (FCS). African green monkey kidney (CV-1) cells were maintained in the same medium containing 10% FCS. Baby hamster kidney 21 clone 13 cells (BHK) were maintained in GEME containing 10% tryptose phosphate broth and 10% newborn calf serum. The construction of recombinant vaccinia virus (VL1RK) expressing the entire HPV 16 L1 open reading frame under the control of a vaccinia late promoter has been described previously. The recombinant virus and the parent WT vaccinia virus (strain WR) were grown in BHK cells and titrated in CV-1 cells.

MOUSE IMMUNISATION AND CELL FUSION
Mice were immunised subcutaneously with 20 μg of a β galactosidase-L1 fusion protein (containing amino acids 198-C terminus of HPV-16 L1) in complete Freund’s adjuvant. After
four weeks a series of subsequent doses of 15 µg protein in Freund’s incomplete adjuvant were given intraperitoneally at two weekly intervals. A final intravenous immunisation of 15 µg was given three days before the fusion. The spleen was removed aseptically and fused to the mouse hybridoma NSO, using the original protocol. Cells were plated at a density of 2 x 10⁶ well in 24-well Linbro plates, and the growth medium changed every two to three days. Positive hybrids were cloned twice by limiting dilution, and clones secreting antibody to L1 were grown in 75 cm² flasks and stored in liquid nitrogen.

SCREENING OF HYBRIDOMA SUPERNATANTS USING ELISA
The supernatants of wells containing hybrids were tested in an ELISA as follows: CV-1 cells were grown in 5 mm wells to confluence and infected with 30 plaque forming units/cell WT or VL1RK vaccinia virus. After 18 hours at 37°C cells were fixed in 5% formaldehyde, 10% sucrose, in phosphate buffered saline (PBS), and air dried. Before use cells were permeabilised with 1% Triton X-100, 10% sucrose in PBS. After blocking with 5% FCS, monolayers were reacted with hybridoma supernatants, followed by incubation with peroxidase-linked protein A, each for 30 minutes at room temperature. The wells were then reacted with O-phenylene diamine and the absorption of the supernatants at 492 nm was measured in an ELISA reader.

IMMUNOFLOUORESCENCE
CV-1 cells were grown on coverslips and infected, fixed, and permeabilised as described for the ELISA method. Cells were reacted with hybridoma supernatant and subsequently incubated with an anti-mouse IgG fluorescein conjugate.

WESTERN BLOT ANALYSIS
BHK-cell monolayers, infected with 30 pfu/cell WT or VL1RK vaccinia virus, were harvested in PBS 20 hours after infection, lysed in sample buffer (24 mm TRIS-HCl, pH 6-18; 100 mm dithiothreitol; 2% sodium dodecyl sulphate (SDS); 20% glycerol; 0-02% bromophenol blue), and sonicated. Samples were denatured at 100°C for three minutes, subjected to SDS-polyacrylamide gel electrophoresis (PAGE) and the proteins transferred to nitrocellulose by electroblotting as described previously. After blocking in 5% dried milk powder in PBS for 30 minutes at room temperature, filters were incubated with 1/2 diluted supernatant for one hour at room temperature, washed three times in PBS-1% NP40 and reacted with affinity purified 125I-protein A (Amersham International, UK) at a concentration of 0.1 µCi/ml. The blots were washed again in PBS-1% NP40, dried, and autoradiographed.

IMMUNOCYTOCHEMICAL STUDIES
Formalin fixed, paraffin wax embedded sections were obtained from diagnostic biopsy specimens taken from patients attending a routine colposcopy clinic. Sections were dewaxed and rehydrated through a xylene alcohol series. Before immunostaining with the Vectastain ABC complex (Vector Laboratories, Peterborough, England) sections were incubated in 0.1% trypsin at 37°C for five minutes and quenched in 0.3% H₂O₂ for 15 minutes.

Routine cervical Papanicoula smears were fixed in 5% formaldehyde in PBS for five minutes, washed in PBS, permeabilised in 1% Triton and subsequently treated with trypsin, quenched, and stained in the same way as the cervical sections.

IN SITU HYBRIDISATION
Sections of cervical lesions embedded in paraffin wax were dewaxed in xylene and alcohol and hybridised with probes containing DNA of HPV 16/18, HPV 31/33, or HPV 6/11, using the Vira Type in situ hybridisation kit (Life Technologies Inc, Gaithersburg, Maryland, USA) according to the manufacturer’s instructions. In certain cases in situ hybridisation was carried out using 35S-labelled DNA, according to the method described by Gowans et al. Briefly, slides were rehydrated, digested with proteinase K and pronase, and treated with acetic anhydride before hybridisation with 35S-labelled HPV-16 DNA. After coating with radiographic emulsion slides were stored for eight days at 4°C and developed, fixed, and stained with haematoxylin.

Results
Mice repeatedly immunised with the β galactosidase-HPV 16 L1 fusion protein developed strong responses to the fusion protein, but the response to the vaccinia recombinant expressing HPV-16 L1 was variable. Some mice also gave high “background” responses to cells infected with wild type vaccinia virus. Spleen cells from mice showing a good differential response in ELISAs to the L1 recombinant vaccinia virus relative to the wild type virus were used in fusions with cell line NSO. The results of a series of three fusions were disappointing in that only two positive hybridoma supernatants were identified by ELISAs on cells infected with VL1RK. Of these, only one retained activity following two rounds of cloning by limiting dilution. This hybridoma was stable to subsequent passage and subcloning and was designated CAMVIR-1.

PROPERTIES OF CAMVIR-1
Tissue culture supernatant from CAMVIR-1 reacted with a 56 kilodalton protein in Western blots of cell lysates infected with VL1RK (fig 1a). A minor species of 64 kilodaltons was also detected. This probably represents a glycosylated form of the protein as it is absent in lysates of cells infected in the presence of tunicamycin, an inhibitor of N-glycosylation. Similar results were reported using a polyclonal rabbit serum raised against an HPV-16 L1 fusion protein. No reactivity was observed against cells infected with wild type vaccinia. The antibody also reacted with the fusion protein used for im-
Reactivity of CAMVIR-1 on tissue sections

The ability of CAMVIR-1 to detect the L1 protein in formalin fixed, paraffin wax embedded tissue sections was examined using a biotin-streptavidin-peroxidase staining method. Sections were obtained from the cervical biopsy specimens of patients with abnormal cervical smears, attending a routine clinic for colposcopy and examination. These lesions were histologically graded as CIN I to III. The sections were tested for the presence of L1 protein using CAMVIR-1 tissue culture supernatant at a concentration of 1/100. Figure 2a shows a lesion in which the nuclei of several cells in the superficial layers are strongly stained. These cells have the morphology of koilocytes. A parallel section of this lesion was subjected to in situ hybridization to establish the presence and type of viral DNA (fig 2b). The cells with the maximum copy number of HPV 16 DNA are found in the same region of the section as the cells containing L1 protein—that is, in the superficial layers of the lesion. We examined a substantial number of lesions (n = 50), and in those cases where staining with CAMVIR-1 was observed, in situ hybridisation showed high copy numbers of HPV-16 DNA in that region of the section. The opposite was not true, however—that is, a minority of lesions were positive for HPV-16 by in situ hybridisation but not with an irrelevant fusion protein containing the N-terminal portion of the HPV-16 L1 protein (amino acids 1-197).

To determine the cellular localisation of the antigen detected by CAMVIR-1, cells were infected with recombinant or WT virus and after formaldehyde fixation reacted with CAMVIR-1 and fluorescein-labelled antimouse immunoglobulin. In cells infected with L1 vaccinia strong nuclear fluorescence was observed, while no reaction could be detected in cell monolayers infected with WT virus (figs 2g and h). An irrelevant monoclonal antibody of the same subclass (IgG2a) did not react with the recombinant infected cells.

Discussion

The absence of a suitable in vitro system for the propagation of human papillomaviruses, together with the fact that in clinical lesions only very low levels of virus are present, has necessitated the use of alternative strategies for the production of immunological reagents to these viruses. In the production of monoclonal antibodies suitable targets for the screening of hybridoma fluids are an important requirement. To date, the monoclonal antibodies reported have been raised and screened against bacterial derived proteins. We used cells infected with a vaccinia virus recombinant expressing the HPV 16 L1 open reading frame as a target for the selection of monoclonal antibodies. This system provided a very convenient and plentiful source of viral protein, free of potentially crossreactive bacterial elements.

The antibody against the HPV-16 L1 pro-
Figure 2
Immunocytochemical staining with CAMVIR-1 of sections of cervical biopsy specimens containing HPV-16 or HPV-33, routine cervical smears, and cells infected with the L1 vaccinia recombinants. (a) Staining with CAMVIR-1 at 1/100 dilution on formalin fixed, paraffin wax embedded section of a cervical lesion containing HPV-16. Sections were dewaxed and rehydrated through a xylene alcohol series, incubated in 0.1% trypsin at 37°C for five minutes, quenched in 0.3% H2O2 for 15 minutes, and stained with the Vectastain ABC complex. (b) In situ hybridisation using 32P-labelled total HPV-16 DNA on the section adjacent to that in fig 2a. (c) Staining with CAMVIR-1 at 1/100 on a cervical lesion containing HPV-33 using the same protocol as in fig 2a. (d) Staining of the same lesion as in fig 2c, using a commercially available polyclonal anti-BPV-16 antibody (Dako) at 1/100, followed by the Vectastain ABC complex. (e and f) Staining with CAMVIR-1 at 1/100 of routine cervical smears fixed in 5% formaldehyde-PBS for five minutes, permeabilised in 1% Triton, and subsequently treated with trypsin, quenched, and stained in the same way as the cervical sections. (g and h) Reactivity of CAMVIR-1 at 1/100 dilution on CV-1 cells infected with 30 pfu/cell L1 vaccinia (g) or wild type vaccinia (h). Cells were fixed in 5% formaldehyde-PBS for five minutes, washed in PBS, and after incubation with CAMVIR-1 for 30 minutes at room temperature, washed again in PBS, and reacted with FITC-labelled anti-mouse IgG at 1/70 (Dako).

Monoclonal antibody to HPV 16

We also obtained nuclear staining of morphologically koilocytic cells on formalin fixed, paraffin wax embedded tissue sections using CAMVIR-1. It was clear from the subsequent typing of these sections using in situ hybridisation that CAMVIR-1 binds strongly to cells containing HPV-16 or -33. Very occasionally, however, a number of cells in the superficial layer of an HPV-6 containing lesion would show nuclear staining. From the results on cervical smears it is clear that cells containing very high copy numbers of HPV-11 also give positive results with CAMVIR-1. A substantial number of lesions containing HPV-6, however, were totally negative when reacted with CAMVIR-1. The antibody probably binds, to a much lesser extent, to cells containing HPV-6 or -11, so that only cells with a very high viral DNA copy number show positive staining. The epitope to which CAMVIR-1 is directed was identified as GFGAMDF (a.a.number 230–236) using overlapping peptides in an ELISA (Cason J, McCance D, personal communication). This region is highly con-
served among the different HPV types, and there is one amino acid substitution between HPV16 and HPV-6 or -11 in this region, namely, D to N (GGFAMMFN), which would explain the weak cross reactivity with these virus types. CAMVIR-1 identifies an epitope on HPV-16 L1 that is strongly reactive against HPV-16 and -33 and, to a much lesser extent, with HPV-6 and -11, and thus is able to identify a subgroup of human papillomaviruses with malignant oncogenic potential.

Expression of the L1 capsid protein, as identified with this monoclonal antibody, is strongly focal, and in any one lesion only a few cells with high copy viral DNA synthesise L1 protein. This situation differs from that seen in some other lesions induced by papillomavirus, particularly cutaneous warts19 and bovine papillomomas20 in which many cells express viral capsid proteins and in which viral particles are present in substantial numbers. The focality of L1 expression in the cervical lesions may be related to the paucity of viral particles in lesions containing HPV 16. Alternatively, it may be related to the stage of infection at which the cervical lesion has been biopsied. The natural history of HPV-16 related disease is virtually unknown. It is evident from studies on other papillomaviruses, particularly CRPV and BPV-1, that the development of a lesion in which viral capsid protein and viral particles are produced requires a considerable time period—of the order of months.21 Anecdotal evidence in human disease suggests that a similar time period is involved in the development of genital condylomata.22 In many patients biopsy specimens may be taken at a stage in the natural history of the disease when viral particles are not produced, either because the lesion is at an early stage or because a quiescent or persistent phase is another feature of the disease process. Whatever the explanation it is evident that L1 capsid protein is synthesised in a few cells containing high numbers of viral DNA copies in the superficial, terminally differentiated surface layers of the epithelium.

Detection of viral DNA only informs about presence of the genome and gives little information on virus replication and pathogenesis. This is particularly important in that papillomaviruses are strictly epitheliotropic, and the available information indicates that permissive virus replication only occurs in keratinocytes and specifically in the keratinocyte undergoing the normal process of differentiation in the epithelium. The intra-epithelial neoplastic lesions of the cervix are characterised by a progressive loss and distortion of the normal differentiation process, and these features are included in the diagnostic criteria of these lesions.23 Changes in the expression of viral proteins may be a reflection of neoplastic phenotype and may give useful information on progression and prognosis, and the antibody CAMVIR-1 may be a useful reagent in this context.

We are grateful to Drs Lynn Kennedy and Ian Frazier of the Department of Medicine and Lions Immunology Laboratory, University of Queensland, Australia, for the provision of HPV16 positive smears; the Histopathology Department, Addenbrooke’s Hospital, Cambridge, for tissue sections of cervical biopsies; and to Mrs Mary Wright for typing the manuscript.