Characterisation of adenoid cystic carcinoma of the breast by immunohistology

W DÜE, H HERBST, V LOY, H STEIN
From the Institute of Pathology, Klinikum Steglitz, Free University of Berlin, West Germany

SUMMARY An adenoid cystic carcinoma of the breast in a 78 year old woman was analysed immunohistologically for the production of type IV collagen, the expression of vimentin, epithelial membrane antigen (EMA) and steroid receptors, and the proliferative activity of the tumour cells. The data were compared with those obtained in eight adenoid cystic carcinomas of salivary glands and in ductal carcinomas of the breast with a cribriform growth pattern. The patients' ages were as follows: 45–80 years (mean 63:2) for the salivary gland carcinomas; 37–69 years (mean 50:6) for the ductal breast carcinomas. In contrast to the cribriform spaces of ductal carcinomas, the pseudocysts in adenoid cystic carcinomas were lined by type IV collagen. The opposite pattern was observed for EMA. Like the myoepithelium of normal breast, the myoepithelium-like cells of adenoid cystic carcinoma stained positive for vimentin while the ductular epithelium-like ones did not. All adenoid cystic carcinomas, including that of the breast, were negative for the oestrogen and progesterone receptors, unlike the ductal carcinomas. Proliferative activity of the adenoid cystic carcinoma of the breast was relatively low.

These data broaden the range of antibodies suitable for differential diagnosis of both tumour types. They may explain the differences in prognosis, and they explain why hormonal treatment or radiotherapy of adenoid cystic carcinoma of the breast are often ineffectual.

While common in salivary glands, adenoid cystic carcinomas of the breast are rare. They are composed of duct lining and myoepithelium-like cells with a dimorphic differentiation into ductular and pseudocystic spaces. Because pseudocystic areas can be more numerous than the ductular ones, adenoid cystic carcinoma of the breast can easily be confused with ductal carcinoma with a cribriform growth pattern. This differential diagnosis, however, is of some interest, because the first of these tumours is associated with a better prognosis. While both can be distinguished by their distribution of laminin and fibronectin, this seems not to be true for type III collagen.

In the present study the production of type IV collagen, as a further component of the basement membrane, and the expression of vimentin were immunohistologically examined in adenoid cystic carcinomas of salivary glands and of the breast, and were compared with those of ductal breast carcinomas with a cribriform growth pattern. From results of previous immunohistological and ultrastructural studies of basement membrane distribution, it was thought that both tumour types should also differ in their pattern of epithelial membrane antigen (EMA) distribution, the determination of which could provide additional information for differential diagnosis. Because this differential diagnosis may be relevant not only for prognosis but also for treatment, the expression of oestrogen and progesterone receptors was also determined immunohistologically. As prognosis may be influenced and explained by differences in the proliferative activity of tumours the growth fraction was examined in both tumour types using the monoclonal antibody Ki-67, which detects an antigen expressed in all stages of the cell cycle except G0.

Material and methods

The adenoid cystic carcinoma of the breast was obtained from a 78 year old woman. The tumour had first been diagnosed five years previously through the resection of a 50 × 30 × 30 mm mass from between the upper quadrants. Histologically, the carcinoma showed an infiltration of the corium and was well circumscribed and surrounded by fibrotic breast tis-
Adenoid cystic carcinoma of the breast

The patient was then admitted for follow up without further treatment. In May 1988 a newly developed nodule with a maximal diameter of 9 mm, located beside the areola, was resected from the right breast and again diagnosed as adenoid cystic carcinoma with infiltration of the corium. Simple mastectomy was carried out and a biopsy specimen showed no remaining tumour, but fibrosis and mild chronic inflammation of the breast. According to the grading system of Szanto et al., which was developed for adenoid cystic carcinomas of salivary glands, the tumour was judged to be grade II. No secondary tumour was present.

For comparison, eight adenoid cystic carcinomas of major and minor salivary glands were randomly selected from the files of our laboratory. According to Szanto et al., three had to be classified as grade I, one as grade II, and four as grade III. Seven ductal carcinomas of the breast were also selected, which contained areas of cribriform growth and which had to be classified as grade II, according to Bloom and Richardson. Six of them were invasive; the seventh was exclusively intraductal. Cryostat sections of all breast tumours were also available.

Immunohistological staining was done using the alkaline phosphatase-antialkaline phosphatase staining procedure, as modified by Stein et al. The sections were examined using two monoclonal antibodies directed against different epitopes of type IV collagen—NC1, directed against the carboxy-terminal cross-linking domain; CIV 22, kindly donated from Dr Odermatt, Zurich, Switzerland, directed against a conformational epitope on type IV collagen. The slides were also stained for vimentin (V9 Dako), EMA (E29 Dako), as well as against the oestrogen receptor (ER-ICA, Abbott) and the progesterone receptor (mPRI, TransBio). The antibody Ki-67, which detects a proliferation-associated antigen like NC1 is only suitable for cryostat sections. For sections fixed in formaldehyde and embedded in paraffin wax, a pre-treatment was necessary for valid detection of the oestrogen receptor.

Results

Adenoid Cystic Carcinoma of the Breast

The adenoid cystic carcinoma of the breast showed the typical differentiation into myoepithelium-like cells and cells resembling ductular epithelium, with a formation of small ductules and predominating areas.

Fig 1 (a) Adenoid cystic carcinoma of the breast composed of pseudocystic and ductular spaces (arrow). Oestrogen receptor is not detected in both tumour components. (ER-ICA.)
(b) The growth fraction, as indicated by the nuclear reaction product, is about 10%. (Ki-67.)
of pseudocystic appearance (fig 1a). Nevertheless, the ductules were easily detectable by routine hematoxylin and eosin staining. In 1985 and 1988, pre-existing ductules were interspersed, partly with mild epitheliosis. Solid areas were present, but they comprised less than 10% of the tumour mass.

Both antibodies against type IV collagen showed a distinct laminar staining pattern which surrounded both the outer border of all tumour growth and the inner "surface" of pseudocystic spaces. The inner surface of the ductules, which by conventional light microscopy seemed to form true spaces, showed no staining for type IV collagen (fig 2a). The reaction for EMA showed the opposite pattern. The pseudocystic spaces and their lining epithelium did not stain while the cells lining true ductules showed a discrete reaction that was accentuated at the luminal cell border. The content of the true lumina also stained slightly (fig 3a).

A dot-like reactivity was observed in a minority of those cells designated as myoepithelium-like. The oestrogen and progesterone receptors were not shown in any of the tumour cells while there was a distinct reaction within the interspersed pre-existing ductular epithelium, which served as an internal control. Vimentin did not stain the ductular epithelium within the tumour. In contrast, nearly all myoepithelium-like tumour cells were intracytoplasmatically stained for vimentin, resulting in a cord-like boundary of the pseudocystic spaces (fig 4a). All these results were identical for cryostat and paraffin wax sections. The immunohistologically determined growth fraction was about 10% (fig 1b); only myoepithelium-like cells were identified as proliferating.

ADENOID CYSTIC CARCINOMAS OF SALIVARY GLANDS

The antigenic pattern was exactly the same for all tumours as that reported for the adenoid cystic carcinoma of the breast. Even those tumour cell groups at the margin, which were clearly infiltrating skeletal muscles, were completely surrounded by type IV collagen, and this was true for all the tumour manifestations forming the outer margin of the whole tumour.

DUCTAL CARCINOMA OF THE BREAST

Both type IV collagen antibodies showed that basal...
Adenoid cystic carcinoma of the breast

lamina surrounded the intraductal and some of the larger invasive tumours at the centre as well as at the border of the carcinomas. Basal lamina were not detected at invasive tumour sites with strong tumour cell differentiation. In contrast to adenoid cystic carcinoma, no corresponding reaction product was seen to line the luminal cells as part of a cribriform growth pattern (fig 2b). Instead, the tumour cells secreted EMA, resulting in a positive staining reaction within the tumour cells, which comprised the cribriform pattern, and this was also seen within the spaces themselves (fig 3b). Vimentin was not expressed in tumour cells (fig 4b). The average fraction of positively staining cells was 72% for oestrogen receptors, 55% for progesterone receptors, and 18% for proliferative activity. One case was negative for both steroid receptors.

The data on the staining of the steroid receptors with Ki-67 for ductal carcinomas are summarised in table 1. The results for both tumour types are shown in table 2.

Discussion

Adenoid cystic carcinoma of the breast is rare. According to Azzopardi, it comprises less than 1% of all breast carcinomas. Although comprising only minor parts of the tumour, the presence of numerous ductular formations allowed this tumour type to be identified easily in our case. This corroborates the statement that most adenoid cystic carcinomas of the breast can be diagnosed by routine light microscopic examination.

Adenoid cystic carcinoma may be confused, however, with ductal carcinoma with a cribriform growth pattern. This resulted in the rejection of some cases suggestive of adenoid cystic carcinomas of the breast. Further investigations are obviously necessary in difficult cases. If immunohistological analysis could help in differential diagnosis this would be more suitable than the use of electron microscopy, suggested by Harris in 1977.

Adenoid cystic carcinomas have been identified...
morphologically at different anatomical sites. In conjunction with the ultrastructural demonstration of basal lamina lining the pseudocysts, immunohistological methods in the breast and in salivary glands have shown these to be continuously surrounded by the basement membrane proteins laminin and fibronectin. The distribution of type IV collagen as a further component of the basement membrane was expected to show the same result. To our knowledge, however, this has not been shown to date. Thus, our results obtained with antibodies against type IV collagen expand the range of antibodies suitable for differential diagnosis of adenoid cystic carcinoma and ductal carcinoma of the breast, and they further substantiate the basement membrane production of those tumour cells lining the pseudocysts. The immunostaining of their content is explained by the presence of basal lamina material in these “spaces”.

In contrast to ductal breast carcinomas, the lining of the outer margin of infiltrating tumour cell complexes was always present in the adenoid cystic carcinomas.
From these data, and the detection of EMA in normal breast epithelium and in malignant breast tissue,\textsuperscript{25-32} it was concluded that the staining for EMA would show an inverse distribution. EMA had been detected in five of 37 adenoid cystic carcinomas of the breast by Sumpio et al.,\textsuperscript{30} but they did not specify to which tumour component this referred. According to our results, immunostaining for EMA is suitable for the identification of adenoid cystic carcinomas of the breast, because the pseudocysts do not react while the epithelium lining the cribriform spaces of ductal carcinomas shows a distinct reaction, particularly at their luminal border. The spaces themselves are filled with immunostained material. The ductular spaces within the adenoid cystic carcinomas are also lined by cells positive for EMA, which lends further support to the hypothesis that there are two distinct cell types in this tumour.

Although the origin of adenoid cystic carcinoma is disputed,\textsuperscript{9,10} that there is a myoepithelial-like differentiation of those tumour cells which line the pseudocystic spaces is widely accepted.\textsuperscript{12,33-34} Correspondingly, the presence of actinomyosin has been shown within these cells by immunohistological methods.\textsuperscript{35} Because the myoepithelium of normal breast tissue stained positive for vimentin in our hands, our data on adenoid cystic carcinomas provide further evidence for the phenotypic similarity of both cell populations. This, however, does not mean that the tumour cells are derived from the myoepithelium. The presence of two epithelial tumour components favours instead the hypothesis that the tumour is derived from a highly potent precursor cell.\textsuperscript{12}

Steroid hormone receptors were not detected in any of the adenoid cystic carcinomas. In contrast, six of our seven ductal carcinomas stained positively with most tumour cells reacting for both receptors (table 2). This corresponds with those data obtained on ductal carcinomas for both immunohistological\textsuperscript{15} and biochemical methods,\textsuperscript{36,37} and it coincides with biochemical data for five cases of previously published cases of adenoid cystic carcinomas of the breast.\textsuperscript{11,12} Furthermore, it explains the negative result of anti-oestrogenic treatment in one case of adenoid cystic carcinoma of the breast.\textsuperscript{13} Only one case has been reported to be positive for oestrogen receptors by the dextran-coated charcoal method,\textsuperscript{35} but determination of the biochemical receptors excludes detailed information about the tissue components. This result could thus be due to pre-existing, receptor positive residual cells of breast tissue, which may be interspersed as seen in our case. Immunohistological determination of hormone receptors is therefore preferable to biochemical assay.

Finally, the adenoid cystic carcinoma of the breast reported here showed low proliferative activity with the antibody Ki-67. This parallels the observation of low mitotic counts in this tumour.\textsuperscript{12} Unfortunately, snap frozen material was not available in any case of adenoid cystic carcinoma of the salivary glands. Because adenoid cystic carcinomas at both sites share a prolonged clinical course,\textsuperscript{1,4,6,30} differences in prognosis\textsuperscript{38} may be due to local conditions for detectability and radical surgery\textsuperscript{4} rather than to proliferating activity alone. Because growth in breast carcinomas depends on the grade of differentiation\textsuperscript{39,40} grade II ductal carcinomas were taken for comparison in this study, although two different classification systems were applied. Our results for the grade II ductal breast carcinomas with cribriform growth patterns show a somewhat larger growth fraction than has been previously reported by Pickartz et al.\textsuperscript{39} for unspecified invasive breast carcinomas with intermediate differentiation, but they are nearly identical with those reported by Barnard et al., as estimated from their fig 3.\textsuperscript{40} The proliferative activity in breast carcinomas shown by Ki-67 has also been determined by Gerdes et al.,\textsuperscript{41} but neither an average growth fraction nor a specification referring to the tumour grade was reported. The relatively high growth activity may partly explain the poorer prognosis of ductal carcinomas compared with adenoid cystic carcinomas of the breast.\textsuperscript{42} More data, however, are necessary to draw definite conclusions in this respect.

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References


Requests for reprints to: Dr W Düe, Institut für Pathologie, Klinikum Steglitz der Freien Universität Berlin, Hindenburgdamm 30, D-1000 Berlin 45, West Germany.