Nitric oxide synthase activity in human breast cancer

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Summary Nitric oxide (NO) is generated by a family of isoenzymes (NO synthases) expressed in a wide range of mammalian cells. We have recently reported NO synthase expression in human gynaecological cancers. In this study we have assessed the activity and distribution of NO synthase in a series of human breast tumours and in normal breast tissue. Calcium-dependent (constitutive) and -independent (inducible) NO synthase activity, as well as NO biosynthesis, was high in invasive tumours compared with benign or normal tissue. Furthermore, for invasive ductal carcinomas, NO biosynthesis was significantly greater for grade III compared with grade II tumours. Immunohistochemical investigations revealed immunolabelling with a monoclonal antibody to murine inducible NO synthase predominantly within tumour-associated macrophages. Immunolabelling with a polyclonal antiserum raised against rat brain NO synthase was also observed in vascular endothelial and myoepithelial cells. Thus NO synthase is expressed in human breast tumours, where its presence correlates with tumour grade.

Keywords: nitric oxide; human; cancer; breast; macrophage

Since the discovery in 1987 that vascular endothelial cells are able to synthesise NO from l-arginine (Palmer et al., 1987), the existence of this biochemical pathway in many other cell types has been thoroughly documented and its relevance in biology is becoming apparent (Moncada et al., 1991). This inorganic free radical gas, synthesised by a family of isoenzymes called NO synthases (Knowles and Moncada, 1994), plays a vital role as a cell signalling molecule in the vascular, nervous and immune systems (Moncada et al., 1991). It is also a cytostatic cytotoxic mediator when generated at higher concentrations by activated macrophages and endothelial cells (Marletta et al., 1988; Li et al., 1991). The role of NO in tumour biology, however, is poorly understood.

Several tumour cell lines express NO synthases (Forstmann et al., 1990; Sherman et al., 1993; Werner-Felmayer et al., 1993; Jenkins et al., 1994), and we have recently reported NO synthase activity in human ovarian and uterine tumour tissue (Thomsen et al., 1994). The correlation observed between enzyme activity and tumour grade suggests that NO plays a role in the biology of these cancers. In the present study we have assessed the activity of NO synthase and its cellular localisation in a series of human breast tumours.

Materials and methods

Materials

All chemicals were from Sigma, Boehringer Mannheim or BDH, unless otherwise indicated. L-[U-14C]Arginine was from Amersham, UK.

Tissue collection

Tissue was collected from 34 breast tumours. Normal breast tissue from six reduction mammoplasty specimens provided a control group. Pieces of tissue (0.5–2.0 g) were divided into two. One was placed immediately into medium (Dulbecco’s MEM, Gibco, Life Technologies, Paisley, UK) and prepared for in vitro culture. The other was snap frozen and stored at −70°C for subsequent assay for NO synthase and immunohistochemistry studies.

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Determination of NO biosynthesis

Tissue was divided into 1 mm3 pieces, washed twice with culture medium and placed into 96-well plates with 200 µl of culture medium with or without L-5-monomethyl-l-arginine (L-NMMA). After 24 h culture at 37°C, in an atmosphere of 95% air 5% carbon dioxide concentration of nitrate + nitrate in supernatants was analysed after reduction of nitrate using cadmium (Thomsen et al., 1991). Nitrate was measured by chemiluminescence (Palmer et al., 1987). NO biosynthesis was determined by the difference in nitrite + nitrate concentrations in culture medium supernatants from wells containing tissue biopsies relative to wells with culture medium alone. Nitrite + nitrate concentration in culture medium alone ranged from 0.5 to 1.0 µM in all experiments. Total protein content of tissue pieces was determined with bichoninic acid reagent after solubilising the tissue with 1 M sodium hydroxide. The limit of detection of nitrate by chemiluminescence was 0.1 µM, equivalent to approximately 0.15 mmol mg−1 protein. Results are expressed as mean ± standard error of the mean (s.e.m.), but data were compared using the non-parametric Mann–Whitney rank test. P < 0.05 was considered statistically significant.

Assay of NO synthase

Frozen tissue was extracted at 0–4°C by homogenisation (with an Ystral homomixer) in 2.5 volumes of a buffer containing 320 mM sucrose, 20 mM EDTA, 1 mM DL-dithiothreitol, 10 µg ml−1 leupeptin, 10 µg ml−1 soybean trypsin inhibitor and 1 µg ml−1 pepstatin brought to pH 7.2 at 20°C with hydrochloric acid. The homogenates were centrifuged at 10 000 g at 0–4°C for 30 min. Supernatants were passed through a 2 ml column of cation-exchange resin (AG 50WX8, Bio-Rad) to remove endogenous arginine and were stored on ice for up to 2 h before use. Nitric oxide synthase in these supernatants (cytosol plus microsomes) was measured by the conversion of L-[U-14C]Arginine to [U-14C]citrulline at 37°C for 10 min as previously described (Salter et al., 1991). The activity of the calcium-dependent enzyme was determined from the difference between the [U-14C]citrulline generated from control samples and samples containing 1 mM (ethylenebis(oxyethylenenitrilo)tetraacetic acid (EGTA); the activity of the calcium-independent enzyme was determined from the difference between samples containing 1 mM EGTA and samples containing both 1 mM EGTA and 1 mM L-NMMA.

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Total protein content of tissue supernatants was determined colorimetrically (Bio-Rad). The limit of detection in this assay was 0.7 pmol min⁻¹ mg⁻¹ protein.

Histology and immunohistochemistry

Tumours were typed and ductal tumours were graded according to Elston and Ellis (1991).

The immunohistochemical studies described below were carried out on all the samples of breast cancer and normal tissue included in this study. Sections (5 μm thick) were cut from frozen tissue, air dried for 30 min then fixed in acetone for 30 min. Two antibodies were used for NO synthase detection: (1) a monoclonal antibody to inducible NO synthase from murine macrophages (ANTI-macNOS; Transduction Laboratories, Lexington, KY, USA) which immunoreact with human inducible NO synthase (I Charles, unpublished observation) and (2) polyclonal antiserum to NO synthase raised in rabbits using purified rat brain NO synthase (Springall et al., 1992). At the dilution used this antiserum has proven reactivity with endothelial and neuronal constitutive NO synthase isoenzymes across several species including humans (Springall et al., 1992; Brave et al., 1993; Terenghi et al., 1993). At a lower dilution (1:100) in Western blots this antiserum also cross-reacts with inducible NO synthase (Thomsen et al., 1994). Sections were incubated in 4% normal swine serum for 30 min before incubation with the monoclonal antibody to inducible NO synthase (2.5 μg ml⁻¹) or the polyclonal antiserum to NO synthase (diluted 1:1000) for 16 h at 4°C. For macrophage detection, adjacent sections were incubated for 30 min with antibodies raised in mouse against the human monocyte/macrophage marker CD68 (3.4 μg ml⁻¹; EBM11; Dako (UK)). Antibody labelling was subsequently visualised using an avidin–biotin complex method.

Results

Nitric oxide biosynthesis by cultured tissue

Nitrite + nitrate accumulated in culture medium during culture of tumour tissue pieces for 24 h (Table I). Nitrite + nitrate generation was significantly higher for the 14 invasive tumours compared with normal controls (P<0.02). A case of benign cystic change yielded concentrations higher than normal controls but less than the mean value of invasive tumours. Interestingly, in a case of in situ ductal carcinoma, nitrite + nitrate was higher than that observed in the normal/benign cases, but again was lower than the mean for invasive tumours. Within the group of ductal carcinomas, nitrite + nitrate generation was greater in grade III than in grade II tumours (P<0.03). Nitrite + nitrate was not detectable in cultures of tissue from the grade I ductal carcinoma. An invasive lobular carcinoma and a phyllodes tumour also showed high concentrations of nitrite + nitrate in culture medium after 24 h culture. Generation of nitrite + nitrate was completely inhibited when tissue pieces were cultured in the presence of 2 mM L-NMMA.

Nitric oxide synthase activity

Nitric oxide synthase activity was detected only in invasive tumour tissue (Table II). For ductal carcinomas, activity was detected in all grade III and in one of two grade II tumours. Nitric oxide synthase was not detectable in tissue from the grade I ductal carcinoma or benign tumours, or in normal breast tissue. For all but two cases (one grade II and one grade III ductal carcinoma) a proportion of the NO synthase activity was found to be calcium independent.

Immunohistochemistry of tissue sections

We used a monoclonal antibody to inducible NO synthase to identify and localise the inducible enzyme and a polyclonal antiserum raised against purified rat brain NO synthase to identify other NO synthase isoenzymes.

The monoclonal antibody to inducible NO synthase labelled peritumoral spindle cells in the tumour stroma with the morphology of macrophages (Figure 1a). These cells were seen in stroma adjacent to invasive islands of tumour cells and also in the stroma intimately associated with ducts involved by in situ carcinoma. Occasional labelled cells were also seen within solid tumour islands of grade III invasive ductal carcinomas and within ducts involved by in situ carcinoma. It was usually possible to recognise that these cells had the morphology of macrophages, and in all cases serial sections stained with the CD68 macrophage marker (EBM/11).

Table I Nitrite + nitrate biosynthesis by breast tissue cultured for 24 h

| Tissue type               | No. of cases | Nitrite + nitrate (nmol mg⁻¹ protein; mean ± s.e.m.) |
|---------------------------|--------------|---------------------------------------------------|
| Normal                    | 4            | 0.0 ± 0.05                                        |
| Benign cystic change      | 1            | 0.3                                               |
| Ductal carcinoma in situ  | 1            | 1.5                                               |
| Invasive tumours          | 14           | 1.9 ± 0.45*                                      |
| Ductal grade I            | 1            | 0.0                                               |
| Ductal grade II           | 7            | 1.0 ± 0.27*                                      |
| Ductal grade III          | 3            | 3.8 ± 0.85*                                      |
| Lobular carcinoma         | 1            | 3.3                                               |
| Phyllodes tumour          | 1            | 1.8                                               |

*P<0.02 compared with normal tissue. *P<0.04 compared with normal tissue. *P<0.03 compared with grade II ductal carcinomas.

Table II NO synthase activity in breast tissue

| Tissue type               | No. of cases | NO synthase (pmol min⁻¹ mg⁻¹ protein; mean ± s.e.m.) | Total | Ca²⁺ independent |
|---------------------------|--------------|------------------------------------------------------|-------|------------------|
| Normal                    | 3            | <0.7                                                | <0.7  |
| Benign lesions*           | 5            | <0.7                                                | <0.7  |
| Invasive tumours          | 15           | 5.1 ± 1.4                                           | 2.3 ± 0.7 |<0.7 |
| Ductal grade I            | 1            | <0.7                                                | <0.7  |
| Ductal grade II           | 2            | 3.9 ± 3.9                                           | <0.7  |
| Ductal grade III          | 7            | 8.5 ± 2.2                                           | 3.7 ± 1.1 |<0.7 |
| Lobular carcinoma         | 4            | 3.3 ± 1.8                                           | 2.0 ± 1.2 |
| Poorly differentiated     | 1            | 4.9                                                 | 0.9    |

*One benign cystic change, one stromal fibrosis and three fibroadenomas.
11) confirmed their macrophage origin (Figure 1b). In sections from normal and benign breast tissue very occasional immunolabelling of stromal macrophages was seen. Few endothelial or myoepithelial cells were also labelled with the monoclonal antibody to inducible NO synthase.

The polyclonal antiserum to NO synthase also labelled peritumoral spindle cells with the morphology of macrophages. In addition, in both benign and malignant breast tissue, immunolabelling of other cells including vascular endothelial (Figure 1c) and some myoepithelial cells (Figure 1d) was also observed. The labelling of these cells was much more evident than that observed with the monoclonal antibody to the inducible enzyme. Some immunopositive cells remain unidentified at present.

Discussion

The present study shows that NO biosynthesis and NO synthase activity are high for tumour tissue obtained from primary human breast cancers compared with benign lesions and normal breast tissue in which activity was low or not detectable. Furthermore, NO biosynthesis was significantly greater for tissue explants from grade III compared with grade II invasive ductal carcinomas. We have previously reported that NO synthase is expressed in human gynaecological tumours and, as with the present study, that there is an association between NO synthase activity and tumour grade (Thomsen et al., 1994).

The biochemical observations of calcium-independent
enzyme activity, and the generation of nitrite and nitrate during culture of viable tumour tissue explants, suggest the presence of inducible NO synthase (Knowles and Moncada, 1994). We have used a monoclonal antibody to inducible NO synthase to determine the presence and localisation of this isoenzyme. Consistent with the biochemical observations, immunohistochemical studies revealed proteins immunoreactive with this monoclonal antibody. This immunoreactivity was localised predominantly within macrophages.

Calcium-dependent activity was also measurable. While calcium-dependent NO synthase activity suggests the threonine to serine irreversible NO synthase isoforms, there are examples in the literature for calcium-dependent inducible NO synthase, and for induction of the neuronal and endothelial NOS isoforms which were originally described as constitutively expressed (reviewed in Knowles and Moncada, 1994). Further immunohistochemical studies using well-characterised antibodies, as well as investigations using molecular biology techniques, are required to determine the specific NO synthase isoform(s) expressed in human breast cancer tissue. However, consistent with the presence of constitutive NO synthase was immunoreactive with the polyclonal antisera to NO synthase in these tumours. This antisera, which immunoreacts with endothelial and neuronal constitutive NO synthases (Springall et al., 1992; Brave et al., 1993; Terenghi et al., 1993), labelled vascular endothelial cells and myoepithelial cells in addition to macrophages. In contrast to gynaecological tumours in which immunolabelling with the antisera to NO synthase was observed in tumour cells (Thomsen et al., 1994), breast tumour cells were not immunolabelled with this antisera or with the monoclonal antibody to inducible NO synthase.

The generation of NO by immune cells is an important aspect of non-specific immunity in animals (Hibbs et al., 1990). Localisation of NO synthase within intratumoral macrophages indicates that NO may also be involved in the immune response in man. Previous investigations have shown that macrophage infiltration is high in invasive ductal carcinomas of the breast regardless of tumour grade (Miles et al., 1994). Interestingly, however, tumour necrosis factor, a cytokine associated with cytotoxic macrophage functions (Sugarmen et al., 1985), is expressed predominantly within tumour-associated macrophages of high-grade breast tumours and expression is grade related (Miles et al., 1994). This cytokine, which as a potent inducer of NO synthase (Hibbs et al., 1990), may regulate the expression of NO synthase in the tumour-infiltrating macrophage population.

While the generation of NO in breast tumours may cause tumour cell cytostasis cytotoxicity, it may also increase tumour blood flow (Andrade et al., 1992; Wood et al., 1993) and promote angiogenesis (Weidner et al., 1992; Jenkins et al., 1993). A balance in favour of the vascular effects may explain the positive correlation between NO biosynthesis and grade of malignancy.

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