Induction of haem oxygenase-1 by nitric oxide and ischaemia in experimental solid tumours and implications for tumour growth

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Summary Induction of haem oxygenase-1 (HO-1) as well as nitric oxide (NO) biosynthesis during tumour growth was investigated in an experimental solid tumour model (AH136B hepatoma) in rats. An immunohistochemical study showed that the inducible isoform of NO synthase (iNOS) was localized in monocyte-derived macrophages, which infiltrated interstitial spaces of solid tumour, but not in the tumour cells. Excessive production of NO in the tumour tissue was unequivocally verified by electron spin resonance spectroscopy. Tumour growth was moderately suppressed by treatment with either N-nitro-l-arginine methyl ester (L-NAME) or S-methylisothiourea sulphate (SMT). In contrast, HO-1 was found only in tumour cells, not in macrophages, by in situ hybridization for HO-1 mRNA. HO-1 expression in AH136B cells in culture was strongly enhanced by an NO (NO+) donor S-nitroso-N-acetyl penicillamine. HO-1 mRNA expression in the solid tumour in vivo decreased significantly after treatment with low doses of NOS inhibitors such as L-NAME and SMT (6–20 mg kg−1). However, the level of HO-1 mRNA in the solid tumour treated with higher doses of NOS inhibitor was similar to that of the solid tumour without NOS inhibitor treatment. Strong induction of HO-1 was also observed in solid tumours after occlusion or embolization of the tumour-feeding artery, indicating that ischaemic stress which may involve oxidative stress triggers HO-1 induction in the solid tumour. Lastly, it is of great importance that an HO inhibitor, zinc protoporphyrin IX injected intra-arterially to the solid tumour suppressed the tumour growth to a great extent. In conclusion, HO-1 expression in the solid tumour may confer resistance of tumour cells to hypoxic stress as well as to NO-mediated cytotoxicity.

Keywords: nitric oxide; NO; NO synthase; haem oxygenase-1; tumour growth; vascular permeability; tumour blood flow

Expression of different isoforms of nitric oxide (NO)3 synthase (NOS) has been documented in various tumour cell lines and solid tumours (Buttery et al, 1993; Bastian et al, 1994; Thomsen et al, 1994; Cobbs et al, 1995). Our previous experiments indicate that enhanced vascular permeability in solid tumour is mediated by at least three factors: NO, bradykinin and prostaglandins (Matsumura et al, 1988; Wu et al, 1998; Maeda et al, 1994). NO production by experimental tumours in mice has been implicated in maintenance of tumour blood flow in the neovasculature, and blockade of NO with Nω-nitro-l-arginine has been shown to reduce blood flow in the tumour-associated neovasculature (Andrade et al, 1992; Tozer et al, 1995, 1997).

Among three types of NOS, the inducible isoenzyme (iNOS) produces a much larger amount of NO for a longer time than do the other two constitutive enzymes, i.e. neuronal NOS and endothelial NOS (Stuehr and Griffith, 1992; Moncada and Higgs, 1993). We recently suggested that NO produced in excess in the local area of the solid tumour seems to sustain rapid tumour growth in an experimental solid tumour (Doi et al, 1996). It has been reported, however, that a particularly high output of NO from iNOS is potentially cytotoxic for various tumour cells (Hibbs et al, 1988; Bastian et al, 1994; Lepoivre et al, 1994). The biological effect of NO in tumour biology is thus still unsettled because of previous inconsistent results, i.e. suppressing effects and promoting effects on tumour growth. These contradictory effects of NO may be explained by a putative protective mechanism in the tumour cells against cytotoxicity of excess NO.

In this respect, of considerable interest is a unique biological function of haem oxygenase (HO), originally involved in disintegration of haem compounds in iron metabolism (Tenhunen et al, 1968). HO degrades haem to release carbon monoxide (CO), iron and biliverdin (Maines, 1997). Two isoforms of HO, i.e. HO-1 and HO-2, are known to exist in eukaryotic systems (Maines and Kappas, 1974). HO-1 is induced by various stimuli, such as pro-inflammatory cytokines and heavy metals (Shibahara et al, 1985; Maines, 1997; Yet et al, 1997). Induction of HO-1 has been suggested to provide an important protective response of cells against oxidative damage, because HO-1 induction may decrease the cellular haem level (pro-oxidant) and elevate the level of bilirubin (antioxidant), which is derived from biliverdin and shows a potent scavenging action against reactive oxygen species (Maines, 1997; Kim et al, 1995a, 1995b). In addition, it is proposed that CO, a product of HO, functions as a gaseous signal transduction molecule in a similar manner to that of NO (Suematsu et al, 1995; Maines, 1997; Prabhakar et al, 1997). It is of interest that HO has been implicated in tumour growth (Lee and Ho, 1994; Hara et al, 1996; Goldman et al, 1996; Takahashi et al, 1996), although details of this mechanism remain unclear.
In the present study, to obtain a better understanding of the role of HO and NO biosynthesis in tumour biology and cancer treatment, we examined the mechanism of induction of HO-1 and iNOS in rat AH136B solid tumours.

MATERIALS AND METHODS

Animals and implantation of AH136B tumour

Donryu rats, weighing from 160 to 180 g, were obtained from a commercial supplier (SLC, Inc., Shizuoka, Japan). They were handled according to the guidelines of the Experimental Animal Center of Kumamoto University.

AH136B tumour (hepatoma) cells were cultured serially in ascitic fluid in rats as described previously (Doi et al, 1996). The cells were implanted subcutaneously (s.c.) in a dorsal site of the foot of Donryu rats with an inoculum size of $1 \times 10^5$ cells per injection site. Tumours were allowed to grow and usually reached 15–20 mm in diameter for 16 days.

Electron spin resonance spectroscopy for analysis of NO production in AH136B solid tumours

Overproduction of NO was examined with the solid tumour obtained 16 days after the tumour cell inoculation by using electron spin resonance (ESR) spectroscopy according to a previously described method (Doi et al, 1996; Setoguchi et al, 1996; Yoshimura et al, 1996). Briefly, a complex of N-(dithiocarboxy)-sarcosine (DTCS, 180 mg kg$^{-1}$; Dojindo Laboratories, Kumamoto, Japan) with FeSO$_4 \cdot 7$H$_2$O (40 mg kg$^{-1}$), which is soluble in water, was injected s.c. ESR was performed 30 min after the administration of the iron complex, with the tumour obtained as described below.

The solid tumours obtained with the rats under sodium pentobarbital anaesthesia were transferred to quartz sample tubes (5 mm in diameter). The samples were immediately frozen in liquid nitrogen and then subjected to ESR by using an X-band ESR spectrometer (Bruker 380E) at 110K. Unless otherwise specified, the conditions for ESR measurement were microwave frequency, 9.39 GHz; microwave power, 4 mW; modulation frequency, 100 kHz; and modulation amplitude, 0.5 mT. The magnetic field was calibrated by using TCNQ-Li salt ($g = 2.00252$).

Simultaneously, to test the effect of NOS inhibitors on the formation of NO in the solid tumour, ESR was performed by using a DTCS-Fe$^{2+}$ complex 1 h after intraperitoneal (i.p.) injection of N$^2$-nitro-1-arginine methyl ester (l-NNAME; Sigma Chemical, St Louis, MO, USA) at 50 mg kg$^{-1}$ or S-methylisothiourea sulphate (SMT; Wako Pure Chemical Co., Ltd, Osaka, Japan) at 50 mg kg$^{-1}$.

Expression and localization of iNOS in AH136B solid tumours

We previously showed that iNOS mRNA expression is highly elevated in AH136B solid tumours implanted and grown in rats (Doi et al, 1996). The localization of iNOS expression was further examined by an immunohistochemical study using a specific antibody for rat iNOS according to our previous method (Setoguchi et al, 1996). Briefly, solid tumour tissue was excised on day 16 after tumour implantation and fixed with 2% periodate–lysine–paraformaldehyde for 6 h. The tissue was then embedded in tissue-embedding medium and frozen in liquid nitrogen, followed by preparation of 6-μm-thick sections by use of a cryostat. After inhibition of endogenous peroxidase activity, the specimens were incubated with an anti-rat iNOS polyclonal antibody (Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) for 1 h at room temperature. After the samples were washed with 0.01 m phosphate-buffered 0.15 m saline (PBS, pH 7.4), they were incubated for 1 h with a sheep antimouse immunoglobulin [F(ab$^\prime$)$_2$] conjugated with peroxidase diluted 1:100. Tissue-bound peroxidase activity was visualized using 3,3$'$-diaminobenzidine as a substrate; haematoxylin was used for nuclear staining.

In addition, tissue sections were stained immunohistochemically with a specific antibody against exudate monocyte-derived macrophages (TRPM-3) (Takeya et al, 1987). As the control, sections were incubated with non-immunized mouse serum or PBS instead of primary antibody, after which they were processed in the manner just described.

Expression and localization of HO-1 in AH136B solid tumours

First, expression of HO-1 mRNA was analysed by Northern blotting. Total RNA was extracted from the tumour tissue obtained at 16 days after tumour inoculation by using the guanidine thiocyionate lysis method with Trizol$^\text{TM}$ reagent (GibcoBRL, Gaithersburg, MD, USA). Each RNA sample (20 μg) underwent electrophoresis on agarose gel and was transferred to the Hybond$^\text{-N+}$ nylon membrane, followed by hybridization of a DNA probe for rat HO-1 mRNA as described previously (Setoguchi et al, 1996; Akaike et al, 1997). The DNA probe was radiolabelled by the random primer technique using [γ$^32$P]-dCTP and the Megaprime labelling system$^\text{TM}$ (Amersham International plc, Buck, UK). An HO-1 cDNA fragment of 882 bp was used for hybridization as reported previously (Takeda et al, 1994), and a cDNA fragment for glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as the control-gene expression (Akaike et al, 1997).

Western blot analysis for HO-1 protein was also performed by using a specific polyclonal anti-rat HO-1 antibody (Stress-Gen Biotechnologies, Victoria, Canada). Tumour tissue obtained 16 days after tumour implantation was homogenized, by use of a Polytron homogenizer (Kinematica GmbH, Luzerne, Switzerland), in PBS (pH 7.4) containing 1 mM ethylenediamine tetraacetic acid (EDTA) and 1 mM phenylmethylsulphonyl fluoride (PMSF).

The microsome fraction was obtained by ultracentrifugation at 105 000 g for 1 h at 4°C. The microsome preparation was then solubilized by adding sodium dodecyl sulphate (SDS), and each aliquot (10 μl; 30 μg of protein content) of the preparation was treated with the buffer for SDS-polyacrylamide gel electrophoresis (PAGE; 10%) under non-reducing conditions as reported earlier (Okamoto et al, 1997). After electrophoresis, protein was transferred to Immobilon$^\text{TM}$ polyvinylidene difluoride membranes (Millipore Co., Ltd., Bedford, MA, USA) followed by incubation with an anti-HO-1 antibody. The protein band that reacted immunologically with the antibody was visualized by using the ECL$^\text{TM}$ system (Amersham), combined with chemiluminescence detection with Kodak XAR-5 film.

To localize HO-1 in solid tumour tissue (on day 16 after tumour inoculation), in situ hybridization was performed with use of a digoxigenin RNA labelling kit (Boehringer Mannheim GmbH, Mannheim, Germany) according to the method described

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Denhardt’s solution, and yeast tRNA (200 ng) in 0.2 m hydrochloric acid (HCl) and were acetylated with 0.25% acetic anhydride in 0.1 m triethanolamine, followed by dehydration through a graded series of ethanol solutions. The sections were then hybridized with either antisense or sense probes for HO-1 mRNA (10 µg ml⁻¹ for each) in 100 µl of hybridization solution containing 50% formamide, 0.6 m sodium chloride, 1 mM EDTA, 10% dextran sulphate, 0.25% SDS, 1 x Denhardt’s solution, and yeast tRNA (200 µg ml⁻¹) in 20 mM Tris–HCl, pH 8.0 at 45°C for 14 h. After hybridization, the sections were washed with 2 x standard saline citrate (SSC) containing 50% formamide at 45°C for 20 min, digested with RNase (5 µg ml⁻¹) at 37°C for 30 min, and washed again with 2 x SSC and then 0.5 x SSC at 45°C. The hybridized digoxigenin-labelled RNA was detected colourimetrically by using a nucleic acid detection kit (Boehringer Mannheim) with a sheep anti-digoxigenin immunoglobulin (Fab’), conjugated with alkaline phosphatase, and nitroblue tetrazolium salt and 5-bromo-4-chloro-3-indolylphosphate for the colour reaction. The phosphatase reaction was carried out in a humidified chamber at 37°C for 1 h.

Measurement of HO activity in tumour tissues

The HO activity in the AH136B solid tumour or various organs such as livers and spleens was assayed as described previously (Maines and Kappas, 1978). Specifically, the tumour tissues, which was obtained 16 days after tumour implantation, or the liver and spleen were homogenized by the Polytron homogenizer in ice-cold 50 mM potassium phosphate buffer (pH 7.4) containing 2 mM EDTA, 2 mM PMSF and 10 µg ml⁻¹ of leupeptin. After the homogenate was centrifuged at 10 000 g for 30 min at 4°C, the resultant supernatant was used to prepare the microsome fraction as described above (105 000 g at 4°C). The microsome fraction was suspended in 0.1 M potassium phosphate buffer (pH 7.4), followed by sonication for 2 s at 4°C and stored at -70°C until use. The reaction mixture for the determination of the HO activity was composed of 2 mM of microsome protein, cytosol fraction of the rat liver (3 mg protein) as a source of biliverdin reductase, 33 µM haemin and 333 µM NADPH in 3 ml of 50 mM potassium phosphate buffer (pH 7.4). The reaction was terminated by addition of 0.1 ml of 0.01 M HCl 15 min after incubation of the reaction mixture. The bilirubin formed in the reaction was extracted with 1 ml of chloroform, and the bilirubin concentration was determined spectroscopically by measuring the difference in absorbance between 464 nm and 530 nm by using a molar extinction coefficient of 40 mm⁻¹ cm⁻¹ (Maines and Kappas, 1978).

Culture of AH136B tumour cells and iNOS and HO-1 induction

AH136B cells obtained from the ascitic form of the tumour were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Gibco, Grand Island, NY, USA) supplemented with non-essential amino acids (Gibco) and 10% fetal bovine serum in 5% carbon dioxide 95% air at 37°C. Cellular iNOS expression was examined by reverse transcriptase-polymerase chain reaction (RT-PCR) and Southern blot analyses. Briefly, total RNA was extracted from a confluent monolayer of the cells in culture on a 6-well polystyrene plate (Falcon, Becton Dickinson Labware, Lincoln Park, NJ, USA; diameter 3.4 cm), and then 3 µg of the RNA were subjected to RT-PCR and Southern blotting for iNOS mRNA. The reaction conditions and oligonucleotide primer used for RT-PCR and a cDNA probe for iNOS are described elsewhere (Doi et al, 1996).

To see whether NO or a related redox form such as nitroso group (NO⁺) induces HO-1 mRNA expression in the tumour cells in vitro, S-nitroso-N-acetyl penicillamine (SNAP; Dojindo Laboratories) was added at various concentrations to the cells in culture (6-well plate) as just described. After a 5 h incubation period at 37°C in the carbon dioxide incubator, total RNA was extracted and 10 µg RNA was used for Northern blotting for HO-1 mRNA in the same manner as for the solid tumour tissues.

Effect of NOS inhibition on HO-1 induction and solid tumour growth in vivo

To explore the mechanism of HO-1 induction in AH136B solid tumour, HO-1 mRNA expression with or without NOS inhibitors was examined by Northern and Western blot analyses. Specifically, tumour-bearing rats received either L-NAME or SMT in 0.2 ml saline i.p. at a dose of 3, 10, 30, or 45 mg kg⁻¹ body weight two times a day for 5 days, beginning 11 days after tumour implantation. The group given only saline served as a control. Solid tumour tissues were obtained 1 day after the last injection of NOS inhibitors, and total RNA was extracted followed by Northern blotting as described above. The microsome fraction of the tissue was isolated by ultracentrifugation and was subjected to Western blotting after SDS-PAGE as noted earlier. The effect of NOS inhibitors on tumour growth was tested by treatment of tumour-bearing rats with L-NAME or SMT with the same protocol as that used for examination of HO-1 expression. Briefly, after administration of NOS inhibitors, tumour volume was assessed by measuring both major (a) and minor (b) axes of the solid tumours with use of a vernier micrometer. Tumour volume was estimated by use of the formula (π × a × b²)/6.

Effect of ischaemic stress on HO-1 expression in AH136B solid tumours in vivo

HO-1 induction after ischaemic treatment was examined by Northern blotting and by measuring its enzyme activity. On day 16 after tumour implantation, the solid tumours were resected at various time points after initiation of surgical occlusion of the iliac artery, which provides the tumour-feeding artery of the tumour-implanted side. Ischaemic stress was also caused in solid tumours by infusion of Lipiodol™ containing styrene-co-maleic acid polymer-conjugated neocarzinostatin (Smancs; 0.1 mg of Smancs in 1.0 ml of Lipiodol per 5 min) via cannula (PE10; Becton Dickinson, Sparks, MD, USA) inserted into the feeding artery (iliac artery) (Yamasaki et al, 1987), and the level of HO-1 mRNA expression was monitored as in the occlusion model.

Treatment of AH136B solid tumours with an HO inhibitor in vivo

To examine the role of HO-1 in the solid tumour growth, zinc protoporphyrin IX (ZnP); (Wako Pure Chemical, Osaka, Japan)
an inhibitor of HO activity (Maines, 1981) was administered to the tumour-bearing animals. Specifically, 0.1 mg of ZnPP in 0.2 ml of 1.0% DMSO in saline was injected intra-arterial to the solid tumour via the feeding artery on day 7 after tumour implantation as just described, and the tumour growth was monitored by measuring the tumour volume in the same manner as the NOS inhibition experiment mentioned above. The group given only 200 μl of 1.0% DMSO in saline served as a control without HO inhibitor treatment.

Statistical analysis

Data are shown as means ± s.e.m. Statistical difference was analysed by the two-tailed unpaired t-test. A P-value of < 0.05 was considered statistically significant.

RESULTS

Excessive production of NO was demonstrated in AH136B solid tumours in rats by ESR spectroscopy: a significant ESR signal of the NO-(DTCS)2-Fe2+ adduct was identified (Figure 1). In contrast, only a negligible ESR signal of the NO adduct was observed when animals were given L-NAME or SMT (Figure 1). Because a strong, steady level of induction of iNOS mRNA was observed, we then examined the localization of iNOS in the tumour tissue.

As demonstrated in Figure 2A, an immunohistochemical study revealed that iNOS is expressed only by cells infiltrating the interstitial space, but not by tumour cells. TRPM-3-positive monocyte-derived macrophages were also found to infiltrate the interstitial space of the solid tumour (Figure 2B). The location of iNOS-positive cells appears to be almost the same as that of TRPM-3-positive cells in the tumour tissue. Therefore, iNOS-positive cells were considered to be interstitial infiltrating macrophages rather than AH136B tumour cells. No appreciable immunostaining was observed by the control staining with non-immunized serum or PBS instead of the primary antibodies (anti-iNOS antibody or TRPM-3) (not shown). This result also suggests that monocyte-derived activated macrophages infiltrating in tumour seem to be primarily responsible for overproduction of NO. The lack of iNOS expression in AH136B tumour cells was further substantiated because no appreciable iNOS mRNA expression was observed with AH136B cells in culture by using RT-PCR (Southern blotting) and Northern blotting (data not shown).

Strong HO-1 expression was observed in the solid tumour as assessed by both Northern and Western blot analyses, and by HO activity measurement as well (Figure 3 A–C). HO-1 mRNA expression in the solid tumour in vivo was higher than that in the tumour cells in vitro (Figure 3A). The level of HO-1 induction in the solid tumours was comparable to that in the spleen, which is known to express HO-1 constitutively under physiological conditions (Figure 3 B, C). In contrast to iNOS, HO-1 was observed only in tumour cells of the solid tumour tissue as identified by in situ hybridization with an HO-1 RNA probe (Figure 4).

Figure 1  ESR spectra obtained with AH136B solid tumour tissue. Thirty minutes before the tissue was obtained, a DTCS-Fe2+ complex (180–40 mg kg−1) was given s.c. to the rats treated (B, C) or untreated (A) with NOS inhibitors (50 mg kg−1, i.p.). All ESR measurements were performed at 110 K. See text for details.
To determine the effect of NO in induction of HO-1, AH136B cells in culture were analysed for expression of HO-1 mRNA after treatment with SNAP in vitro. As shown in Figure 5, induction of HO-1 mRNA was clearly increased in a dose-dependent manner when SNAP was added to the cell culture system. This result indicates that NO or its related redox isoform, NO+, has a potent HO-1-inducing potential in AH136B tumour cells.

The level of HO-1 mRNA in solid tumours in vivo decreased significantly after treatment with low doses (6 and 20 mg kg\(^{-1}\)) of NOS inhibitors such as L-NAME and SMT. However, as the NOS inhibitor dose increased up to 90 mg kg\(^{-1}\) day\(^{-1}\), HO-1 mRNA expression gradually recovered to a level similar to that of the solid tumour without treatment (Figure 6). Similar results for the effect of NOS inhibition on HO-1 protein expression were obtained by Western blotting for rat HO-1 (data not shown). The efficacy of NOS inhibition by L-NAME and SMT was tested by an ESR study, and treatment with either L-NAME or SMT abrogated NO production completely in solid tumour tissues as just mentioned (Figure 1 B, C).

To explain the inconsistency in the effect of NO on HO-1 induction in vitro and in vivo, we examined the effect of ischaemic stress on the solid tumours in rats. Ischaemia caused by occlusion of the tumour-feeding artery and by infusion of Smancs/Lipiodol resulted in strong up-regulation of HO-1 mRNA expression (Figure 7A and B). Although there was about a 24-h gap between HO-1 mRNA up-regulation and HO activity induction, the HO activity in the solid tumour was apparently enhanced by the ischaemic stress (Figure 7C).

Moreover, when tumour growth in vivo was investigated with the use of NOS inhibitors, moderate retardation of tumour growth was observed with both L-NAME and SMT (Figure 8), but this effect was not as remarkable as their strong inhibition of NO biosynthesis, as evidenced by ESR spectroscopy (Figure 1 B, C). In contrast, the treatment of HO inhibitor (ZnPP), which significantly inhibited HO activity in the solid tumour, showed a great suppression of the tumour growth in vivo (\(P < 0.01\)) (Figure 9). In a separate experiment, NO production in the solid tumour with or without ZnPP treatment was investigated by using ESR spectroscopy as just described. ZnPP administered intraarterially, however, did not affect NO biosynthesis by iNOS expressed in the tumour tissue (data not shown).

**DISCUSSION**

In our earlier study, excessive production of NO as assessed by ESR spectroscopy correlated well with the rate of growth of AH136B solid tumours in rats (Doi et al, 1996). The present study was performed to further explore this observation and to clarify the role of NO in solid tumour biology, by focusing on the induction and function of HO-1 in AH136B solid tumours.

Rapid tumour growth is sustained by a number of factors derived from tumour cells or a host’s tissue in solid tumours (Matsumura et al, 1988; Vaupel et al, 1989; Maeda et al, 1994; Nakano et al, 1996; Suzuki et al, 1996; Wu et al, 1998). Maintenance of regional blood flow in solid tumour tissues appears to be most important for tumour cell growth, for a sufficient supply of molecular oxygen and various nutrients. It is now well known that NO released from vascular endothelial cells plays an important physiological role in regulation of blood flow in systemic circulation via its potent vasodilating action (Moncada...
and Higgs, 1993). Growth of solid tumour, therefore, may be sustained by increased blood flow mediated by prolonged and excessive production of NO.

Our previous experiment showed that enhanced vascular permeability to tumour tissues was mediated by NO (Maeda et al., 1994; Wu et al., 1998). Leibovich et al (1994) reported that an L-arginine-dependent NO pathway mediates angiogenic activity. Interestingly, Ziche et al (1997) recently reported that vascular endothelial growth factor, VEGF (also called vascular permeability factor, VPF), induces angiogenesis via formation of NO.

Figure 3  Northern blot (A) and Western blot (B) analyses for HO-1 expression and HO activity (C) in AH136B tumours. Northern hybridization was performed with AH136B tumours obtained in vitro and in vivo with use of a cDNA probe for rat liver HO-1 (A). The expression of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA in each tissue was determined as for HO-1, as shown in the lower panel. Western blotting was done by using a specific antibody for rat HO-1 with AH136B solid tumour tissue, liver, and spleen (B). The HO activity was assayed spectroscopically based on the bilirubin formation from haem in the reaction mixture of the microsome fraction of various tissues in the presence of the rat liver cytosol protein as a source of biliverdin reductase (C). Data are shown as means ± s.e.m. (n = 4; *P < 0.05). Note that the HO activity in each tissue determined was strongly inhibited by addition of 30 μM ZnPP mg⁻¹ of microsome protein to the assay mixture. The microsome fraction of the tissue was isolated by ultracentrifugation and was subjected to Western blot analysis after SDS-PAGE and to the measurement for HO activity. See text for details.
Angiogenesis leading to hypervascularization is often observed during rapid tumour growth. Consequently, it is plausible that the angiogenic potential as well as the vascular permeability enhancing effect of NO may facilitate rapid growth of solid tumours, which have great demands for various nutrient factors. A similar beneficial effect of NO on tumour growth was clearly demonstrated recently in tumour-bearing mice, in which human adenocarcinoma cells, which overexpressed iNOS, were implanted (Jenkins et al, 1995). A promoting effect of NO on tumour progression and metastasis is also suggested by Lala and Orucevic (1998). These results are supported by the finding that inhibition of NO synthesis by an NOS inhibitor given systemically reduced tumour growth in mice (Andrade et al, 1992; Thomsen et al, 1997; our present results).

In contrast, it has been reported that NO is an effector molecule in macrophage-mediated cytotoxicity (Hibbs et al, 1988; Bastian et al, 1994; Lepoivre et al, 1994). We showed in the present experiment that the tumour-infiltrating activated macrophage seems to be a major contributor to overproduction of NO in the solid tumour. This observation appears to conflict with the above-mentioned result. To solve these contradictory issues, we probed the mechanism of efficient tumour growth in the presence of an excess amount of NO in the solid tumours.

In this context, Lancaster’s group reported that pretreatment of rat hepatocytes with a low dose of NO donor confers resistance to oxidative damage of the cells, through induction of HO-1 (Kim et al, 1995a). In their report, it is also demonstrated that HO-1 induction protected the cells from NO-mediated cytotoxicity. HO-1, which catalyses the conversion of haem to biliverdin and CO, has been shown to be constitutively expressed in the liver and spleen and in some tumour cells (Tenhunen et al, 1968; Shibahara et al,

Figure 4 In situ hybridization for HO-1 expression in AH136B solid tumour. Sections of solid tumour were hybridized with antisense (A) or sense (B) RNA probes for HO-1 labelled with digoxigenin. AH136B cells are positively stained with the antisense probe for HO-1. Magnification × 130. See text for details.

Figure 5 Northern blot analysis for HO-1 mRNA expression in AH136B tumour cells in culture treated with SNAP. After treatment of cells with various concentrations of SNAP for 5 h, Northern blotting for HO-1 mRNA expression was performed in the same manner as described in Figure 3. In the lower panel, the HO-1 mRNA signals were quantified by densitometric analysis, after normalization with GAPDH mRNA signals as a standard mRNA expression, using a Macintosh computer with an Image Scanner (GT6500, Epson Co., Ltd, Tokyo, Japan) and the public domain NIH Image Program. See text for details.
Figure 6  Northern blot analysis for HO-1 mRNA expression in AH136B solid tumours treated with NOS inhibitors. Tumour-bearing rats received either L-NAME or SMT i.p. at a dose of 3, 10, 30, or 45 mg kg\(^{-1}\) two times a day for 5 days, starting 11 days after tumour implantation. Northern blotting for HO-1 mRNA expression was performed in the same manner as described in Figure 3. In the lower panels, quantitative image analysis for HO-1 mRNA was performed in a same manner as in Figure 5. See text for details.

Figure 7  HO-1 mRNA expression in AH136B solid tumours under ischaemic conditions caused by occlusion of the feeding artery (A) or embolization with a Smancs-Lipiodol emulsion (0.1 mg of Smancs in 0.1 ml of Lipiodol) (B). Lower panels in (A) and (B) show the relative signal intensities of HO-1 mRNA quantified similar to Figure 5. (C) Induction of HO activity in the solid tumour was also examined after occlusion of the feeding artery in the same manner as in (A). Data are expressed means ± s.e.m. (n = 4; *P < 0.05). Northern blotting was carried out with the solid tumour obtained at various time points after occlusion of the feeding artery or treatment with Smancs-Lipiodol. In a control study (sham operation), HO-1 mRNA expression was examined after laparotomy without occlusion of the feeding artery. See text for details.
45 mg kg⁻¹ two times a day for 5 days, starting 11 days after tumour control. Tumour volumes was estimated by use of the formula (πd₁²d₂)/6. Data are shown as means ± s.e.m. (n = 8); asterisks show significant differences versus the control group by the t-test for unpaired data (*P < 0.05; **P < 0.01).

![Figure 8](image) Effect of HO inhibitor treatment on tumour growth. Tumour-bearing rats received either l-NAME or SMT i.p. at a dose of 3, 10, 30, or 45 mg kg⁻¹ two times a day for 5 days, starting 11 days after tumour implantation. The rate of gain in tumour volume was assessed by comparing sizes of both major (a) and minor (b) axes of the solid tumours at 11 and 16 days after tumour cell implantation. The group given only saline served as control. Tumour volumes was estimated by use of the formula (πd₁²d₂)/6. Data are shown as means ± s.e.m. (n = 8); asterisks show significant differences versus the control group by the t-test for unpaired data (*P < 0.05; **P < 0.01).

![Figure 9](image) Effect of HO inhibitor treatment on tumour growth. ZnPP (0.1 mg) in 0.2 ml saline (1.0% DMSO) was administered to the solid tumour via the feeding artery (iliac artery) involving laparotomy at 7 days after tumour implantation. The tumour growth was assessed in the same manner as described in Figure 8. The control group received only 0.2 ml of 1.0% DMSO in saline. Data are shown as means ± s.e.m. (n = 4); asterisks show significant differences versus the control group by the t-test for unpaired data (*P < 0.05; **P < 0.01). The inset shows the HO activity in the solid tumour, which was assessed before (7 days after tumour inoculation) and after ZnPP treatment (n = 4 except that n = 1 for day 7 after ZnPP treatment).

In conclusion, our current study indicates that HO-1 expression in solid tumour appears to be regulated by NO and ischaemic stress, which should often occur in tumour tissue during rapid tumour growth in vivo. The up-regulation of HO-1 may have protective and beneficial effects for tumour cells against anti-tumour actions of the host as well as many anticancer agents.

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