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**Design and evaluation of S-nitrosylated human serum albumin (SNO-HSA) as a novel anticancer drug**

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**Running Title Page:**

NO-HSA induces tumor cell apoptosis

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The number of text pages: 33

The number of tables: 1

The number of figures: 6

The number of references: 40

The number of words in the Abstract: 242

The number of words in the Introduction: 571

The number of words in the Discussion: 1461

ABBREVIATIONS: NO, nitric oxide; NO-HSA, nitrosated human serum albumin; rHSA, recombinant human serum albumin; ROS, reactive oxygen species; NO-NSAID, nitric oxide-donating non-steroidal anti-inflammatory drug; NO-ASA, nitric oxide-donating aspirin; DTPA, diethylenetriaminepentaacetic acid; HBSS, Hanks' balanced salt solution; KPB, potassium phosphate buffer; C26, murine colon 26 carcinoma; FCS, fetal calf serum; CM-H<sub>2</sub>DCFDA, 5-(and-6)-chloromethyl-2',7'-dichlorodihydrofluorescein diacetate, acetyl ester; FITC, fluorescein isothiocyanate; HRP, Horseradish Peroxidase; TdT, terminal

deoxynucleotidyl transferase; DAB, 3,3-diaminobenzidine; Cr, serum creatinine; BUN, blood urea nitrogen; ALT, alanine aminotransferase; AST, aspartate aminotransferase; ALP, alkaline phosphatase; CD, circular dichroism; PAGE, polyacrylamide gel electrophoresis; BSA, bovine serum albumin; GSH, glutathione; GSNO, S-nitrosoglutathione; R410C, genetic variant of human serum albumin mutated at position 410.

## Abstract

Recently, the cytotoxic activity of NO has been investigated for its potential use in anticancer therapies. Nitrosated human serum albumin (NO-HSA) may act as a reservoir of NO *in vivo*. However, there are no published reports regarding the effects of NO-HSA on cancer. Therefore, the present study investigated the anti-tumor activity of NO-HSA. NO-HSA was prepared by incubating HSA, which had been sulfhydrylated using iminothiolane, with isopentyl nitrite (6.64 mol NO/mol HSA). Anti-tumor activity was examined *in vitro* using murine colon 26 carcinoma (C26) cells and *in vivo* using C26 tumor-bearing mice. Exposure to NO-HSA increased the production of reactive oxygen species (ROS) in C26 cells. Flow cytometric analysis using rhodamine 123 showed that NO-HSA caused mitochondrial depolarization. Activation of caspase-3 and DNA fragmentation were observed in C26 cells after incubation with 100  $\mu$ M NO-HSA for 24 h, and NO-HSA inhibited the growth of C26 cells in a concentration-dependent manner. The growth of C26 tumors in mice was significantly inhibited by administration of NO-HSA compared with saline- and HSA-treatment. Immunohistochemical analysis of tumor tissues demonstrated an increase in TUNEL-positive cells in NO-HSA-treated mice, suggesting that inhibition of tumor growth by NO-HSA was mediated through induction of apoptosis. Biochemical parameters (such as Cr, BUN, AST and ALT) showed no significant differences among the three treatment groups, indicating that NO-HSA did not cause hepatic or renal damage. These results suggest that NO-HSA has the potential for chemopreventive and/or chemotherapeutic activity with few side effects.

## Introduction

Although cancer primarily arises from disorders of cell proliferation, it also may arise from disruptions in programmed cell death signaling pathways, resulting in decreased apoptosis of cancerous cells (Okada et al., 2004). Therefore, induction of apoptosis in neoplastic cells is a very effective therapy for tumor eradication (Meng et al., 2006). However, this type of chemotherapy often has negative side effects, such as transient cell cycle arrest, senescence, and autophagy. Drug delivery systems that facilitate selective apoptosis of neoplastic cells have been suggested as a way of overcoming this problem (Kaufmann et al., 2000; Kondo et al., 2005).

Nitric oxide (NO) is a unique diffusible molecular messenger that occupies a central role in mammalian pathophysiology (Brune et al., 1998). Its multiple actions include vascular smooth muscle relaxation (Moncada et al., 1986; Ignarro et al., 1989), inhibition of platelet aggregation (Azuma et al., 1986), effects on neurotransmission (Garthweite, 1991), and regulation of immune function (Marletta et al., 1988). Alternatively, under some circumstances, NO is cytotoxic (Laval et al., 1994). NO causes cellular iron losses and inhibits DNA synthesis, mitochondrial respiration, and aconitase activity in L10 hepatoma cells (Hibbs et al., 1988). In addition, NO reacts with superoxide anion (which is produced by activated macrophages and other cells), to form peroxynitrite. This byproduct of NO is a potent chemical oxidant, which alters protein function and damages DNA (Beckman et al., 1993). These effects are part of the nonspecific host defense, which facilitates killing of tumor cells and intracellular pathogens. In addition, the cytotoxicity arising from long-lasting NO generation has been attributed to induction of apoptosis (Brune et al., 1998).

Recently, the cytotoxic activity of NO has been studied to assess its therapeutic

potential in cancer treatment. NO-donating non-steroidal anti-inflammatory drugs (NO-NSAIDs), especially NO-aspirin (NO-ASA), have been investigated as promising chemopreventive agents (Fabbri et al., 2005; Kashifi et al., 2002; Williams et al., 2001). NO-ASA consists of traditional ASA to which a NO-releasing moiety is bound via a spacer. This agent induces oxidative stress by increasing intracellular peroxide and  $O_2^-$ , thereby inducing apoptosis via activation of the intrinsic apoptosis pathway (Gao et al., 2005). JS-K is a pro-drug designed to release NO after reacting with glutathione S-transferase, which induces double-stranded DNA breaks, activates DNA-damage-response pathways, and induces apoptosis in human multiple myeloma cells both *in vitro* and *in vivo* (Kiziltepe et al., 2007).

Human serum albumin (HSA) is an abundant circulating protein and the nitrosated form serves as a reservoir of NO (Stamler et al., 1992). Therefore, NO-HSA is an NO donor that is currently being investigated for its potential therapeutic applications. For example, administration of NO-HSA to animals with ischemia-reperfusion injury minimizes the tissue damage that occurs after reperfusion (Semisroth et al., 2005). In a balloon-injured rabbit femoral artery model, locally delivered NO-HSA preferentially binds to sites of vessel injury and inhibits both platelet accumulation and the subsequent development of neointimal hyperplasia (Marks et al., 1995). NO-HSA also shows potent antibacterial activity and inhibits the proliferation of cultured human vascular smooth muscle cells (Ishima et al., 2007). However, there are no reports describing the effects of NO-HSA on cancer.

Accordingly, the present study evaluated the anti-tumor activity of NO bound to HSA (NO-HSA) via an S-nitrosothiol linkage using iminothiolane as a spacer. The molecular events related to induction of apoptosis by NO-HSA were studied *in vitro* and

the antitumor activity of NO-HSA was studied *in vivo* using a murine model of C26 colon carcinoma.

## Methods

### *Chemicals*

Traut's Reagent (2-imminothiolane) was purchased from Pierce Chemical Co. (Rockford, IL, USA). Isopentyl nitrite, diethylenetriaminepentaacetic acid (DTPA) and Cell Counting Kit-8 were purchased from Wako Pure Chemical Industries, Ltd. (Osaka, Japan). RPMI-1640 medium, Hanks' balanced salt solution (HBSS) and RNase A was obtained from Sigma Chemical (St Louis, MO, USA). Proteinase K was obtained from Roche Applied Science (Indianapolis, IN, USA). All other reagents used were of the highest grade available from commercial sources.

### *Expression and purification of recombinant human serum albumin (rHSA)*

rHSA was produced using a yeast expression system as described previously (Matsushita et al., 2004). Briefly, for constructing the HSA expression vector pPIC9-HSA, native HSA coding region was incorporated into the methanol-inducible pPIC9 vector (Invitrogen Co., San Diego, CA, U.S.A.). The resulting vector was introduced into the yeast species *P. pastoris* (strain GS115) to express rHSA. Secreted rHSA was isolated from the growth medium by a combination of precipitation with 60% (w/v)  $(\text{NH}_4)_2\text{SO}_4$  and purification on a Blue Sepharose CL-6B column (GE Healthcare) followed by Phenyl HP column (GE Healthcare). Isolated protein was defatted by using the charcoal procedure described by Chen (Chen, 1967), deionized, freeze-dried and then stored at  $-20^\circ\text{C}$  until use. The resulting rHSA (treated with dithiothreitol) exhibited a single band on SDS/PAGE. Density analysis of protein bands stained with Coomassie Brilliant Blue showed that its purity was more than 97%.

### ***Synthesis of nitroso-albumin (NO-HSA)***

Terminal sulfhydryl groups were added to the HSA molecule by incubating 0.15 mM rHSA with 3 mM Traut's Reagent (2-iminothiolane) in 100 mM potassium phosphate buffer (KPB) containing 0.5 mM DTPA (pH 7.8) for 1 h at room temperature. The resultant modified rHSA then was S-nitrosylated by 3 h incubation with 15 mM isopentyl nitrite at room temperature (Figure 1). The resulting NO-HSA was concentrated, exchanged with saline using a PelliconXL filtration device (Millipore Corporation, Billerica, MA, USA), and the final concentration adjusted to 2 mM NO-HSA. The sample was stored at -80 °C until use.

### ***Determination of S-nitrosylation efficiency***

The amount of the S-nitroso moieties of NO-HSA was quantified using a 96-well plate. First, 20- $\mu$ l aliquots of NO-HSA solution and NaNO<sub>2</sub> (standard) were incubated with 0.2 ml of 10 mM sodium acetate buffer (pH 5.5) containing 100 mM NaCl, 0.5 mM DTPA, 0.015% N-1-naphthylstyrene-diamide and 0.15% sulfanilamide with or without 0.09 mM HgCl<sub>2</sub>, for 30 min at room temperature. Then, the absorbance was measured at 540 nm. The number of moles of NO per mole of HSA, was obtained by subtracting the values in the absence of HgCl<sub>2</sub> from values in the presence of HgCl<sub>2</sub>; the value thus obtained, was  $6.64 \pm 0.54$  mol NO/mol HSA.

### ***Cellular experiments with Colon-26 (C26) cells***

C26 cells, which were donated by the Institute of Development, Aging and Cancer, at Tohoku University (Sendai, Miyagi, Japan), were cultured at 37°C in RPMI 1640

medium containing 10% fetal calf serum (FCS), 100 U/ml penicillin, and 10 µg/ml streptomycin (medium A). Trypsin (0.25%) EDTA solution was used to detach the cells from the culture flask for plating and passing the cells. All cell culture experiments were performed at 37°C in a humidified atmosphere of 5% CO<sub>2</sub> in air.

For detection of reactive oxygen species (ROS), C26 cells (1.0 x 10<sup>4</sup> cells/well) were cultured in 96-well plates in medium A for 12 h, washed twice with PBS, and incubated for an additional 30 min in HBSS containing 5 µM 5-(and-6)-chloromethyl-2',7'- dichlorodihydrofluorescein diacetate, acetyl ester (CM-H<sub>2</sub>DCFDA) (Molecular probes). After washing twice with HBSS, the cells were cultured in HBSS containing 5% FCS for 15 min followed by the addition of PBS, 50 µM HSA or 50 µM NO-HSA. After incubation, fluorescence was measured using a plate reader (excitation wavelength, 485 nm; emission wavelength, 535 nm). The change in fluorescence was calculated by subtracting the fluorescence at 0 h from the fluorescence measured at the indicated times. The fluorescence intensities of cells incubated with PBS, 50 µM HSA and 50 µM NO-HSA at 0 h were 201.3, 166.1 and 181.3, respectively.

Changes in the mitochondrial membrane potential of C26 cells were monitored using flow cytometric analysis with rhodamine 123 staining. C26 cells (1.0 x 10<sup>6</sup> cells/well) were cultured in 6-well plates for 12 h, washed twice with PBS and incubated with PBS and either 100 µM HSA or various concentrations of NO-HSA in medium A for 24 h. The cells were then trypsinized, washed twice with PBS, and incubated for 15 min with 5 µg/ml rhodamine 123. The mean fluorescence intensity of rhodamine 123 in the cells was measured using a flow cytometer (FACS Calibur; Becton Dickinson Biosciences, Franklin Lakes, USA).

For the determination of caspase-3 activity, cells were cultured to confluence in 6-well plates, washed twice with PBS, and incubated with medium A containing 100  $\mu$ M HSA or various concentrations of NO-HSA. Cells were incubated for 24 h, trypsinized, and washed with 0.2 ml of ice-cold PBS. The cell pellet was resuspended in 15  $\mu$ l cell lysis buffer, lysed by freeze-thawing and then incubated on ice for 15 min. The cell lysates were centrifuged at 15,000 rpm for 20 minutes at 4 °C, and the supernatant fraction was collected (cell extract). The caspase-3 activity in the cell extract was assessed using the colorimetric CaspASE<sup>TM</sup> Assay System (Promega), according to the manufacturer's instructions.

For the detection of DNA degradation (DNA ladder), C26 cells ( $1 \times 10^6$  cell/well) were cultured in 6-well plates. Cells were cultured for 12 h, washed twice with PBS, and incubated with PBS and either 100  $\mu$ M HSA or various concentrations of NO-HSA for 24 h. The cells then were trypsinized, collected, and centrifuged at 4000 rpm for 10 min. After removing the supernatant, the cell pellet was resuspended in 0.2 ml of PBS and centrifuged at 4000 rpm for 10 min. The supernatant was again removed and the remaining pellet was incubated in 20  $\mu$ l 10 mM Tris-HCl buffer (pH 7.8) containing 2 mM EDTA and 0.5 % SDS (cell lysis buffer) for 10 min at 4 °C, followed by centrifugation at 15000 rpm for 5 min. The resulting supernatant (cell extract) was collected and incubated with 1  $\mu$ l RNase A (10  $\mu$ g/ml) for 30 min at 50 °C. 1  $\mu$ l proteinase K (10  $\mu$ g/ml) was added to the cell extract, followed by a 1 h incubation at 50 °C. The resulting DNA extract was electrophoresed in a 2.0% agarose gel, followed by staining of the gel with ethidium bromide and visualization of the DNA bands using ultraviolet illumination.

The cell viability assay was performed using the Cell Counting Kit-8 (WST-8),

which is based on the MTT assay. C26 cells were plated in 96-well plates at  $1.0 \times 10^4$  cells/well and were cultured for 32 h in medium A. Then, the cells were washed twice with PBS and incubated in a total volume of 0.2 ml medium A containing various concentrations of HSA or NO-HSA. After incubating the cultures for various lengths of time, 5  $\mu$ l WST-8 solution was added to each well and the cells were incubated for an additional 2 h at 37°C. The number of surviving cells was determined by measuring the absorbance at 450 nm. Cell viability was calculated as the percentage of the control value (without HSA or NO-HSA) (Ishiyama et al., 1996).

### ***Animal experiments***

Five-week-old male BALB/c AnNCrIcrlj mice (17 - 20 g), were purchased from Charles River (Calco, Como, Italy). The mice were housed in a 12/12 h light/dark cycle in a humidity-controlled room. Mice were acclimated for at least 5 days before being used in experiments.

For tumor induction, mice were inoculated with C26 cells ( $1 \times 10^6$  cells/mouse) by a subcutaneous injection into the dorsal skin. Three days after inoculation, C26 carcinoma-bearing mice were randomly divided into three groups: control, HSA, and NO-HSA. The mice received a daily intravenous injection of saline, HSA (10  $\mu$ mol/kg), or NO-HSA (10  $\mu$ mol/kg), for 10 days from day 3 to 12 post-inoculation. Tumor volume was calculated using the formula  $0.4(a \times b^2)$ , where 'a' is the largest, and 'b' is the smallest, diameter of the tumor (Shimizu et al., 2005), and monitored from day 7 to 17 after inoculation. Variance in each group was evaluated using the Bartlett test, and differences in mean tumor volume were evaluated using the Tukey-Kramer test.

Some animals in each of the three treatment groups were used for

immunohistochemical analysis and serum biochemistry. When the mice received five times per day on days from 3 to 7 post-inoculation (the tumors in each group reached approximately 5 mm in diameter), blood samples were collected from the abdominal vena cava under diethylether anesthesia approximately 2 h after the daily injection and then the mice were sacrificed.

Tumors were dissected, fixed immediately with 2% periodate/lysine/ paraformaldehyde fixative at 4°C for 5 h and washed with a graded series of sucrose solutions in PBS (10, 15 and 20%). After immersion in 20% sucrose in PBS to inhibit ice crystal formation, the tissues were embedded in O.C.T compound (Sakura Fine Technochemical, Tokyo, Japan), frozen in liquid nitrogen, and stored at -80°C. Five- $\mu$ m tumor sections were prepared using a cryostatic microtome (HM500M, Microm, Walldorf, Germany), and were mounted on poly-L-lysine-coated slides. The slides were stained using the TUNEL method and an *in situ* Apoptosis Detection Kit (Takara-Bio Co.Ltd, shiga, Japan). The slides were washed 3 times with 0.01 M phosphate buffer (pH 7.4) containing 0.9 % NaCl, followed by application of methanol containing 0.3% H<sub>2</sub>O<sub>2</sub> to inactivate endogenous peroxidase and incubation at room temperature for 30 min. The slides were washed 3 times with 0.01 M phosphate buffered saline, and then were incubated in 100 ml of permeabilization buffer on ice for 5 min. The slides were washed 3 times with 0.01 M phosphate buffered saline, and then were incubated with 50 ml of freshly prepared terminal deoxynucleotidyl transferase(TdT) reaction mixture (TdT enzyme 5 ml + Labeling Safe buffer 45 ml) at 37°C for 60 min. After the slides were washed 3 times with 0.01 M PBS, they were incubated in 70 ml of anti-FITC-HRP conjugate antibody at 37°C for 30 min. After the slides were washed 3 times with 0.01 M PBS, they were incubated in DAB-H<sub>2</sub>O<sub>2</sub> reaction buffer at room temperature for 10

min. After the slides were washed 3 times with distilled water, they were stained with 3% methyl green for 10 min, dehydrated, penetrated and sealed (Gravrieli et al., 1992). Each slide then was visualized under a light microscope (Olympus, Tokyo, Japan), at a magnification of x 400.

Serum was separated by centrifugation. Routine clinical laboratory techniques were used to determine the concentrations of total protein, serum creatinine (Cr), blood urea nitrogen (BUN), alanine aminotransferase (ALT), aspartate aminotransferase (AST) and alkaline phosphatase (ALP) in serum. Variance in each group was evaluated using the Bartlett test, and differences were evaluated using the Tukey-Kramer test.

## Results

### *NO-HSA induces cell death via apoptosis in vitro*

Apoptosis is induced by a variety of factors. Among them, it is well-known that intracellular accumulation of ROS, such as  $\text{H}_2\text{O}_2$ ,  $\text{O}_2^-$  and peroxynitrite, causes apoptosis. Moreover, production of ROS also plays a major role in NO-ASA-induced apoptosis. To examine whether NO-HSA promoted ROS production in C26 cells, a fluorescent probe (CM- $\text{H}_2\text{DCFDA}$ ), which undergoes conversion to 2', 7'-dichlorofluorescein (DCF) in the presence of intracellular ROS, was used. Addition of NO-HSA to C26 cells increased the amount of ROS compared with treatment with vehicle or HSA (Figure 2). In addition, the ROS in the C26 cell culture medium increased with time after addition of NO-HSA. This result suggests that NO-HSA promotes a signal cascade leading to apoptosis by increasing intracellular production of ROS.

To evaluate the effect of NO-HSA on mitochondrial function, C26 cells were loaded with a mitochondria-selective fluorescent cation (rhodamine 123) to monitor the mitochondrial membrane potential. Compared with vehicle, in cells treated with 50 or 100  $\mu\text{M}$  NO-HSA rhodamine 123, fluorescence was decreased by 75%; whereas treatment with 25  $\mu\text{M}$  NO-HSA or 100  $\mu\text{M}$  HSA did not affect rhodamine 123 fluorescence compared with vehicle (Figure 3A). These observations indicate that NO-HSA induces depolarization of the mitochondrial membrane.

Caspase-3 is a cell-death protease that is involved in the downstream execution phase of apoptosis. During this phase of apoptosis, cells undergo morphological changes, such as DNA fragmentation, chromatin condensation, and formation of

apoptotic bodies. Compared with the effect of vehicle, cells treated with 25 or 50  $\mu\text{M}$  NO-HSA had slightly increased caspase-3 activity, and cells treated with 100  $\mu\text{M}$  NO-HSA showed a larger increase in caspase-3 activity (Figure 3B). Cells treated with HSA had the same level of caspase-3 activity as cells treated with PBS.

To further confirm that NO-HSA induced apoptosis in C26 cells, DNA fragmentation, which is a typical morphological change in the execution phase of apoptosis, was examined. DNA fragmentation was observed in C26 cells treated with 100  $\mu\text{M}$  NO-HSA, but not in C26 cells treated with 25 or 50  $\mu\text{M}$  NO-HSA (Figure 3C). Neither vehicle nor 100  $\mu\text{M}$  HSA induced DNA fragmentation in C26 cells (Figure 3C). These findings suggest that NO-HSA induces apoptosis by increasing ROS production, activating caspase-3, and hyperpolarizing the mitochondrial membrane potential.

To determine the effect of NO-HSA on cell growth, the viability of C26 cells was examined after treatment with HSA or various concentrations of NO-HSA. NO-HSA inhibited growth of C26 cells in a concentration-dependent manner; cell growth was suppressed by 71, 80, and 85% after a 48-h incubation with 25, 50, and 100  $\mu\text{M}$  NO-HSA, respectively (Figure 4A). The viability of C26 cells incubated with 50  $\mu\text{M}$  NO-HSA significantly decreased with increasing incubation times (Figure 4B). NO-HSA inhibited growth to a greater extent than did HSA, which had only a weak inhibitory effect. These results suggest that NO-HSA inhibits cell growth of C26 cells by inducing apoptosis.

### ***NO-HSA exerts anti-tumor effect via the apoptotic pathway in vivo***

To investigate the anti-tumor effect of NO-HSA *in vivo*, C26 tumor-bearing mice were intravenously injected with saline, HSA or NO-HSA. Mean tumor area increased

with time in the saline-treated group. In the HSA-treated group, tumor growth was suppressed, compared with that in the control group, but the difference was not statistically significant. In contrast, tumor growth was significantly inhibited by administration of NO-HSA (Figure 5).

To clarify whether the suppression of tumor growth by NO-HSA is mediated via apoptosis, tumor tissues from C26 tumor-bearing mice injected with NO-HSA were examined using immunohistochemistry. In NO-HSA-treated mice, there were more TUNEL-positive cells than in the saline- and HSA-treated animals. In addition, the tumor tissue architecture was less defined in animals treated with NO-HSA than in the other groups, suggesting that NO-HSA induced apoptosis in C26 tumor cells and thus exerted an anti-tumor effect *in vivo* (Figure 6).

To evaluate the side effects of NO-HSA treatment, several serum biochemical parameters were measured in tumor-bearing mice treated with saline, HSA or NO-HSA (Table 1). There were no significant differences in total protein, Cr, BUN, AST or ALT among the three groups, suggesting that NO-HSA did not cause kidney or liver damage. However, compared with the control group, mice treated with HSA had significantly lower serum levels of ALP ( $345.3 \pm 4.7$  U/l versus  $385.3 \pm 18.3$  U/l). Moreover, the serum concentration of ALP in mice treated with NO-HSA was  $300.5 \pm 23.0$  U/l, which was significantly lower than the control ( $p < 0.01$ ) and HSA ( $p < 0.05$ ) groups. In general, ALP levels increase in several types of cancer, such as liver, lung and bone cancer; thus, the present findings suggest that NO-HSA is an effective anti-cancer agent. The vasodilating effect of NO-HSA was also evaluated in rats after intravenous injection at a dose of  $10 \mu\text{mol/kg}$  ( $66 \mu\text{mol}$  of NO per kilogram). NO-HSA induced a decrease in the mean arterial blood pressure immediately after intravenous injection and the maximum

reduction effect was  $32.8 \pm 7.3$  mmHg. In contrast, HSA had no significant effect on the blood pressure. The fall in pressure returned to the initial levels in 30 min (data not shown).

## Discussion

There have been many trials of NO as a therapeutic agent, because of its powerful biological activity (Moncada et al., 1993). However, the *in vivo* half-life of NO (~0.1 s) is often too short to capitalize on its potential biological actions. The half-life of NO can be prolonged by adding S-nitrosothiols moieties with cysteine residues of proteins. For example, nitrosated HSA appears to act as a reservoir of NO *in vivo* (Stamler et al., 1992). Simon et al. incubated bovine serum albumin (BSA) with 200-fold excess concentration of NaNO<sub>2</sub> under acidic condition to synthesize polynitrosylated BSA that is highly modified at the thiol group of cysteine, hydroxyl group of tyrosine and amines (38 mol NO/mol BSA). The polynitrosylated BSA has been shown to exhibit antiplatelet activity. However polynitrosylated S-NO-BSA, a NO-BSA conjugate prepared with the same method except that the BSA has been reduced with dithiothreitol and contains 19 mol of “S-NO” per mol of BSA, was a significantly more potent platelet inhibitor than polynitrosylated BSA described above (Simon et al., 1996). These findings show that nitrosylated BSA behave as a NO donor, in particular, the poly(S-nitroso) derivative could be by far the most potent compound. One molecule of HSA contains 35 cysteine residues, 34 of which form 17 non- reactive disulfide bonds, and one of which (Cys-34) forms a reactive free thiol (Peters, 1985). Thus, the number of NO molecules that can be bound to HSA is limited because only one free cysteine per HSA molecule is available for conjugation. Ewing et al. increased the number of free sulfhydryl groups on bovine serum albumin (BSA) by reduction with dithiothreitol and thiolation with N-acetylhomocysteine, thereby preparing poly-nitrosated BSA (12-15 mol NO/mol BSA) (Ewing et al., 1997). Marks et al. produced poly-nitrosated BSA (5.9

mol NO/mol BSA) by adding free sulfhydryl groups to the molecule and by treating the BSA with N-acetylhomocysteine thiolactone (Marks et al., 1995). However, the poly-nitrosated BSAs prepared in these studies formed aggregates as a result of intermolecular disulfide formation. Aggregate formation results in molecular heterogeneity, which limits the therapeutic application of S-nitroso residues. In the present study, iminothiolane, which reacts with primary amines to introduce sulfhydryl groups while maintaining charge properties similar to the original amino groups, was selected as the thiolation reagent. Iminothiolane was used to produce poly-nitrosated HSA (NO-HSA) (6.6 mol NO/mol HSA), which did not form aggregates after non-reducing SDS-PAGE or native-PAGE (data not shown). Moreover, the far-UV spectra of NO-HSA were nearly identical to those of HSA (data not shown). Therefore, NO-HSA is expected to be clinically applicable as a biocompatible pharmacological agent, although further study is required to clarify other potential issues, including the antigenicity of this protein.

NO-NSAIDs have been extensively investigated as therapeutic agents for cancer due to their ability to release NO, thereby promoting apoptosis. NO-NSAIDs are categorized as organic nitrate esters, which are readily reduced to organic nitrite esters by cytosolic enzymes. Subsequently, GSH reacts with organic nitrite esters to form GSNO, indicating that NO-NSAIDs release NO via S-nitrosothiol (Wong et al., 1999). Alternatively, the transfer of NO from NO-HSA to the cytosol could be inferred from a study by Ramachandran et al. The authors reported that NO is released from extracellular S-nitrosothiols by a cell-surface enzyme (protein disulfide isomerase) and accumulates in the cell membrane where it reacts with O<sub>2</sub> to produce N<sub>2</sub>O<sub>3</sub>, which is then available for nitrosation reactions with intracellular thiols at the membrane-cytosol

interface (Ramachandran et al., 2001). Therefore, it is likely that NO-HSA also releases NO by the intracellular formation of S-nitrosothiol, suggesting that the species of NO released within the cell by S-nitrosothiols, as well as the reactive substances (such as ROS) derived from the released NO, would not differ significantly between NO-NSAIDs and NO-HSA. In support of this hypothesis, NO-HSA caused depolarization of the mitochondrial membrane potential, activation of caspase-3 and DNA fragmentation in the present study, consistent with the effects of NO-NSAIDs. Additional studies are needed to determine the details of the molecular events and the systematic pathways affected by NO-HSA, but the mechanism of action should be similar to that of NO-NSAIDs. Recently, Gao J et al. elucidated the detailed mechanism of apoptosis induced by NO-ASA. Intracellular accumulation of ROS is a key proximal event in NO-ASA-induced apoptosis and correlates with the effect on tumor cell growth (Gao et al., 2005). In the present study, NO-HSA induced accumulation of ROS in tumor cells, suggesting that increased ROS production may be an important proximal event leading to induction of apoptosis.

The results of the *in vivo* study showed that NO-HSA significantly suppressed tumor growth by inducing apoptosis, without adverse changes in serum biochemical parameters in treated mice. Recently, Trachootham et al., using immortalized cell lines and their oncogenic progeny transfected with *H-Ras*<sup>V12</sup>, demonstrated that cancer cells typically produce more ROS than normal cells. Moreover, the pro-oxidant status of cancer cells increases their susceptibility to treatment with agents that cause oxidative stress, as demonstrated in a study using  $\beta$ -phenylethyl isothiocyanate (Trachootham et al., 2006). In addition, Feng et al. reported that cyaniding-3-rutinoside selectively induces accumulation of peroxides in HL-60 human leukemic cells, but not in normal

PBMC (Feng et al., 2007). Schumacker has proposed that ROS toxicity induced by certain chemo-therapeutic agents may be an effective means of selectively eradicating malignant cells (Shumacker, 2006). In the present study, we presumed that although NO reacts with superoxide anion to form peroxynitrite (a potent oxidant and nitrating agent), these highly reactive oxidant species are likely produced at higher levels in C26 cells compared with normal cells. Therefore, NO-HSA may show selective cytotoxicity for tumor cells and not affect normal cells. These findings strongly suggest that NO-HSA is a promising therapeutic anti-cancer agent, given the unusual redox conditions typical of malignant cells.

Anti-apoptotic effects of NO have been observed in a variety of cells, including T cells, hepatocytes, endothelial cells, neurons, ovarian follicle cells, eosinophils, thymocytes and embryonic kidney cells (Liu et al., 1999). In a recent study using U937 human promonocytic cells, NO-R410C (a genetic variant of HSA) had anti-apoptotic activity (Ishima et al., 2007). Whether NO ultimately inhibits or promotes apoptosis likely depends on the cell, the signal, the source, the molecule, the amount, and the presence or absence of co-reactants. For example, the amount of NO bound to the carrier molecule appears to account for the discrepant results between previous investigations and the present study. The NO content of NO-R410C (in previous study) and NO-HSA (in this study) were 1.3 and 6.6 mol NO/mol HSA, respectively, and the S-nitroso moiety concentrations *in vitro* were 26-130 and 165-660  $\mu\text{M}$ , respectively. Mohr et al. reported that 10-100  $\mu\text{M}$  GSNO inhibits the activation of caspase-3 induced by actinomycin D in U937 cells (Mohr et al., 1997). In contrast, apoptosis (characterized by DNA fragmentation and morphological changes) was observed in U937 cells treated with GSNO at concentrations in excess of 250  $\mu\text{M}$  (Mebmer et al.,

1995). Based on the results of the present study and those of previously published investigations, the critical NO threshold concentration between promotion and inhibition of apoptosis appears to be 100-200  $\mu$ M.

Matsumura et al. examined the accumulation of differently sized proteins within tumor tissues of tumor-bearing mice. Macromolecules containing HSA tended to accumulate in tumor tissues, apparently due to hypervascularization and enhanced vascular permeability (even to macromolecules) of the tumors, with little export of macromolecules from the tumor tissue via blood or lymphatic vessels (Matsumura et al., 1987). Therefore, NO-HSA may be a useful agent for targeting chemotherapeutics to tumor tissue. However, the short half-life of NO has been one of the greatest obstacles to therapeutic application of NO donors. Consequently, the pharmacokinetic properties of NO-HSA in mice were measured to determine the biological fate of NO. The apparent half-life of S-nitroso moieties in NO-HSA was estimated to be 18.9 min (data not shown), which is similar to that of NO-R410C, but much longer than that of the low-molecular-weight NO donor, GSNO (4.2 min) (Ishima et al., 2007). In the present study, the difference in NO half-life between NO-HSA and GSNO appeared to be due to reduced renal excretion of HSA compared with GSH due to its larger molecular size, suggesting that HSA may be a useful NO carrier *in vivo*.

In summary, NO-HSA was synthesized by inducing S-nitrosothiol linkages using iminothiolane as a spacer. NO-HSA generated ROS in C26 cells, induced intrinsic apoptotic events, such as depolarization of mitochondria membrane potentials, activation of caspase-3 and induction of DNA fragmentation. Moreover, NO-HSA inhibited proliferation of tumor cells *in vitro* in a concentration-dependent manner. In the *in vivo* experiments, NO-HSA also strongly inhibited tumor growth by inducing

apoptosis, with no side-effects. The results of the present study suggest that NO-HSA has promise as a new-generation anti-cancer agent with few side effects.

## References

- Azuma H, Ishikawa M, and Sekizaki S (1986) Endothelium-dependent inhibition of platelet aggregation. *Br J Pharmacol* **88**: 411-415.
- Beckman JS and Crow JP (1993) Pathological implication of nitric oxide, superoxide and peroxynitrite formation. *Biochem Soc Trans* **21**: 330-334.
- Brune B, Sandau K, and Knethen AV (1998) Apoptotic cell death and nitric oxide: activating and antagonistic Transducing Pathways. *Biochemistry (Moscow)* **63**: 817-825.
- Chen RF (1967) Removal of fatty acids from serum albumin by charcoal Treatment. *J Biol Chem* **242**: 173-181.
- Ewing JF, Young DV, Janero DR, Garvey DS, and Grinnell TA (1997) Nitrosylated bovine serum albumin derivatives as pharmacologically active nitric oxide congeners. *J Pharmacol Exp Ther* **283**: 947-954.
- Fabbri F, Brigliadori G, Ulivi P, Tesei A, Vannini I, Rosetti M, Bravaccini S, Amadori D, Bolla M, and Zoli W (2005) Pro-apoptotic effect of a nitric oxide-donating NSAID, NCX4040, on bladder carcinoma cells. *Apoptosis* **10**: 1095-1103.
- Feng R, Ni HM, Wang SY, Tourkova IL, Shulin MR, Harada H, and Yin XM (2007) Cyanidin-3-rutinoside, a natural polyphenol antioxidant, selectively kills leukemic cells by induction of oxidative stress. *J Biol Chem* **282**: 13468-13476.
- Gao J, Liu X, and Rigas B (2005) Nitric oxide-donating aspirin induces apoptosis in human colon cancer cells through induction of oxidative stress. *Proc Natl Acad Sci USA* **102**: 17207-17212.
- Garthweite J (1991) Glutamate, nitric oxide and cell-cell signaling in the nervous system. *Trends Neurosci* **14**: 60-67.

- Gavrieli Y, Sherman Y, and Ben-Sasson SA (1992) Identification of programmed cell death in Situ via specific labeling of nuclear DNA fragmentation. *J Cell Biol* **119**: 493-501.
- Hibbs JB Jr, Taintor RR, Vavrin Z, and Rachlin EM (1988) Nitric oxide: A cytotoxic activated macrophage effector molecule. *Biochem Biophys Res Commun* **157**: 87-94.
- Ignarro LJ (1989) Endothelium-derived nitric oxide - Pharmacology and relationship to the actions of organic nitrate esters. *Pharmacol Res* **6**: 651-659.
- Ishima Y, Sawa T, Kragh-Hansen U, Miyamoto Y, Matsushita S, Akaike T, and Otagiri M (2007) S-nitrosylation of human variant albumin Liprizzi (R410C) confers potent antibacterial and cytoprotective properties. *J Pharmacol Exp Ther* **320**: 969-977.
- Ishiyama M, Tominaga H, Shiga M, Sasamoto K, Ohkura Y, and Ueno K (1996) A combined assay of cell viability and in vitro cytotoxicity with a highly water-soluble tetrazolium salt, neutral red and crystal violet. *Biol Pharm Bull* **19(11)**: 1518-1520.
- Kashifi K, Ryann Y, Qiao LL, Williams JL, Chen J, Soldato PD, Traganos FT, and Rigas B (2002) Nitric oxide-donating nonsteroidal anti-inflammatory drugs inhibit the growth of various cultured human cancer cells: Evidence of a tissue type-independent effect. *J Pharmacol Exp Ther* **303**: 1273-1282.
- Kaufmann SH and Gores GJ (2000) Apoptosis in cancer: cause and cure. *Bioessays* **22**: 1007-1017.
- Kiziltepe T, Hideshima T, Ishitsuka K, Ocio EM, Raje N, Catley L, Li CQ, Trudel LJ, Yasi H, Vallet S, Kutok JL, Chauhan D, Mitsiades CS, Saavedra JE, Wogan GN, Keefer LK, Shami PJ, and Anderson KC (2007) JS-K, a GST-activated nitric oxide generator, induces DNA double-strand breaks, activates DNA damage response pathways, and induces apoptosis in vitro and in vivo in human multiple myeloma cells.

*Blood* **110**: 709-718.

Kondo Y, Kanzawa T, Sawaya R, and Kondo S (2005) The role of autophagy in cancer development and response to therapy. *Nat Rev Cancer* **5**: 726-734.

Laval F and Wink DA (1994) Inhibition by nitric oxide of the repair protein, O6-methylgluanine-DNA-methyltransferase. *Carcinogenesis* **15**: 443-447.

Liu L and Stamler JS (1999) NO: an inhibitor of cell death. *Cell Death and Differentiation* **6**: 937-942.

Marks DS, Vita JA, Folts JD, Keaney JF Jr, and Welch GN (1995) Loscalzo J. Inhibition of neointimal proliferation intimal proliferation in rabbits after vascular injury by a single treatment with a protein adduct of nitric oxide. *J Clin Investig* **96**: 2630-2638.

Marletta MA, Yoon PS, Iyengar R, Leaf CD, and Wishnok JS (1988) Macrophage oxidation of L-arginine to nitrite and nitrate - Nitric oxide is an *intermediate*. *Biochemistry* **27**: 8706-8711.

Matsumura Y, Oda T, and Maeda H (1987) General mechanism of intratumor accumulation of macromolecules: Advantage of macromolecular therapeutics. *Jpn J Cancer Chemother* **14**: PART II 821-829.

Matsushita S, Ishima Y, Chuang VT, Watanabe H, Tanase S, Maruyama T, and Otagiri M (2004) Functional analysis of recombinant human serum albumin domains for pharmaceutical applications. *Pharm Res* **10**: 1924-1932.

Mebmer UK, Reimer DM, and Brune B (1995) Nitric oxide-induced apoptosis: p53-dependent and p53-indeoendent signaling pathways. *Biochem J* **319**: 299-305.

Meng XW, Lee SH, and Kaufmann SH (2006) Apoptosis in the treatment of cancer: a promise kept? *Curr Opin Cell Biol* **18**: 668-676.

Mohr S, Zech B, Lapetina EG, and Brune B (1997) Inhibition of caspase-3 by

- S-nitrosation and oxidation caused by Nitric oxide. *Biochem Biophys Res Com* **238**: 387-391.
- Moncada S and Higgs A (1993) The L-arginine-nitric oxide pathway. *N Engl J Med* **329**: 2002-2012.
- Moncada S, Palmer RMJ, and Gryglewski RJ (1986) Mechanism of action of some inhibitors of endothelium-derived relaxing factor. *Proc Natl Acad Sci USA* **83**: 9164-9168.
- Okada H and Mak TW (2004) Pathways of apoptotic and non-apoptotic death in tumor cells. *Nat Rev Cancer* **4**: 592-603.
- Peters T Jr (1985) Serum albumin. *Adv Protein Chem* **37**: 161-245.
- Ramachandran N, Root P, Jiang XM, Hogg OJ, and Mutus B (2001) Mechanism of transfer of NO from extracellular S nitrosothiols into the cytosol by cell-surface protein disulfide isomerase. *Proc Natl Acad Sci USA* **98**: 9539-9544.
- Schumacker PT (2006) Reactive oxygen species in cancer cells: Live by sword, die by the sword. *Cancer Cell* **10**: 175-176.
- Semisroth S, Fellner B, Trescher K, Bernecker OY, Kalinowski L, Gasser H, Hallstrom S, Malinski T, and Podesser BK (2005) S-Nitroso human serum albumin attenuates ischemia/reperfusion injury after cardioplegic arrest in isolated rabbit hearts. *J Heart Lung Transplant* **24**: 2226-2234.
- Shimizu K, Asai T, Fuse C, Sadzuka Y, Sonobe T, Ogino K, Taki T, Tanaka T, and Oku N (2005) Applicability of anti-neovascular therapy to drug-resistant tumor: Suppression of drug-resistant P388 tumor growth with neovessel-targeted liposomal adriamycin. *Int J Pharm* **296**: 133-141.
- Simon DI, Mullins ME, Jia L, Gaston B, Singel DJ, and Stampler JS (1996)

Polynitrosylated proteins: Characterization, Bioactivity, and functional consequences.

*Proc Natl Acad Sci USA* **93**: 4736-4741.

Stamler JS, Jaraki O, Osborne J, Simon DI, Keaney J, Vita J, Singel D, Valeri CR, and

Loscalzo J (1992) Nitric oxide circulates in mammalian plasma primarily as an

S-nitroso adduct of serum albumin. *Proc Natl Acad Sci USA* **89**: 7674-7677.

Trachootham D, Zhou Y, Zhang H, Demizu Y, Chen Z, Pelicano H, Chiao PJ, Achanta G,

Arlinghaus RB, Liu J, and Huang P (2006) Selective killing of oncogenically

transformed cells through a ROS-mediated mechanism by  $\beta$ -phenylethyl

isothiocyanate. *Cancer Cell* **10**: 241-252.

Williams JL, Borgo S, Hasan I, Castillo E, Traganos F, and Rigas B (2001) Nitric

oxide-releasing nonsteroidal anti-inflammatory drugs (NSAIDs) alter the kinetics of

human colon cancer cell lines more effectively than traditional NSAIDs: Implication

for colon cancer chemoprevention. *Cancer Res* **61**: 3285-3289.

Wong PS and Fukuto JM (1999) Reaction of organic nitrate esters and S-nitrosothiols

with reduced flavins: A possible mechanism of bioactivation. *Drug Metab Dispos*

**2704**: 502-509.

### **Legends for Figures:**

**Figure 1. The scheme for the reaction of 2-iminothiolane with primary amines followed by S-nitrosylation**

**Figure 2. Production of ROS in C26 cells after NO-HSA treatment**

C26 cells were pre-treated with H<sub>2</sub>DCFDA for uptake into C26 cells and hydrolysis by cellular esterase, followed by addition of either PBS (*open circles*), 50 μM HSA (*open squares*) or 50 μM NO-HSA (*closed circles*). Excitation of the probes was done at 485 nm and emission was measured at 535 nm. Change in fluorescence was calculated by subtracting the fluorescence at 0 h from that at the indicated times. The fluorescence intensities of the PBS, 50 μM HSA and 50 μM NO-HSA groups at 0 h were 201.3, 166.1 and 181.3, respectively. Results are the mean ± S.D. of three separate experiments.

**Figure 3. Induction of apoptosis of C26 cells after NO-HSA treatment**

**(A) Alteration in the mitochondrial membrane potential after NO-HSA treatment**

C26 cells were cultured with PBS, 100 μM HSA or various concentrations of NO-HSA for 24 h, followed by addition of rhodamine 123. The amounts of cell-associated rhodamine 123 were determined as described in Materials and Methods. Results are the mean ± S.D. of three separate experiments.

**(B) Activation of caspase-3 after NO-HSA treatment**

C26 cells were incubated with PBS, 100 μM HSA or various concentrations of NO-HSA for 24 h. Caspase-3 activity was estimated by monitoring *p*-nitroaniline (absorbance at 405 nm) released from the substrate upon cleavage by caspase-3. Change

in absorbance was calculated by subtracting absorbance after incubation with caspase inhibitor (Z-VAD-FMK), from absorbance after incubation without caspase inhibitor. The absorbances among PBS-, HSA- and NO-HSA-treated cells incubated with Z-VAD-FMK were almost identical ( $0.170 \pm 0.17$ ). Results are the means  $\pm$  S.D. of three separate experiments.

### (C) DNA fragmentation after NO-HSA treatment

C26 cells were incubated with PBS, 100  $\mu$ M HSA or various concentrations of NO-HSA for 24 h. DNA fragmentation was detected as described in the Materials and Methods.

### Figure 4. Effect of NO-HSA on C26 cell viability

(A) C26 cells were treated for 48 h with various concentrations of HSA (*open squares*) or NO-HSA (*closed circles*). (B) C26 cells were incubated for the indicated times with 100  $\mu$ M HSA (*open squares*) or 100  $\mu$ M NO-HSA (*closed circles*). Cell viability was determined as described in the Methods. Results are the mean  $\pm$  S.D. of three separate experiments.

### Figure 5. Effect of NO-HSA on tumor growth in C26 tumor-bearing mice

C26 tumor-bearing mice were given daily intravenous injections of saline (5 ml/kg), HSA (10  $\mu$ mol/5 ml/kg) or NO-HSA (10  $\mu$ mol/5 ml/kg) for ten days from day 3 to day 12 after inoculation with tumor cells. Tumor size was measured and tumor volume was calculated according to the formula:  $V = 0.4a^2b$ , where  $a$  is the smallest, and  $b$  is the largest, superficial diameter. Results are means  $\pm$  S.D;  $n = 10$  animals per

experimental group. \*Statistically significant reduction compared with saline ( $P<0.01$ ) or HSA ( $P<0.05$ ) at the corresponding time. \*\*Statistically significant reduction compared with saline ( $P<0.01$ ) or HSA ( $P<0.01$ ) at the corresponding time.

**Figure 6. Immunohistochemical staining of tumor tissues of C26 tumor-bearing mice intravenously injected with NO-HSA using the TUNEL method.**

C26 tumor-bearing mice were given daily intravenous injections of saline (5 ml/kg) (A), HSA (10  $\mu$ mol/5 ml/kg) (B) or NO-HSA (10  $\mu$ mol/5 ml/kg) (C) for five days from day 3 to day 7 after inoculation with tumor cells. TUNEL-staining, performed as described in the Materials and Methods, shows apoptotic cells in the tumor tissue of mice treated with NO-HSA.

**Table**

**Table 1. Serum biochemical parameters of C26 tumor-bearing mice treated with saline, HSA or NO-HSA**

C26 tumor-bearing mice were intravenously injected five times per day with saline (5 ml/kg), HSA (10  $\mu$ mol/5 ml/kg) or NO-HSA (10  $\mu$ mol/5 ml/kg) on days from 3 to 7 after inoculation with tumor cells. Blood samples were collected from the abdominal vena cava under anesthesia with diethylether approximately 2 h after the last treatment injection and mice were sacrificed. Serum biochemical parameters were measured using routine clinical laboratory techniques.

	Saline	HSA	NO-HSA
Total protein (g/dl)	5.23 $\pm$ 0.21	5.33 $\pm$ 0.06	5.55 $\pm$ 0.24
Cr (mg/dl)	0.13 $\pm$ 0.04	0.14 $\pm$ 0.01	0.16 $\pm$ 0.01
BUN (mg/dl)	15.25 $\pm$ 0.96	15.33 $\pm$ 1.53	13.75 $\pm$ 1.26
AST (U/l)	143.5 $\pm$ 62.9	91.7 $\pm$ 39.4	116.3 $\pm$ 69.1
ALT (U/l)	169.8 $\pm$ 111.6	110.0 $\pm$ 81.2	161.3 $\pm$ 126.2
ALP (U/l)	385.3 $\pm$ 18.3	345.3 $\pm$ 4.7 <sup>a</sup>	300.5 $\pm$ 23.0 <sup>b,c</sup>

a; p<0.05, Saline vs HSA, b; p<0.01, Saline vs NO-HSA, c; p<0.05, HSA vs NO-HSA















