Therapeutic effect of lecithinized superoxide dismutase (PC-SOD) against colitis

TOMOAKI ISHIHARA, KEN-ICHIRO TANAKA, YUICHI TASAKA, TAKUSHI NAMBA, JUN SUZUKI, TSUTOMU ISHIHARA, SUSUMU OKAMOTO, TOSHIFUMI HIBI, MITSUKO TAKENAGA, RIE IGARASHI, KEIZO SATO, YUTAKA MIZUSHIMA and TOHRU MIZUSHIMA

Graduate School of Medical and Pharmaceutical Sciences, Kumamoto University, Kumamoto 862-0973, Japan (T.I., K-I.T., Y.T., T.N., J.S., T.I., K.S., T.M)

DDS Institute, The Jikei University School of Medicine, Tokyo 105-8461, Japan (T.I., Y.M)

Department of Internal Medicine, Keio University School of Medicine, Tokyo 160-8582, Japan (S.O., T.H)

Division of Drug Delivery System, Institute of Medical Science, St. Marianna University, Kawasaki 216-8512, Japan (M.T., R.I)
Running title: PC-SOD and DSS-induced colitis

Address correspondence to: Dr. Tohru Mizushima, Graduate School of Medical and Pharmaceutical Sciences, Kumamoto University, 5-1 Oe-honmachi, Kumamoto 862-0973, Japan. TEL & FAX: 81-96-371-4323, E-mail: mizu@gpo.kumamoto-u.ac.jp

The number of text pages: 52

The number of tables: 4

The number of figures: 7

The number of references: 39

The number of words in the abstract: 245

The number of words in the introduction: 745

The number of words in the discussion: 1196

Abbreviations: ANOVA, analysis of variance; 5-ASA, 5-aminosalicylic acid; CL, chemiluminescence; DAI, disease activity index; DMPO, 5,5-dimethyl-1-pyrroline-N-oxide; DSS, dextran sulfate sodium; DTPA, Diethylenetriamine-N, N, N’, N’’, N’’’-pentaactic acid; ELISA, enzyme-linked immunosorbent assay; ESR, electron spin resonance; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; IBD, inflammatory bowel disease; MPO, myeloperoxidase; NF-κB, nuclear factor kappa B; RT, room temperature; PC, phosphatidylcholine; PC-SOD, lecithinized superoxide dismutase; PMA, phorbol 12-myristate 13-acetate; POBN, α-(4-pyridyl-1-oxide)-N-tert-butyl nitron; QOL, quality of life; ROS, reactive oxygen species; S.E.M., standard error of the mean; SOD, superoxide dismutase; TNF, tumor necrosis factor; UC, ulcerative colitis; U-SOD, unmodified superoxide dismutase.
Abstract

Ulcerative colitis (UC) involves intestinal mucosal damage induced by reactive oxygen species (ROS), in particular superoxide anion. Superoxide dismutase (SOD) catalyses dismutation of superoxide anion to hydrogen peroxide, which is subsequently detoxified by catalase. Lecithinized SOD (PC-SOD) is a new modified form of SOD which has overcome previous clinical limitations of SOD. In this study, we examined the action of PC-SOD using an animal model of UC, dextran sulfate sodium (DSS)-induced colitis. DSS-induced colitis was ameliorated by daily intravenous administration of PC-SOD. Unmodified SOD (U-SOD) produced a similar effect, but only at more than 30 times the concentration of PC-SOD. In vivo electron spin resonance analysis confirmed that the increase in the colonic level of ROS associated with development of colitis was suppressed by PC-SOD administration. The dose-response profile of PC-SOD was bell-shaped, but simultaneous administration of catalase restored the ameliorative effect at high doses of PC-SOD. Accumulation of hydrogen peroxide was observed with the administration of high doses of PC-SOD, an effect which was suppressed by the simultaneous administration of catalase. We also found that either a weekly intravenous administration or daily oral administration of PC-SOD conferred
protection. These results suggest that PC-SOD achieves its ameliorative effect against colitis through decreasing the colonic level of ROS, and that its ineffectiveness at higher doses is due to the accumulation of hydrogen peroxide. Furthermore, we consider that intermittent or oral administration of PC-SOD can be applied clinically to improve the quality of life of UC patients.
Introduction

Inflammatory bowel disease (IBD), Crohn’s disease and ulcerative colitis (UC), have become substantial health problems (Cuzzocrea, 2003). Recent studies suggest that IBD is a chronic inflammatory disorder occurs in the intestine due to “a vicious cycle”: infiltration of leukocytes into intestinal tissues causes mucosal damage induced by reactive oxygen species (ROS) that are released from the activated leukocytes, and this damage further stimulates the infiltration of leukocytes through induction of pro-inflammatory cytokines, in particular tumor necrosis factor (TNF)-α (Podolsky, 2002). Among the various ROS, superoxide anion is particularly important, as it has a potent ability to damage cells and leads to the formation of other ROS, such as hydroxy radicals (Kruidenier and Verspaget, 2002). A positive correlation between the severity of IBD and the intestinal level of ROS has been reported (Simmonds et al., 1992). Thus, anti-oxidant molecules (radical scavengers) have attracted considerable attention as therapeutic candidates for the treatment of IBD.

Superoxide dismutase (SOD) is one such anti-oxidant protein. SOD catalyses the dismutation of superoxide anion to hydrogen peroxide, which is subsequently detoxified.
to oxygen and water by catalase or glutathione peroxidase (Kruidenier and Verspaget, 2002). Among three isoforms of human SOD, Cu/Zn-SOD mainly contributes to the SOD activity in IBD patients (Kruidenier et al., 2003a). Decreased expression of SOD, especially Cu/Zn-SOD, has been observed in IBD patients (Kruidenier et al., 2003a; Kruidenier et al., 2003b). Furthermore, administration of Cu/Zn-SOD suppresses the development of IBD-related colitis in the experimental animal models (Keshavarzian et al., 1990; Segui et al., 2004). These findings raised the prospect that SOD could be of therapeutic benefit in the treatment of IBD. However, subsequent clinical trials of Cu/Zn-SOD have proven unsuccessful, mostly due to its low affinity to the cell membrane where superoxide anion is produced, and its low stability in plasma, with a half-life of only a few minutes (Greenwald, 1990; Tsao et al., 1991; Igarashi et al., 1992; Igarashi et al., 1994). Therefore, various drug delivery systems have been applied to SOD to overcome these limitations (Keshavarzian et al., 1990; Igarashi et al., 1992; Igarashi et al., 1994; Yasui and Baba, 2006).

Among these applications, lecithinized SOD (PC-SOD) is potentially beneficial for clinical treatment of IBD, especially UC. PC-SOD is lecithinized human Cu/Zn-SOD
in which four phosphatidylcholine (PC) derivative molecules are covalently bound to each SOD dimer (Igarashi et al., 1992). *In vitro* experiments using cultured cells have shown that this modification drastically improves the cell membrane affinity of SOD without decreasing its SOD activity (Igarashi et al., 1992; Igarashi et al., 1994), while *in vivo* experiments using rats have demonstrated that it also greatly improves plasma stability (Igarashi et al., 1992). In phase I clinical studies, intravenously administered PC-SOD (40 - 160 mg) had a terminal half-life of more than 24 h, with good safety and tolerability (Broeyer et al., 2008; Suzuki et al., 2008a). Furthermore, intravenously administered PC-SOD ameliorated dextran sulfate sodium (DSS)-induced colitis in rats, an IBD-related colitis animal model (Hori et al., 1997), suggesting that PC-SOD is effective for the treatment of IBD patients. In fact, recent published results of phase II clinical study have shown that intravenously administered PC-SOD (40 or 80 mg) significantly improved the disease activity index (DAI) scores of UC patients (Suzuki et al., 2008b). However, the comparison of PC-SOD with unmodified SOD (U-SOD) based on pharmacological activity against colitis has not been undertaken and a decrease in the ROS level with PC-SOD administration has not been demonstrated *in vivo*. As well as
U-SOD, a bell-shaped dose-response profile of PC-SOD has been reported for various pharmacological activities, including anti-colitis activity (Mao et al., 1993; Hori et al., 1997; Tamagawa et al., 2000; Tsubokawa et al., 2007). However, its underlying mechanism has remained unknown. Furthermore, when considering the quality of life (QOL) of patients, the present clinical protocol of PC-SOD administration (intravenous infusion once daily for 4 weeks) is expected to be improved. In this study, we compared PC-SOD and U-SOD for their pharmacological activity against DSS-induced colitis and found that PC-SOD has more than 30 times higher activity. In vivo electron spin resonance (ESR) analysis showed that administration of PC-SOD suppressed the increase in the ROS level induced by DSS treatment. We also provide evidence that the ineffectiveness of higher doses of PC-SOD is due to accumulation of hydrogen peroxide at the intestine. Furthermore, based on results obtained here, we propose that intermittent administration or oral administration of PC-SOD is a clinically viable option to improve the QOL of UC patients.
Methods

Chemicals and Animals

Paraformaldehyde, o-dianisidine, phorbol 12-myristate 13-acetate (PMA), fetal bovine serum (FBS) and catalase from bovine liver (1340 U/mg) were obtained from Sigma (St. Louis, MO). RPMI 1640 was from Nissui Pharmaceutical (Tokyo, Japan). Enzymatic digest of animal tissue (Proteose peptone) was from BD Biosciences (San Jose, CA). LPS was from List Biological Laboratories (Campbell, CA). Alexa Fluor 488 goat anti-rabbit immunoglobulin G was purchased from Invitrogen (Carlsbad, CA). Mounting medium for immunohistochemical analysis (VECTASHIELD) was from Vector Laboratories (Burlingame, CA). 5,5-dimethyl-1-pyrroline-N-oxide (DMPO) was purchased from LABOTEC LTD (Tokyo, Japan), lymphocyte isolation sterile solution (Ficoll-Paque PLUS) from GE Healthcare (Buckinghamshire, UK), DSS (M.W. 5000, 15–20% sulfur content) and luminol from WAKO Pure Chemicals (Tokyo, Japan), and Mayer’s hematoxylin, 1% eosin alcohol solution and mounting medium for histological examination (Malinol) from MUTO Pure Chemicals (Tokyo, Japan). The RNeasy kit
was obtained from Qiagen (Valencia, CA), the PrimeScript® 1st strand cDNA Synthesis Kit was purchased from TAKARA Bio (Ohtsu, Japan), and mix for real time RT-PCR (iQ SYBR Green Supermix) was from Bio-Rad Laboratories (Hercules, CA). α-(4-pyridyl-1-oxide)-N-tert-butynitrone (POBN) was from Alexis (San Diego, CA). U-SOD (5190 U/mg) and PC-SOD (3000 U/mg) were from our laboratory stock (Igarashi et al., 1992). SODs were dissolved in 5% xylitol and administered intravenously (tail vein) or orally. Diethylenetriamine-N, N, N’, N”’, N”’-pentaactic acid (DTPA) and 4’, 6-diamidino-2-phenylindole, dihydrochloride (DAPI) were from Dojindo (Kumamoto, Japan). An antibody against phospho-nuclear factor kappa B (NF-κB) p65 (Ser536) was from Cell Signaling Technology (Danvers, MA). Wild-type mice (8-week-old, ICR, male) were used throughout. The experiments and procedures described here were carried out in accordance with the Guide for the Care and Use of Laboratory Animals as adopted and promulgated by the National Institute of Health, and were approved by the Animal Care Committee of Kumamoto University.

**Development of DSS-induced Colitis and Measurement of Colon Length and DAI**
DSS-induced colitis was induced in mice by the addition of 4% DSS (w/v, final concentration) to their drinking water as described previously (Tanaka et al., 2007). The first administration of PC-SOD was done just before the start of DSS administration. The animals were allowed free access to the DSS-containing water for 7 days. For measurement of myeloperoxidase (MPO) activity, expression of mRNAs and the ROS level, we used rectum and distal colon tissue.

After 7 days, animals were placed under deep ether anesthesia and sacrificed, the colons were dissected and their length measured from the ileocecal junction to the anal verge.

The DAI was determined macroscopically by an observer unaware of the treatment the mice had received, according to previously reported criteria (Tanaka et al., 2007). Briefly, the DAI was calculated as the sum of the diarrheal stool score (0: normal stool; 1: mildly soft stool; 2: very soft stool; 3: watery stool) and the bloody stool score (0: normal colored stool; 1: brown stool; 2: reddish stool; 3: bloody stool).
MPO Activity

MPO activity in the colonic tissues was measured as previously described (Tanaka et al., 2007). After DSS treatment, colons were dissected, rinsed with cold saline and cut into small pieces. Samples were homogenized and protein concentrations of the samples were determined using the Bradford method. MPO activity was determined in 10 mM phosphate buffer with 0.5 mM o-dianisidine, 0.00005% (w/v) hydrogen peroxide and 20 µg protein. MPO activity was obtained from the slope of the reaction curve and specific activity was expressed as the number of hydrogen peroxide molecules converted per min per mg protein.

Real-time RT-PCR Analysis

Real-time RT-PCR was performed as previously described (Mima et al., 2005) with some modifications. Total RNA was extracted from intestinal tissues or mouse peritoneal macrophages using an RNeasy kit according to the manufacturer’s protocol. Samples (2.5 µg RNA) were reverse-transcribed using a first-strand cDNA synthesis kit. Synthesized cDNA was used in real-time RT-PCR (Chromo 4 instrument (Bio-Rad...
Laboratories, Hercules, CA) experiments using mix for real time RT-PCR, and analyzed with Opticon Monitor Software. Specificity was confirmed by electrophoretic analysis of the reaction products and by inclusion of template- or reverse transcriptase-free controls.

To normalize the amount of total RNA present in each reaction, glyceraldehyde-3-phosphate dehydrogenase (GAPDH) cDNA was used as an internal standard.

Primers were designed using the Primer3 website (http://frodo.wi.mit.edu/cgi-bin/primer3/primer3_www.cgi). The primers used were (name: forward primer, reverse primer): 

- **Tnf-α**: 5’-cgtcagccgatttgctatct-3’, 5’-cggactccgcaaagtctaag-3’;
- **Gapdh**: 5’-aactttggcattgtggaagg-3’, 5’-acacattgagggaaca-3’;
- **Il-1β**: 5’-gatcccaagcaatacccaaa-3’, 5’-ggggaactctgcagactcaa-3’;
- **Il-6**: 5’-ctggagtcacagaaggagtgg-3’, 5’-ggagtcacagaagtgcag-3’;
- **Il-23p19**: 5’-gccccgtatccagtgaag-3’, 5’-ggtaagtcacagaagttt-3’;
- **Il-23p19**: 5’-gccccgtatccagtgaag-3’, 5’-ggtaagtcacagaagttt-3’.

**Histological and Immunohistochemical Analysis**
Colonic tissue samples were fixed in 4% buffered paraformaldehyde, then embedded in paraffin before being cut into 4 µm sections.

For histological examination, sections were stained first with Mayer’s hematoxylin and then with 1% eosin alcohol solution. Samples were mounted with mounting medium and inspected with the aid of an Olympus BX51 microscope.

For immunohistochemical analysis, sections were blocked with 3% BSA for 30 min, incubated for 12 h with antibody against phospho-NF-κB (1:100 dilution) in the presence of 2.5% BSA, and finally incubated for 1 h with Alexa Fluor 488 goat anti-mouse immunoglobulin G in the presence of DAPI (5 µg/ml). Samples were mounted with mounting medium and inspected using fluorescence microscopy (Olympus BX51).

**Measurement of ROS in Neutrophils in Vitro**

Human neutrophils were prepared as previously described (Karakawa et al., 2008). Briefly, polymorphonuclear leukocytes and mononuclear cells were separated
using a gradient of lymphocyte isolation sterile solution. Red blood cells remaining in the polymorphonuclear leukocyte fractions were lysed with 0.2% NaCl.

The chemiluminescence (CL) response induced by the superoxide anion released from neutrophils was measured as described (Muranaka et al., 1997). Prepared neutrophils were mixed with 25 ng/ml PMA in RPMI medium containing 10 µM luminol and 500 µM DTPA. The CL response was continuously recorded for 10 min at room temperature (RT) using a luminometer (Advantec Co., Tokyo, Japan).

The level of superoxide anion was also assayed by ESR spin trapping with DMPO as previously described (Karakawa et al., 2008). Prepared neutrophils were incubated with 10 ng/ml PMA in RPMI medium containing 500 µM DTPA and 25 mM DMPO for 5 min at RT. ESR spectra were recorded at RT on a JES-TE200 ESR spectrometer (JEOL, Tokyo, Japan) under the following conditions: modulation frequency, 100 kHz; microwave frequency, 9.43 GHz; microwave power, 40 mW; scanning field, 335.2 ± 5 mT; sweep time, 2 min; field modulation width, 0.25 mT; receiver gain, 100; time count, 0.3 sec. After recording the ESR spectra, the signal
intensities of the DMPO–OOH adducts were normalized against that of a manganese oxide marker.

**Determination of ROS Level and the Amount of Hydrogen Peroxide in Vivo**

*In vivo* ESR analysis was performed as previously described (Sato et al., 1992; Sato et al., 2002), with some modifications. After DSS administration for 7 days, animals were placed under deep anesthesia with chloral hydrate (250 mg/kg) and injected with POBN intraperitoneally (4 mmol/kg). After 1 h, mice were sacrificed, the colons were dissected and the lipid phase from the samples was extracted as described elsewhere (Sato et al., 1992; Sato et al., 2002). After evaporating the sample, ESR spectra were immediately recorded at RT in a JES-TE200 spectrometer under the following conditions: modulation frequency, 100 kHz; microwave frequency, 9.43 GHz; microwave power, 40 mW; scanning field, 335.2 ± 5 mT; sweep time, 2 min; field modulation width, 0.25 mT; receiver gain, 630; time count, 0.3 sec. Every buffer and solutions of the reaction mixture used for ESR measurement was treated with chelex100 resin (Bio-rad Laboratories, Hercules, CA) before use to remove metals.
For determination of hydrogen peroxide levels, colons were dissected, cut into small pieces, suspended in PBS and incubated for 30 min at RT with rotation. After centrifugation, the supernatants were applied to the NWLSS™ NWK-HYP01 assay kit (Northwest Life Science Specialties, Vancouver, WA).

**Determination of the Amount of PC-SOD and TNF-α in Vivo.**

Determination of the amount of PC-SOD was carried out as previously described (Igarashi et al., 1992). After administration of PC-SOD, the blood was collected and serum samples were obtained by centrifugation. On the other hand, colons were dissected, cut into small pieces, homogenized and centrifuged to obtain the supernatants. Samples were analyzed using a human Cu/ZnSOD enzyme-linked immunosorbent assay (ELISA) kit (Bender MedSystem, Burlingame, CA). We used PC-SOD (or U-SOD) for drawing the standard curve of ELISA and determined the amount of PC-SOD (or U-SOD). The amount of TNF-α in serum was determined similarly by use of its ELISA kit from Pierce (Rockford, IL).
Preparation of Mouse Peritoneal Macrophages

Mouse peritoneal macrophages were prepared as described previously (Salimuddin et al., 1999). Mice were given 2 ml of 10% Enzymatic digest of animal tissue by intraperitoneal injection and peritoneal cells were harvested 3 days later. The cells were seeded in 60 mm culture dishes at 4 X 10^6 cells/dish in RPMI 1640 medium supplemented with 10% heat-inactivated FBS. After incubation for 4 h, non-adherent cells were removed and the adherent cells were cultured for use in the experiments. Virtually all of the adherent cells were macrophages, as previously described (Salimuddin et al., 1999).

Statistical Analysis

All values are expressed as the mean ± standard error of the mean (S.E.M.). Two-way analysis of variance (ANOVA) followed by the Tukey test or the Student's \( t \)-test for unpaired results were used to evaluate differences between more than three groups or between two groups, respectively. Differences were considered to be significant for values of \( P < 0.05 \).
Results

A Comparison of the Effect of PC-SOD and U-SOD on DSS-induced Colitis

The severity of DSS-induced colitis can be monitored by various indices, such as DAI, length of colon, MPO activity, and histological analysis. We compared PC-SOD and U-SOD for their effect on the development of colitis induced by 4% DSS administration. The clinical study was performed with 40 and 80 mg PC-SOD (Suzuki et al., 2008b), which corresponds to 2 and 4 kU/kg and therefore, we chose the dose of 3 kU/kg for the following experiments. PC-SOD and U-SOD were intravenously administered once daily. There was no significant difference in the volume of water consumed by each group of mice (data not shown). Administration of 4% DSS increased the DAI and this increase was significantly suppressed by the administration of PC-SOD (3 kU/kg) but not U-SOD (3 kU/kg) (Fig. 1A). DSS-induced colon shortening, used as a morphometric measure for the degree of inflammation, was significantly ameliorated in the PC-SOD-treated animals (Fig. 1B), as was colonic MPO activity, an indicator of
leukocyte infiltration (Fig. 1C). Figure 1D shows the results of histological analyses of colonic tissues. Crypt loss and infiltration of leukocytes were observed in DSS-treated mice and these phenotypes were improved by administration of PC-SOD and, to a lesser extent, U-SOD (Fig. 1D). Taken together, these findings demonstrate that PC-SOD is more effective than U-SOD for the amelioration of DSS-induced colitis.

To compare the specific activity of PC-SOD and U-SOD, we determined their dose-response profiles. As shown in Fig. 2A, PC-SOD produced the maximum beneficial effect at 1.5 - 3 kU/kg, whereas higher doses (6 – 12 kU/kg) had no significant effect on DAI. A similar bell-shaped profile has also been reported in a rat model of DSS-induced colitis (Hori et al., 1997). In the case of colon shortening and colonic MPO activation, the maximal effect was again observed in response to 1.5 - 3 kU/kg PC-SOD (Fig. 2B and C). In contrast, U-SOD at the much higher concentration of 48 kU/kg only ameliorated DSS-induced colitis to a similar extent to that obtained with 1.5 kU/kg PC-SOD (Fig. 2D-F), indicating that the specific activity of PC-SOD is more than 30 times that of U-SOD. We next used an ELISA assay to compare the level of PC-SOD and U-SOD in serum and colonic tissues after their intravenous administration. As shown in Table 1,
after daily intravenous administration for 7 days, and 6 h after the final injection, PC-SOD was detected in serum and colonic tissues at a concentration of 3.3 µg/ml and 0.23 ng/mg, respectively. However, U-SOD was not detected in either preparation, suggesting that the enhanced activity of PC-SOD is partly due to its greater stability \textit{in vivo}.

We also examined the effect of DSS-treatment on the level of PC-SOD after intravenous administration. As shown in Table 2, DSS-treatment significantly increased the level of PC-SOD in colonic tissues, but not so clearly in serum, suggesting that PC-SOD can be enriched in damaged or inflamed tissues, which is an advantage for its clinical application for UC.

\textit{Involvement of ROS in Amelioration of DSS-induced Colitis by PC-SOD}

Previous studies have suggested that PC-SOD is more potent than U-SOD for decreasing the superoxide anion released from activated neutrophils \textit{in vitro} by showing that PC-SOD showed higher activity than U-SOD for the suppression of endothelial cell damage induced by activated neutrophils and that more PC-SOD than U-SOD remains on
the surface of neutrophils after washing (Igarashi et al., 1994). In this study, we investigated this directly by measuring the superoxide anion by ESR. As shown in Fig. 3A, a radical spin adduct of the ESR spectrum corresponding to superoxide anion (DMPO–OOH adduct) was observed for PMA-activated human neutrophils. Pre-incubation of neutrophils with PC-SOD lowered the peak of DMPO–OOH adduct in a dose-dependent manner, showing that PC-SOD decreases the level of superoxide anion. U-SOD produced a similar, but less pronounced, effect (Fig. 3A and B). We also quantitatively measured the activity of PC-SOD and U-SOD based on CL analysis. An increase in CL, in other words, the amount of superoxide anion released from activated neutrophils, decreased in the presence of PC-SOD and U-SOD. Again, PC-SOD was more potent than U-SOD (Fig. 3C and D), perhaps due to its higher cell membrane affinity, as previously reported (Igarashi et al., 1994).

Although it is generally believed that administration of PC-SOD decreases the level of ROS in vivo, no direct evidence in support of this idea has been reported. In this study, we examined the effect of PC-SOD administration on the intestinal level of ROS measuring the lipid-derived free radical spin adduct with ESR spectroscopy and spin trap
POBN, which reacts with ROS to form a radical spin adduct. We have recently reported that this method, in vivo free radical production and ex vivo detection, is effective for monitoring ROS level in the intestine (Namba et al., submitted). The hyperfine coupling constants for the POBN radical adducts were $g^N = 14.92 \pm 0.06$ G and $g^H = 2.44 \pm 0.05$ G, which are similar to previous data (Namba et al., submitted). Therefore, our detected radical was determined as a lipid-derived free radical. As shown in Fig. 4A and B, the intestinal level of ROS (the height of the ESR peak shown by the bar) was increased by DSS administration, an effect that was suppressed by the administration of PC-SOD. This is the first direct evidence that PC-SOD decreases the level of ROS in vivo.

We also examined the effect of PC-SOD on the mRNA expression of various cytokine (TNF-α, IL-1β, IL-6 and IL-23) in the intestine by real-time RT-PCR analysis. The mRNA expression of Tnf-α and Il-1β was up-regulated by DSS treatment and this up-regulation was suppressed by administration of PC-SOD (3 kU/kg) (Figs. 4C and 5E). On the other hand, administration of neither DSS nor PC-SOD affected the mRNA expression of Il-6 and Il-23p19 (Fig. 4C). The effect of PC-SOD on mRNA expression of Tnf-α and Il-1β was also examined in vitro. Treatment of peritoneal macrophages
prepared from wild-type mice with LPS induced the mRNA expression of \textit{Tnf-}\(\alpha\) and \textit{Il-1}\(\beta\) and this induction was partially suppressed by simultaneous treatment of cells with PC-SOD (Fig. 4D). These results suggest that PC-SOD suppresses the expression of TNF-\(\alpha\) and IL-1\(\beta\), by lowering the intestinal level of ROS, resulting in the amelioration of DSS-induced colitis.

As described in the introduction, a bell-shaped dose-response profile has been observed for various pharmacological activities of PC-SOD (Hori et al., 1997; Tamagawa et al., 2000; Tsubokawa et al., 2007). One possible explanation for the ineffectiveness of high doses of PC-SOD is the accumulation of hydrogen peroxide due to the relatively higher activity of SOD compared with catalase (Mao et al., 1993), however, this idea has not been proved. In this study, we tested this idea by examining the effect of simultaneous administration of catalase; if the ineffectiveness of higher doses of PC-SOD is due to the accumulation of hydrogen peroxide, the effect would be restored by simultaneous administration of catalase that detoxifies hydrogen peroxide into oxygen and water. Administration of a high dose of PC-SOD (6 kU/kg) improved the DAI score in DSS-treated mice in the presence of simultaneous intravenous administration of
catalase but not in its absence (Fig. 5A). Administration of catalase alone tended to improve the DAI, but this effect was not statistically significant (Fig. 5A). Similar results were observed for DSS-induced colon shortening and colonic MPO activation (Fig. 5B and C). These findings suggest that the ineffectiveness of high doses of PC-SOD on DSS-induced colitis is caused by accumulation of hydrogen peroxide.

We further tested this idea by direct measurement of the colonic level of hydrogen peroxide. As shown in Fig. 5D, DSS treatment increased the colonic level of hydrogen peroxide and administration of a low dose (3 kU/kg) of PC-SOD suppressed this effect. However, in the case of a high dose of PC-SOD (12 kU/kg), a significant effect was only observed in the presence of simultaneous administration of catalase (Fig. 5D), which alone did not significantly suppress the increase (Fig. 5D). Similar, but less pronounced, results were obtained with 6 kU/kg PC-SOD as compared with a 12 kU/kg dose (data not shown).

We also examined the effect of catalase on the mRNA expression of Tnf-α in the presence of high dose of PC-SOD. The up-regulated mRNA expression of Tnf-α by DSS treatment was not suppressed by administration of a high (6 kU/kg) dose of...
PC-SOD (Fig. 5E). However, simultaneous administration of catalase with the high dose of PC-SOD significantly suppressed the DSS-induced mRNA expression of Tnf-α (Fig. 5E). We also determined the serum level of TNF-α and found that the level was increased by DSS-treatment and this increase was suppressed by administration of a low (3 kU/kg) dose of PC-SOD and a high (6 kU/kg) dose of PC-SOD with catalase (Fig. 5F). Being different from the case of mRNA expression in the intestine (Fig. 5E), the serum level of TNF-α was decreased by administration of catalase alone. These results suggest that TNF-α plays an important role in the ameliorative effect of PC-SOD against DSS-induced colitis.

Immunohistochemical analysis with antibody against phosphorylated form of NF-κB p65 at Ser536 (active form of NF-κB p65) demonstrated that DSS-administration increases the level of active NF-κB in the colonic mucosa (Fig. 5G), suggesting that the inflammatory response occurs in epithelial cells and infiltrated leukocytes. Again, this increase in the level of active NF-κB expression was suppressed by administration of a low (3 kU/kg) dose of PC-SOD and a high (6 kU/kg) dose of PC-SOD with catalase.
Effect of Modified Methods of PC-SOD Administration

To obtain some useful clues for refining the clinical guidelines for administration of PC-SOD, we tested the outcome of other protocols and routes of administration in the treatment of DSS-induced colitis. As illustrated in Fig. 6A, we first intravenously administered PC-SOD once at the start of DSS treatment (day 0), then monitored the DAI for 7 days. Although the dose found to be effective with daily administration (3 kU/kg) did not improve the DAI score, a higher dose (6 – 24 kU/kg) produced a significant improvement (Fig. 6A). However, higher doses of 48 – 96 kU/kg worsened the DAI during the early stage of colitis development (Fig. 6A). The effectiveness of a one-shot administration of PC-SOD (12 – 24 kU/kg) was also shown by measuring colon shortening and colonic MPO activation (Fig. 6B and C). The findings suggest that intermittent (for example once weekly) administration of PC-SOD, a regime that greatly improves the QOL of UC patients, is a viable clinical protocol.-When we monitored the level of PC-SOD after this single-dose administration (12 kU/kg) we found that it dropped below detectable limits 72 h (in serum) or 24 h (in colonic tissues) after the injection (Table 3).
We also examined the effect of oral administration (once daily) of PC-SOD. As shown in Fig. 7A, significant improvement in the DAI score was observed at most of the doses tested. The ameliorative effect of oral administration of PC-SOD was also observed in terms of colon shortening and colonic MPO activation (Fig. 7B and C). This suggests that oral administration of PC-SOD, a regime that greatly improves the QOL of UC patients, is a viable clinical protocol, is also clinically viable. We found that the level of PC-SOD in serum did not increase at any time-points (6 – 48 h) after the oral administration of PC-SOD (48 kU/kg) (Table 4), suggesting that orally administered PC-SOD is not absorbed and reaches the intestinal mucosa directly. By employing an ELISA assay, we detected the PC-SOD in the colonic tissues 24 h after its oral administration (48 kU/kg) (Table 4). However, low doses (0.75 – 1.5 kU/kg) proved undetectable (data not shown).
Discussion

The efficacy of PC-SOD for the treatment of UC patients has already been demonstrated by a phase II clinical study (Suzuki et al., 2008b). However, the mechanism of its action is not fully understood. Given that determining the underlying mechanism is important to advance the further development of this drug, in the present study, we examined the action of PC-SOD in an animal model of UC, DSS-induced colitis. Furthermore, as the current clinical protocol for the administration of PC-SOD (once daily intravenous infusion for 4 weeks) does not provide patients with good QOL, we also tested other dosing regimes in our animal model.

The superior character of PC-SOD to U-SOD has been shown both in vitro (high cell membrane affinity) and in vivo (high stability in plasma). However, a direct comparison of their pharmacological activity has not been reported. In this study, we have demonstrated that the ameliorative effect of PC-SOD against DSS-induced colitis is more than 30 times higher than that of U-SOD. The higher stability in serum (Table 1) and higher activity for decreasing superoxide anion (Fig. 3) of PC-SOD could contribute
to this effect. Analysis of intestinal ROS level in vivo is difficult, and thus the decrease in the level of ROS by PC-SOD has not been directly shown. In this study, we have demonstrated this by use of radical spin adduct ESR spectrum analysis. This analysis should also be useful for detecting the alteration in the intestinal level of ROS associated with various other diseases, and for evaluation of drugs for the treatment of such conditions.

The bell-shaped dose-response profile of PC-SOD is of clinical concern, as this may reflect side-effects of the drug. In this study, we have revealed that the efficacy of higher doses of PC-SOD is restored by simultaneous administration of catalase that converts hydrogen peroxide to water and oxygen. Furthermore, we have directly determined the colonic level of hydrogen peroxide and found that low doses of PC-SOD suppress the DSS-induced increase in the intestinal level of hydrogen peroxide, and that simultaneous administration of catalase with high doses of PC-SOD but not the PC-SOD alone produces a similar effect. These results suggest that the ineffectiveness of high doses of PC-SOD on DSS-induced colitis is caused by accumulation of hydrogen peroxide. Although catalase is abundant, recent studies have suggested that its activity
fluctuates during the development of colitis (Kruidenier et al., 2003b; Kruidenier et al., 2003c; Mahgoub et al., 2003). This may affect the clinical efficacy of PC-SOD, and individual examination of catalase activity before the administration of the drug may result in safer and more effective treatment.

Hydrogen peroxide is not itself major cause of ROS-mediated cell damage, but it does react with Fe^{2+} to produce a potent hydroxy radical according to the Fenton reaction (Mao et al., 1993). Furthermore, it has been reported that among various ROS, hydrogen peroxide is the most potent activator of NF-κB, a strong inducer of inflammation through induction of pro-inflammatory cytokines, especially TNF-α (Schmidt et al., 1995; Marikovsky et al., 2003), and that NF-κB plays an important role in intestinal colitis (Schreiber et al., 1998; Herfarth et al., 2000). Here we have shown that activation of NF-κB, the mRNA expression of Tnf-α and serum level of TNF-α are induced when the colonic level of hydrogen peroxide increases, suggesting that hydrogen peroxide damages the intestinal mucosa both through induction of inflammation via activation of NF-κB and through direct cell damage mediated in conjunction with hydroxy radical formation.
In the present study, we have also demonstrated that administration of a single high-dose of PC-SOD at the start of DSS treatment significantly suppresses colitis. The serum or intestinal level of PC-SOD dropped below detectable limits 72 h or 24 h, respectively, after the injection (Table 3), suggesting that the existence of PC-SOD in the early stages of development of colitis is important for its ameliorative effect. The present clinical protocol for PC-SOD administration to UC patients enforces their hospitalization. Our results suggest that intermittent (for example once weekly) administration of PC-SOD, thereby allowing ambulatory administration of the drug, may be an effective and preferable treatment for UC patients.

We also found that oral administration of PC-SOD (0.75 – 48 kU/kg) significantly suppressed DSS-induced colitis. The serum level of PC-SOD did not increase after the oral administration, suggesting that orally administered PC-SOD is not absorbed but reaches the colonic mucosa directly. However, based on ELISA assay, we only detected PC-SOD in colonic tissues after administration of the highest dose tested (48 kU/kg), the tissue level being similar to that obtained after intravenous administration of a 3 kU/kg dose (Table 1). Thus, it seems that a very small amount of PC-SOD (under the limit of
detection) is effective when administered orally. The distribution of PC-SOD at the intestine may differ depending on the route of administration and this may contribute to the higher specific activity of PC-SOD following oral treatment. The fact that a bell-shaped dose-response profile was not observed upon oral administration of PC-SOD (Fig. 7) supports this idea. On the other hand, it is also possible that PC-SOD is modified at the gastric and intestinal mucosa in such a way that it is not recognized by the ELISA assay although its activity is maintained. If oral administration of PC-SOD is applied clinically, it should greatly improve the QOL of patients treated with the drug. We used human SOD, not mouse protein for examination of effect of oral administration of PC-SOD on DSS-induced colitis in mice. Human SOD may modify the mucosal immunological reaction in mice and this modification may stimulate the beneficial effect of PC-SOD against colitis in mice. In this case, oral administration of PC-SOD may not give beneficial effects for human.

Glucocorticoids, 5-aminosalicylic acid (5-ASA) and immunosuppressive drugs are currently used for the clinical treatment of IBD (Podolsky, 2002; Baumgart and Sandborn, 2007). Although some new types of drugs, such as infliximab, have been
developed recently, a number of clinical problems, such as side-effects, are yet to be addressed (Keane et al., 2001; Bongartz et al., 2006; Baumgart and Sandborn, 2007). Thus, IBD remains a disease for which the development of new types of drugs remains therapeutically important. Clinical studies have suggested that PC-SOD is a relatively safe drug. Furthermore, it has a mechanism of action that is different from that of other currently used drugs as well as those that are under clinical development. Thus, PC-SOD remains a therapeutically beneficial option for UC patients either through a single administration or in combination with other types of drugs.

ROS play an important role in the progression of not only IBD but also various other diseases, such as focal cerebral ischemic injury, pulmonary fibrosis, chemotherapy-induced cardiotoxicity, and motor dysfunction after spinal cord injury. It is therefore worth noting in conclusion that a number of animal studies have suggested that PC-SOD is also effective in the treatment of these conditions (Tamagawa et al., 2000; Chikawa et al., 2001; Nakajima et al., 2001; den Hartog et al., 2004; Tsubokawa et al., 2007). Thus, the results obtained in this paper should provide useful evidence when
considering the most appropriate clinical protocol for therapeutic administration of PC-SOD against these diseases.
References


Footnotes

This work was supported by Grants-in-Aid for Scientific Research from the Ministry of Health, Labour, and Welfare of Japan, as well as Japan Science and Technology Agency and Grants-in-Aid for Scientific Research from the Ministry of Education, Culture, Sports, Science and Technology, Japan.
Legends for Figures

**Fig. 1.** Effect of PC-SOD and U-SOD on development of DSS-induced colitis. Mice treated with 4% DSS for 7 days, as described in the materials and methods, were intravenously administered PC-SOD or U-SOD once daily. DAI was measured daily (A). The length of the colon (B) and colonic MPO activity (C) were determined at the end of the experimental period. Sections of colonic tissue were also prepared and subjected to histological examination by hematoxylin and eosin staining (D). CTRL: control without DSS treatment. Values are mean ± S.E.M. * or # P<0.05; ** or ## P<0.01.

**Fig. 2.** Dose-response profile of effect of PC-SOD and U-SOD on development of DSS-induced colitis. Mice were treated with DSS and PC-SOD (A-C) or U-SOD (D-F) and colitis was assessed as described in the legend of Fig. 1. Values are mean ± S.E.M. *P<0.05; **P<0.01 (versus without PC-SOD or U-SOD).

**Fig. 3.** Effect of PC-SOD or U-SOD on the amount of superoxide anion *in vitro*. Human neutrophils were pre-incubated with PC-SOD or U-SOD for 1 h and washed with medium.
Neutrophils were then activated with PMA and the amount of superoxide anion was measured by ESR (A, B) or CL analysis (C, D). The intensity of the ESR signal of the superoxide anion adduct (DMPO–OOH adduct shown by the bar in (A)) was determined (B). The area under the graph (C) was also determined and is presented as a relative measure in (D). Values are given as the mean ± S.E.M. (n=3). #P<0.05; ** or ##P<0.01.

**Fig. 4.** Effect of PC-SOD on the level of ROS and expression of cytokines. DSS and PC-SOD were administered to mice, as described in the legend of Fig. 1 (A-C). After 7 days, POBN was administered and the colons were dissected and subjected to radical adduct ESR spectrum analysis (A). The intensity of the ESR signal (shown by the bar in (A)) was determined (B). Samples of colonic mRNA were subjected to real-time RT-PCR, using a specific primer set for *Il-1β, Il-6 and Il-23p19*. Values were normalized to *Gapdh*, expressed relative to the control sample (i.e. without DSS treatment) (C). Mouse peritoneal macrophages were preincubated with or without indicated concentration of PC-SOD for 1 h and further treated with LPS (1 µg/ml) in the presence of same concentration of PC-SOD as in the preincubation step for 3 h. The mRNA
fractions were prepared and analysed by real-time RT-PCR as described above (D).

Values are given as the mean ± S.E.M. (n=3) (B, C) or S.D. (D). * or #P<0.05; ** or ##P<0.01; n.s.: not significant; CTRL: control.

**Fig. 5.** The effect of simultaneous administration of catalase on the ameliorative effect of PC-SOD on DSS-induced colitis, colonic level of hydrogen peroxide, TNF-α and NF-κB. Mice were treated with DSS and PC-SOD (A-G) and colitis was assessed (A-C), as described in the legend of Fig. 1. Catalase was intravenously administered once daily (A-G). Colonic tissues were removed and the amount of hydrogen peroxide was determined (D). Samples of colonic mRNA were subjected to real-time RT-PCR as described in the legend of Fig. 4 (E). The serum levels of TNF-α were determined by ELISA (F). Sections of intestinal tissues were prepared and subjected to immunohistochemical analysis with an antibody against phospho-NF-κB p65 and DAPI staining as described in materials and methods (G). Values are mean ± S.E.M. *, # or †P<0.05; **, ## or ††P<0.01; n.s.: not significant.
**Fig. 6.** Effect of a single dose of PC-SOD on the development of DSS-induced colitis.

PC-SOD was intravenously administered to mice once at the start of DSS treatment (day 0). Colitis was subsequently assessed, as described in the legend of Fig. 1. Values are mean ± S.E.M. *P<0.05; **P<0.01.

**Fig. 7.** Effect of oral administration of PC-SOD on development of DSS-induced colitis.

PC-SOD was orally administered to mice once daily. Colitis was assessed, as described in the legend of Fig. 1. Values are mean ± S.E.M. *P<0.05; **P<0.01.
Table 1

Serum and colonic levels of PC-SOD and U-SOD.

Mice were intravenously administered indicated dose of PC-SOD or U-SOD once daily for 7 days. They were also treated with DSS, as described in the legend of Fig. 1. Blood and colonic tissues were taken 6 h after the final administration. The levels of PC-SOD or U-SOD in the samples were determined by ELISA. Values are mean ± S.E.M.

<table>
<thead>
<tr>
<th></th>
<th>DSS + PC-SOD 3kU/kg</th>
<th>DSS + U-SOD 3 kU/kg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum (µg/mL)</td>
<td>3.3 ± 0.42</td>
<td>&lt;0.063</td>
</tr>
<tr>
<td>Tissue (ng/mg)</td>
<td>0.23 ± 0.038</td>
<td>&lt;0.013</td>
</tr>
</tbody>
</table>
Table 2

Serum and colonic levels of PC-SOD.

Mice were intravenously administered indicated dose of PC-SOD once daily for 7 days. They were also treated with or without DSS, as described in the legend of Fig. 1. Blood and colonic tissues were taken 6 h after the final administration. The levels of PC-SOD in the samples were determined by ELISA. Values are mean ± S.E.M. *P<0.05.

<table>
<thead>
<tr>
<th></th>
<th>PC-SOD 3 kU/kg</th>
<th>DSS + PC-SOD 3 kU/kg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum (µg/mL)</td>
<td>3.5 ± 0.21</td>
<td>4.0 ± 0.61</td>
</tr>
<tr>
<td>Tissue (ng/mg)</td>
<td>0.066 ± 0.029</td>
<td>0.24 ± 0.043 *</td>
</tr>
</tbody>
</table>
Table 3

Serum and colonic levels of PC-SOD.

Mice were intravenously administered 12 kU/kg PC-SOD once on day 0 and treated with DSS for 3 days. Blood and colonic tissue samples were taken periodically. The levels of PC-SOD in the samples were determined by ELISA. Values are mean ± S.E.M.

<table>
<thead>
<tr>
<th></th>
<th>15 min</th>
<th>24 h</th>
<th>48 h</th>
<th>72 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum (µg/mL)</td>
<td>170 ± 5.7</td>
<td>2.3 ± 0.67</td>
<td>0.41 ± 0.12</td>
<td>&lt;0.16</td>
</tr>
<tr>
<td>Tissue (ng/mg)</td>
<td>1.6 ± 0.69</td>
<td>&lt;0.031</td>
<td>&lt;0.031</td>
<td>&lt;0.031</td>
</tr>
</tbody>
</table>
Table 4

Serum and colonic levels of PC-SOD.

Mice were orally administered 48 kU/kg PC-SOD once and treated with DSS for 2 days.

Blood and colonic tissue samples were taken periodically. The levels of PC-SOD in the samples were determined by ELISA. Values are mean ± S.E.M.

<table>
<thead>
<tr>
<th></th>
<th>6 h</th>
<th>12 h</th>
<th>24 h</th>
<th>48 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum (µg/mL)</td>
<td>&lt;0.16</td>
<td>&lt;0.16</td>
<td>&lt;0.16</td>
<td>&lt;0.16</td>
</tr>
<tr>
<td>Tissue (ng/mg)</td>
<td>&lt;0.031</td>
<td>&lt;0.031</td>
<td>0.46 ± 0.24</td>
<td>&lt;0.031</td>
</tr>
</tbody>
</table>
Figure 2

A

Disease activity index

DSS (n=11)
+ PC-SOD 0.38 kU/kg (n=8)
+ PC-SOD 0.75 kU/kg (n=6)
+ PC-SOD 1.5 kU/kg (n=9)
+ PC-SOD 3 kU/kg (n=9)
+ PC-SOD 6 kU/kg (n=10)
+ PC-SOD 12 kU/kg (n=11)

Day

0 1 2 3 4 5 6 7

B

Length of colon (cm)

PC-SOD (kU/kg)
0 0.38 0.75 1.5 3 6 12
(n=7) (n=8) (n=9) (n=11)

C

MPO activity
(umol H_2O_2/mg protein)

PC-SOD (kU/kg)
0 0.38 0.75 1.5 3 6 12
(n=11) (n=8) (n=9) (n=10) (n=7) (n=8)
Figure 2

D

Disease activity index

Day

0 1 2 3 4 5 6 7

DSS (n=11)
+ U-SOD 1.5 kU/kg (n=12)
+ U-SOD 3 kU/kg (n=10)
+ U-SOD 6 kU/kg (n=7)
+ U-SOD 12 kU/kg (n=6)
+ U-SOD 24 kU/kg (n=6)
+ U-SOD 48 kU/kg (n=9)

E

Length of colon (cm)

U-SOD (kU/kg)

0 1.5 3 6 12 24 48

(n=7) (n=5) (n=5) (n=6) (n=6) (n=9)

F

MPO activity (μmol H₂O₂/mg protein)

U-SOD (kU/kg)

0 1.5 3 6 12 24 48

(n=11) (n=10) (n=5) (n=6) (n=6) (n=9)
Figure 3

A

+ PC-SOD 0.25 U/mL
+ U-SOD 0.25 U/mL
+ PC-SOD 2.5 U/mL
+ U-SOD 2.5 U/mL
+ PC-SOD 25 U/mL
+ U-SOD 25 U/mL

B

Relative intensity

0 1 2 3

PMA  PMA  PMA
+ PC-SOD 2.5 U/mL  + U-SOD 2.5 U/mL

C

Chemiluminescence (cpm)

0 2000 4000 6000 8000 10000

PMA  + PC-SOD 2.5 U/mL  + U-SOD 2.5 U/mL  + PC-SOD 25 U/mL  + U-SOD 25 U/mL  + PC-SOD 250 U/mL  + U-SOD 250 U/mL

D

% of control

0 25 50 75 100

PMA  PMA  PMA
+ PC-SOD 25 U/mL  + U-SOD 25 U/mL

** #

## **

** #

Downloaded from Jpet.aspetjournals.org at AS/PEJournals on July 8, 2017.
Figure 5

G

<table>
<thead>
<tr>
<th>CTRL</th>
<th>DSS</th>
<th>DSS + PC-SOD 3 kU/kg</th>
<th>DSS + PC-SOD 6 kU/kg</th>
<th>DSS + Catalase 6 kU/kg</th>
<th>DSS + PC-SOD 6 kU/kg</th>
</tr>
</thead>
</table>

Phospho-NF-κB

DAPI