

Role of Cu,Zn-SOD in the synthesis of endogenous vasodilator hydrogen peroxide during reactive hyperemia in mouse mesenteric microcirculation in vivo

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Yada T, Shimokawa H, Morikawa K, Takaki A, Shinozaki Y, Mori H, Goto M, Ogasawara Y, Kajiya F. Role of Cu,Zn-SOD in the synthesis of endogenous vasodilator hydrogen peroxide during reactive hyperemia in mouse mesenteric microcirculation in vivo. *Am J Physiol Heart Circ Physiol* 294: H441–H448, 2008. First published November 16, 2007; doi:10.1152/ajpheart.01021.2007.—We have recently demonstrated that endothelium-derived hydrogen peroxide (H_2O_2) is an endothelium-derived hyperpolarizing factor and that endothelial Cu/Zn-superoxide dismutase (SOD) plays an important role in the synthesis of endogenous H_2O_2 in both animals and humans. We examined whether SOD plays a role in the synthesis of endogenous H_2O_2 during in vivo reactive hyperemia (RH), an important regulatory mechanism. Mesenteric arterioles from wild-type and Cu,Zn-SOD^{-/-} mice were continuously observed by a pencil-type charge-coupled device (CCD) intravital microscope during RH (reperfusion after 20 and 60 s of mesenteric artery occlusion) in the cyclooxygenase blockade under the following four conditions: control, catalase alone, N^ω-monomethyl-L-arginine (L-NMMA) alone, and L-NMMA + catalase. Vasodilation during RH was significantly decreased by catalase or L-NMMA alone and was almost completely inhibited by L-NMMA + catalase in wild-type mice, whereas it was inhibited by L-NMMA and L-NMMA + catalase in the Cu,Zn-SOD^{-/-} mice. RH-induced increase in blood flow after L-NMMA was significantly increased in the wild-type mice, whereas it was significantly reduced in the Cu,Zn-SOD^{-/-} mice. In mesenteric arterioles of the Cu,Zn-SOD^{-/-} mice, Tempol, an SOD mimetic, significantly increased the ACh-induced vasodilatation, and the enhancing effect of Tempol was decreased by catalase. Vascular H_2O_2 production by fluorescent microscopy in mesenteric arterioles after RH was significantly increased in response to ACh in wild-type mice but markedly impaired in Cu,Zn-SOD^{-/-} mice. Endothelial Cu,Zn-SOD plays an important role in the synthesis of endogenous H_2O_2 that contributes to RH in mouse mesenteric smaller arterioles.

nitric oxide; endothelium-derived hyperpolarizing factor; arteriole; vasodilatation

THE ENDOTHELIUM SYNTHESIZES and releases endothelium-derived relaxing factors (EDRFs), including vasodilator prostaglandins, nitric oxide (NO), and as yet unidentified endothelium-derived hyperpolarizing factor (EDHF). Since the first reports on the existence of EDHFs (4, 8), several candidates for EDHF

have been proposed (9), including cytochrome P-450 metabolites (2, 3), endothelium-derived K^+ channel (7), and electrical communications through gap junctions between endothelial cells and vascular smooth muscle cells (34). Matoba et al. (19a, 19b, 20) previously identified that endothelium-derived hydrogen peroxide (H_2O_2) is a primary EDHF in mesenteric arteries of mice, pigs, and humans. Morikawa et al. (24a, 25) subsequently confirmed that endothelial Cu/Zn-superoxide dismutase (SOD) plays an important role in synthesizing EDHF/ H_2O_2 in mice and humans. Recently, our laboratory (41a, 42) confirmed that endogenous H_2O_2 plays an important role for autoregulation and protection against reperfusion injury in canine coronary microcirculation.

Reactive hyperemia (RH) is an important regulatory mechanism of the cardiovascular system in response to a temporal reduction in blood flow for which both mechanosensitive (e.g., myogenic and shear mediated) and metabolic regulatory processes may be involved (6, 14a, 28). For the RH response of canine coronary microcirculation, NO, ATP-sensitive K^+ channels, and adenosine may all be involved (11, 41). Shear stress plays a crucial role in modulating vascular tone by stimulating the release of EDRFs (8, 32), and all three EDRFs (PGI₂, NO, and EDHF) are involved in flow-induced vasodilatation (15, 18, 33, 44).

However, it remains to be examined whether endogenous H_2O_2 is involved in the vasodilator mechanism of RH and, if so, whether endothelial Cu,Zn-SOD plays a role in the synthesis of endogenous H_2O_2 during RH. The present study was thus designed to address these important issues in mice. Our laboratory (42, 44) previously reported that the contribution of EDHF to the vasodilatory mechanisms increases as the diameter of the vessel decreases. Thus, by employing a pencil-type charge-coupled device (CCD) intravital microscope with a high resolution, we focused on the arterioles with a diameter of <50 μ m in vivo.

METHODS

The present study was approved by the Animal Care and Use Committee of Kawasaki Medical School and conformed to the guidelines on animal experiments of Kawasaki Medical School and the

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Animal preparation. Male Cu,Zn-SOD^{-/-} and control mice (10–16 wk of age) derived from breeding pairs of heterozygous (Cu,Zn-SOD^{+/-}) mice (Jackson Laboratory, Bar Harbor, ME) were used (25). They were placed on a heating blanket to maintain body temperature at 37°C throughout the experiment. The animals were anesthetized with 1% inhalational anesthesia of isoflurane. After tracheal intubation, they were ventilated with a mixture of room air and oxygen by a ventilator. The abdomen was opened, and a 24-Fr catheter was inserted into the abdominal aorta to measure aortic pressure. Mesenteric arterioles were continuously observed by a pencil-type intravital microscope (Nihon Kohden, Tokyo, Japan) (13).

Measurements of diameter by pencil-type intravital microscope. Mesenteric arterioles were visualized using a pencil-type intravital microscope (13). The system was modified for the visualization of microcirculation from our previous needle-probe CCD videomicroscope system (40). The microscopic images were monitored and recorded on a digital videocassette recorder (Sony, Tokyo, Japan) every 33 ms (30 frames/s). The spatial resolution of a static image of this system is 0.5 μm for ×600 magnification. The field of view is 367 × 248 μm, and the focal depth is 50 μm.

Measurements of regional blood flow in mesenteric arteries. Regional blood flow in mesenteric arteries was measured by the nonradioactive microsphere (15 μm; Sekisui Plastic, Tokyo, Japan) technique at the end of the experiments, as previously described (24). Briefly, a bolus (50 μl) of the microspheres suspension (5 × 10⁵ spheres; Ce and Ba) were injected into the abdominal aorta at baseline and 5 s after the reperfusion of the mesenteric artery with confirming changes of the blood flow of the mesenteric artery by a CCD intravital microscope and without inducing hemodynamic changes (14). Mice were euthanized, and the mesenterium was extracted. The X-ray fluorescence of the stable heavy elements was measured by a wavelength-dispersive spectrometer (model PW 1480; Phillips, Eindhoven, the Netherlands). The relative increase in blood flow of mesenterium [microsphere count/tissue weight (g)] during RH from baseline was calculated.

Detection of H₂O₂ and NO production in mesenteric microvessels. 2',7'-Dichlorodihydrofluorescein diacetate (DCF-DA; Molecular Probes, Eugene, OR) and diaminorhodamine-4M AM (DAR; Daiichi Pure Chemicals, Tokyo, Japan) were used to detect H₂O₂ and NO production in mesenteric microvessels, respectively, as previously described (41a). Briefly, fresh and unfixed mesenteric tissue was cut into several blocks and immediately frozen in an optimal cutting temperature compound (Tissue-Tek; Sakura Fine Chemical, Tokyo, Japan). After washout of the mesenteric tissue with phosphate-buffered solution under a normal temperature, fluorescent images of the microvessels

were obtained 3 min after application of acetylcholine (ACh) by using a fluorescence microscope (Olympus BX51) (41a). We defined the baseline fluorescent intensity as the response in the vascular endothelium just after the injection of NO or H₂O₂ fluorescent dye. The fluorescence data at baseline (both DCF-DA and DAR) were obtained after the RH.

Experimental protocol. We performed four protocols. First, mesenteric arterioles in wild-type and Cu,Zn-SOD^{-/-} mice were continuously observed by a pencil-type intravital microscope during RH (reperfusion after 20 and 60 s of mesenteric artery occlusion) with cyclooxygenase blockade [indomethacin, 5 × 10⁻⁵ mol/l topical administration (ta)] with the following four conditions: control, catalase alone [1,500 U·min⁻¹·100 g body wt⁻¹ intra-arterial administration (ia)] polyethylene glycol-catalase, a specific decomposer of H₂O₂, NO synthase inhibitor alone (10⁻⁸ mol/l ta L-NMMA), and L-NMMA + catalase (17). In the presence of indomethacin and L-NMMA, microspheres were administered at baseline and 5 s after the reperfusion into the abdominal aorta by bolus injection because RH peaked within 20–60 s after release from 20- and 60-s occlusion (29). Maximal vascular diameter was measured within 20 and 60 s after the reperfusion. Second, ACh (10⁻⁷ to 10⁻⁵ mol/l ta)-induced endothelium-dependent vasodilatation was examined under the control conditions and in the presence of Tempol, a SOD mimetic 4-hydroxy-2,2,6,6-tetramethylpiperidine-N-oxyl (50 μg·min⁻¹·100 g body wt⁻¹ ia) (17), and Tempol + catalase. In the combined infusion protocol (Tempol or Tempol + catalase) in the presence of cyclooxygenase blockade + L-NMMA, the combined infusion was performed simultaneously for 20 min. ACh was infused for 10 min, and the vascular diameter was measured. Third, sodium nitroprusside (SNP; 10⁻⁷ to 10⁻⁵ mol/l ta, each 10 min)-induced endothelium-independent vasodilatation was examined in wild-type and Cu,Zn-SOD^{-/-} mice. Fourth, fresh and unfixed mesenteric tissue was then cut into several blocks and immediately frozen in optimal cutting temperature compound.

Statistical analysis. The results are expressed as means ± SE. Dose-response curves were analyzed by two-way ANOVA followed by the Scheffé's post hoc test for multiple comparisons. Vascular responses were analyzed by one-way ANOVA followed by the Scheffé's post hoc test for multiple comparisons. *P* < 0.05 was considered to be statistically significant.

RESULTS

Hemodynamics and blood gases during RH. Throughout the experiments, mean aortic pressure and heart rate were constant and comparable (Tables 1 and 2), and P_{O₂}, P_{CO₂}, and pH were maintained within the physiological ranges (>70 mmHg P_{O₂}, 25–40 mmHg P_{CO₂}, and pH 7.35–7.45). Baseline mesenteric

Table 1. Hemodynamics during RH

	n	Control		Catalase		L-NMMA		L-NMMA + Catalase	
		B	RH	B	RH	B	RH	B	RH
MBP, RH 20, mmHg									
WT	10	83 ± 7	85 ± 9	81 ± 7	82 ± 7	82 ± 8	81 ± 6	83 ± 8	82 ± 6
Cu,Zn-SOD ^{-/-}	10	85 ± 12	87 ± 10	83 ± 8	82 ± 8	82 ± 8	82 ± 8	82 ± 8	80 ± 9
MBP, RH 60, mmHg									
WT	5	86 ± 8	88 ± 7	87 ± 7	88 ± 7	88 ± 8	87 ± 7	88 ± 7	87 ± 7
Cu,Zn-SOD ^{-/-}	5	88 ± 8	86 ± 8	87 ± 6	89 ± 9	88 ± 7	88 ± 8	89 ± 6	90 ± 11
HR, RH 20, beats/min									
WT	10	346 ± 14	348 ± 14	335 ± 15	333 ± 17	315 ± 15	310 ± 17	330 ± 18	330 ± 18
Cu,Zn-SOD ^{-/-}	10	364 ± 27	354 ± 22	350 ± 18	351 ± 15	355 ± 15	340 ± 17	355 ± 15	335 ± 17
HR, RH 60, beats/min									
WT	5	351 ± 31	361 ± 9	353 ± 21	356 ± 13	358 ± 10	364 ± 37	354 ± 22	355 ± 15
Cu,Zn-SOD ^{-/-}	5	346 ± 18	356 ± 25	356 ± 15	361 ± 19	351 ± 31	361 ± 9	358 ± 10	364 ± 27

Values are means ± SE; n, number of rats. RH, reactive hyperemia; L-NMMA, N^G-monomethyl-L-arginine; B, baseline; MBP, mean blood pressure; HR, heart rate; WT, wild-type.

Table 2. Hemodynamics during administration of ACh and SNP

MBP, mmHg	n	Control							Tempol							Tempol + Catalase							SNP						
		B		ACh 10 ⁻⁷	ACh 10 ⁻⁶	ACh 10 ⁻⁵	ACh 10 ⁻⁴	ACh 10 ⁻³	ACh 10 ⁻⁶	ACh 10 ⁻⁵	ACh 10 ⁻⁴	ACh 10 ⁻³	ACh 10 ⁻²	ACh 10 ⁻⁵	ACh 10 ⁻⁴	ACh 10 ⁻³	ACh 10 ⁻²	ACh 10 ⁻⁵	ACh 10 ⁻⁴	ACh 10 ⁻³	ACh 10 ⁻²	B	SNP 10 ⁻⁷	SNP 10 ⁻⁶	SNP 10 ⁻⁵	SNP 10 ⁻⁴	SNP 10 ⁻³		
		n	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	MBP	HR	
WT	10	90±7	88±7	87±7	89±6	91±10	90±11	84±11	91±8	86±8	86±8	86±8	84±11	91±8	86±8	86±8	84±11	91±8	86±8	86±8	78±13	77±13	76±13	76±13	76±13	76±13	76±13		
Cu,Zn-SOD ^{-/-}	10	93±11	92±10	94±10	97±10	98±16	97±15	99±15	91±8	91±8	91±8	91±8	99±15	91±8	91±8	91±8	99±15	91±8	91±8	91±8	81±8	77±10	75±11	75±11	75±11	75±11	75±11		
WT	10	361±9	340±17	340±17	340±17	342±24	336±25	336±25	320±19	330±27	330±27	330±27	336±25	320±19	330±27	330±27	336±25	320±19	330±27	330±27	370±57	364±37	351±31	351±31	351±31	351±31	351±31		
Cu,Zn-SOD ^{-/-}	10	386±11	381±24	371±21	396±13	386±11	377±15	377±15	362±22	356±25	356±25	356±25	377±15	362±22	356±25	356±25	377±15	362±22	356±25	356±25	372±21	369±6	358±10	358±10	358±10	358±10	358±10		

Values are means ± SE; n, number of rats. SNP, sodium nitroprusside; ACh, acetylcholine.

arteriolar diameter was comparable in the absence and presence of inhibitors under the four different experimental conditions (Tables 3 and 4). Those different inhibitors (L-NMMA, catalase, and Tempol) did not affect basal diameter.

Mesenteric vasodilatation during RH. We were able to observe EDHF-sensitive smaller arterioles (18–66 μ m) by using a newly developed pencil-type CCD intravital microscope with a higher resolution. In the mesenteric arterioles of wild-type mice, vasodilatation during RH to 20- and 60-s arterial occlusion was decreased by catalase or L-NMMA alone and was almost completely inhibited by L-NMMA + catalase (Fig. 1). In contrast, in mesenteric arterioles of Cu,Zn-SOD^{-/-} mice, vasodilatation during RH to 20- and 60-s arterial occlusion was decreased by catalase alone and was almost completely inhibited by L-NMMA alone or L-NMMA + catalase (Fig. 1). Blood flow measurement by microsphere technique showed that in the presence of indomethacin and L-NMMA, RH-induced increase in blood flow was 232 ± 4% (20 s) and 331 ± 4% (60 s) of baseline in control and was sensitive to catalase (137 ± 4%, 20 s; and 147 ± 17%, 60 s) in the wild-type mice, whereas in the Cu,Zn-SOD^{-/-} mice, the vasodilator response was significantly reduced to 125 ± 19% (20 s) and 145 ± 23% (60 s) in control and was insensitive to catalase (120 ± 24%, 20 s; and 139 ± 19%, 60 s) (Fig. 2). With the longer occlusion of the mesenteric artery, the shear stimulus for H₂O₂ release was significantly increased in the control condition and was significantly decreased by catalase.

Endothelium-dependent vasodilatation. In mesenteric arterioles of wild-type mice, endothelium-dependent vasodilatation to ACh (10⁻⁷ to 10⁻⁵ mol/l) in the presence of indomethacin and L-NMMA was unchanged with Tempol but significantly inhibited by the addition of catalase (Fig. 3). In contrast, in the mesenteric arterioles of the Cu,Zn-SOD^{-/-} mice, the response to ACh was significantly enhanced with Tempol, a response that was sensitive to the addition of catalase (Fig. 3).

Endothelium-independent vasodilatation. Endothelium-independent vasodilatation to SNP (10⁻⁷ to 10⁻⁵ mol/l) in the presence of L-NMMA + catalase was comparable between the two strains (Table 4).

Detection of H₂O₂ and NO production in the mesenteric artery. Fluorescent microscopy with DCF-DA showed that vascular H₂O₂ production in mesenteric arterioles was significantly increased in response to ACh in wild-type mice compared with baseline but markedly impaired in Cu,Zn-SOD^{-/-} mice (Fig. 4). In contrast, vascular NO production in mesenteric arterioles, as assessed by DAR fluorescent intensity, was significantly increased in response to ACh in wild-type mice compared with baseline and was unaltered in Cu,Zn-SOD^{-/-} mice (Fig. 5).

DISCUSSION

The novel finding of the present study with a newly developed pencil-type CCD intravital microscope *in vivo* is that Cu,Zn-SOD plays an important role in the synthesis of endogenous H₂O₂, which is substantially involved in the mechanisms of RH-induced vasodilatation in mouse mesenteric circulation.

Impaired EDHF-mediated vasodilatation in Cu,Zn-SOD^{-/-} mice *in vivo*. Matoba et al. (19a, 20) have previously identified that endothelium-derived H₂O₂ is an EDHF in mouse and

Table 3. Diameter change during RH

	Control		Catalase		L-NMMA		L-NMMA + Catalase	
	B	RH	B	RH	B	RH	B	RH
RH 20, μm								
WT	36 \pm 4	49 \pm 4 \dagger	36 \pm 4	44 \pm 4 \dagger	36 \pm 3	42 \pm 3*	36 \pm 4	38 \pm 3
Cu,Zn-SOD ^{-/-}	36 \pm 4	48 \pm 5 \dagger	36 \pm 4	42 \pm 5 \dagger	36 \pm 4	38 \pm 4	36 \pm 3	38 \pm 4
RH 60, μm								
WT	40 \pm 4	55 \pm 4 \dagger	40 \pm 4	51 \pm 4 \dagger	40 \pm 5	47 \pm 4*	39 \pm 4	41 \pm 4
Cu,Zn-SOD ^{-/-}	40 \pm 3	56 \pm 4 \dagger	40 \pm 3	51 \pm 4 \dagger	39 \pm 4	42 \pm 3	40 \pm 5	42 \pm 3

Values are means \pm SE; n, number of arterioles per animal. * $P < 0.01$ vs. B.

human mesenteric microvessels. Subsequently, our laboratory (42) and others (23) have confirmed that endogenous H_2O_2 exerts important vasodilator effects in canine coronary microcirculation in vivo and in isolated human coronary microvessels, respectively. H_2O_2 can be formed from superoxide anions derived from several sources in endothelial cells, including endothelial NO synthase (eNOS), cyclooxygenase, lipoxygenase, cytochrome P-450 enzymes, and reduced NADP [NAD(P)H] oxidases. Gupte et al. (10) demonstrated that cytosolic NADH redox and Cu,Zn-SOD activity have important roles in controlling the inhibitory effects of superoxide anions derived from NADH oxidase. Morikawa et al. (24a, 25) have also demonstrated that endothelial Cu,Zn-SOD plays an important role in the synthesis of H_2O_2 in mouse and human mesenteric arteries in vitro.

In the present study, catalase or L-NMMA alone significantly, but not completely, inhibited the RH-induced vasodilatation of mesenteric arterioles in wild-type mice in vivo, whereas L-NMMA + catalase markedly attenuated the remaining vasodilatation. In contrast, in Cu,Zn-SOD^{-/-} mice, L-NMMA alone significantly decreased the vasodilatation and blood flow in response to 20- and 60-s arterial occlusion (Figs. 1 and 2). These results obtained using a pencil-type CCD intravital microscope indicate that H_2O_2 exerts important vasodilator effects on mesenteric smaller arterioles during RH and that Cu,Zn-SOD plays an important role in the synthesis of endogenous H_2O_2 during RH in vivo. Koller and Bagi (14a) showed that RH in rat isolated coronary arterioles was sensitive to pressure/stretch and flow/shear stress. Miura et al. (23) also showed the important role of endogenous H_2O_2 in flow-induced vasodilatation of human coronary arterioles. Koller and Bagi (14a) also suggested that H_2O_2 contributes to the development of the early peak phase of RH but not the duration of reactive vasodilatation, whereas NO prolongs the later phase of RH in rat isolated coronary arterioles, suggesting that H_2O_2 released endogenously within the vascular wall changes hemodynamic forces. In the present study, peak blood flow was significantly decreased after catalase (Fig. 2), suggesting that flow-induced vasodilatation during the early phase of RH is

indeed mediated by H_2O_2 in mouse mesenteric arterioles in vivo.

Compensatory vasodilator mechanism between H_2O_2 and NO. It is well known that coronary vascular tone is regulated by the interactions among hemodynamic forces and several endogenous vasodilators, including NO, H_2O_2 , and adenosine (41a, 42). Koller and Bagi (14a) demonstrated that mechanosensitive mechanisms were activated by changes in pressure and flow/shear stress during RH in isolated coronary arterioles. A superoxide anion is dismutated to H_2O_2 by manganese SOD (Mn-SOD, mitochondrial matrix) and Cu,Zn-SOD. H_2O_2 diffuses across the mitochondrial membrane to act on vascular smooth muscle (45). Tsunoda et al. (35) demonstrated that Mn-SOD augmented RH during 60-s canine coronary ischemia and reperfusion. H_2O_2 generated in the arteriolar smooth muscle could cause the response of activation of cGMP in rat skeletal muscle arterioles (38). Kitakaze et al. (12) indicated that the augmentation of reactive hyperemic flow caused by SOD is attributed to the enhanced release of adenosine in canine coronary circulation. These endogenous vasodilators may play an important role in causing the compensatory vasodilatation of coronary microvessels during myocardial ischemia.

In the present study, endothelium-dependent vasodilatation during RH (in the presence of L-NMMA) was almost completely inhibited by catalase in wild-type mice. In the Cu,Zn-SOD^{-/-} mice, vasodilatation during RH remained under the control condition but was almost completely inhibited by L-NMMA (Fig. 1). The RH-induced increase in blood flow (in the presence of indomethacin and L-NMMA) was significantly inhibited by catalase in the wild-type mice but not in Cu,Zn-SOD^{-/-} mice (Fig. 2). RH-induced increase in blood flow (in the presence of indomethacin and L-NMMA) remained in Cu,Zn-SOD^{-/-} mice (Fig. 2). H_2O_2 may compensate for the loss of action of NO. H_2O_2 produced by SOD other than Cu,Zn-SOD may compensate for the loss of action of Cu,Zn-SOD-derived H_2O_2 .

Table 4. Diameter change during administration of ACh and SNP

	ACh			SNP			
	B	ACh 10^{-7}	ACh 10^{-6}	B	SNP 10^{-7}	SNP 10^{-6}	SNP 10^{-5}
WT, μm	36 \pm 3	41 \pm 3*	45 \pm 3 \dagger	35 \pm 4	39 \pm 4*	43 \pm 5 \dagger	46 \pm 5 \dagger
Cu,Zn-SOD ^{-/-} , μm	36 \pm 4	38 \pm 4	41 \pm 4*	34 \pm 4	38 \pm 4*	41 \pm 4 \dagger	44 \pm 4 \dagger

Values are means \pm SE; n, number of arterioles per animal. * $P < 0.05$; $\dagger P < 0.01$ vs. B.

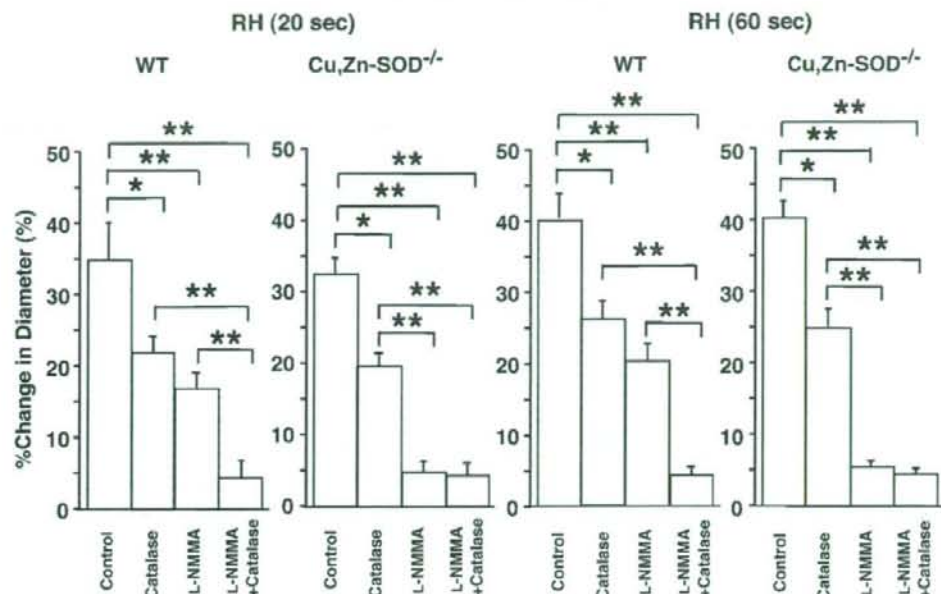


Fig. 1. Mesenteric vasodilation during reactive hyperemia (RH). In the wild-type (WT) mice, vasodilation during RH was inhibited by catalase or *N*³-monomethyl-L-arginine (L-NMMA) and further inhibited by L-NMMA + catalase. In the *Cu,Zn-SOD*^{-/-} mice (*Cu,Zn-SOD*^{-/-}), vasodilation during RH was inhibited by catalase and markedly inhibited by L-NMMA, and the remaining response was not inhibited by catalase. The number of arterioles per animals used was 10/5 for each group. **P* < 0.05; ***P* < 0.01.

Improvement of ACh-induced vasodilatation by Tempol in Cu,Zn-SOD^{-/-} mice. It was previously reported that Tempol, a cell membrane-permeable SOD mimetic 4-hydroxy-2,2,6,6-tetramethylpiperidine-*N*-oxyl, decreased oxidative stress in the spontaneously hypertensive rat (31). In the present study, Tempol significantly improved the ACh-induced vasodilatation in *Cu,Zn-SOD*^{-/-} mice, whereas catalase abolished the beneficial effect of Tempol (Fig. 3), indicating that the effect of Tempol was mediated by endogenous H₂O₂ in vivo. In contrast, Tempol had no enhancing effect on the ACh-induced vasodilatation in control mice (Fig. 3), suggesting that a sufficient amount of SOD is present in this strain. In *Cu,Zn-*

SOD^{-/-} mice, L-NMMA did not abolish the ACh-induced vasodilatation, and the DCF-DA stain showed remaining fluorescent intensity (Fig. 4). Thus the residual vasodilatation could be caused by the following possible mechanisms. First, NO may also be synthesized in a nonenzymatic manner (27). Nonenzymatic synthesis of NO could occur in the presence of NADPH, glutathione, and L-cysteine, etc., opposing the effects of NOS inhibition (27). Second, the effects of L-NMMA may be limited since it is known that L-NMMA does not abolish NO production (1). H₂O₂ produced from vascular smooth muscle cells and other tissues may also contribute to the residual vasodilatation (5, 30). Third, the contribution of other proposed

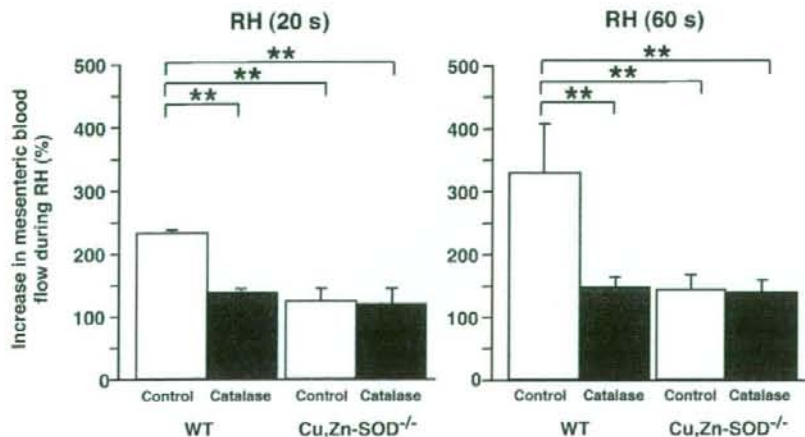
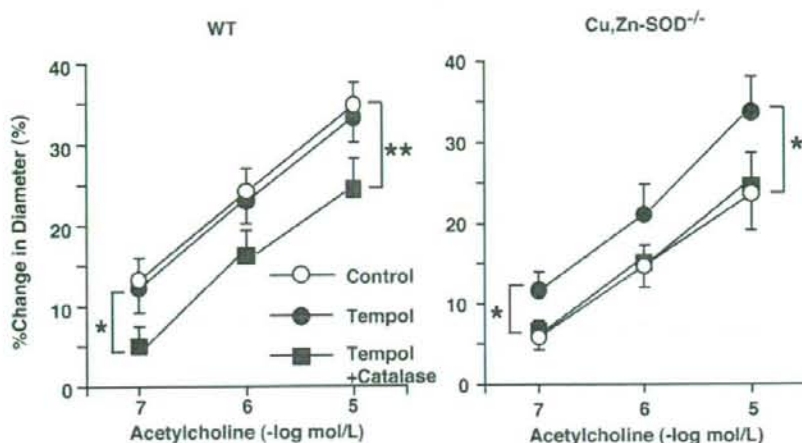


Fig. 2. The increase in mesenteric blood flow during RH. In the presence of indomethacin and L-NMMA, RH-induced increase in blood flow was sensitive to catalase in the WT mice, whereas in the *Cu,Zn-SOD*^{-/-}, the vasodilatation was significantly reduced in control and was insensitive to catalase. The number of animals used was 5 for each group. ***P* < 0.01.

Fig. 3. Endothelium-dependent relaxations to ACh. In the WT mice, endothelium-dependent vasodilatation to ACh (in the presence of indomethacin and L-NMMA) was unchanged with Tempol but significantly inhibited by the addition of catalase. In the Cu,Zn-SOD^{-/-}, the vasodilation was significantly enhanced with Tempol, where the response was sensitive to the addition of catalase. The number of arterioles per animals used was 10/5 for each group. **P* < 0.05; ***P* < 0.01.



EDHF candidates, such as *P*-450 metabolites (2, 3) and potassium ion (7), may contribute to the residual vasodilatation. Although RH and ACh have different mechanisms of vasodilator effects, they also share the same flow-induced vasodilator mechanism.

Endothelium-independent vasodilatation in Cu,Zn-SOD^{-/-} mice. Microvascular dysfunction in hypercholesterolemic rats was confined to the endothelium because the dilator response to SNP and adenosine was unchanged (37). In the present study, endothelium-independent vasodilatation in response to SNP was comparable between the two genotypes, suggesting

that vasodilatation properties of vascular smooth muscle cells were preserved in the Cu,Zn-SOD^{-/-} mice in vivo.

Detection of vascular H₂O₂ and NO production. Our laboratory (41a) has recently demonstrated that vascular production of H₂O₂ and NO after ischemia-reperfusion is enhanced in small coronary arteries and arterioles in vivo, respectively. It was previously shown that a ACh-induced increase in fluorescence intensity in endothelial cells of the mesenteric artery is significantly reduced in Cu,Zn-SOD^{-/-} mice (25). In the present study, vascular H₂O₂ production, as assessed by DCF-DA fluorescent intensity in mesenteric arterioles, was markedly impaired

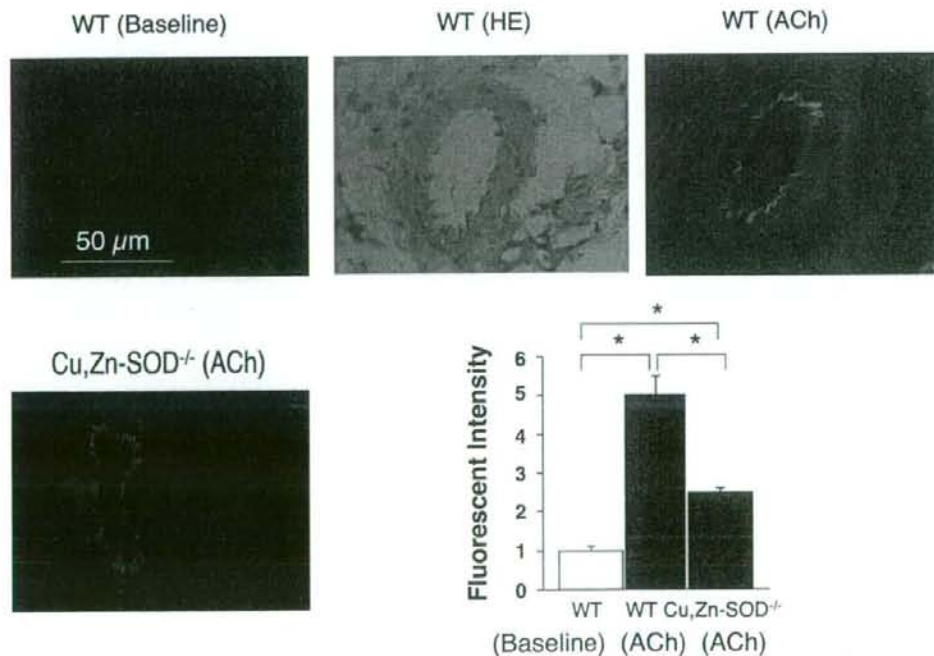


Fig. 4. Detection of vascular H₂O₂ production. Vascular H₂O₂ production in mesenteric arterioles was significantly increased in response to ACh in WT mice but markedly impaired in Cu,Zn-SOD^{-/-}. The number of arterioles per animals used was 10/5 for each group. **P* < 0.05. HE, Hematoxylin eosin.

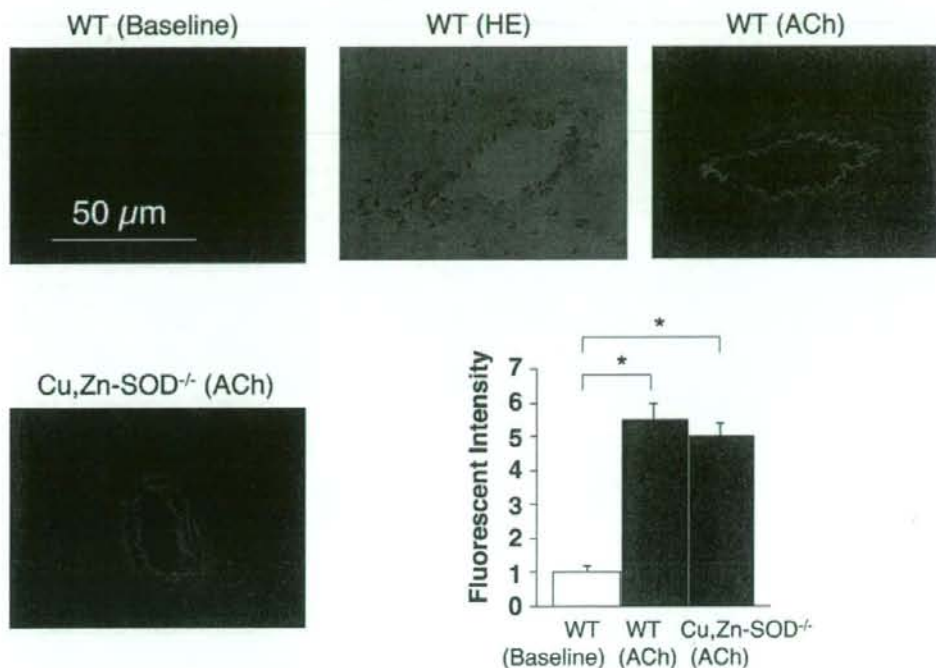


Fig. 5. Detection of vascular nitric oxide (NO) production. Vascular NO production in mesenteric arterioles was significantly increased in response to ACh in WT mice and unaltered in Cu,Zn-SOD^{-/-}. The number of arterioles per animals used was 10/5 for each group. * $P < 0.05$.

in Cu,Zn-SOD^{-/-} mice (Fig. 4). These findings indicate that endothelial production of H₂O₂ is significantly impaired in Cu,Zn-SOD^{-/-} mice, confirming the importance of the enzyme in endothelial synthesis of H₂O₂.

In the previous study by Morikawa et al. (25), eNOS protein expression was comparable between Cu,Zn-SOD^{-/-} and wild-type mice. In the present study, vascular NO production in small mesenteric artery was unaltered in Cu,Zn-SOD^{-/-} mice compared with wild-type mice (Fig. 5). NO could compensate for the loss of action of H₂O₂, although there are still many uncertainties about the local cellular dynamics of superoxide anions and NO.

Study limitations. Several limitations should be mentioned for the present study. First, we estimated blood flow in the mesenteric circulation using microspheres. We were unable to calculate the absolute values of local blood flow or shear stress because of the methodological limitations. However, since the flow measurement with microspheres was performed at the end of the experiments, it should not have influenced other results. Second, we used Cu,Zn-SOD^{-/-} mice in the present study, where unknown compensatory mechanisms may be operative, and we were unable to elucidate the mechanism(s) for the remaining EDHF-mediated responses in those mice.

Clinical implications. RH is an important regulatory mechanism of the cardiovascular system, reflecting the flow reserve in response to a brief period of cessation of flow. An impaired flow reserve in resistance vessels is a hallmark of microvascular dysfunction with coronary risk factors. Hypertension is associated with structural alterations in the microcirculation and a reduced endothelium-dependent dilation in conduit ar-

teries (19). It is well known that abnormality in Cu,Zn-SOD is noted in several diseases, including hypertension and diabetes mellitus (36, 39).

In conclusion, endogenous H₂O₂ exerts important vasodilator effects of mesenteric smaller arterioles during RH, especially at the low level of NO, and that Cu,Zn-SOD plays an important role in the synthesis of endogenous H₂O₂ during RH in vivo.

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研究論文

X-ray Spectra from a Brass-target Plasma Triode

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Characteristic x-rays, Absorption of zinc K β rays

Abstract

In this paper, we describe a recently developed table-top plasma x-ray generator utilizing a brass-target triode, which we used in preliminary experiments for the superposition of K-series characteristic x-rays in weakly ionized plasma and for producing their higher harmonics. In the plasma flash x-ray generator, a 200 nF condenser was charged, and flash x-rays were produced by discharging. The x-ray tube was a demountable triode with a brass target containing 65% copper and 35% zinc by weight, and a turbomolecular pump evacuated air from the tube with a pressure of 1 mPa. Target evaporation led to the formation of weakly ionized linear plasma, consisting of metal ions and electrons, around the rod target, and intense characteristic x-rays were produced. At a charging voltage of 50 kV, the maximum tube voltage was almost equal to the charging voltage of the main condenser, and the peak current was 15 kA. When the charging voltage was increased, the linear plasma formed, and the K-series characteristic x-ray intensities of zinc K α , copper K β , and copper K β lines increased substantially. On the other hand, hardly any zinc K β lines were detected. In particular, we confirmed the irradiation of the second unsharp harmonics of the fundamental K α lines of copper and zinc. The x-ray pulse widths were 700 ns, and the time-integrated x-ray intensity was 1.2 mGy at 1.0 m per pulse with a charging voltage of 50 kV.

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1. Introduction

Synchrotrons produce monochromatic parallel beams using single crystals, and the photon energy is selected by Bragg's angle. In particular, the beams with photon energies of approximately 35 keV have been employed for enhanced K-edge angiography¹⁻³⁾ using iodine contrast media, because iodine media with a K-edge of 33.2 keV absorb the beams effectively. Thus, we have developed a steady state cerium x-ray generator⁴⁻⁶⁾ and have succeeded in carrying out cone-beam enhanced angiography using cerium K-series characteristic x-rays.

Conventional flash x-ray generators⁷⁾ utilize high-voltage condensers and cold-cathode x-ray tubes, and produce extremely short x-ray pulses with durations of less than 1 μ s. For use in biomedical radiography, we have developed several different flash x-ray generators⁸⁻¹⁰⁾ corresponding to specific radiographic objectives, and have succeeded in producing clean K-series characteristic x-rays of nickel and copper from a weakly ionized linear plasma using a plasma triode¹¹⁻¹⁴⁾. We have confirmed the irradiations of clean K-series characteristic x-rays of molybdenum using a compact flash x-ray generator with a disk-cathode diode^{15,16)}, and we have developed an intense plasma diode to produce high-photon-energy characteristic x-rays of molybdenum, cerium¹⁷⁾, gadolinium, tantalum¹⁸⁾ and tungsten¹⁹⁾. In particular, the tantalum and tungsten K α rays have been applied to perform high-speed K-edge angiography using gadolinium-based contrast media.

In order to carry out wide-photon-energy or energy subtraction radiography, we are very interested in the variations of characteristic x-

ray intensities from evaporating plasma target. In particular, the absorption of K-series characteristic x-rays in the plasma consisting of electrons and two-element metal ions requires investigations, since all K-rays of two elements are produced from the two-element solid alloy target. Furthermore, because we have confirmed the irradiation of higher harmonic hard x-rays using nickel and copper targets, the x-ray spectra with photon energies beyond the K-edges should be measured.

In this paper, we describe a recently developed table-top plasma x-ray generator utilizing a brass-target triode. We used it to carry out preliminary experiments for the superposition of K-series characteristic x-rays in weakly ionized plasma and for producing their higher harmonics.

2. Generator

Figure 1 shows a block diagram of a high-intensity plasma flash x-ray generator. This generator consists of the following essential components: a high-voltage power supply, a high-voltage condenser with a capacity of 200 nF, a turbomolecular pump, a krytron pulse generator as a trigger device, and a flash x-ray tube. A low-impedance transmission line is employed in order to increase the maximum tube current in the generator. The high-voltage main condenser is charged to 50 kV by the power supply, and electric charges in the condenser are discharged to the tube after triggering the cathode electrode with the trigger device. The plasma flash x-rays are then produced.

The x-ray tube is a demountable cold-cathode triode that is connected to the turbomolecular pump with a pressure of approximately 1

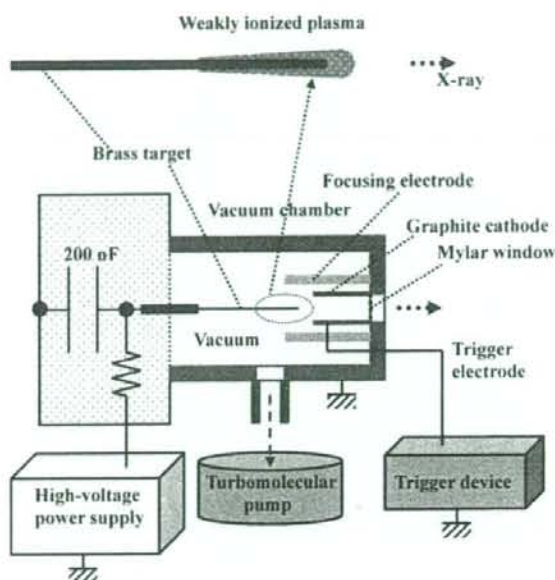


Fig. 1 Block diagram including the electric circuit of the high-intensity plasma flash x-ray generator.

mPa. This tube consists of the following major parts: a pipe-shaped graphite cathode with a bore diameter of 10.0 mm, a trigger electrode made from copper wire, a brass focusing electrode, a stainless steel vacuum chamber, a nylon insulator, a polyethylene terephthalate (Mylar) x-ray window 0.25 mm in thickness, and a 4.0-mm-diameter rod brass target containing 65% copper and 35% zinc by weight. The distance between the target and cathode is 20 mm, and the trigger electrode is set in the cathode electrode. As electron beams from the cathode are roughly converged to the target by the focusing electrode, evaporation leads to the formation of a weakly ionized linear plasma, consisting of metal ions and electrons, around the target.

3. Characteristics

3.1. Tube voltage and current

Tube voltage and current were measured by

a high-voltage divider with an input impedance of $1\text{ G}\Omega$ and a current transformer, respectively. Figure 2 shows the time relation between the tube voltage and current. The tube voltage and current roughly displayed damped oscillations at the indicated charging voltages. When the charging voltage was increased, both the maximum tube voltage and current increased. At a charging voltage of 50 kV, the maximum tube voltage was almost equal to the charging voltage of the main condenser, and the maximum tube current was 15 kA.

3.2. X-ray output

X-ray output pulse was detected using a combination of a plastic scintillator and a photomultiplier (Fig. 3). The x-ray pulse height substantially increased with corresponding increases in the charging voltage. The x-ray pulse widths were 700 ns, and the time-integrated x-ray intensity measured by a thermolu-

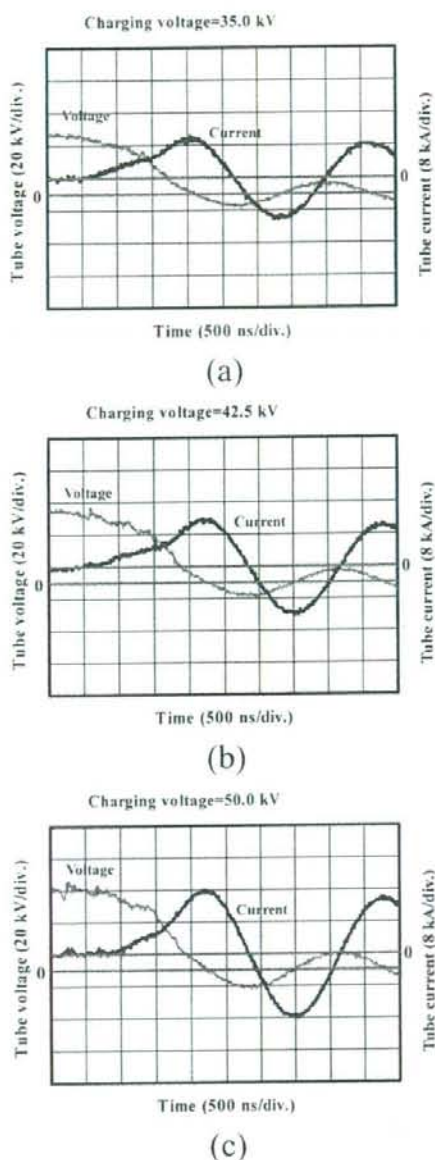


Fig. 2 Tube voltages and currents with a charging voltage of (a) 35.0 kV, (b) 42.5 kV and (c) 50.0 kV.

minescence dosimeter (Kyokko TLD Reader 1500 with MSO-S elements without energy compensation) had a value of 1.2 mGy at 1.0 m per pulse with a charging voltage of 50 kV.

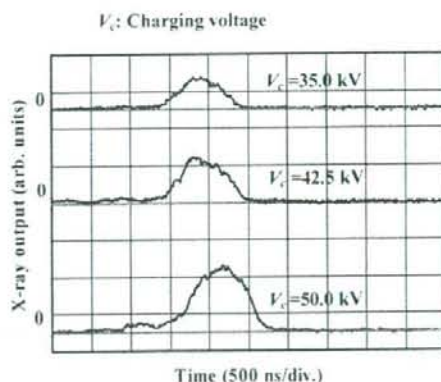


Fig. 3 X-ray outputs at the indicated conditions.

3.3. X-ray source

In order to roughly observe images of the plasma x-ray source in the detector plane, we employed a 100- μ m-diameter pinhole camera and an x-ray film (Polaroid XR-7) (Fig. 4). When the charging voltage was increased, the plasma x-ray source grew, and both spot dimension and intensity increased. Because the x-ray intensity was the highest at the center of the target, both the dimension and intensity decreased according to both increases in the thickness of the filter for absorbing x-rays and decreases in the pinhole diameter. The minimum dimension was equal to the target diameter of 3.0 mm.

3.4. X-ray spectra

X-ray spectra from the plasma source were measured by a transmission-type spectrometer with a lithium fluoride curved crystal 0.5 mm in thickness. The spectra were taken by a computed radiography (CR) system (Konica Regius 150)²⁰⁾ with a wide dynamic range, and relative x-ray intensity was calculated from Dicom digital data.

Figure 5 shows measured spectra from

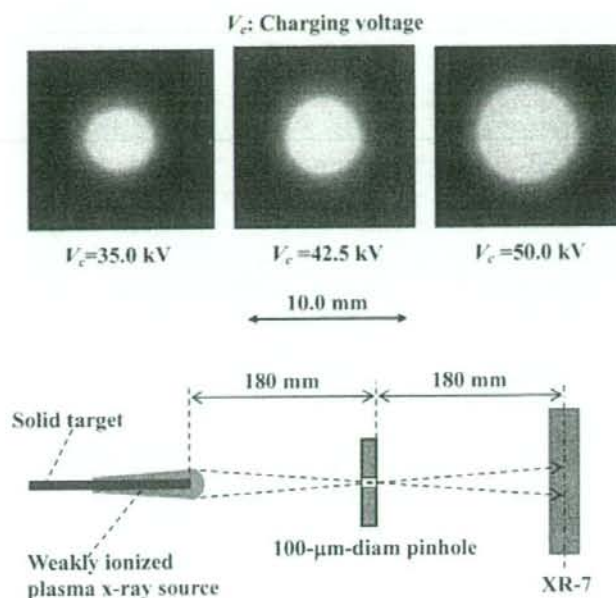


Fig. 4 Images of plasma x-ray source, and the experimental setup for imaging the source.

weakly ionized metal plasma. We observed clean copper $K\alpha$ copper $K\beta$ and zinc $K\alpha$ lines. However, zinc $K\beta$ and bremsstrahlung rays were barely detected. The characteristic x-ray intensity substantially increased with increases in the charging voltage. In the high-photon-energy region, we confirmed the irradiation of the second unsharp harmonics of the fundamental K-lines of copper and zinc elements. The x-ray intensities of the harmonics increased with increases in the charging voltage, and the harmonic bremsstrahlung rays survived due to the x-ray resonance in the plasma.

4. Radiography

Plasma radiography was carried out with the CR system, and the charging voltage and the distance were 50 kV and 1.2 m, respectively.

Figure 6 shows radiograms of tungsten

wires coiled around a pipe made of polymethyl methacrylate. Although the image contrast increased with increases in the wire diameter, a 50- μm -diameter wire could be observed. Next, the image of aluminum grains falling into a polypropylene beaker from a glass test tube is shown in Fig. 7. Because the x-ray duration was approximately 700 ns, the stop-motion image of grains could be obtained. Figure 8 shows angiograms of a rabbit thigh obtained utilizing iodine-based microspheres of 15 μm in diameter. In the angiography, a 100- μm -diam tungsten wire was used for determining the diameter of blood vessels, and fine blood vessels of about 100 μm were clearly visible.

5. Conclusions and Outlook

Regarding the spectrum measurement, although we confirmed clean copper $K\alpha$ copper $K\beta$ and zinc $K\alpha$ lines, zinc $K\beta$ lines were

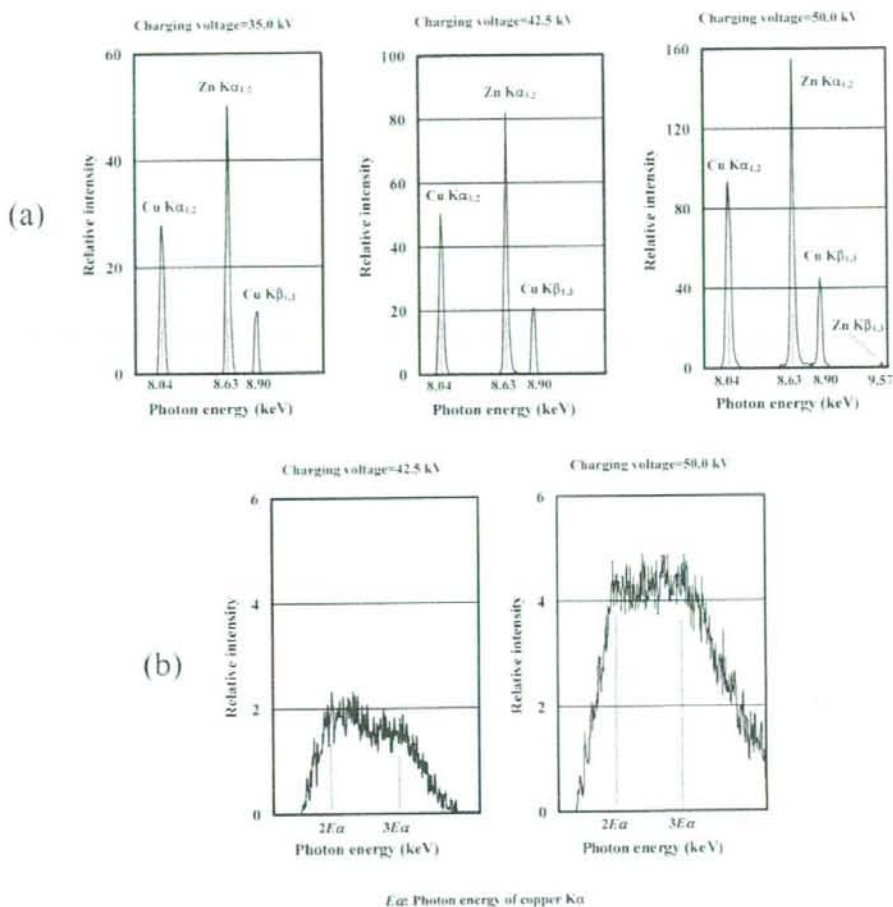


Fig. 5 X-ray spectra at the indicated conditions. (a) Characteristic x-rays and (b) bremsstrahlung x-rays.

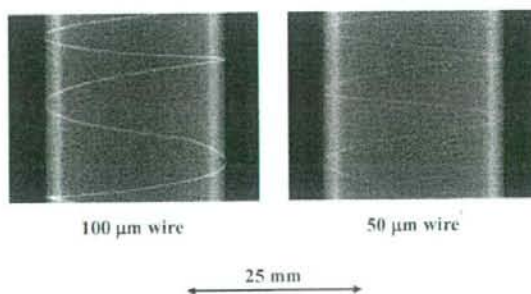


Fig. 6 Radiograms of tungsten wires coiled around pipes made of polymethyl methacrylate.

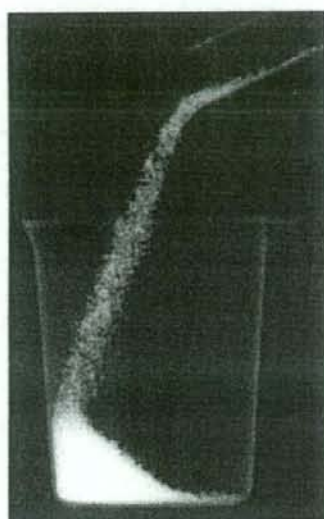


Fig. 7 Radiogram of aluminum grains falling from a glass test tube.

barely observed. Because the weakly ionized zinc plasma transmitted zinc $K\beta$ lines easily, these lines were absorbed by the copper plasma. In the high-photon-energy region, we could not observe clean higher harmonics, and bremsstrahlung x-rays with photon energies of approximately $2E_{\alpha}$ were left in cases where high charging voltages beyond 40 kV were applied.

From the experimental results, we saw that the x-ray spectra with photon energies just beyond the copper K-edge were absorbed effectively by the copper plasma, and zinc $K\beta$ rays would be useful for producing copper fluorescent rays. Next, we would like to obtain results using a capillary-type target for forming weakly ionized linear plasma.

In this research, we obtained sufficient characteristic x-ray intensity per pulse for CR radiography, and the generator produced number

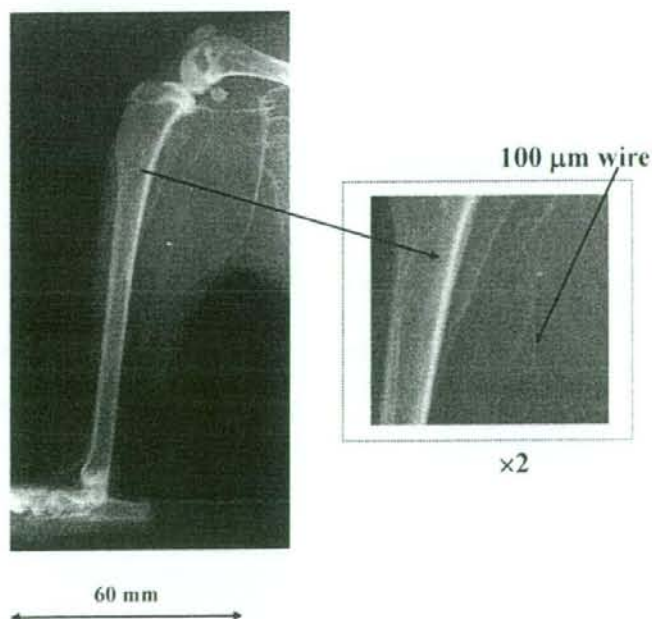


Fig. 8 Angiogram of a rabbit thigh.

of characteristic photons was approximately 1×10^8 photons \cdot cm⁻² at 1.0 m per pulse. In addition, since the photon energy of characteristic x-rays can be controlled by changing target elements, various quasi-monochromatic high-speed radiographies, such as flash energy subtraction radiography using a metal filter and wide-photon-energy radiography, will be possible.

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Magnification K-Edge Angiography Utilizing 100- μm -Focus Tungsten Tube and Gadolinium-Based Contrast Media

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A microfocus X-ray tube is useful for performing magnification digital radiography. The 100- μm -focus X-ray generator consists of a main controller, for regulating tube voltage and current, and a tube unit with a high-voltage circuit and a fixed anode X-ray tube. The maximum tube voltage, current, and electric power were 106 kV, 0.5 mA, and 50 W, respectively. Using a 100- μm -thick tungsten filter, the X-ray intensity was 19.8 $\mu\text{Gy/s}$ at 1.0 m from the source with a tube voltage of 100 kV and a current of 0.50 mA. Since K α rays from tungsten targets are absorbed effectively by gadolinium-based contrast media, these rays are useful for performing enhanced angiography. Magnification angiography was performed by threefold magnification imaging using a computed radiography system and the filter at a tube voltage of 100 kV. In angiography of nonliving animals, we observed fine blood vessels of approximately 100 μm with high contrasts.

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KEYWORDS: angiography, gadolinium-based contrast media, 100- μm -focus X-ray tube, tungsten K-rays, magnification radiography

1. Introduction

Energy-selective imaging has been performed using monochromatic X-rays, and extremely clean monochromatic parallel beams have been formed using synchrotrons in conjunction with single silicon crystals. In particular, because these beams with photon energies just beyond the iodine K-absorption edge 33.2 keV are absorbed effectively by iodine-based contrast media, enhanced K-edge angiography^{1–3} has been carried out.

To produce extremely high-dose-rate pulse X-rays, various flash X-ray generators have been developed.^{4,5} In particular, weakly ionized plasma flash X-ray generators^{6–8} are useful for producing intense clean K-series characteristic X-rays because bremsstrahlung X-rays are absorbed effectively by the plasma. However, it is difficult to generate steady-state X-rays because conventional flash tubes utilize cold cathodes.

Recently, a steady-state X-ray generator utilizing a cerium-target tube^{9,10} has been developed and has been applied to enhanced K-edge angiography achieved with cerium K α rays and iodine-based contrast media, since the K α rays (34.6 keV) are absorbed effectively by iodine-based media. When a computed radiography (CR) system (Konica Minolta Regius 150)¹¹ with a sampling pitch of 87.5 μm is used for angiography, blood vessels of 100 μm in diameter are observed. However, the spatial resolution measured using a lead test chart is approximately 200 μm .

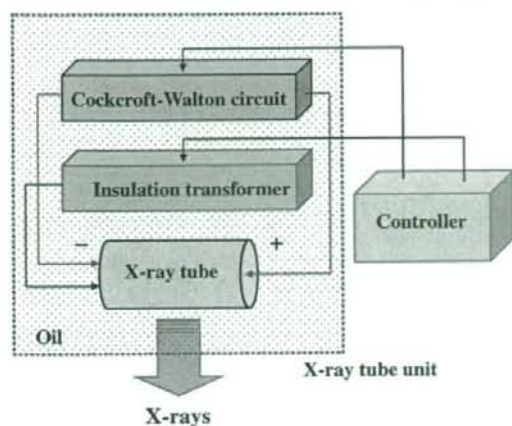
K α rays from ytterbium, tantalum, and tungsten targets are absorbed effectively by gadolinium-based contrast media, and these flash K α rays are useful for performing high-speed enhanced K-edge angiography^{12,13} with X-ray durations less than 1 μs . As compared with iodine K-edge

angiography, the absorbed dose can be decreased easily in cases in which gadolinium media are employed. Although X-ray angiographic techniques utilizing gadolinium media have been performed,^{14,15} gadolinium K-edge angiography is useful for decreasing the absorbed dose using effective X-ray spectra.

The maximum densities of the gadolinium and iodine elements in liquid contrast media are 150 and 370 mg/mL, respectively, and it is difficult to improve the image contrast of blood vessels using gadolinium media. Lately, we have developed a highly sensitive X-ray computed tomography (CT) system¹⁶ utilizing a cadmium telluride detector and have performed iodine K-edge CT angiography (CTA) with an iodine density of 4 mg/mL using optimum X-rays. To decrease the absorbed dose for patients in angiographic techniques including CTA,^{17,18} we are very interested in gadolinium K-edge angiography achieved using steady-state tungsten K α rays and low-density contrast media, and unnecessary bremsstrahlung X-rays should be absorbed using a metal filter in enhanced angiography.

Microfocus X-ray tubes have been applied to phase contrast radiography utilizing magnification imaging.¹⁹ Furthermore, the magnification is useful for improving spatial resolution^{20,21} in digital radiography, and the image quality is improved by decreasing scattering X-ray intensity from an object. Because most microfocus X-ray tubes employ tungsten targets, gadolinium K-edge angiography can be performed using tungsten K α rays.

In the present research, we employed a 100- μm -focus X-ray generator to perform a preliminary study on enhanced magnification angiography utilizing gadolinium media by decreasing bremsstrahlung X-ray intensity.

Fig. 1. Block diagram of 100- μm -focus X-ray generator.

2. X-ray Generator

Figure 1 shows the block diagram of a microfocuss X-ray generator used in this experiment, and the generator consists of a main controller and an X-ray tube unit with a 100- μm -focus tube. Tube voltage, current, and exposure time can be controlled using the controller. The high-voltage line employs the Cockcroft-Walton circuit in order to decrease the dimensions of the tube unit. In the X-ray tube unit, the positive and negative high voltages are applied to the anode and cathode electrodes, respectively. The filament heating current is supplied by an AC power supply in the controller in conjunction with an insulation transformer, which is used for isolation from the high voltage from the Cockcroft-Walton circuit. In this experiment, the tube voltage applied was from 80 to 106 kV, and the tube current was regulated to be within 0.50 mA (maximum current) by filament temperature. Exposure time is controlled in order to obtain optimum X-ray intensity, and K-series characteristic X-rays are selected out using a 100- μm -thick tungsten filter.

To design a tungsten K-edge filter, the transmittance T of K α rays is given by

$$T = I/I_0 = \exp[-\mu(\varepsilon) \cdot \xi] \approx 0.5, \quad (1)$$

where I is the transmission K α intensity, I_0 is the incident K α intensity, $\mu(\varepsilon)$ is the linear absorption coefficient at an average photon energy ε of K α rays, and ξ is the filter thickness (half-value layer). Thus, ξ is calculated as approximately 100 μm and is written as

$$\xi = \log 2 / [\mu_w(\varepsilon) \cdot \rho] \approx 100 \mu\text{m}, \quad (2)$$

where $\mu_w(\varepsilon)$ is the mass absorption coefficient of tungsten at an ε , and ρ is the tungsten density.

3. Results and Discussion

3.1 X-ray intensity

X-ray intensity was measured by a Victoreen 660 ionization chamber at 1.0 m from the X-ray source. At a constant tube current of 0.50 mA, X-ray intensity increased when tube voltage was increased. At a tube voltage of 100 kV, the intensity was substantially decreased using the filter, and the intensities without filtering and with filtering were 449 and 19.8 $\mu\text{Gy/s}$, respectively (Fig. 2). Therefore, the absorbed dose in patients can be decreased using the filter for gadolinium angiography.

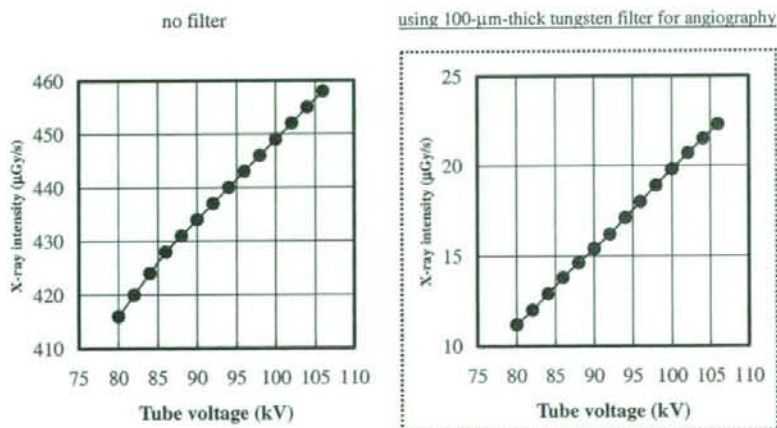
3.2 X-ray spectra

In order to measure X-ray spectra, we employed a cadmium telluride detector (Amptek XR-100T) using the filter (Fig. 3). When tube voltage was increased, the characteristic X-ray intensity increased, and both the maximum photon energy and the spectrum peak energy increased.

For gadolinium K-edge angiography, tungsten K α rays are useful, and bremsstrahlung X-rays with energies below the gadolinium K-edge decrease the image contrast. Using this filter, because bremsstrahlung rays with energies higher than 100 keV were not absorbed easily, the tube voltage for angiography was determined as 100 kV by considering the filtering effect of an object.

3.3 Magnification radiography

Magnification radiography was performed by threefold magnification imaging using the CR system at a tube voltage

Fig. 2. X-ray intensities ($\mu\text{Gy/s}$) at 1.0 m from source as function of tube voltage (kV) at tube current of 0.50 mA.

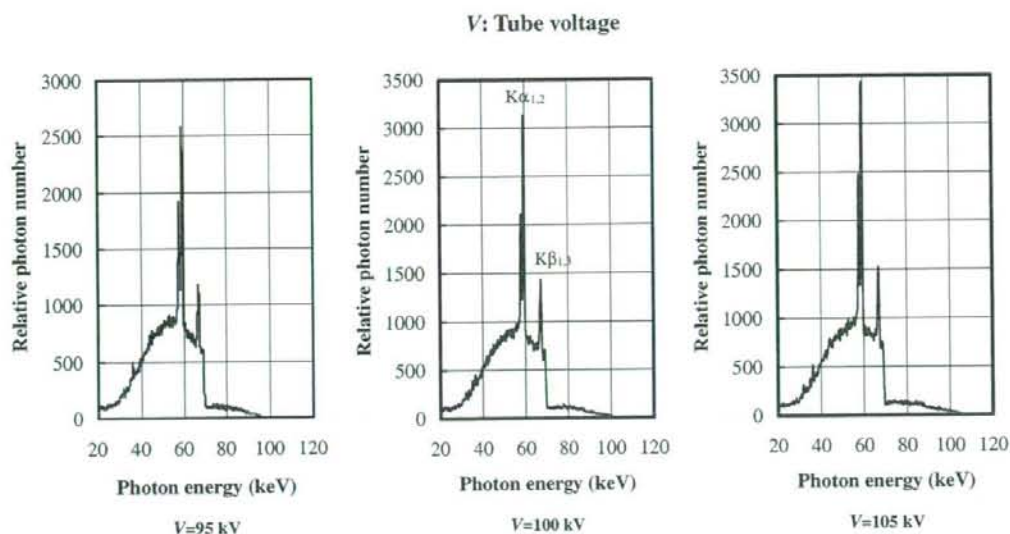


Fig. 3. X-ray spectra measured using 100- μm -thick tungsten filter with changes in tube voltage.

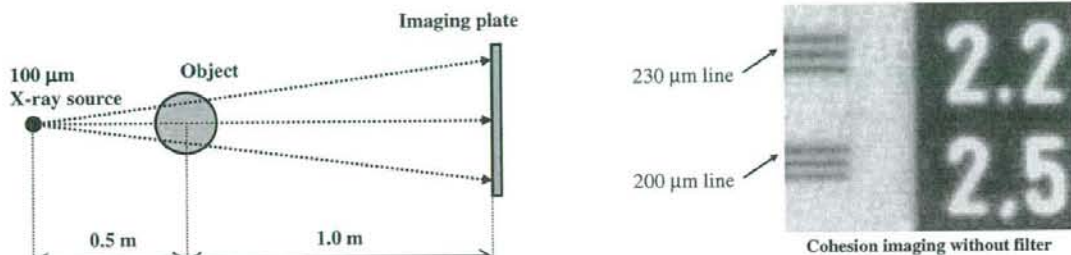


Fig. 4. Threefold magnification imaging using imaging plate.

of 100 kV, a tube current of 0.50 mA, and a distance of 1.5 m between the X-ray source and the imaging plate (Fig. 4). First, spatial resolution was measured using a lead test chart at an exposure time of 10 s. Although the spatial resolution of cohesion radiography was approximately 200 μm , 71- μm -thick lines (seven line pairs) were visible in magnification radiography (Fig. 5).

Figure 6 shows radiographs of a 50- μm -diameter tungsten wire coiled around a 30-mm-diameter rod made of poly(methyl methacrylate) (PMMA) at an exposure time of 10 s. Owing to the blurring of the image caused by a sampling pitch of 87.5 μm , the wire diameter in radiography was decreased utilizing magnification imaging. Although the tungsten wire transmitted tungsten K-rays easily, the contrast of the wire increased using the filter for absorbing bremsstrahlung rays, because low-photon-energy bremsstrahlung rays with energies below the K-edge were absorbed easily by the rod.

3.4 Enhanced magnification angiography

Figure 7 shows the mass attenuation coefficients of gadolinium at the selected energies; the coefficient curve is discontinuous at the gadolinium K-edge. The average photon energy of the tungsten $K\alpha$ lines is shown above the gadolinium K-edge. The average photon energy of tungsten

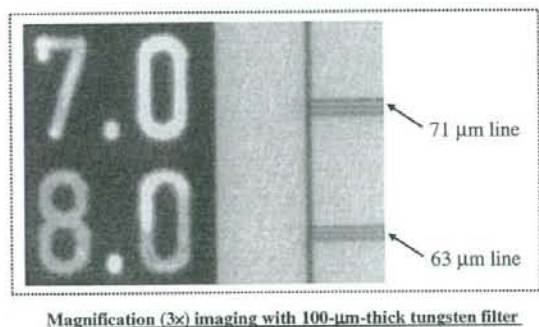


Fig. 5. Radiographs of test chart for measuring spatial resolution using filter.

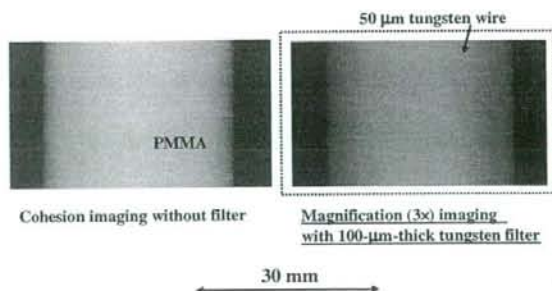


Fig. 6. Radiographs of tungsten wire coiled around PMMA rod.