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Carbon Monoxide: An Endogenous Modulator of Sinusoidal Tone in the Perfused Rat Liver

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Abstract

Heme oxygenase is a heme-oxidizing enzyme which generates biliverdin and carbon monoxide (CO). The present study was designed to elucidate whether CO endogenously produced by this enzyme serves as an active vasorelaxant in the hepatic microcirculation. Microvasculature of the isolated perfused rat liver was visualized by dual-color digital microfluorography to alternately monitor sinusoidal lining and fat-storing Ito cells. In the control liver, the CO flux in the venous effluent ranged at 0.7 nmol/min per gram of liver. Administration of a heme oxygenase inhibitor zinc protoporphyrin IX (1 μM) eliminated the baseline CO generation, and the vascular resistance exhibited a 30% elevation concurrent with discrete patterns of constriction in sinusoids and reduction of the sinusoidal perfusion velocity. The major sites of the constriction corresponded to local sinusoidal segments colocalized with Ito cells which were identified by imaging their vitamin A autofluorescence. The increase in the vascular resistance and sinusoidal constriction were attenuated significantly by adding CO (1 μM) or a cGMP analogue 8-bromo-cGMP (1 μM) in the perfusate. From these findings, we propose that CO can function as an endogenous modulator of hepatic sinusoidal perfusion through a relaxing mechanism involving Ito cells. (J. Clin. Invest. 1995, 96:2431–2437.) Key words: heme oxygenase • nitric oxide • cGMP • Ito cells • microcirculation

Introduction

Carbon monoxide, a product of heme oxygenase (EC 1.14.99.3) (1, 2) is thought to up-regulate cGMP via activation of guanylate cyclase, and thereby shares several biological actions with nitric oxide (NO), such as smooth muscle relaxation (3) or inhibition of platelet aggregation (4). There is, however, little experimental evidence suggesting that CO generated endogenously by heme oxygenase activity participates in modulation of cell function under physiologic conditions, except for recent observations which provide evidence for the contribution of CO as a neural messenger in the brain (5, 6). As to its vascular actions, no definite evidence has been provided to establish CO as an endogenous modulator of vascular tone.

Liver is, like the brain and spleen, one of the most abundant sources of the heme oxygenase among organs (1). We have recently reported that zinc protoporphyrin IX (ZnPp), a potent inhibitor of heme oxygenase, elicited an increase in the perfusion pressure in the rat liver, suggesting that constitutive levels of CO are necessary to maintain its low vascular resistance (7). The present study was undertaken to explore the microvascular mechanisms through which CO generated by heme oxygenase modulates the vascular resistance. We have examined the role of Ito cells, liver-specific microvascular pericytes, as a putative CO-sensitive machinery to maintain sinusoidal tone in a relaxing state.

Methods

Animal preparation. Male Wistar rats (300–320 grams) allowed free access to laboratory chow and tap water were fasted 24 h before experiments. After the animals were anesthetized with pentobarbital sodium (40 mg/kg intraperitoneally), the livers were perfused with hemoglobin- and albumin-free Krebs-Henseleit bicarbonate-buffered solution (pH 7.4, 37°C) gassed with carbogen (95% O2, 5% CO2) as described previously (8, 9). The common bile duct was cannulated by a PE-10 polyethylene catheter to monitor the bile output every 5 min after the start of experiments. The perfusate was pumped through the liver with a peristaltic pump at a constant flow rate of 3.4 ml/min per gram of liver weight in a single pass mode under monitoring the inlet perfusion pressure and the portal and venous oxygen concentrations using oxygen microelectrodes (PO2-100; Inter Medical Co., Tokyo, Japan) (10).

Digital microfluorography of hepatic microcirculation. The isolated perfused liver was placed on a plastic stage in which the bottom of the liver was supported by an elastic nylon net to minimize local damages of preparation. The upper surface of the left lobe was covered with a spheroid-shaped cover glass and was observed through an orthostatic fluorescence microscope (TMD; Nikon, Tokyo, Japan) equipped with a 16 × quartz objective lens and an intensified charged couple device camera (TEC-470, Optronics Engineering, Goleta, CA), a gray density filter (ND16; Nikon), and a programing electric shutter (Shutter Controller; Sankei, Inc., Tokyo, Japan), which allowed us to limit unnecessary exposure to epi-illumination by a xenon lamp (100 W). To determine the diameter of hepatic sinusoids and to confirm the state of microvascular perfusion, 2.0% solution of FITC-BSA was injected into the perfusion circuit at a rate of 250 μl/s for 4 s. The intralobular movement of the fluorescence tracer was visualized by epi-illuminating at 480 nm with passage through a 530-nm emission filter (BAS520: Nikon), and was recorded continuously to a digital videorecorder (S-VHS-HQ; Victor-JVC, Tokyo, Japan). By analyzing the replayed videotapes using an off-line digital videodensitometer (Capiflow; IM-Capi-
flow, Kista, Sweden) (9, 10), the intralobular perfusion velocity (Vp-c) was estimated as follows:

\[ V_{p-c} = \frac{L_{oc}/ \Delta T_{p-c}}{s}, \]

where \(L_{oc}\) is the distance between the centers of the optical measurement windows (50 \( \times \) 50 \( \mu \)m) in periportal and pericentral regions, and \(\Delta T_{p-c}\) is the time delay of the peak intensity between the periportal and pericentral regions which was estimated by the videodensitometer. Since the \(V_{p-c}\) values varied among different livers or lobules, ranging from 300 \( \mu \)m/s to 800 \( \mu \)m/s, the data were divided by the control velocity at time 0, and expressed as the normalized velocity as compared with the control (%Vp-c).

The perfusion rate or the nitrogen correlation between Ito cells and sinusoidal constriction. Intralobular distribution of Ito cells was identified by our previous method on the basis of visualization of vitamin A autofluorescence (10, 11). Briefly, the liver surface was epiluminatized at 360 nm by using a narrow band pass filter (BA400; Nikon), and the autofluorescence at 450 nm was visualized by the intensified charged couple device camera. The camera controller was able to average eight videos divided into six subgroups (control, 1 \( \mu \)M ZnPP, 1 \( \mu \)M CO + 1 \( \mu \)M ZnPP, 1 \( \mu \)M 8-bromo-cGMP [8BrcGMP] + 1 \( \mu \)M ZnPP, 1 \( \mu \)M CO alone, 1 \( \mu \)M 8BrcGMP alone; \( n = 8 \) for each group). 20 min after completing the preparation of the perfused liver, FITC-BSA-associate microangiographs and the corresponding vitamin A autofluorescences were recorded in the same microscopic field as control microangiographs. ZnPP was added to the perfusate reservoir to give a final concentration at CO- or 8BrcGMP-containing Krebs-Henseleit buffer was infused into the liver 5 min before the start of the ZnPP administration. Every 10 min after the start of the ZnPP administration, both FITC-BSA and vitamin A microangiographs were recorded. The total exposure time to record Ito cell images (vitamin A autofluorescence) during the entire course (30 min) of experiments was within 4 s. Under these conditions, the autofluorescence derived from Ito cells was well maintained with a minimal photobleaching phenomenon until the end of experiment. By epi-illuminating at 360 nm continuously at the end of each experiment, the autofluorescence in Ito cells was totally eliminated, confirming the rapid photobleaching property of vitamin A (10, 11).

Finally, we evaluated time history of oxygen consumption and bile output to confirm the viability of each perfused liver. Our experimental results showed that neither administration of 8 \( \mu \)M CO nor its combination with 1 \( \mu \)M ZnPP did elicit a significant reduction of the baseline bile output or oxygen consumption, suggesting the absence of possible noxious effects of these reagents on the perfused liver. When the bile output was reduced to < 1.0 \( \mu \)l/min per gram liver or the oxygen consumption ranged at < 1.8 \( \mu \)mol/min per gram liver during the observation period, such an experiment was discarded to avoid data from low quality preparations which mainly resulted from local perfusion failure.

In another set of experiments, the \( \alpha_{1}\)-adrenergic receptor agonist phenylephrine (PE) was used as the positive control, since adrenergic nerve endings (12) and catecholamine-mediated construction are localized in the portal venules in the rat (13). The whole organ vascular resistance in the liver was estimated as the ratio of the pressure gradient between the portal and hepatic veins versus flow volume according to the previous method (14). In some experiments, heterogeneity of the lobular perfusion was macroscopically evaluated to check the difference in the vasocostructive mechanism between the ZnPP- and the PE-treated livers. To that effect, 2 ml of the trypan blue solution (3%) was infused at the rate of 400 \( \mu \)l/min at the end of experiments (15). Other reagents used in the present study such as N\(^{\text{N}}\)-nitro-L-arginine methyl ester (L-NAME) and aminoguanidine (AG) were purchased from Sigma Chemical Co. (St. Louis, MO).

**Measurements of CO in the effluent.** Concentrations of CO in the effluent of the perfused liver were determined by measuring spectrophotometrically the formation of ferrous-CO complex of myoglobin (Mb) (horse heart; Sigma Chemical Co.). Approximately 20 ml of the effluent buffer was collected from the hepatic venous outlet using a gas-tight syringe 20 min after the start of the control perfusion. The glass cuvette with a gas-tight rubber cap was prepared for spectrophotometry of the samples. A 27-gauge long needle was inserted down to the bottom of the cuvette. After a 30-gauge short needle was inserted for air evacuation, the cuvette was filled with the sample by injecting an aliquot of the effluent (about 4.0 ml) through the long needle, and the air bubbles were carefully removed by purging the cuvette with injection of the rest of the effluent sample. The ferric Mb, which was oxidized by pretreatment with potassium ferricyanide, was purified using carboxymethyl cellulose chromatography (pH 7.2), and was stored at ~80°C until the day of experiments. 200 \( \mu \)l of the Mb solution (60 \( \mu \)M) was injected into the cuvette (3.0 \( \mu \)M at a final concentration) through the long needle so that the whole amount of Mb was placed at the bottom of the cuvette. After eliminating the needle and stirring the sample, the absorption spectrum was measured by a recording spectrophotometer (400–600 nm, MPS-2000; Shimazu Inc., Kyoto, Japan) equipped with a personal computer (PC 9801VX; NEC, Tokyo, Japan) which allows us to digitally process the data. The ferrous CO complex of Mb was yielded by the addition of an excess amount of dithionite (40 \( \mu \)l of the 0.5-M solution which gives ~5 mM at a final concentration in the cuvette), and the absorption spectrum was again recorded. Finally, 40 \( \mu \)l of the CO-saturated buffer which was kept in a gas-tight syringe (~1 mm) was injected into the reaction cuvette to saturate ferrous Mb with CO, and the spectrum was recorded. Difference spectrum between the samples before and after adding dithionite was then digitally estimated. Under low concentrations of CO, Mb in the sample after adding dithionite is a mixture of ferrous and ferrous–CO forms, where the difference in absorbance observed at \( \lambda \) (\( \Delta A_{\lambda} \)) is given as below:

\[ \Delta A_{\lambda} = \left[ \left[ \text{Fe}^{2+}\text{CO} \right]_{\lambda} \text{CO} + \left[ \text{Mb} \right]_{\lambda} - \left[ \text{Fe}^{2+}\text{CO} \right]_{\lambda} \text{CO} \right] \text{trans} - \left[ \text{Mb} \right]_{\lambda} \text{trans} \]

where, \( \left[ \text{Fe}^{2+}\text{CO} \right]_{\lambda} \text{CO} \) and \( \left[ \text{Mb} \right]_{\lambda} \text{trans} \) are concentrations of ferrous–CO form, the initial concentration of Mb added in the cuvette (3.0 \( \mu \)M), molar extinction coefficients at \( \lambda \) of ferrous–CO, ferrous, and ferric forms, respectively. Since one of the isobestic point between ferrous and ferric Mb exists at 419.1 nm in which \( \epsilon_{419.1} \text{trans} = \epsilon_{419.1} \text{trans} = 0 \), the formula can be arranged as follows:

\[ \left[ \text{Fe}^{2+}\text{CO} \right] = \Delta A_{419.1}(1/\epsilon_{419.1} \text{CO} + \epsilon_{419.1} \text{trans}), \]

indicating that \( \left[ \text{Fe}^{2+}\text{CO} \right] \) is a linear function of \( \Delta A_{419.1} \). The values of \( \epsilon_{419.1} \text{CO} \) and \( \epsilon_{419.1} \text{trans} \) were determined by examining the absorption spectrum of the same concentration of ferrous Mb in the presence and absence of saturated amount of CO, respectively. To obtain the CO-saturated ferrous Mb solution, 40 \( \mu \)l of the CO-saturated buffer (>1 mM) was injected into the cuvette to give a final concentration greater than 10 \( \mu \)M. Since the dissociation constant of ferrous-CO form of Mb is 34.5

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nM (16) at room temperature (pH 7.4), ~98% of the total amounts of CO in the Mb-containing sample mixture binds to ferrous Mb, and only ~2% is present as a free form, when CO concentrations in the samples range at ~3.0 μM. We therefore determined the concentration of CO as that of ferrous-CO complex of Mb in samples.

Primary culture of Ito cells and cGMP measurements. Isolation of Ito cells was carried out according to the previous method by Ito et al. (17). Briefly, livers of male Wistar rats (400–450 grams) fed ad lib. on a stock diet, were treated with collagenase perfusion, and the nonparenchymal cells were pelleted by centrifugation (450 g, 4°C for 10 min). An Ito cell–enriched fraction was obtained by the use of centrifugation with a triple-layered (9, 11, and 17%) Nycodenz cushion (1,400 g, 4°C for 20 min). After plating the cells, the culture medium was changed every other day. Ito cells in 5-d culture (5 × 10^5 cells) inoculated onto the 35-mm collagen-coated culture dishes were used to determine the contents of cGMP. The cultured cells were washed twice with the buffer (137 mM NaCl, 2.7 mM KCl, 8.1 mM NaHPO4, 2H2O, 1.5 mM KH2PO4, 0.5 mM MgCl2, and 0.9 mM CaCl2·2H2O, pH 7.4). After incubation with 1 ml of the same buffer containing 0.5 mM 3-isobutyl-1-methylxanthine, an inhibitor of phosphodiesterase, for 10 min, the cells were treated with desired concentrations of CO for 30 min at 37°C. CO concentrations in the culture medium after completing the reaction were determined by detecting the ferrous–CO complex of Mb as was described earlier in this study for the measurements of CO in the effluent from the perfused liver. To correct diffusion of CO in the medium to the atmosphere during the 30-min incubation time, the decline of CO concentration in the culture dish were estimated as a function of time (every 10 min after the start of incubation). The mean value for four measurements (0, 10, 20, and 30 min) of CO concentrations was then defined as the actual CO concentration in the corresponding system. The reaction was terminated by adding 1 mM HCl and 95% ethanol. The cGMP contents in the extracts were measured by an enzyme-linked immunoassay using assay kits (RPN225; Amersham Corp., Arlington Heights, IL). Differences in the data among the groups were analyzed by one-way ANOVA combined with Scheffe-type multiple comparison test. A value of P < 0.05 was considered significant.

Results

ZnPP-induced increase in the hepatic vascular resistance. Fig. 1 illustrates the effects of ZnPP on the whole organ vascular resistance and the concentration of CO in the effluent of the perfused liver. In the control liver, the vascular resistance was maintained in a steady state during the whole course of observation, and the concentration of CO in the effluent ranged at 0.23±0.09 μM. Accordingly, the actual flux of CO in the effluent ranged approximately at 0.7 nmol/min per gram of liver. In the liver treated with 1 μM ZnPP, the resistance exhibited a time-dependent increase and reached ~30% higher levels than that of the control at 30 min. Under these circumstances, the CO concentration in the effluent decreased to an undetectable level. The same concentration of hematin, however, did not alter the perfusion resistance (0.25±0.05 cm H2O/min per ml at 30 min, n = 6). Pretreatment with the perfusate containing 1 μM CO significantly attenuated the ZnPP-induced elevation of the vascular resistance by ~80%. Furthermore, the CO concentration in the effluent ranged at 0.62±0.08 μM (n = 8) which is high enough to recover the CO levels in the control effluent. Since this concentration of CO alone did not alter the vascular resistance in the control (data not shown), the effects of exogenous CO supplement seems to result from its modulatory action on the ZnPP-elicted vascular response. Pretreatment with 1 μM 8Br-cGMP exhibited ~60% inhibition of the ZnPP-induced changes in the resistance, while the CO concentrations in the effluent ranged at an undetectable level.

Since NO is known to up-regulate cGMP, we have examined whether the hepatic vascular resistance might be elevated by NO synthase inhibitors such as L-NAME and AG, which block constitutive and inducible forms of the enzyme, respectively. Neither L-NAME nor AG at 1 mM altered the vascular resistance, suggesting a minor role of endogenous NO production in modulation of the vascular resistance in the steady state, if any.

Inverse correlation between the vascular resistance and CO concentrations in the hepatic venous efflux. The finding that CO concentration in the effluent varied among the individual livers tempted us to examine the relationship between the vascular resistance and CO levels in the effluent in each experiment. Fig. 2 illustrates the correlation between the CO concentrations in the effluent and the whole organ vascular resistance, showing a significant inverse correlation between the level of CO in the effluent and the vascular resistance (P < 0.05, r^2 = 0.71).

ZnPP-induced sinusoidal constriction colocalized with Ito cells. The intrahepatic compartment responsible for the ZnPP-induced increase in the vascular resistance seems to be in microvessels according to the digital microfluorography using FITC-BSA. The early event in ZnPP-treated hepatic microvasculature (10 min after the start of the ZnPP administration) is a discrete pattern of sinusoidal constriction (Fig. 3, arrows 1–9). At the same time, the sinusoidal perfusion velocity exhibited a slight increase and then gradually decreased in a time-dependent manner, showing a 25% reduction at 30 min after the start of ZnPP administration (Fig. 4). Pretreatment with CO (1 μM) or with 8Br-cGMP (1 μM) significantly attenuated the ZnPP-
elicitated reduction of the sinusoidal perfusion velocity. During the observation period, no significant changes in the inner diameter of central venules were observed, suggesting little possibility that postsinusoidal elements achieve major roles in the contractile mechanisms of this model.

We investigated a topographic correlation between the sites of sinusoidal constriction and the distribution of Ito cells, inasmuch as the regional constricting sites seem to correspond to the localization of Ito cells (Fig. 3, C and D). Fig. 5 illustrates ZnPP-induced changes in the diameter measured at local sinusoidal segments colocalized with Ito cells and at those without Ito cells. ZnPP administration (1 μM) induced an ~20% decrease in the diameter, preferentially at the local sinusoidal segments involving Ito cell-associated vitamin A autofluorescence. The ZnPP-induced sinusoidal constriction was abolished by a continuous infusion of CO (1 μM), and was partly inhibited by pretreatment with 8Br-cGMP (1 μM).

To differentiate the direct vasoactive effects of ZnPP at the sinusoidal level from its effects at the level of portal venules, we investigated the effects of PE on sinusoidal hemodynamics in situ as well as on the whole organ perfusion. Administration of PE (500 nM) induced a significant increase in the vascular resistance (0.35±0.09 cmH2O/min per ml, n = 6) which was equivalent to that elicited by 1 μM ZnPP (Fig. 1). However, the sinusoidal response exhibited quite different pictures between PE- and ZnPP-treated livers. Namely, as far as the perfused lobules were observed by FITC-BSA-assisted digital microfluorangiography, there were no sinusoidal narrowing patterns which were typically observed in the ZnPP-treated liver (Fig. 5). These results imply that the vascular sites responsible for the PE-induced elevation of vascular resistance may be the proximal portal veins or venules rather than sinusoids, confirming the previous observation by Zhang et al. (18). Such a difference between the two groups was observed in the state of macroscopic perfusion which was assessed by the trypan blue technique. In the ZnPP-treated liver, the intrahepatic distribution of the dye color exhibited uniform and homogeneous patterns. By contrast, the PE-treated liver was stained in patches and the fluorescence intensity exhibited a marked heterogeneity even among the perfused lobules, indicating nonuniform and heterogenous liver perfusion. Taken together, the present findings suggest that sinusoidal constriction, rather than presinusoidal constriction, contributes to the ZnPP-induced increase in the hepatic vascular resistance.

Up-regulation of cGMP in cultured Ito cells in response to CO. Colocalization of Ito cells with the regional sites of the ZnPP-elicited sinusoidal constriction and modulatory effects of CO and 8Br-cGMP lead us to hypothesize that Ito cells might involve the local vasorelaxation mechanism operated by the CO-dependent activation of guanylate cyclase. We therefore investigated the CO-elicited response of cGMP, a cyclic nucleotide which relaxes cytosome of Ito cells (17). Fig. 6 illustrates the effects of CO which were applied exogenously to primary cultured Ito cells on the amounts of cGMP. The application of the CO-containing buffer induced a dose-dependent increase in cGMP contents in Ito cells. Since the elevation of cGMP has been considered a relaxing mechanism for Ito cells (17), the present results suggest that these cells could be relaxed in response to extracellularly applied CO. The minimum concentration of CO which evoked significant increase in cGMP ranged at 20 μM, which was ~100-fold higher than that detected in the effluent of the control perfused liver. CO at 50 μM elicited a greater increase in cGMP that was almost equivalent to that elicited by 100 μM sodium nitroprusside. On the other hand, ZnPP at 1 μM did not alter the levels of cGMP in cultured Ito cells, suggesting that an inhibitory action of this reagent on guanylate cyclase (19) in Ito cells may not contribute to the sinusoidal constriction under the current experimental conditions.

Discussion

The present study together with our previous communication (7) has first provided substantial evidence that constitutive levels of CO generation play a crucial role in maintenance of hepatic sinusoidal tone in a relaxing state. In agreement with the previous observations (1, 20), the major source of CO in the liver is heme oxygenase, since CO detected in the effluent of the liver were abolished by the administration of 1 μM ZnPP, a potent inhibitor of the enzyme. In the control liver, parenchymal cells are considered the major cellular compartment of heme oxygenase reaction which yields >85% of the total flux of CO and biliverdin in vivo (21). According to the current results obtained from the perfused liver preparation, the output of CO released from the whole liver into circulatory system ranged approximately at 2.2 nmol/min per 100 gram of rat weight, assuming that the liver weight ranges at 2.8% of the rat body weight (22). This estimation matches roughly the rate of spontaneous generation of bilirubin, an end product of heme degradation pathway, in fresh isolated hepatocytes (3.0 nmol/min per 100 gram of rat weight), based on the assumption that rat liver contains an average of 5 × 10⁸ parenchymal cells/100.
grams of body weight as previously described (23). Taking into account these findings, our results suggest that CO constitutively released from hepatic parenchyma may influence local vascular resistance serving as a tissue-derived vasorelaxant in the liver.

The local vascular sites responsible for the ZnPP-induced elevation of vascular resistance seems to be sinusoidal microvessels rather than the proximal portal vessels with large sizes to which PE exerts vasoconstriction and a resultant heterogeneous patterns of macroscopic vascular perfusion (18). It should be noted that ZnPP administration evoked discrete patterns of sinusoidal constriction, the sites of which corresponded to the distribution of Ito cells, the liver-specific microvascular pericytes (24). Since the ZnPP-induced elevation of the vascular resistance was attenuated by supplement with CO or with 8Br-cGMP, we hypothesized that Ito cells may be responsible for lowering the sinusoidal resistance via the vasorelaxing mechanism involving CO-dependent activation of guanylate cyclase. There are several lines of evidence in the present study which support the hypothesis: exogenously administered cGMP partially attenuated the ZnPP-induced elevation of the vascular resistance. In addition, exogenous CO application at millimolar levels significantly up-regulated cGMP in cultured Ito cells. Furthermore, attenuation of the ZnPP-elicited regional sinusoidal constriction concomitant with the elevation of vascular resistance by 8Br-cGMP occurred without increasing the CO concentration in the efflux solution. There results suggest that CO endogenously generated by heme oxygenase may activate guanylate cyclase in Ito cells and thereby maintain sinusoids in a dilated state.

Of interest is that endogenous NO production is unlikely to be involved in the vasorelaxing mechanism in hepatic sinusoids, inasmuch as neither constitutive nor inducible NO synthase inhibitors did alter the vascular resistance and the sinusoidal hemodynamics. Since the reactivity of NO with heme proteins is much higher than that of CO in vitro (25), there is a possibility that the concentration of NO required for sinusoidal constriction in the local compartment (e.g., Disse space) might be

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**Figure 3.** Representative pictures of ZnPP-induced sinusoidal constriction and its topographic relationship with the localization of Ito cells. Parts A and B illustrate microfluorographs during the injection of FITC-labeled albumin before, and 10 min after the start of ZnPP administration at 1 μM, respectively. Intrahepatic localization of Ito cells can be identified by vitamin A autofluorescence, which is visualized at 450 nm emission under epi-illumination at 360 nm. Since vitamin A autofluorescence is rapidly photobleached, the exposure time to visualize Ito cells was automatically controlled to 1 s using an electric shutter. Vitamin A microfluorographs were stored in the computer and were digitally overlaid to the corresponding microvascular images. Parts C and D show such superimposed images before and after the start of ZnPP administration at 1 μM, respectively. Note that individual sinusoids (arrows 1-9) represent marked constriction, p, periportal regions; c, central venules. Bar denotes 100 μm.
extremely low as compared with that of CO. Recent observation by Bautista et al. showing the interaction between superoxide anion and NO in the perfused liver (26) could give a possible explanation for this hypothesis. Namely, under normal conditions, NO or superoxide concentrations in the liver may virtually be negligible, presumably because they counteract each other to maintain a normal balance between them. Under these circumstances, a nonradical monoanion such as CO may serve as an effective agonist for cGMP up-regulation. The actual sinusoidal perfusion in vivo is influenced by diverse factors such as neural regulation, the hepatic arterial tone, and the dynamics of intestinal circulation that are not involved in the perfused system. It is therefore likely that perturbation of endogenous NO generation may alter the sinusoidal tone in vivo, as was shown by Nishida et al. (27). Although such an intravital study should further be evaluated, the current results shed light on the possible physiologic role of tissue-derived CO as an alternative cGMP-dependent microvascular modulator in the liver.

Another hypothesis to be taken into account as a CO-dependent vasorelaxing mechanism in sinusoids is the possible involvement of the cGMP-independent pathway. Several findings in the current study support this possibility: first, the minimum CO concentration which was required to elicit a significant increase in cGMP in cultured Ito cells was at least 100-fold higher than that detected in the effluent of the control perfused liver. Secondly, the effects of 8Br-cGMP on the ZnPP-induced increase in the vascular resistance was limited only to the 60% inhibition, suggesting the presence of alternative mechanisms for CO-mediated vasorelaxation. Such an unknown action of CO which involves cGMP-independent signal transduction mechanism has recently been postulated by Kourembanas et al. (28) who reported that exogenously applied CO can modulate the expression of endothelin and growth factors independently of cGMP. Recent observation by Rich et al. has provided evidence that CO elicits an increase in potassium current and thereby hyperpolarizes the resting membrane potential in rabbit corneal epithelial cells (29). Since the hyperpolarizing effects of CO has also been reported in other types of cells such as intestinal smooth muscle cells (30), electrophysiological properties of cultured Ito cells and the effect of CO on transmembrane currents should be evaluated to reveal a novel vasorelaxing mechanism of CO. Moreover, since sinusoidal endothelial cells per se may serve as another possible contractile element which alters fluid filtration via regulation of tone in their sieve plates (31), it is also necessary to elucidate involvement of such Ito-cell-independent mechanisms for CO-mediated regulation of sinusoidal resistance. Although the CO-mediated mechanisms for sinusoidal relaxation have not fully been revealed at the present moment, our observation that CO endogenously generated by heme oxygenase reaction exerts a modulatory action on sinusoidal relaxation not only gives a clue to the mechanism by which the liver can maintain an extremely low level of vascular resistance as compared to that in other organs, but may also provide an insight into a novel physiologic role of heme oxygenase as an active vasorelaxing machinery in addition to its established role for heme degradation.

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