Copper-dependent activation of hypoxia-inducible factor (HIF)–1: implications for ceruloplasmin regulation

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Cellular oxygen partial pressure is sensed by a family of prolyl-4-hydroxylase domain (PHD) enzymes that modify hypoxia-inducible factor (HIF)α subunits. Upon hydroxylation under normoxic conditions, HIFα is bound by the von Hippel–Lindau tumor suppressor protein and targeted for proteasomal destruction. Since PHD activity is dependent on oxygen and ferrous iron, HIF-1 mediates not only oxygen- but also iron-regulated transcriptional gene expression. Here we show that copper (CuCl₂) stabilizes nuclear HIF-1α under normoxic conditions, resulting in hypoxia-response element (HRE)-dependent reporter gene expression. In vitro hydroxylation assays CuCl₂ inhibited prolyl-4-hydroxylation independently of the iron concentration. Ceruloplasmin, the main copper transport protein in the plasma and a known HIF-1 target in vitro, was also induced in vivo in the liver of hypoxic mice. Both hypoxia and CuCl₂ increased ceruloplasmin (as well as vascular endothelial growth factor [VEGF] and glucose transporter 1 [Glut-1]) mRNA levels in hepatoma cells, which was due to transcriptional induction of the ceruloplasmin gene (CP) promoter. In conclusion, our data suggest that PHD/HIF/HRE-dependent gene regulation can serve as a sensory system not only for oxygen and iron but also for copper metabolism, regulating the oxygen-, iron- and copper-binding transport proteins hemoglobin, transferrin, and ceruloplasmin, respectively. (Blood. 2005;105:4613-4619)

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Introduction

The hypoxia-inducible factor 1 (HIF-1) is an ubiquitously expressed transcriptional master regulator of many genes regulating mammalian oxygen homeostasis. Among others, the corresponding gene products are involved in erythropoiesis, iron metabolism, angiogenesis, control of blood flow, glucose uptake and glycolysis, pH regulation, and cell-cycle control. HIF-1 is a αβ heterodimer specifically recognizing the HIF-binding site within cis-regulatory hypoxia response elements. Under normoxic conditions, the von Hippel–Lindau tumor suppressor protein (pVHL) targets the HIF-1α subunit for rapid ubiquitination and proteasomal degradation. Binding of the pVHL tumor suppressor protein requires the modification of HIF-1α by prolyl-4-hydroxylation at prolines 402 and 564 of human HIF-1α. A family of 3 oxygen- and iron-dependent prolyl-4-hydroxylases called PHD1, PHD2, PHD3, or HPH3, HPH2, HPH1, respectively, has been shown to hydroxylate HIFα. A fourth member, called PH-4, regulates HIF-1α in overexpression conditions only. Thus, limited oxygen supply prevents HIFα hydroxylation and degradation. This unusual mechanism of protein regulation provides the basis for the very rapid HIF-1α response to hypoxia. In addition to protein stability, oxygen-dependent C-terminal asparagine hydroxylation of HIF-1α by factor inhibiting HIF (FIH) prevents transcriptional cofactor recruitment, thereby fine-tuning HIF-1 activity following a further decrease in oxygen availability.

Among the HIF-1 targets are the genes encoding transferrin, transferrin receptor, heme oxygenase-1, and ceruloplasmin, which coordinate regulation iron metabolism. Increased iron uptake, release from the liver, plasma transport, and uptake in the bone marrow are essential to sustain the erythropoietic function of erythropoietin, the prototype HIF-1 target. Ceruloplasmin is a multicopper plasma protein containing ferroxidase activity necessary for Fe²⁺ saturation of transferrin. Hereditary aceruloplasminemia in humans as well as targeted deletion of the ceruloplasmin gene (Cp) in mice results in iron metabolism disorders characterized by anemia, hepatic iron overload, and neurodegeneration, demonstrating a tight connection between copper and iron metabolism.

Iron deficiency has been known for more than a decade to induce erythropoietin gene expression and HIF-1α protein stabilization. Nowadays, these results are most likely explained by inactivation of the iron-dependent protein hydroxylases PHD1 to 3 and FIH. Iron deficiency also results in mRNA induction of ceruloplasmin by HIF-1–dependent promoter activation and subsequent transcriptional up-regulation of the Cp gene. These results suggest that PHDs and FIH not only function as oxygen...
sensors but also display iron-sensing properties. Whereas transition metal ions such as Co\(^{2+}\) and Ni\(^{2+}\) stabilize HIF-1\(\alpha\) by inhibiting PHD function, it is unknown whether copper salts also interfere with PHD function. In a screen for agents that modulate HIF-1 transcriptional activity, we found that Cu\(^{2+}\) can induce HIF-1–dependent reporter gene induction. Detailed analysis of this finding suggests that free Cu\(^{2+}\) can induce ceruloplasmin synthesis in a HIF-1–dependent way.

**Materials and methods**

**Cell lines and cell culture**

All cell lines were cultured in Dulbecco modified Eagle medium (high glucose) as described previously.\(^{20}\) Oxygen partial pressures in the hypoxic workstation (InVivoX2-400; Ruskin Technology, Leeds, United Kingdom) or in the incubator (Model 3319; Forma Scientific, Illkirch, France) were either 140 mmHg (20% O\(_2\) vol/vol, normoxia) or 7 mmHg (1% O\(_2\) vol/vol, hypoxia). Stable transfection with the plasmid pH3SVL led to HIF-1–dependent reporter cell lines designated HRCHO5 (derived from CHO Chinese hamster ovary cells) or HRB5 (derived from Hep3B human hepatoma cells), respectively, as described before.\(^{30,32}\) The luciferase reporter gene pH5SVL is driven by a simian virus 40 (SV40) promoter and contains a total of 6 HIF-1 DNA-binding sites derived from the transferrin gene.\(^{16}\)

**Cell proliferation/viability assays**

Cell proliferation/viability was assessed by the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) conversion assay as described.\(^{29}\) Absorbances at 570 nm were determined in a 96-well photometer and noncellular background was subtracted. Data were normalized to the MTT conversion activity of solvent-treated normoxic control cells, which were arbitrarily defined as 1.

**Reporter gene assays**

Stably transfected hypoxia reporter cell lines were seeded in triplicates into 96-well dishes at a density of 10\(^4\)/well. After incubation overnight, freshly dissolved CuCl\(_2\) or CuCl in phosphate-buffered saline (PBS) was added at the concentrations indicated and the cells were exposed to normoxic or hypoxic conditions for 24 hours. In some experiments, the kinase inhibitors LY294002, PD98059, or PD169316 (Calbiochem, VWR International, Schwalbach, Germany) dissolved in dimethyl sulfoxide (DMSO), or DMSO alone, were added in addition to 100 \(\mu\)M CuCl\(_2\). Firefly luciferase reporter gene activity was determined as described by the manufacturer (Promega, Madison, WI) using a microplate luminometer (Berthold, Hamburg, Germany). Results were normalized to the solvent-treated normoxic control values, which were arbitrarily defined as 1.

**Transient transfections**

Hep3B cells were transiently transfected with the CP promoter reporter construct pGL-Cp4774\(^{20}\) by the calcium phosphate coprecipitation method.\(^{34}\) CP enhancer constructs, either containing a wild-type or a mutant HRE, were cloned into pGL3Prom (Promega) as described previously.\(^{20}\) Semiconfluent cells on a 10-cm dish were cotransfected with 20 \(\mu\)g of the renilla luciferase control construct pRL-SV40 (Promega) for 16 hours. Cells were detached with 2 mM EDTA (ethylenediaminetetraacetic acid; pH 8.0) in PBS, distributed onto 2 24-well dishes, and allowed to recover for 8 hours. Freshly prepared CuCl\(_2\) in PBS or PBS alone was added and the cells were incubated for 24 hours under normoxic or hypoxic conditions. Luciferase activities were determined using the dual-luciferase kit (Promega) as described previously.

**Protein extraction and immunoblot analyses**

Cells were treated with CuCl\(_2\) or the iron chelator ciclopirox olamine (CPX)\(^{32}\) at the concentrations indicated for 6 or 24 hours. For stability experiments, 20 \(\mu\)g/mL cycloheximide was added following 6 hours hypoxic induction with or without CuCl\(_2\) and the cells were allowed to reoxygenate. Combined cytoplasmic and nuclear extracts of cultured cells were prepared using 0.4 M NaCl and 0.1% Nonidet P-40 (NP-40) in extraction buffer as described previously.\(^{35}\) Protein concentrations were determined by the Bradford method using bovine serum albumin (BSA) as a standard.\(^{36}\) For immunoblot analysis, cellular protein (100 \(\mu\)g or 50 \(\mu\)g in stability experiments) was electrophoresed through 5% sodium dodecyl sulfate (SDS)–polyacrylamide gels and electrotransferred onto nitrocellulose membranes (Amersham, Freiburg, Germany) by semidy blotting (BioRad, München, Germany). Membranes were stained with Ponceau S (Sigma, Buchs, Switzerland) to confirm equal protein loading and transfer. HIF-1\(\alpha\) was detected using a mouse monoclonal anti-HIF-1\(\alpha\) immunoglobulin G\(_1\) (IgG\(_1\); Transduction Laboratories, Heidelberg, Germany) followed by a goat anti-mouse secondary antibody (Santa Cruz Biotechnology, Santa Cruz, CA). Anti–\(\beta\)-actin antibodies were from Sigma. Chemiluminescence detection of horseradish peroxidase–coupled secondary antibodies was performed by incubation of the membranes with 100 mM Tris (tris(hydroxymethyl)aminomethane)–HCl (pH 8.5), 2.65 mM H\(_2\)O\(_2\), 0.45 mM luminol, and 0.625 mM coumaric acid (all purchased from Sigma) for 1 minute before exposure to X-ray films (Fuji, Düsseldorf, Germany).

**HIF-1\(\alpha\) immunofluorescence**

Immunofluorescence analysis was performed as described before.\(^{29}\) Briefly, cells were fixed with formaldehyde for 10 minutes, washed with PBS, permeabilized with Triton X-100, and rinsed again with PBS. After blocking nonspecific binding sites with 3% BSA in PBS for 30 minutes, the cells were incubated for 1 hour with mouse monoclonal anti-HIF-1\(\alpha\) IgG\(_1\) (Transduction Laboratories) diluted 1:10 in 3% BSA in PBS, followed by a fluorescein isothiocyanate–coupled secondary anti-mouse (Dako, Copenhagen, Denmark) antibody diluted 1:100 with 3% BSA in PBS. After extensive washings with PBS, the slides were mounted in mowiol and analyzed by fluorescence microscopy (Axioplan 2000, equipped with an Axiosview digital camera and Axioview software; Carl Zeiss Vision, Mannheim, Germany).

**In vitro prolyl-4-hydroxylation assay**

Prolyl-4-hydroxylation activity was determined as recently described in detail elsewhere.\(^{26}\) Briefly, a biotinylated peptide containing Pro564 of HIF-1\(\alpha\) was bound to NeutraVidin-coated 96-well plates and incubated with partially purified PHD1, PHD2, or PHD3 in the presence of 2-oxoglutarate, FeSO\(_4\), and ascorbate for 1 hour. After washing, thioredoxin-tagged pVHL in complex with elongins B and C was allowed to bind to the hydroxylated peptide, followed by detection of the thioredoxin tag by primary antithioredoxin antibodies and secondary horseradish peroxidase–coupled anti-mouse antibodies (Sigma) using the TMB (3,3′,5,5′-tetramethylbenzidine) substrate kit (Pierce, Bonn, Germany). MBP-PHD2 (amino acids 196-426) was expressed and purified by using the pMALT system according to the instructions provided by the manufacturer (New England Biolabs, Frankfurt, Germany). PHD1 and PHD3 were expressed with an N-terminal 6His-Tag in S\(\phi\)P insect cells as described elsewhere.\(^{36,37}\)

**mRNA quantification**

Exposure of mice in triplicate to 7.5% oxygen for 0, 24, 48, and 72 hours and excision of their livers was described earlier.\(^{38}\) Total RNA from frozen mouse liver tissue or cultured Hep3B cells was isolated and analyzed by Northern blotting as described previously.\(^{16}\) Hybridization probes were obtained by restriction digestion and gel isolation followed by labeling with the random-primed method.\(^{16}\) The mouse ceruloplasmin cDNA\(^{39}\) was a kind gift of J. Gitlin (St Louis, MO) and the ribosomal protein L28 cDNA was cloned from a HepG2 cDNA library.\(^{40}\) Radioactive signals were detected by phosphoimaging and quantitated using QuantityOne software (Bio-Rad). Human ceruloplasmin, vascular endothelial growth factor (VEGF), glucose transporter-1 (Glut-1) and L28 were determined by real-time polymerase chain reaction (PCR). Briefly, 2 \(\mu\)g of total RNA was reverse transcribed with Superscript III according to the manufacturer’s
We previously established a Chinese hamster ovary (CHO) cell line, called HRCHOS, which was stably transfected with an HRE-containing luciferase reporter gene containing multiple hypoxia-response elements (HREs). Apart from hypoxia, reporter activity in these cells could also be induced by cobalt and nickel salts as well as by iron chelation. Unexpectedly, we found that CuCl₂ dose-dependently induced reporter gene induction approximately 4.1-fold at 500 μM CuCl₂ for 24 hours. The cells were prepared for indirect immunofluorescence analysis as described in “Materials and methods.” Original magnification, ×630.

**Results**

**CuCl₂ can activate a hypoxia-response element-containing reporter gene**

We previously established a Chinese hamster ovary (CHO) cell line, called HRCHOS, which was stably transfected with a HIF-1α–dependent luciferase reporter gene containing multiple hypoxia-response elements (HREs). Apart from hypoxia, reporter activity in these cells could also be induced by cobalt and nickel salts as well as by iron chelation. Unexpectedly, we found that CuCl₂ dose-dependently induced reporter gene induction approximately 4.1-fold at 500 μM CuCl₂ in CHO cells (Figure 1A). The human hepatoma Hep3B cell line is known to synthesize the plasma proteins transferrin and ceruloplasmin, both of which are induced in a HIF-dependent manner upon iron depletion of these cells. We thus stably transfected Hep3B cells with the HIF-dependent reporter gene like previously used to generate the HRCHO5 cell line. When the resulting cell line (termed HRB5) was treated with various doses of CuCl₂ for 24 hours, reporter gene expression increased up to 14-fold at 125 μM CuCl₂ (Figure 1B, top graph). Hypoxia alone (24 hours of 1% O₂) induced reporter gene activity 11-fold (Figure 1B, bottom graph) and the combined treatment with 1% O₂ and 125 μM CuCl₂ resulted in a 118-fold reporter gene induction. The CuCl₂ dose-response relationship revealed a relatively narrow window of active CuCl₂ concentrations. Since we suspected CuCl₂ cytotoxicity at higher concentrations, MTT conversion assays were performed to analyze cell viability (ie, the combined decrease in cell growth and/or the increase in cell death). A dose-dependent reduction of MTT conversion activity was found in both cell lines (Figure 1A-B). At CuCl₂ concentrations that maximally induced reporter gene expression, MTT conversion was still 60% in CHO and 67% in Hep3B cells, respectively, when compared with untreated cells. Hypoxic conditions did not significantly alter MTT conversion activity in Hep3B cells. In addition to Cu²⁺, Cu²⁺ also was able to induce reporter gene activity (Figure 1C), which might be due to the disproportionation of cuprous ions to cupric ions and copper in aqueous solutions. Other metal ions tested (Al³⁺, Ca²⁺, Fe²⁺, Fe³⁺, Gd²⁺, Mg²⁺, Mn²⁺, Se⁴⁺, and Zn²⁺) did not alter reporter gene activity in this type of assay (data not shown).

**Nuclear HIF-1α protein induction by CuCl₂**

The most likely explanation for the induction of reporter gene expression by CuCl₂ is the activation of the HIF system. We therefore analyzed whether CuCl₂ treatment leads to the stabilization of the HIF-1α protein. As shown by immunoblotting (Figure 2A), CuCl₂ treatment strongly induced HIF-1α protein stability in Hep3B cells after 6 and 24 hours. Both the range of CuCl₂ concentrations and the combined effects with hypoxia are consistent with the reporter gene data, especially after 24 hours of induction. To examine whether Cu²⁺ also stabilizes HIF-1α in a nonhepatic cell line, human cervical carcinoma HeLa cells were...
treated with CuCl₂. As shown in Figure 2B, HIF-1α was also stabilized in HeLa cells, though at a considerably higher CuCl₂ following hypoxia or CuCl₂ treatment but not in normoxic control accumulation exclusively within the nucleus, excluding the nucleoli, shown by indirect immunofluorescence analysis, HIF-1α is activated in a HIF-1-dependent manner. Thus, we further analyzed ceruloplasmin regulation, a known HIF-1 target gene inducible by hypoxia and iron depletion in vitro cultured cells.20,28 First, we determined ceruloplasmin mRNA levels in livers of mice exposed to 7.5% O₂ for up to 3 days by Northern blotting (Figure 5A). Liver ceruloplasmin mRNA levels significantly (p < .05) increased 1.7-fold and 1.5-fold from days 1 and 2 to day 3, respectively. Only nonsignificant changes occurred during the first 2 days of exposure (Figure 5B). These data provide the first in vivo evidence that prolonged hypoxia can induce the mouse Cp gene.

CuCl₂ stabilizes HIF-1α under normoxic conditions

To gain more insight into the molecular mechanisms by which CuCl₂ induces and activates HIF-1α, the phosphatidylinositol 3 (PI3)–kinase and mitogen-activated protein kinase (MAPK)–kinase signaling pathways, which are known to be responsible for normoxic HIF-1α induction by a large number of stimuli, were investigated. Therefore, the PI3-kinase inhibitor LY294002, the MAPK kinase-1 (MEK-1) inhibitor PD98059, and the p38 kinase inhibitor PD169316 were used to specifically block these pathways. However, as shown in Figure 3A, these 3 kinase inhibitors had no significant impact on reporter gene induction by 100 μM CuCl₂, suggesting that other mechanisms are responsible for CuCl₂-dependent HIF-1α induction.

In contrast to the kinase pathways that induce HIF-1α protein by translational up-regulation, hypoxia is known to induce HIF-1α by preventing its normoxic degradation. Thus, we induced HIF-1α accumulation by hypoxia in Hep3B cells for 6 hours in the presence or absence of 100 μM CuCl₂. Thereafter, cycloheximide was added to block de novo translation of HIF-1α and reoxygenation was allowed for up to 135 minutes. As shown in Figure 3B, the continued presence of CuCl₂ clearly delayed HIF-1α degradation, suggesting that CuCl₂ stabilizes HIF-1α protein rather than inducing its production. In contrast, β-actin protein levels remained unaffected by this treatment, demonstrating a specific inhibition of HIF-1α degradation.

CuCl₂ inhibits HIF-1α prolyl-4-hydroxylation independent of Fe²⁺ concentration

CuCl₂-dependent stabilization of HIF-1α might be due to inhibition of HIF-1α prolyl-4-hydroxylation by PHD isoenzymes. Indeed, using a recently developed cell-free in vitro assay, we found that CuCl₂ strongly inhibited hydroxylation of a peptide containing HIF-1α Pro564 by PHD1, PHD2, and PHD3, respectively (Figure 4A).

To gain more insight into the mechanism by which CuCl₂ inhibits PHD activity, we repeated the prolyl-4-hydroxylation assays in the presence of limited concentrations (1 μM) or a vast excess (100 μM) of FeSO₄. As shown in Figure 4B, PHD3 activity was inhibited by CuCl₂ with an IC₅₀ value of approximately 2.3 μM. However, the inhibitory concentration at 50% (IC₅₀) value did not significantly change when Fe²⁺ was present in excess, suggesting that iron availability or/and competitive replacement from the active center is not the cause for CuCl₂-dependent inhibition of PHD activity.

Induction of ceruloplasmin mRNA by hypoxia and Cu²⁺

Cu²⁺-dependent induction of HIF-1α offers the intriguing possibility that the Cu²⁺–binding protein ceruloplasmin itself is regulated by Cu²⁺ in a HIF-1–dependent manner. Thus, we further analyzed ceruloplasmin regulation, a known HIF-1 target gene inducible by hypoxia and iron depletion in vitro cultured cells.20,28 First, we determined ceruloplasmin mRNA levels in livers of mice exposed to 7.5% O₂ for up to 3 days by Northern blotting (Figure 5A). Liver ceruloplasmin mRNA levels significantly (t test, p < .05) increased 1.7-fold and 1.5-fold from days 1 and 2 to day 3, respectively. Only nonsignificant changes occurred during the first 2 days of exposure (Figure 5B). These data provide the first in vivo evidence that prolonged hypoxia can induce the mouse Cp gene.

Figure 3. Molecular mechanism of HIF-1α protein stabilization by CuCl₂. (A) Hep3B hepatoma reporter cells (subline HRB5) were treated with or without 100 μM CuCl₂ for 24 hours in the presence or absence of the indicated kinase inhibitors under normoxic conditions. Shown are mean induction factors of luciferase activity in the presence or absence of 150 μM CuCl₂ before the addition of 20 μM cycloheximide (CHX) to block translation. Following reoxygenation for the indicated time periods, HIF-1α levels were determined by immunoblots as in Figure 2. Subsequent detection of β-actin served as control for equal loading.

Figure 4. Inhibition of PHD activity by CuCl₂ in vitro. PHD activity was measured in the presence of oxygen, 2-oxoglutarate, and ascorbate by the binding of a purified pVHL-elonginB-elonginC complex to a HIF-1α–derived peptide bound onto microtiter plates as described in "Materials and methods." (A) PHD enzyme–free assays using a nonhydroxylated peptide or 40 nM of a hydroxyproline-containing peptide served as negative and positive controls (black bars labeled with HIF-P564 and HIF-P564OH, respectively). PHD1 and PHD2 were obtained from insect cells as 6His fusion proteins and PHD2 was expressed in bacteria as MBP fusion protein. Representative experiments performed in triplicate are shown as mean values ± SD. (B) PHD3–dependent HIF-1α Pro⁴⁰⁴ hydroxylation activity was determined in the presence of either 1 or 100 μM FeSO₄. Shown are mean values ± SD of n = 3 independent experiments. OD indicates optical density.
We next determined mRNA concentrations in cultured Hep3B cells (subline HRB5) exposed to hypoxic conditions for up to 3 days by real-time reverse transcription (RT)–PCR. As shown in Figure 6, ceruloplasmin as well as the prototype oxygen-regulated VEGF and Glut-1, but not L28 control mRNA, are induced by hypoxia at all time points examined, confirming that ceruloplasmin is hypoxia-inducible in Hep3B cells. Of note, 150 μM CuCl2 also induced ceruloplasmin, VEGF, and Glut-1, but not L28 mRNA, in Hep3B cells and the combination with hypoxia further increased these mRNA levels (Figure 6).

Maximal ceruloplasmin induction by CuCl2 was 5.5-fold after 72 hours of treatment under normoxic conditions.

The ceruloplasmin gene promoter is activated by Cu2+.

Cu2+-dependent HIF-1α protein and ceruloplasmin mRNA induction suggested that ceruloplasmin gene expression is regulated by HIF-1–dependent promoter activation. We thus transiently transfected Hep3B cells with a ceruloplasmin promoter–firefly luciferase reporter gene containing a functional HRE within a 4774 base pair (bp) fragment upstream of the ATG translational start codon. A cotransfected constitutive renilla luciferase reporter gene served to normalize for differences in transfection efficiency. Following 24 hours of treatment with CuCl2, reporter gene activity dose-dependently increased up to 4-fold at 200 μM CuCl2 (Figure 7A). Hypoxia alone induced reporter gene activity 3.5-fold and the combination of CuCl2 with hypoxia further increased reporter gene activity. However, CuCl2 concentrations greater than 200 μM were considerably more toxic under hypoxic conditions as could be judged from the decrease in cell density and renilla reporter gene activity (not shown).

The involvement of HIF-1 was further analyzed by using the HRE-containing fragment of the ceruloplasmin upstream region linked to the constitutive SV40 promoter. As shown in Figure 7B, the presence of this fragment was sufficient to induce firefly luciferase activity approximately 5-fold by 150 μM CuCl2 or 1% O2 after 24 hours of treatment. Point mutation of this HRE reduced but did not completely inhibit induction by CuCl2 or hypoxia in these experiments, suggesting that other elements might also be involved in CuCl2-dependent activation of ceruloplasmin gene expression.
Discussion

Molecular mimicry of hypoxia by distinct transition metal ions has been known since 1988, when Bunn and coworkers reported that HepG2 and Hep3B hepatoma cell lines increase erythropoietin gene expression not only under hypoxic conditions but also following treatment with CoOCl2 or NiCl2, and to a lesser extent MnOCl2 salts.43 However, CuOCl2 has not been found to induce erythropoietin gene expression in these early experiments. In our study, we demonstrated that CuOCl2 can functionally stabilize HIF-1α, leading to target gene induction in hepatoma cells by a mechanism likely to involve the inhibition of prolyl-4-hydroxylation.

Apart from CoOCl2, NiOCl2, or CuOCl2, several other transition metals have been reported to induce HIF-1α, including vanadate,42 arsenite,42 and chromium.44 A common mechanism by which these transition metals interfere with PHD function is unknown. Salminen and colleagues reported that intracellular ascorbate required for PHD activity might be depleted by CoOCl2 and NiOCl2.45 Reactive oxygen species, PI3K/Akt, p38, p70S6K,1 PKCδ, and adenosine monophosphate (AMP) kinases have been reported to be involved in transition metal-dependent HIF-1α induction.42-44,46-48 However, several kinase pathway inhibitors did not affect CuOCl2-dependent HIF-1 induction in our hands. Moreover, all of these mechanisms are based on an intact cellular environment, but in our experiments we could demonstrate a direct inhibition of PHD activity in a cell-free system in vitro with an IC50 of approximately 2.3 μM and full inhibition at approximately 10 μM CuCl2. These values are somewhat lower than the approximately 150 μM CuCl2 required to obtain maximal induction of HIF-1α in hepatoma cells. However, cell culture conditions, including the CuOCl2-binding capacity of serum proteins present in the fetal calf serum (FCS), cannot be directly compared with in vitro assays performed with different kinectics and purified proteins in a cell-free environment. Over all, our data suggest that inhibition of PHD activity is the most likely mechanism by which CuCl2 induced HIF-1α.

Spectroscopic and structural studies on 2-oxoglutarate-dependent dioxygenases showed that CuOCl2 can substitute active-site FeOCl2 in this class of enzymes.12 However, under our experimental conditions CuOCl2 does probably not compete for the active site of HIF-PHD3 because even the presence of a vast excess of ferrous iron did not reverse the inhibitory function of CuCl2. Although it cannot be excluded that CuOCl2 binds to the active site of PHD with considerably higher affinity compared with FeOCl2, it appears to be more likely that other, yet unknown mechanisms are responsible for the inhibition of HIF PHDs by transition metal ions.

Recently, van Heerden and coworkers demonstrated that rainbow trout exposed to CuSO4 induce HIF-1α protein in their gills.49 These results confirm that CuOCl2 salts can also induce HIF-1α protein in vivo. van Heerden and coworkers attributed their results to an increased oxygen consumption leading to regional tissue hypoxia. However, our data suggest that a direct inhibition of HIF-1α prolyl-4-hydroxylation by CuOCl2 salts might also contribute to the accumulation of HIF-1α protein. Whereas the fish were exposed to water containing 1.65 μM CuSO4,49 optimal CuCl2 concentrations in our experiments were between 100 and 200 μM for hepatoma cells. This concentration range is very similar to the widely used CoCl2 that generally stabilizes HIF-1α in the range between 50-150 μM.41,50 Of note, under normal conditions the “free” intracellular copper ion concentration is less than 1 atom per cell.51 Due to its toxicity, intracellular copper is tightly regulated by copper-scavenging metallochaperones.52 Their copper-binding capacity either needs to be saturated before “free” copper is available to interfere with the PHDs or the copper-bound metallochaperones might be directly involved in PHD regulation. However, the precise molecular mechanisms acting between extracellular transition metal concentrations and intracellular PHD target protein regulation remain unknown.

Liver-derived ceruloplasmin is the main copper-binding plasma transport protein and functions as a ferroxidase essential for the oxidation of Fe2+ before it can be bound to transferrin as Fe3+.21 Previous studies demonstrated that ceruloplasmin is transcriptionally regulated by HIF-1 under hypoxic conditions in vitro.20 Here, we demonstrated that ceruloplasmin mRNA expression is also induced in the liver of hypoxic mice in vivo. Because both ceruloplasmin and apotransferrin facilitate iron release from liver cells and macrophages,53-56 the coordinate induction of their transcription under hypoxic conditions is likely required to match the increased iron demand of erythropoietic precursor cells in the bone marrow following stimulation by erythropoietin.

The finding that CuOCl2 inhibits PHD activity provides the intriguing possibility that the concentration of free CuOCl2 regulates the expression of its binding protein in the liver in a HIF-dependent manner. Indeed, when CuSO4 was administered intravenously to copper-deficient pigs, an increase in plasma ceruloplasmin was observed which preceded the increase in plasma iron.57 Other known HIF-1 target genes induced by CuOCl2 in vivo include VEGF which is increased at wound sites topicaly treated with CuSO4.58 These data are consistent with our findings and support a physiological role of HIF-1 in CuOCl2-dependent gene regulation, extending the sensory functions of the PHD family beyond oxygen sensing.

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References


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