Oxyleghemoglobin scavenges nitrogen monoxide and peroxynitrite

A possible role in functioning nodules?

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Oxyleghemoglobin scavenges nitrogen monoxide and peroxynitrite: a possible role in functioning nodules?

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Abstract It has been demonstrated that the $\text{NO}^\bullet$ produced by nitric oxide synthase or by the reduction of nitrite by nitrate reductase plays an important role in plants' defense against microbial pathogens. The detection of nitrosyl Lb in nodules strongly suggests that $\text{NO}^\bullet$ is also formed in functional nodules. Moreover, $\text{NO}^\bullet$ may react with superoxide (which has been shown to be produced in nodules by various processes), leading to the formation of peroxynitrite. We have determined the second-order rate constants of the reactions of soybean oxyleghemoglobin with nitrogen monoxide and peroxynitrite. At pH 7.3 and 20 °C, the values are on the order of 10^8 and $10^4 \text{ M}^{-1} \text{ s}^{-1}$, respectively. In the presence of physiological amounts of CO_2 (1.2 mM), the second-order rate constant of the reaction of oxyleghemoglobin peroxynitrite is even larger ($10^5 \text{ M}^{-1} \text{ s}^{-1}$). The results presented here clearly show that oxyleghemoglobin is able to scavenge any $\text{NO}^\bullet$ and peroxynitrite formed in functional nodules. This may help to stop $\text{NO}^\bullet$ triggering a plant defense reaction.

Keywords Leghemoglobin · Nitric oxide · Peroxynitrite · Hemoglobin · Kinetics · Mechanism

Abbreviation EPR: Electron paramagnetic resonance · Hb: Human hemoglobin · Lb: Leghemoglobin · LbFeO_2 (oxyLb): Oxyleghemoglobin ·

$\text{LbFe}^{\text{IV}}=\text{O}$ (ferrylLb): Oxoiron(IV)-leghemoglobin · MetLb: Iron(III)leghemoglobin · Mb: Myoglobin · NOS: Nitric oxide synthase

Introduction

Hemoglobins, heme-containing proteins that display a conserved three-on-three arrangement of α -helices (globin fold), have been found ubiquitously in eukaryotes and in many bacteria. Plant hemoglobins were first identified in the root nodules of legumes [1], and these proteins were therefore called leghemoglobins (Lbs). For many years it was believed that Lbs were the only hemoglobins present in plants. However, in the 1980s a new class of hemoglobins was discovered in both nodulating and non-nodulating plants [2]. At least three types of hemoglobins are now believed to be present in plants [3]. Symbiotic Lbs are found in high concentrations in the root nodules of legumes that establish a symbiosis with nitrogen-fixing bacteria [4]. The primary function of these hemoglobins is to facilitate the transport of O_2 to bacteroids for ATP generation while protecting nitrogenase in the bacteroids from O_2 -mediated inactivation. Nonsymbiotic hemoglobins are thought to be present in all plants, albeit at much lower concentrations. These proteins have been divided into two groups, class-1 and class-2 hemoglobins, according to their phylogeny, protein sequences, biochemical properties, and gene expression profiles. The diverse properties of the hemoglobins belonging to these two classes suggest that class-1 and the class-2 proteins have different functions [5]. Class-1 hemoglobins have been proposed to have important functions in plant growth and ATP metabolism under hypoxic conditions, and are expressed in plant tissue in response to specific metabolic stresses [3]. Class-2 hemoglobins are induced in response to microorganisms and, contrary to class-1 hemoglobins, by cytokinin treatment [5]. Compared to class-1 hemoglobins, these proteins display sequences closer to those of Lbs. Finally, the third type of plant hemoglobins

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consist of truncated hemoglobins, hemoproteins that display amino acid sequences 20–40 residues shorter than normal hemoglobins [6, 7].

In animals, nitrogen monoxide ($\text{NO}^\bullet$, nitric oxide) was first identified as an endothelium-derived relaxing factor (EDRF), but is now recognized to regulate a variety of different functions [8]. Three isoforms of nitric oxide synthase (NOS) are responsible for the generation of $\text{NO}^\bullet$ in mammalian systems: constitutive endothelial and neuronal NOSs, and inducible NOS, which is associated with immune and inflammatory responses [9]. $\text{NO}^\bullet$ is also produced in plants, but much less is known about the mechanism of its generation and its functions [10, 11]. Evidence for the existence of a mammalian-type NOS in plants, mostly based on cross-reactions with mammalian NOS antibodies, has repeatedly been reported [12]. However, it has also been demonstrated that some of these antibodies interact with plant proteins other than NOS [13]. Alternative $\text{NO}^\bullet$ sources may occur in plants, and one candidate is considered to be nitrate reductase [14–16]. This enzyme has been shown to transform nitrite into $\text{NO}^\bullet$ in a NADH-dependent process, which is operative when photosynthetic activity is absent or inhibited, and when nitrite can be accumulated [14, 17].

Depending on the concentration applied, both cytotoxic and protecting properties have been attributed to $\text{NO}^\bullet$ in plants [18]. It has been demonstrated that $\text{NO}^\bullet$ plays an important role in the mechanism of plant defenses against microbial pathogens [18–20]. Furthermore, $\text{NO}^\bullet$ was demonstrated to be essential for the regulation of normal plant physiological processes [21]. As for mammalian [22] and bacterial [23] hemoglobins, the reaction of plant hemoglobins with $\text{NO}^\bullet$ may also have a significant physiological function. In this context, it has been shown that both class-I hemoglobins and $\text{NO}^\bullet$ are produced during hypoxic stress [24]. Consequently, it has been suggested that these hemoglobins play a crucial role in modulating the levels of $\text{NO}^\bullet$. Hemoglobin can act as a $\text{NO}^\bullet$ dioxygenase [25] and thus, by lowering the $\text{NO}^\bullet$ concentration, prevent $\text{NO}^\bullet$ -mediated inhibition of cytochrome c oxidase in mitochondria [26, 27].

Interestingly, it has been reported that a NOS-like enzyme is present in the roots and nodules of a leguminous plant (*Lupinus albus*) [28]. Since $\text{NO}^\bullet$ is an inhibitor of nitrogenase from nitrogen-fixing bacteria [29], it is not known whether $\text{NO}^\bullet$ has a biological role in nodules. However, a nitrosyl-leghemoglobin complex ($\text{LbFe}^{\text{II}}\text{NO}$) has been detected in soybean [30], cowpea [31], and alfalfa [32] nodules, an observation that suggests that $\text{NO}^\bullet$ is generated in functional nodules and that Lb may indeed react with it. Moreover, since superoxide may be produced in nodules by various processes [33], including Lb autoxidation [34], it is also conceivable that also peroxynitrite is formed in these organs. In animals, peroxynitrite generated from the diffusion-controlled reaction between $\text{NO}^\bullet$ and $\text{O}_2^{\bullet-}$ ($(1.6 \pm 0.3) \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$ [35]) causes protein nitration and oxidative tissue damage [36].

We have recently reported that human methemoglobin (metHb) and horse heart metmyoglobin (metMb) catalyze the isomerization of peroxynitrite to nitrate, albeit not very efficiently [37]. Moreover, oxyHb and oxyMb have also been shown to scavenge peroxynitrite and to prevent nitration of their own tyrosine residues as well as externally added Tyr [38–40]. We have started to investigate the reactivity of soybean leghemoglobin (Lb) toward so-called reactive nitrogen species. In this work, we have mainly focused on oxyLb, which is the physiologically active form in the nodules. Our data show that this form can efficiently scavenge $\text{NO}^\bullet$ and peroxynitrite; the possible roles of these reactions in the nodulation process are discussed.

Materials and methods

Reagents

Buffer solutions were prepared from $\text{K}_2\text{HPO}_4/\text{KH}_2\text{PO}_4$ (Fluka, Deisenhofen, Germany) with deionized Milli-Q water. Sodium dithionite, potassium superoxide, and hydrogen peroxide were obtained from Fluka. Sodium bicarbonate was purchased from Merck (Darmstadt, Germany). Nitrogen monoxide (Linde, Höllriegelskreuth, Germany) was passed through a NaOH solution and a column of NaOH pellets before use in order to remove higher nitrogen oxides. Nitrogen monoxide solutions were prepared as described previously [41].

Peroxynitrite, carbon dioxide, and protein solutions

Peroxynitrite was prepared either from KO_2 and gaseous nitrogen monoxide according to Koppenol et al [42], or by treating tetramethylammonium superoxide with nitrogen monoxide at -77°C in liquid ammonia, followed by isolation as a crystalline solid via removal of the ammonia [43]. No difference was observed between the reactivity of peroxynitrite prepared according to the two procedures. The peroxynitrite solutions, which contained variable amounts of nitrite (maximally 50% relative to the peroxynitrite concentration) and no hydrogen peroxide, were stored in small aliquots at -80°C . Nitrite did not interfere with our studies, since the reactions of nitrite with oxyLb or with $\text{LbFe}^{\text{IV}}=\text{O}$ proceed at a significantly slower rate than the corresponding reactions with peroxynitrite [44, 45]. The stock solutions were diluted either with 0.01 M NaOH or with water, and the concentration of peroxynitrite was determined spectrophotometrically prior to each experiment by measuring the absorbance at 302 nm ($\epsilon_{302} = 1,705 \text{ M}^{-1} \text{ cm}^{-1}$) [43].

For the experiments carried out in the absence of added CO_2 , the buffers and the 0.01 M NaOH solutions were prepared fresh daily and thoroughly degassed. Experiments in the presence of CO_2 were carried out by adding the required amount of a freshly prepared 0.5 M

sodium bicarbonate solution to the protein solutions as described in detail in [40]. The values for the constant of the hydration–dehydration equilibrium $\text{CO}_2 + \text{H}_2\text{O} \rightleftharpoons \text{H}^+ + \text{HCO}_3^-$ were taken from [46], by considering the ionic strengths of the solutions. After addition of CO_2 or bicarbonate, the protein solutions were allowed to equilibrate at room temperature for at least 5 min.

Soybeans (*Glycine max*) were grown in a glasshouse and the Lb components purified from the root nodules as described previously [47]. All of the experiments reported here were carried out with Lb c_2 . The metLb concentration was determined by measuring the absorbance in 0.1 M phosphate buffer (pH 7.0–7.3) at 404 nm ($\epsilon_{404} = 141 \text{ mM}^{-1} \text{ cm}^{-1}$). OxyLb was prepared by reducing a concentrated metLb solutions with a slight excess of sodium dithionite. The solution was purified chromatographically on a Sephadex G25 column by using a 0.1 M phosphate buffer solution pH 7.0 as the eluant. OxyLb solutions were prepared by diluting this stock solution with buffer, and concentrations were determined by measuring the absorbance at 411, 542 and/or 575 nm ($\epsilon_{411} = 125 \text{ mM}^{-1} \text{ cm}^{-1}$, $\epsilon_{542} = 15.0 \text{ mM}^{-1} \text{ cm}^{-1}$ and $\epsilon_{575} = 14.8 \text{ mM}^{-1} \text{ cm}^{-1}$). FerrylLb solutions were prepared as described in [48], and their concentrations were determined by measuring the absorbance at 416 or 543 nm ($\epsilon_{416} = 97.2 \text{ mM}^{-1} \text{ cm}^{-1}$ and $\epsilon_{543} = 10.4 \text{ mM}^{-1} \text{ cm}^{-1}$) [48].

UV/visible spectroscopy

Absorption spectra were collected in 1 cm cells with a UVIKON 820 spectrophotometer (Kontron, Basel, Switzerland).

Stopped-flow kinetics analysis

Single-wavelength stopped-flow studies were carried out with an Applied Photophysics (Leatherhead, UK) SX17MV or a SX18MV-R instrument. The measured reaction time courses were analyzed with Kaleidagraph (Synergy Software, Reading, PA), version 3.6.2. Rapid-scan spectrophotometric studies were performed with an On-Line Instrument Systems Inc. (OLIS; Bogart, GA) stopped-flow instrument equipped with an OLIS RSM 1000 rapid scanning monochromator. The length of the cell in each of the three spectrophotometers was 1 cm. The mixing time of the OLIS instrument is ~ 4 ms, whereas those of the single-wavelength instruments are ~ 2 ms. All of the measurements were carried out at 20 °C.

Kinetics studies of the reaction of oxyLb with $\text{NO}^\bullet$ were carried out under pseudo-first order conditions with oxyLb in excess with the Applied Photophysics instruments. Reaction time courses used for second-order rate constant determination were collected at 405 nm. For this purpose, the $\text{NO}^\bullet$ solutions were prepared by diluting the aqueous stock solution (2 mM)

with degassed 0.1 M phosphate buffer (pH 7.3) directly in a Hamilton (Martinsried, Germany) SampleLock syringe and the oxyLb solutions were prepared by diluting the stock solution with degassed 0.1 M phosphate buffer (pH 7.3). The concentrations of the oxyLb solutions were determined spectrophotometrically prior to each experiment. Detection of the metLb–peroxynitrite intermediate was performed with the OLIS instrument at pH 9.5. Both the protein and the $\text{NO}^\bullet$ solutions were prepared in 0.1 M borate buffer (pH 9.5).

The reactions of peroxynitrite with oxyLb and with ferrylLb were first studied by rapid-scan stopped-flow spectroscopy, both in the presence and in the absence of CO_2 . The second-order rate constants were then determined by single-wavelength measurements at 413 (or 415) and 430 nm. The protein solutions were prepared by diluting the oxyLb and the ferrylLb stock solutions to the desired concentration with 0.1 M phosphate buffer (pH 7.0–7.3) under aerobic conditions, in the absence or presence of added bicarbonate. Peroxynitrite solutions were prepared by diluting the stock solution immediately before use with 0.01 M NaOH to achieve the required concentration. Kinetics studies of peroxynitrite decay in the presence of different amounts of metLb were performed by single-wavelength stopped-flow spectroscopy by following the absorbance decrease at 302 nm. For all reactions with peroxynitrite, the protein was dissolved in a 0.1 M buffer at a pH slightly more acidic (0.2–0.4 pH units lower) than the desired final pH, which was always measured at the end of the reaction for the control.

Statistics

The experiments reported in this article were carried out in triplicate at least, on independent days. The results are given as mean values of at least three experiments $\pm$ the corresponding standard deviation.

Results

Kinetics studies of the reaction of oxyLb with nitrogen monoxide

The reaction between $\text{NO}^\bullet$ and oxyLb was first studied under anaerobic conditions by rapid-scan stopped-flow spectroscopy at pH 7.3 and 20 °C by following the change in absorbance in the range 300–650 nm. Upon reaction with a slight excess of $\text{NO}^\bullet$ (3–10 μM) the characteristic absorbance maxima of oxyLb (2–9 μM) at 411, 542, and 575 nm shifted within the mixing time essentially to those of metLb ($\lambda_{\text{max}} = 404, 532$, and 623 nm) (data not shown). To determine the second-order rate constant of this reaction, we therefore used the single-wavelength stopped-flow instrument, which has a shorter mixing time and is more suited to measuring extremely fast reactions. Because of the difficulties linked with accurately determining the

concentrations of $\text{NO}^\bullet$ solutions, we chose to keep the concentration of oxyLb at a large excess compared to that of $\text{NO}^\bullet$, to maintain pseudo first-order conditions. The reaction time courses were collected at 405 nm, one of the wavelengths at which the difference in absorbance between oxyLb and metLb is the largest ($\Delta\epsilon_{405} \approx 30 \text{ mM}^{-1} \text{ cm}^{-1}$). As shown in Fig. 1, the traces could all be fitted well to a single-exponential expression. The second-order rate constant, obtained from the linear fit of the plot of the observed pseudo first-order rate constants versus oxyLb concentration (Fig. 1) is $(8.2 \pm 0.5) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$. This value is very similar to that reported for the corresponding reactions of $\text{NO}^\bullet$ with human oxyHb and horse heart oxyMb (Table 1) [49].

The reactions of $\text{NO}^\bullet$ with oxyHb and oxyMb have been shown to proceed via the corresponding peroxynitrite iron(III) complexes ($\text{Fe}^{\text{III}}\text{OONO}$), which have been characterized by UV/vis spectroscopy under alkaline conditions [49, 50]. Moreover, the hemoglobin intermediate $\text{HbFe}^{\text{III}}\text{OONO}$ has recently been characterized under similar conditions by rapid-freeze EPR spectroscopy [50]. As observed for Hb and Mb, rapid-scan spectroscopic studies of the reaction of $\text{NO}^\bullet$ with oxyLb did not show the presence of a detectable intermediate under neutral conditions (pH 7.3). Interestingly, as shown in Fig. 2, the reaction of oxyLb with $\text{NO}^\bullet$ yields the alkaline form of metLb ($\lambda_{\text{max}} = 539$ and 570 nm) without the formation of a detectable intermediate, even under alkaline conditions (0.1 M borate buffer pH 9.5). Under identical conditions, $\text{MbFe}^{\text{III}}\text{OONO}$ has been shown to decay to metMb and nitrate at a rate of $205 \pm 5 \text{ s}^{-1}$ [49]. The $\text{HbFe}^{\text{III}}\text{OONO}$

complex is more stable and the two subunits decay at a rate of 47 ± 3 and $6.9 \pm 0.1 \text{ s}^{-1}$ [49]. Recent reinvestigation of the kinetics of the decay of $\text{HbFe}^{\text{III}}\text{OONO}$ in 0.1 M borate buffer (pH 9.5) showed that the decay rate of the fast component that we reported previously ($36 \pm 5 \text{ s}^{-1}$, [49]) is incorrect.

Kinetics studies of the reaction of oxyLb with peroxynitrite

The reaction between oxyLb and peroxynitrite was studied by rapid-scan UV/vis spectroscopy at pH 7.3 and 20 °C. As shown in Fig. 3a, upon reaction with peroxynitrite, the intensities of the two absorbance bands characteristic for oxyLb ($\lambda_{\text{max}} = 542$ and 575 nm, thin line) decreased and the spectrum of metLb ($\lambda_{\text{max}} = 535$ nm, bold line) appeared. Two sets of isosbestic points were identified during the reaction, at 525/584 nm (Fig. 3a) and 525/611 nm (Fig. 3b), respectively.

This observation indicates that the reaction of peroxynitrite with oxyLb proceeds via an intermediate. The isosbestic points observed at the beginning of the reaction (Fig. 3a) correspond to those between the spectra of oxyLb and ferrylLb, whereas those present at the end of the reaction (Fig. 3b) correspond approximately to those between the spectra of ferrylLb and metLb. In analogy to the reaction of peroxynitrite with human oxyHb and horse heart oxyMb [39, 40, 51], our results strongly suggest that the reaction between peroxynitrite and oxyLb also proceeds in two steps via the formation of ferrylLb (Reactions 1 and 2).

It is worth noting that the absorbance spectrum of the reaction product (the bold line in Fig. 3b) is not completely identical to that of metLb. The small differences—the slightly lower absorbance around 500 nm and the absorbance maximum at 539 instead of 535 nm—indicate that the final product is a mixture of metLb and its nitrite complex $\text{LbFe}^{\text{III}}\text{NO}_2$. Nitrite, present as an impurity in the peroxynitrite solutions, is in excess relative to metLb and thus partly binds to the product but does not influence the rate of the reaction between oxyLb and peroxynitrite. At pH 7.3 and 20 °C, the second-order rate constant for nitrite binding to metLb is $(4.7 \pm 0.1) \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$, and the absorbance maxima of $\text{LbFe}^{\text{III}}\text{NO}_2$ are at 410 and 538 nm [45].

As shown in Fig. 4, the absorbance changes observed for the Soret band in the course of the reaction of oxyLb with an excess of peroxynitrite are also consistent with the proposed mechanism. The characteristic band for oxyLb ($\lambda_{\text{max}} = 411$ nm, thin line) decreased and the absorbance maximum of the final product ($\lambda_{\text{max}} = 405$ –406 nm, bold line) is very close to that of metLb ($\lambda_{\text{max}} = 405$ nm). As discussed above, the absorbance

Fig. 1 Determination of the second-order rate constant for the $\text{NO}^\bullet$ -mediated oxidation of oxyLb. Plot of k_{obs} versus oxyLb concentration for the reaction between $\text{NO}^\bullet$ (0.1–0.3 μM) and variable amounts of oxyLb, measured at 20 °C in 0.1 M phosphate buffer (pH 7.3). *Inset* Reaction time course (average of ten traces) measured at 405 nm for the reaction of 1.65 μM oxyLb with $\sim 0.16 \mu\text{M}$ $\text{NO}^\bullet$ in 0.1 M phosphate buffer pH 7.3. The dotted bold line represents the experimental data, whereas the full line corresponds to the best fit to the data ($k_{\text{obs}} = 146 \pm 2 \text{ s}^{-1}$)

Table 1 Comparison of rate constants for reactions of different forms of leghemoglobin (Lb), human hemoglobin (Hb), and horse heart myoglobin (Mb) with $\text{NO}^\bullet$ and peroxynitrite at pH 7.0–7.3 and 20 °C

Protein	Reaction	Rate constants ($\text{M}^{-1} \text{s}^{-1}$)	Reference
Lb	$\text{Fe(II)O}_2 + \text{NO}^\bullet + \text{H}_2\text{O} \rightarrow \text{Fe(III)OH}_2 + \text{NO}_3^-$	$k = (8.2 \pm 0.5) \times 10^7$	This study
Lb	$\text{Fe(II)O}_2 + \text{HOONO/ONOO}^- \rightarrow \text{Fe(IV)=O}$	$k = (5.5 \pm 0.5) \times 10^4$	This study
Lb	In the presence of 1.2 mM CO_2	$k = (8.8 \pm 0.9) \times 10^5$	
Lb	$\text{Fe(IV)=O} + \text{HOONO/ONOO}^- \rightarrow \text{Fe(III)OH}_2$	$k = (2.1 \pm 0.2) \times 10^4$	This study
Lb	In the presence of 1.2 mM CO_2	$(3.6 \pm 0.5) \times 10^5$	
Lb	$\text{HOONO/ONOO}^- \text{Fe(III)NO}_3^- (+ \text{H}^+)$	$k_{\text{cat}} = (1.45 \pm 0.02) \times 10^5$	This study
Hb	$\text{Fe(II)O}_2 + \text{NO}^\bullet + \text{H}_2\text{O} \rightarrow \text{Fe(III)OH}_2 + \text{NO}_3^-$	$k = (8.9 \pm 0.3) \times 10^7$	[66]
Hb	$\text{Fe(II)O}_2 + \text{HOONO/ONOO}^- \rightarrow \text{Fe(IV)=O}$	$k = (3.3 \pm 0.1) \times 10^4$	[51]
Hb	In the presence of 1.2 mM CO_2	$k = (3.5 \pm 0.3) \times 10^5$	
Hb	$\text{Fe(IV)=O} + \text{HOONO/ONOO}^- \rightarrow \text{Fe(III)OH}_2$	$k = (3.3 \pm 0.4) \times 10^4$	[51]
Hb	In the presence of 1.2 mM CO_2	$k = (1.1 \pm 0.1) \times 10^5$	
Hb	$\text{HOONO/ONOO}^- \text{Fe(III)NO}_3^- (+ \text{H}^+)$	$k_{\text{cat}} = (1.2 \pm 0.1) \times 10^4$	[37]
Mb	$\text{Fe(II)O}_2 + \text{NO}^\bullet + \text{H}_2\text{O} \rightarrow \text{Fe(III)OH}_2 + \text{NO}_3^-$	$k = (4.4 \pm 0.1) \times 10^7$	[49]
Mb	$\text{Fe(II)O}_2 + \text{HOONO/ONOO}^- \rightarrow \text{Fe(IV)=O}$	$k = (5.4 \pm 0.2) \times 10^4$	[39]
Mb	In the presence of 1.2 mM CO_2	$k = (4.1 \pm 0.7) \times 10^5$	[40]
Mb	$\text{Fe(IV)=O} + \text{HOONO/ONOO}^- \rightarrow \text{Fe(III)OH}_2$	$k = (2.2 \pm 0.1) \times 10^4$	[39]
Mb	In the presence of 1.2 mM CO_2	$k = (3.2 \pm 0.2) \times 10^4$	[40]
Mb	$\text{HOONO/ONOO}^- \text{Fe(III)NO}_3^- (+ \text{H}^+)$	$k_{\text{cat}} = (2.9 \pm 0.1) \times 10^4$	[37]

changes in the visible part of the spectrum suggest the formation of small amounts of $\text{LbFe}^{\text{III}}\text{NO}_2$ ($\lambda_{\text{max}} = 410 \text{ nm}$), which may be responsible for the slight shift observed for the Soret band. The intermediate band with a maximum at 409 nm arises from a mixture of mainly ferrylLb and metLb.

The kinetics of the reaction between oxyLb and an excess of peroxynitrite were studied by single-wavelength stopped-flow spectroscopy under pseudo first-order conditions. The two steps of the reaction were studied separately. The reaction time courses of the first step were measured at 413 or 414 nm and those of the second step at 399 or 430 nm, close to the isosbestic points for the second and the first reaction step, respectively. As shown in Fig. 5, both time courses could be fitted to a single-exponential expression. However, the fits for the second traces had to start from the point when the first step finished (see vertical line in Fig. 5b).

Fig. 2 Rapid-scan UV/vis spectra of the reaction of oxyLb (10.6 μM) with $\text{NO}^\bullet$ ($\sim 50 \mu\text{M}$) in 0.1 M borate buffer pH 9.5, 20 °C. Conversion of oxyLb (*thin line*) to the alkaline form of metLb (*bold line*), collected $\sim 2 \text{ ms}$ after mixing

Fig. 3a–b Rapid-scan UV/vis spectra of the reaction of oxyLb (13 μM) with peroxynitrite (170 μM) in 0.05 M phosphate buffer pH 7.3, 20 °C. Conversion of oxyLb (*thin line*) to a mixture of $\text{LbFe}^{\text{III}}\text{NO}_2$ and metLb (*bold line*). **a** The shown spectra were collected with a time interval of 100 ms. **b** The shown traces were collected at 400, 500, 600, 900, 1,200, and 1,800 ms

As shown in Fig. 6, we obtained a linear dependence of the observed rate constants on the peroxynitrite concentration for both reaction steps. The second-order rate constants obtained from the linear fits of the data for the two reaction steps are $(5.5 \pm 0.5) \times 10^4$ and $(2.1 \pm 0.2) \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$, respectively. These values are very close to those obtained for the reaction of peroxynitrite with human oxyHb and with horse heart oxyMb under similar conditions (Table 1).

To confirm that the second step of the reaction between peroxynitrite and oxyLb corresponds to the reduction of ferrylLb to metLb, we studied the reaction of peroxynitrite with ferrylLb, prepared separately by the addition of four equivalents of H_2O_2 to deoxyLb [48]. In contrast to ferrylHb [52], it has been shown that ferrylLb is rather stable and (among other species) does not react with H_2S [48]. Figure 7a shows that upon

addition of an excess of peroxynitrite to ferrylLb, the maximum of the Soret band shifts from 415 to 416 nm (characteristic for ferrylLb) to 405 nm (λ_{max} of metLb). This conversion proceeds with a single isosbestic point at 414 nm, the same wavelength of the isosbestic point found for the second step of the reaction between oxyLb and peroxynitrite (Fig. 4b). The second-order rate constant of the reaction between ferrylLb and peroxynitrite was determined by fitting the reaction time courses at 420 nm to a single-exponential expression (Fig. 7b, inset). From a linear fit of the plot of the observed rate constants versus peroxynitrite concentration (Fig. 7b), we obtain a value of $(3.4 \pm 0.7) \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$, very close to that for the second step of the reaction between peroxynitrite and oxyLb. Taken together, our results suggest that the reaction of peroxynitrite with oxyLb proceeds according to reactions 1 and 2; in other words,

Fig. 4a–b Rapid-scan UV/vis spectra of the reaction of oxyLb (3 μM) with peroxynitrite (87 μM) in 0.05 M phosphate buffer pH 7.3, 20 °C. Conversion of oxyLb (*thin line*) to the intermediate (*bold dotted line*) and to a mixture of $\text{LbFe}^{\text{III}}\text{NO}_2$ metLb (*bold line*). **a** The spectra shown were collected over a time interval of 50 ms. **b** The spectra shown were collected starting from 200 ms (*dotted bold line*) with a time interval of 150 ms

Fig. 5a–b Reaction time courses measured at **a** 413 and **b** 399 nm for the two steps of the reaction of 3 μM oxyLb with 270 μM peroxynitrite in 0.05 M phosphate buffer pH 7.3 and 20 °C. The *dotted bold lines* represent the experimental data, extracted from the rapid-scan UV/vis spectra. In **(b)**, the *vertical line* indicates the starting point for the fit of the second step; that is, the point when the first step is finished. The two observed rate constants resulting from the best fits to the data (*thin lines*) are 17.4 ± 0.5 and $6.9 \pm 0.3 \text{ s}^{-1}$, for the first (**a**) and the second (**b**) reaction step, respectively

Fig. 6 Determination of the second-order rate constants for the two steps of the peroxynitrite-mediated oxidation of oxyLb. Plots of k_{obs} versus peroxynitrite concentration for the two steps of the reaction between oxyLb (3 μM) and peroxynitrite, followed at 413/414 or 399/430 nm, respectively (in 0.05 M phosphate buffer pH 7.3 at 20 °C)

analogously to the reaction of peroxynitrite with human oxyHb [51] and horse heart oxyMb [39].

The reaction between oxyLb and peroxynitrite was also studied in the presence of 1.2 mM CO_2 . Rapid-scan UV/vis spectroscopic studies showed that the absorbance changes of the Soret band and of the visible part of the spectrum that occurred in the course of this reaction are essentially identical to those discussed above for the reaction in the absence of CO_2 (data not shown). The kinetics of the two reaction steps were also studied separately by following the absorbance changes on two isobestic points, 414 and 430 nm, for the first and second step, respectively. As shown in Fig. 8, in the presence of 1.2 mM CO_2 the observed rate constants increased linearly with increasing peroxynitrite concentration. The second-order rate constants obtained for the two steps from the linear fits shown are $(8.8 \pm 0.9) \times 10^5$ and $(3.6 \pm 0.5) \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$, respectively. These values are one order of magnitude larger than those obtained for the two steps of the reaction that took place in the absence of CO_2 (Table 1). Similar increases in the values of the rate constants were also found for the analogous reaction between human oxyHb and peroxynitrite (Table 1).

The second reaction step, the reduction of ferrylLb by peroxynitrite, was studied with separately prepared ferrylLb, again in the presence of 1.2 mM CO_2 . The reaction time courses were measured at 420 nm (data not shown). For all peroxynitrite concentrations studied, the traces could be fitted to a single-exponential expression and the observed rate constants were very similar to those of the second step of the reaction between peroxynitrite and oxyLb measured under similar conditions. The second-order rate constant obtained from a linear fit of the plot of the observed rate constant versus peroxynitrite concentration (data not shown) is $(2.3 \pm 0.5) \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$.

Catalysis of the decay of peroxynitrite by metLb

The reaction of metLb with peroxynitrite was studied by stopped-flow spectroscopy by following the absorbance changes at 302 nm, the absorbance maximum of peroxynitrite. As shown in Fig. 9, at pH 7.3 and 20 °C, addition of $\sim 6 \mu\text{M}$ of metLb reduced the half-life of the decay of (100 μM) peroxynitrite from $t_{1/2} = 3.5 \text{ s}$ (in the absence of the protein) to $t_{1/2} = 0.6 \text{ s}$. Interestingly, under identical experimental conditions metLb was significantly more efficient than human metHb at catalyzing the decay of peroxynitrite (Fig. 9). The half-life for the

Fig. 7a–b Reaction of ferrylLb with peroxynitrite. **a** Rapid-scan UV/vis spectra of the reaction of ferrylLb (2.2 μM) with peroxynitrite (130 μM) in 0.05 M phosphate buffer pH 7.3, 20 °C. Conversion of ferrylLb (thin line) to metLb (bold line). The spectra shown were collected over a time interval of 50 ms. **b** Determination of the second-order rate constant for the peroxynitrite-mediated reduction of ferrylLb. Plot of k_{obs} versus peroxynitrite concentration for the reaction between ferrylLb (2–4 μM) and variable amounts of peroxynitrite, measured at 20 °C in 0.05 M phosphate buffer pH 7.3. *Inset* Reaction time course measured at 420 nm for the reaction of 4 μM ferrylLb with 160 μM peroxynitrite in 0.05 M phosphate buffer pH 7.3 and 20 °C. The dotted bold line represents the experimental data, extracted from the rapid-scan UV/vis spectra, and the thin line the best fit to the data ($k_{\text{obs}} = 7.3 \pm 0.3 \text{ s}^{-1}$)

Fig. 8 Determination of the second-order rate constants for the two steps of the peroxynitrite-mediated oxidation of oxyLb in the presence of CO₂. Plots of k_{obs} versus peroxynitrite concentration for the reaction between oxyLb (3 μM) and peroxynitrite in the presence of 1.2 mM CO₂, followed at 414 or 430 nm, for the first and the second reaction steps, respectively (in 0.05 M phosphate buffer, pH 7.4–7.5 at 20 °C)

reaction performed under identical conditions but with ~ 6 μM metHb was $t_{1/2} = 2.8$ s.

Experiments in the presence of variable amounts of metLb showed that the observed decay rate of peroxynitrite increased linearly with increasing protein concentration (Fig. 10). At pH 7.3 and 20 °C, the catalytic rate constant obtained from the linear fit shown in Fig. 10 was $(1.45 \pm 0.02) \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$. This value is one order of magnitude larger than that previously reported for human metHb and horse heart metMb (Table 1) [37].

Fig. 9 Comparison of the efficiencies of human metHb and metLb at catalyzing the decay of peroxynitrite under identical experimental conditions. Absorbance decrease measured at 302 nm for the decay of 100 μM peroxynitrite in the absence and presence of 5.9 μM metLb or 5.9 μM metHb, in 0.05 M phosphate buffer pH 7.3, at 20 °C

Analogous studies were carried out at pH 6.5. As shown in Fig. 10, at lower pH the rate constants observed for the decay of peroxynitrite were significantly larger, both in the absence and in the presence of metLb. Moreover, the efficiency of metLb to catalyze the decay of peroxynitrite also increased. Indeed, the catalytic rate constant obtained from the linear fit at pH 6.5 and 20 °C was $(5.15 \pm 0.02) \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$. This result suggests that, analogous with the reaction with human metHb [37], peroxynitrous acid (HOONO) is the species that reacts with metLb. Interestingly, under acidic conditions the catalytic rate constant for metLb is one order of magnitude larger than that for human metHb, which is $(2.3 \pm 0.1) \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ (at pH 6.5 and 20 °C [37]).

The reaction of peroxynitrite with metLb was also studied by following the absorbance changes at different wavelengths on the Soret band (404, 411, and 415 nm). In analogy with the reaction with human metHb, the time courses did not only show the decay of peroxynitrite, which still absorbs in this range (because of its very broad band with $\lambda_{\text{max}} = 302$ nm). In most cases, one or two additional processes were observed, which took place at a faster rate than that of peroxynitrite decay (data not shown). The absorbance changes linked with these processes are small and it is possible that they represent the binding of peroxynitrite to metLb.

In addition, further absorbance changes were detected over a much longer timescale (up to 500 s). The directions of these changes (increase/decrease) indicated that they did not arise from the reaction of metLb with nitrite, always present as a contaminant in our peroxynitrite solutions (see below). Because of the large excess of peroxynitrite over metLb used in these experiments, it is conceivable that a fraction of peroxynitrite eludes the reaction with the heme and thus leads to modification of the amino acid residues of Lb. Indeed, the analogous

Fig. 10 Determination of the catalytic rate constant for the metLb-mediated decay of peroxynitrite. Plot of k_{obs} versus metLb concentration for the protein-catalyzed decay of peroxynitrite (100 μM), measured at 20 °C in 0.05 M phosphate buffer (pH 6.5 and 7.3)

reaction of metHb with an excess of peroxynitrite leads to the formation of a long-lived radical characterized by EPR spectroscopy and assigned as a tyrosyl radical [53]. Moreover, similar studies carried out with distal histidine sperm whale metMb mutants also showed the formation of small amounts of nitrated tyrosine residues in those mutants that displayed similar or even larger values of catalytic rate constants for peroxynitrite isomerization [54]. Thus, the absorbance changes detected in our experiments with metLb when peroxynitrite had already completely decayed may arise from subsequent reactions of reactive (radical) intermediates generated on the globin, which slightly modify the absorbance features of the Soret band. Nevertheless, the λ_{max} of metLb was unchanged at the end of the reaction, an observation that suggests that these subsequent reactions are minor processes. In conclusion, since these reactions are certainly due to the large quantities of peroxynitrite used in our experiments, which are significantly larger than those possibly generated in plants, we decided not to investigate them further.

Discussion

Leghemoglobins are the most extensively studied oxygen-binding hemoproteins in plants, as Hb and Mb are for vertebrates. The major function of Lb is undoubtedly to guarantee oxygen supply to respiring nitrogen-fixing microsymbionts at very low oxygen concentrations (10 nM) [4]. However, the discovery that NO• is also produced in plants may reveal further roles for Lb, related to the biochemistry of so-called reactive nitrogen species, as has happened for Hb and Mb [22, 55, 56]. Indeed, it has recently been demonstrated that besides O₂ uptake and storage, Hb and Mb also act as NO• and peroxynitrite scavengers [38, 39, 49].

In plants, one of the functions of NO• is to act as an essential messenger in defense signaling against pathogens [11]. The detection of nitrosyl Lb in nodules [30–32] strongly suggests that NO• is also formed in functional nodules. This has recently been confirmed by experiments performed in our laboratories with a fluorescent probe (A. Puppo, unpublished results). Moreover, NO• may react with superoxide, which may be produced in nodules by various processes [33, 34, 57], thus leading to the formation of peroxynitrite. Since hemoproteins are among the major targets of NO• and peroxynitrite in vivo, in this work we studied the reactions of different forms of soybean Lb with these two so-called reactive nitrogen species, focusing particularly on the physiologically active oxyLb form.

The kinetics studies presented here show that the reaction of NO• with oxyLb is comparable to those with oxyHb and oxyMb. Indeed, the reaction proceeds via ferrylLb, the final product is metLb, and the values of the second-order rate constant are all of the same order of magnitude (Table 1). Detailed studies with Mb

mutants have revealed that, because of the high chemical reactivity of NO• towards the oxygenated heme center, diffusion of NO• into the distal pocket is very likely to be the rate-limiting step for this reaction [58–60]. The same argument holds for the binding of NO• to deoxygenated forms of hemoglobins and myoglobins. Indeed, a good correlation has been found between the values of the rate constants for NO•-mediated oxidations of several oxyMb mutants in which residues of the distal pocket were replaced and those of NO• binding to the corresponding deoxygenated proteins [58]. This observation supports the hypothesis of a common rate-limiting step. The value of the second-order rate constant for NO• binding to deoxyLb ($1.2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, [61]) is four times larger than that of the reaction between the two subunits of deoxyHb and NO• ($3.0 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$, [62]). According to this hypothesis, the observation that the NO•-mediated oxidations of oxyLb and oxyHb proceed at comparable rates is unexpected. Indeed, it would have been predicted that the second-order rate constant for the NO•-mediated oxidation of oxyLb is four times larger than that for the corresponding reaction with oxyHb.

The amino acid sequence of soybean leghemoglobin shows a high degree of homology with those of vertebrate myoglobins and hemoglobins [4]. Also, its tertiary folding pattern is very similar to that of Mb and of the α - and the β -subunits of human Hb [63, 64]. Moreover, soybean Lb contains two histidine residues in positions analogous to those of the “proximal” and “distal” histidine of mammalian Hb and Mb. In oxyHb and oxyMb, the distal histidine is strongly hydrogen bonded to the coordinated oxygen, a feature that will lead to stabilization of the $\text{Fe}^{\text{III}}\text{O}_2^{\cdot -}$ resonance structure (vs. $\text{Fe}^{\text{II}}\text{O}_2$). In contrast, kinetics studies with distal histidine Lb mutants (His61 in soybean Lb) [63] as well as spin echo EPR studies [65] showed that the hydrogen bond between the distal histidine and the coordinated O₂ in oxyLb is very weak at neutral pH and becomes strong only at low pH. Consequently, in oxyLb the $\text{Fe}^{\text{III}}\text{O}_2^{\cdot -}$ resonance structure is likely to be less stabilized and so radical recombination reaction with NO• may proceed at a slightly slower rate.

In analogy to the reaction of NO• with oxyHb and oxyMb [49, 66], it is conceivable that the metLb–peroxynitrite complex ($\text{LbFe}^{\text{III}}\text{OONO}$) is formed in the first step of the corresponding reaction with oxyLb. However, no intermediate could be detected in our stopped-flow studies, even under conditions that allowed for the spectroscopic characterization of the corresponding human Hb complex (pH 9.5) [50, 66]. Our previous studies with Hb and Mb suggested that the rate of decay of the intermediate iron(III) peroxynitrite complex is strongly influenced by the protein environment surrounding the heme [49, 66]. Indeed, significant differences were observed among the decay rates of the peroxynitrite complexes of metMb and the iron(III) forms of the two subunits of Hb: at pH 9.5 and 20 °C, the values are 205, 47, and 6.9 s^{-1} , respectively [49].

A significant difference between the structures of Hb/Mb and Lb is the presence of a wider and more flexible heme pocket in Lb [63, 64]. Thus, the observation that $\text{LbFe}^{\text{III}}\text{OONO}$ decays at a considerably larger rate than the corresponding Mb and Hb complexes seems to suggest that isomerization to nitrate is facilitated when the distal pocket is wider. Studies with distal histidine mutants of Mb (or Lb) would help to corroborate this hypothesis.

The average concentration of Lb over the entire nodule is $\sim 300\ \mu\text{M}$. However, in very active nodules its local concentration can reach 2–3 mM [4]. In biological systems, Lb is mainly present in the reduced LbFe^{II} form. Indeed, because of the low oxygen tension in root nodules, only $\sim 35\%$ of Lb is found in the oxygenated form (oxyLb) [67]. Even if a large part of $\text{NO}^\bullet$ did bind to deoxyLb, the $\text{NO}^\bullet$ scavenging activity of oxyLb would be noticeable. Both processes would protect nitrogenase, which is rapidly inhibited by $\text{NO}^\bullet$ [29]. Moreover, it has been shown that some Lb genes are induced early in the symbiotic interaction [68], before nitrogenase is expressed. At this stage, the oxygen concentration may still be high and larger amounts of Lb may be in the oxygenated form. Under these conditions, the $\text{NO}^\bullet$ scavenging activity of oxyLb would preclude any triggering of a plant defense reaction by this reactive species.

The protective role of oxyLb may be even greater against the damaging effects of peroxynitrite. In nodules, superoxide formation may occur from Lb autoxidation [34] and as a result of the strong reducing conditions required for nitrogen fixation and the action of several proteins including ferredoxin, uricase and hydrogenase [33]. Thus, peroxynitrite is also likely to be produced in functional nodules. Our kinetics studies show that oxyLb is a good scavenger for peroxynitrite. Indeed, the half-life of peroxynitrite in the presence of 1 mM oxyLb is ~ 10 ms. Moreover, it has been reported that the CO_2 concentration in active nodules is about 1.3 mM [69]. Our studies in the presence of 1.2 mM CO_2 showed that oxyLb is even more efficient under these conditions: 1 mM oxyLb reacts with peroxynitrite with a half-life of ~ 1 ms. The mechanism of the reaction between oxyLb and peroxynitrite seems to be identical to that between oxyHb and peroxynitrite. Thus, our work shows that, particularly in the presence of CO_2 , oxyLb can not only scavenge peroxynitrite but also the radicals derived from its decomposition: $\text{CO}_3^{\bullet -}$ and $\text{NO}_2^\bullet$.

Because of the high affinity of deoxyLb for O_2 , with the stopped-flow instruments at our disposal it was not possible to determine the rate constant of its reaction with peroxynitrite. To get qualitative information, we added 5 equiv. of peroxynitrite to deoxyLb under strictly anaerobic conditions in a UV/vis cell, and we have confirmed that metLb is produced from this reaction (data not shown). We have previously shown that the reactions of peroxynitrite with deoxyMb and deoxyHb proceed at a similar rate (Hb) or faster (Mb) than those with the oxygenated forms of these proteins

[39, 51]. Thus, even if Lb was mostly in its deoxygenated form, it may still be an efficient peroxynitrite scavenger.

Interestingly, most extracts from root nodules contain some amount of the oxidized iron(III) form of Lb (metLb). It has been argued that Lb may be oxidized during the extraction process [4, 67]. Nevertheless, direct evidence for metLb occurrence in soybean nodules, in particular in old nodules, has been presented by carrying out UV/vis spectroscopic measurements of intact nodules attached to roots [70].

Here, we have shown that metLb catalyzes the isomerization of peroxynitrite to nitrate, analogously to metHb or metMb. Nevertheless, metLb is ten times more active. We have previously shown that the efficiencies of metHb and metMb as peroxynitrite isomerization catalysts are strongly affected by the presence of the hydrogen bond between the distal histidine and the H_2O molecule bound to the iron(III) center [37, 54, 71]. This strong hydrogen bond has to be cleaved in the rate-determining step of the catalytic process: peroxynitrite binding to the iron(III) center. Thus, metMb mutants in which this hydrogen bond is absent are much more efficient catalysts for the isomerization of peroxynitrite to nitrate [54, 71].

Studies with distal histidine metLb mutants have suggested that this residue undergoes hydrogen bonding to the water molecule coordinated to the iron(III) center in Lb as well [63, 72]. However, it has been shown that this hydrogen bond is weaker in metLb than those in metMb and metHb [63, 72]. Moreover, the position of the Soret absorbance maximum of metLb (404 nm) indicates that the water molecule is only weakly bound to the iron(III). Taken together, the wider, more flexible heme pocket and the weaker hydrogen bond offer an explanation for the larger value of the catalytic rate constant obtained for metLb. Nevertheless, the value of k_{cat} is still more than one order of magnitude smaller than those of the H64A- and H64D-metMb mutants [54], proteins in which this hydrogen bond is absent.

It is interesting to note that the reactivity of metLb towards peroxynitrite is very different to that towards another oxidant generated in plants, H_2O_2 . Indeed, exposure of metLb to H_2O_2 results in the formation of the highly valent oxoiron(IV) form ($\text{LbFe}^{\text{IV}}=\text{O}$) and one or more radicals on the globin [48]. In addition, a green compound is produced, which is believed to be a cross-linked species produced from the reaction of a globin radical with the heme [73].

The results presented here clearly show that oxyLb is able to scavenge any $\text{NO}^\bullet$ formed in functional nodules. This may contribute to the protection of nitrogenase, which is rapidly inactivated by this reactive species [29]. OxyLb is also able to scavenge peroxynitrite, which is likely to be produced from the reaction of $\text{NO}^\bullet$ with superoxide formed in nodules [33, 34, 57], precluding any damaging effect of this species. In both cases, the metLb generated can be reduced by a metLb reductase [74] to its ferrous form, which can give rise to oxyLb. Thus, oxyLb may have a protective role against these

reactive species in functional nodules. On the other hand, taking into account the early inductions of some Lb genes [68], a possible role in the nodulation process cannot be excluded.

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