Kinetic studies on the oxidation of nitrite by horseradish peroxidase and lactoperoxidase

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The reaction of nitrite (NO$_2^-$) with horseradish peroxidase and lactoperoxidase was studied. Sequential mixing stopped-flow measurements gave the following values for the rate constants of the reaction of nitrite with compounds II (oxoferryl heme intermediates) of horseradish peroxidase and lactoperoxidase at pH 7.0, 13.3 ± 0.07 mol$^{-1}$ dm$^3$ s$^{-1}$ and 3.5 ± 0.05 × 10$^{-1}$ mol$^{-1}$ dm$^3$ s$^{-1}$, respectively. Nitrite, at neutral pH, influenced measurements of activity of lactoperoxidase with typical substrates like 2,2'-azino-bis[ethyl-benzothiazoline-(6)-sulphonic acid] (ABTS), guaiacol or thiocyanate (SCN$^-\$). The rate of ABTS and guaiacol oxidation increased linearly with nitrite concentration up to 2.5-5 mmol dm$^{-3}$. On the other hand, two-electron SCN$^-$ oxidation was inhibited in the presence of nitrite. Thus, nitrite competed with the investigated substrates of lactoperoxidase. The intermediate, most probably nitrogen dioxide (NO$_2^*$), reacted more rapidly with ABTS or guaiacol than did lactoperoxidase compound II. It did not, however, effectively oxidize SCN$^-$ to OSCI. NO$_2^*$ did not influence the activity measurements of horseradish peroxidase by ABTS or guaiacol method.

The heme-containing peroxidases catalyze one-electron oxidation of a wide range of substrates with concomitant reduction of peroxides:

Peroxidase + H$_2$O$_2$ → Compound I + H$_2$O (1)

Compound I + SH → Compound II + S$^*$ (2)

Compound II + SH → Peroxidase + S$^*$ (3)

where compound I is an oxoferryl porphyrin $\pi$-cation radical, Fe(IV)-O(Por$^{3+}$), which, in the case of lactoperoxidase (LPO) (and some other peroxidases, but not horseradish peroxidase (HRP)), spontaneously transforms

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Abbreviations: ABTS, 2,2'-azinobis[3-ethyl-benzothiazoline-(6)-sulphonic acid]; HRP, horseradish peroxidase; LPO, lactoperoxidase; MPO, myeloperoxidase; TNB, 5-thio-2-nitrobenzoic acid.
to oxoferryl protein radical, Fe(IV)=O(R*); compound II is an oxoferryl heme intermediate, Fe(IV)=O; SH is a substrate to be oxidized. Many organic and inorganic compounds are known to serve as substrates of the peroxidase reaction [1].

Since about ten years, i.e., since the discovery that nitric oxide (NO) plays a key role in several processes, both physiologically and pathophysiological, reactions of reactive nitrogen intermediates with biologically important molecules have drawn considerable interest. It is known that NO is a short-lived molecule the content of which is nearly 0.01–0.1 μmol dm⁻³ in mammalian tissues, and is significantly increased in pathophysiological processes [2]. Its autooxidation in aqueous solution is a very complex process which can be summarized as follows:

\[ 4 \text{NO} + O_2 + 2 H_2O \rightarrow 4 H^+ + 4 \text{NO}_2^- \]  (4)

*NO reacts rapidly with O₂ to yield peroxynitrite (ONOOCO⁻), a powerful oxidizing agent. Isomerization of peroxynitrite gives NO₃⁻, but oxidation by ONOO⁻/ONOOCO⁻ (pKₐ = 6.8), results in concomitant production of NO₂⁻. Thus, nitrite is an important end product of NO metabolism. The amount of nitrite found in healthy human subject ranges from hundreds of nanomoles in plasma up to hundreds of micromoles in saliva [3]. The level of nitrite concentration reflects not only endogenous *NO production but also NO₃⁻/NO₂⁻ ingestion [4]. It is worth noting that nitrite is also a significant contaminant in solutions of peroxynitrite synthesized on the laboratory scale. NO₂⁻ may be present at millimolar concentration in peroxynitrite samples [5].

It has been shown that nitrite is oxidized by heme proteins: oxyhemoglobin [6], catalase [7, 8] and peroxidases [9, 10]. More recently, it was reported that catalase was inhibited by NO₂⁻ [11]. In the case of peroxidases, nitrite peroxidation occurs in the normal peroxidase cycle (reactions (1)–(3)) and the formation of nitrogen dioxide (*NO₂) as an intermediate has been postulated [9]. The second-order rate constants for the reactions of compounds I and II of horseradish peroxidase (HRP) with nitrite increased with increasing acidity over the pH range 6–8 [9]. Shibata et al. [10] reported that nitrite added to a mixture containing HRP and H₂O₂ bleached chlorophyll and nitrated tyrosine. The optimum pH for bleaching of chlorophyll was 4.0. Chance [8] suggested that protonated nitrite (HNO₂, pKₐ = 3.3) could be the active form of the electron donor. Klebanoff [12] showed that myeloperoxidase (MPO) together with H₂O₂ and nitrite was toxic to E. coli at pH 5–7. Recently van der Vliet et al. [3] and Sampson et al. [13] reported that HRP, LPO and MPO oxidize nitrite in the presence of H₂O₂ to catalyze nitration of tyrosine and tyrosine residues in proteins at neutral pH. The efficiency of tyrosine nitration by investigated enzymes was found to be as follows: HRP < LPO < MPO [13].

Rzeszka et al. [14, 15] found that the presence of nitrite dramatically enhanced oxidation of some substrates by the LPO/H₂O₂ system at neutral pH. The same authors also reported that the anticancer agent, mitoxantrone, which was not metabolized by LPO/H₂O₂ underwent oxidation to the biologically active quinone/diiminoquinone form of the drug by a nitrite-derived metabolite of LPO/H₂O₂, presumably the *NO₂⁻ radical [16].

The aim of this work was to compare the reactivity of two peroxidases: HRP and LPO towards nitrite and to investigate the influence of the intermediate species of nitrite oxidation, most likely *NO₂⁻, on reactions catalyzed by these enzymes.

MATERIALS AND METHODS

Horseradish peroxidase (type VI) (EC 1.11.1.7) with the purity number RZ = A₄₀₀/ A₂₈₀ of 3.0 (a measure of hemin content using content of the aromatic amino acids as reference), and lactoperoxidase (EC 1.11.1.7) with RZ = A₄₁₂/ A₂₈₀ of 0.8 were obtained from
Sigma and used as received. Extinction coefficients of 1.02 \cdot 10^5 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1} at 403 nm [17] and 1.12 \cdot 10^5 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1} at 412 nm [18] were used for spectrophotometric determinations of HRP and LPO concentrations, respectively. Sodium nitrite (NaNO₂), 2,2'-azino-bis[ethyl-benzothiazoline-(6)-sulphonic acid] (ABTS), guaiacol, sodium thiocyanate (NaSCN), hydrogen peroxide (H₂O₂) 30% were also obtained from Sigma. All other reagents were of analytical grade. All experiments were carried out at ambient temperature and at pH = 7.0. (0.01 \text{ mol dm}^{-3} phosphate buffer). We checked that reaction kinetics in the absence or presence of 0.1 \text{ mol dm}^{-3} NaNO₃ was the same, i.e. that ion strength did not influence the reactions investigated. Nano-pure water from MilliQ (Millipore) was used throughout.

In order to determine the dissociation constant, K_D, for the LPO/NO₂ complex, difference absorption spectra (spectrum of the enzyme–substrate versus spectrum of the enzyme) were taken. Titration were carried out by adding nitrite at concentrations of 1-50 mmol dm⁻³ to 4 µmol dm⁻³ LPO solution and to the buffer (pH 7.0) in the reference cell. K_D value was calculated according to [19].

Kinetic measurements for reactions taking place within milliseconds or seconds were performed using the SX 17 MV stopped-flow spectrofluorimeter (Applied Photophysics) with 1-cm cell. The kinetics of the reaction of compounds I or II of HRP with nitrite were studied using the sequential mixing mode of the stopped-flow apparatus. HRP solution (8 µmol dm⁻³) placed in syringe 1, was premixed with an equal volume of the 10 µmol dm⁻³ H₂O₂ solution, placed in syringe 2. After a delay of 100 ms, the compound I formed in the aging loop was mixed with the solution containing nitrite, the concentration of which was chosen to ensure at least 10-fold excess over that of compound I. Reactions of HRP compounds I or II with nitrite were followed by changes in absorption at 412 nm (isosbestic point between native enzyme and compound II) or 398 nm (isosbestic point between compound I and compound II), respectively.

The reaction of compound II of LPO with nitrite was studied in a similar way: 4.5 µmol dm⁻³ solution of LPO was premixed with 20 µmol dm⁻³ H₂O₂ (concentrations before mixing are given). Delay time was 2 s. We checked that after that time LPO compound I was converted to the species with absorption maximum at 430 nm, characteristic for compound II. Now it is accepted that LPO compound I, which occurs primarily in the form of oxoferryl porphyrin π-cation radical, spontaneously transforms to the form of oxoferryl protein radical, the absorption spectrum of which is indistinguishable from that of compound II [20, 21]. Nevertheless, reduction of compound I by an internal donor to compound II cannot be excluded. The formed LPO intermediate species (compound I in the form of protein radical and/or compound II) was then mixed with varying concentrations of nitrite (kept in at least 10-fold excess over LPO species). The rate of the decrease of absorbance at 430 nm was taken as a measure of the rate of the reaction of compound II with substrate. Note that, when compound I, existing in the form of a protein radical, reacts with substrate to form compound II, no changes in absorption at 430 nm should be seen.

Five determinations of rate constants were performed for each nitrite concentration and the mean values were plotted against the substrate concentration. The apparent second-order rate constants of the reactions of compounds I and II of HRP and for compound II of LPO with nitrite were calculated from the respective slopes using least-squares regression.

The kinetic measurements for the reactions occurring at a longer time-scale were performed using Hewlett-Packard 8452A diode array spectrophotometer.
RESULTS AND DISCUSSION

It was shown that nitrite binds to heme proteins at the heme iron [22], but the concentration of nitrite as high as mmol dm\(^{-3}\) is required to change the absorption spectrum. Ligation of NO\(_2^-\) to HRP is very sensitive to pH. The optical spectrum of HRP in the presence of 20 mmol dm\(^{-3}\) NO\(_2^-\) showed no change after 13 h at pH 7.7. However, at pH 5.4, in the presence of 100 mmol dm\(^{-3}\) NO\(_2^-\), the absorption maximum shifted to 416 nm within 1 min [22]. We have checked whether nitrite used at the concentrations required to determine the rate constants with HRP compounds I and II and in activity measurements does influence absorption spectrum of HRP. Our experiments have shown that absorption spectrum of HRP in the presence of nitrite at concentrations from \(10^{-5}\) to \(5 \cdot 10^{-2}\) mol dm\(^{-3}\) at pH 7.0 remained unchanged even after several days (storing at 8°C, measurement at room temperature). This means that under our experimental conditions most of nitrite was dissociated from the enzyme.

Spectral scans of the reaction between HRP and H\(_2\)O\(_2\) in the presence of nitrite are shown in Fig. 1. The presence of isosbestic points during formation of compound II (Fig. 1a) and restoration of native enzyme from compound II (Fig. 1b) clearly indicate that the reaction occurs in two kinetically separated steps, involving a nitrite induced one-electron reduction of compounds I and II, respectively. The respective second-order rate constants for the reactions of HRP compound I and II with NO\(_2^-\) at pH 7.0 determined by us are 4.5 ± 0.05 \(\cdot\) \(10^2\) mol\(^{-1}\) dm\(^3\) s\(^{-1}\) and 13.3 ± 0.07 mol\(^{-1}\) dm\(^3\) s\(^{-1}\). These values fall into the range reported earlier [9]. In order to check whether \(^{\ast}\)NO\(_2^-\), most probably formed as an reaction intermediate, does inactivate the enzyme, a solution containing 2 \(\mu\)mol dm\(^{-3}\) HRP was incubated with 2.5 \(\mu\)mol dm\(^{-3}\) H\(_2\)O\(_2\) and with different amounts of NO\(_2^-\) (from \(10^{-5}\) to \(5 \cdot 10^{-2}\) mol dm\(^{-3}\)) for the time ranging from 10 min to 24 h. Then the aliquots of the solution were diluted with the buffer in proportion 1:500 and the rates of ABTS (2.5–10 mmol dm\(^{-3}\)) or guaiacol (2.5–10 mmol dm\(^{-3}\)) oxidation were measured. We found that activity of the samples incubated with NO\(_2^-\), irrespective of nitrite concentration and incubation time, was the same as in the absence of nitrite (not shown). We have also measured the rates of ABTS or guaiacol oxidation by adding nitrite at concentrations of \(10^{-3}\)–5 \(\cdot\) \(10^{-2}\) mol dm\(^{-3}\) directly to the cuvette. Again, no change of reaction kinetics has been observed, irrespective of substrate concentration.

It has been shown that nitrite interacts with LPO [23]. The value of dissociation constant for the LPO/NO\(_2^-\) adduct, \(K_D = 2.43\) mmol dm\(^{-3}\) at pH 5.5 was reported [24]. When nitrite concentration equals \(K_D\), half of the enzyme present in the solution is complexed with NO\(_2^-\) (assuming that nitrite is in a great excess with respect to the enzyme). The ab-

![Figure 1. Absorption spectra of compound II formation (A, spectra taken every 10 s) and decay (B, spectra taken every 3 min), [HRP] = 2 \(\mu\)mol dm\(^{-3}\), [H\(_2\)O\(_2\)] = 5 \(\mu\)mol dm\(^{-3}\), [NaNO\(_2\)] = 5 \(\cdot\) \(10^{-4}\) mol dm\(^{-3}\), pH = 7.0.](image-url)
absorption maximum for the LPO/NO$_2^-$ complex at pH 5.5 was found at 422 nm [24]. We have found that $K_D$ value for the LPO/NO$_2^-$ complex is $20 \pm 0.5$ mmol dm$^{-3}$ at pH 7.0. This indicates that for LPO, similarly to HRP, the interaction between NO$_2^-$ and ferric iron is weaker at neutral pH than in acidic solution.

In a similar manner as in the experiments with HRP, i.e. by the method of sequential mixing, we have tried to determine the rate constant of the reaction of LPO compound I and compound II with nitrite. It is known that LPO compound I, occurring in the form of porphyrin π-cation radical is very unstable ($t_{1/2} = 0.2$ s) and spontaneously transforms to the species showing an absorption spectrum typical for compound II [25]. However, it has been suggested, that the species formed could be compound I in the o xo ferryl protein radical form. In fact, it is spectroscopically indistinguishable from compound II [21]. After a delay of 50 ms, i.e. at a time when LPO compound I in the form of porphyrin π-cation radical is being formed, the solution was mixed with nitrite and an increase of absorbance at 430 nm (maximum of compound II absorption band) was detected. Unfortunately, the spontaneous transformation of porphyrin radical form of compound I into protein radical form ($k = 3.5 \pm 0.05$ s$^{-1}$ (our unpublished results)), strongly disturbs observations of the reaction of compound I with nitrite and prevents precise determination of the rate constant for this reaction. The absorbance at 430 nm observed after LPO compound II formation is by about 30% lower in the presence of both H$_2$O$_2$ and nitrite than in the presence of H$_2$O$_2$ alone. The lowered concentration of LPO compound II is most probably due to a smaller, in comparison with HRP, difference in the rate constants for the reactions of LPO compounds I and II with nitrite. A similar observation were made by Metodiewa & Dunford [26] who studied the reactions of LPO compounds I and II with 3-aminotriazole.

Using sequential mixing with a delay time of 2 s (see Materials and Methods) it was possible to measure the rate constant of LPO compound II with nitrite. We obtained a value of $3.5 \cdot 10^4 \pm 0.05$ mol$^{-1}$ dm$^3$ s$^{-1}$ (pH 7.0). It is three orders of magnitude higher than that for HRP.

Bactericidal activity of lactoperoxidase in vitro is connected with two-electron oxidation of thiocyanate to hypothyocyanite anion, SCN$^-$ (at pH 7.0), the formation of which may be followed spectrophotometrically at 235 nm [27]. The rate of thiocyanate oxidation ([SCN$^-1$] = 1 mmol dm$^{-3}$) by LPO in the presence of 1 mmol dm$^{-3}$ H$_2$O$_2$ decreased with increasing nitrite concentration and was completely inhibited when 2.5 mmol dm$^{-3}$ nitrite was present in the sample (not shown). The reported values of the rate constant of LPO compound I with SCN$^-$ at pH 7.5 are in the range $4 \cdot 10^4 - 1.1 \cdot 10^5$ mol$^{-1}$ dm$^3$ s$^{-1}$ [28, 29] i.e. they are comparable with the rate constant of LPO compound II with nitrite obtained by us. Note that the rate constant for the reaction of LPO compound I with nitrite estimated by us was only several times higher (also of the order of $10^6$ mol$^{-1}$ dm$^3$ s$^{-1}$). This may suggest that SCN$^-$ and NO$_2^-$ compete with each other for LPO compound I. The other possibility is that NO$_2^-$ blocks the binding site for SCN$^-$ in the enzyme. It is generally accepted that peroxidase substrates bind to the enzyme prior to the catalytic reaction. The dissociation constant, $K_D$, for LPO/SCN$^-$ complex was reported to be 20.3 mmol dm$^{-3}$ at pH 5.5 [24] but $90 \pm 0.5$ mmol dm$^{-3}$ at pH 6.1 [30]. Nitrite ions strongly compete with SCN$^-$ in binding to the enzyme and the $K_D$ value for the LPO/SCN$^-$ complex in the presence of NO$_2^-$ (at a concentration four times as high as $K_D$) is 88.1 mmol dm$^{-3}$ [24] at pH 5.5. However, as we carried out our measurements at ion concentrations lower than $K_D$, i.e. under conditions when more than half of the amount of enzyme remained in the uncomplexed form, it seems unlikely that inhibition of SCN$^-$ oxidation by nitrite was caused by difficulties in SCN$^-$ binding to protein prior to the catalytic reaction.
Van der Vliet et al. [3] found that the oxidation of 5-thio-2-nitrobenzoic acid (TNB) by the LPO/H$_2$O$_2$/SCN$^-$ system was enhanced by NO$_2^-$ when [NO$_2^-$] was two orders of magnitude lower than [SCN$^-$]. Thus, when the ratio [SCN$^-$/[NO$_2^-$] = 100, the competition between those two anions with LPO compound I seems unlikely. It is possible that LPO compound I, in addition to reacting with SCN$^-$, is reduced to compound II by an internal donor. Compound II is unable to oxidize SCN$^-$ but can react with nitrite to produce •NO$_2^-$, which in turn oxidizes TNB, additionally to OSCN$^-$. Incubation of a solution containing 2 µmol of NO$_2^-$ (Fig. 2-inset). We have observed that the rate of ABTS and guaiacol oxidation increases linearly up to about 2.5 - 5 mmol dm$^{-3}$ NO$_2^-$ . Above these concentrations, the influence of NO$_2^-$ becomes less and less pronounced to reach saturation at about 12 mmol dm$^{-3}$ and 20 mmol dm$^{-3}$ nitrite, respectively, for guaiacol and ABTS. On comparing the rate constants for the reactions of LPO compound II with nitrite with those for the investigated substrates one can conclude that nitrite is able to compete with LPO substrates. The intermediate, most probably •NO$_2^-$, reacts more rapidly with ABTS and guaiacol than does

dm$^{-3}$ LPO, 20 µmol dm$^{-3}$ H$_2$O$_2$ and nitrite at concentrations up to 0.1 mmol dm$^{-3}$ for different time intervals (from 10 min to 24 h) followed by 500-fold dilution of the samples and determination of activity by the ABTS method indicates that •NO$_2^-$ does not inactivate LPO.

The presence of nitrite at a concentration ≥ 0.1 mmol dm$^{-3}$ caused an increase of the rate of oxidation of ABTS (not shown) or guaiacol (Fig. 2). At 2.5 mmol dm$^{-3}$ concentration, nitrite accelerated oxidation of 2.5 mmol dm$^{-3}$ ABTS approximately fivefold (not shown) and the oxidation of guaiacol even more efficiently. In the presence of 2.5 mmol dm$^{-3}$ nitrite, 5 mmol dm$^{-3}$ guaiacol was oxidized sixteen times faster than in the absence of LPO compound II ([31, 32] and our unpublished results), thus their oxidation in the presence of nitrite occurs faster. On the other hand, •NO$_2^-$ seems to be unable to oxidize SCN$^-$ to OSCN$^-$. When NO$_2^-$ concentration is comparable with $K_D$, its significant part is complexed with ferric iron and, in consequence, the reaction with hydrogen peroxide is blocked and enzyme becomes inactivated. Thus the observed deviations from linearity in the plot of the reaction rate for substrate oxidation vs [NO$_2^-$], observed for nitrite concentrations above 2.5 - 5 mmol dm$^{-3}$ (Fig. 2), is most probably connected with partial enzyme inactivation caused by nitrite binding to the ferric iron.
It has been shown that nitrite is able to inhibit catalase activity. In the presence of 1 mmol dm$^{-3}$ nitrite, at pH 7.0, only 23% of the initial activity (the rate of H$_2$O$_2$ decomposition) was detected ([H$_2$O$_2$] used at this experiment was 10.5 mmol dm$^{-3}$) [11]. The mechanism of this inhibition is not clear. The rate constant for the reaction of compound I of catalase with nitrite and with H$_2$O$_2$ is of the order of $10^{-3}$ mol$^{-1}$ dm$^3$ s$^{-1}$ at pH 7.0 [7, 32] and of $10^{-7}$ mol$^{-1}$ dm$^3$ s$^{-1}$ [33], respectively. Thus, the observed inhibition could not be due to the competition between nitrite and H$_2$O$_2$ for compound I, but was rather due to the modification of enzyme caused by nitrite.

CONCLUSIONS

Nitrite reacts with lactoperoxidase with a rate constant comparable with those for typical peroxidase substrates at pH 7.0. The intermediate, most probably *NO$_2$, does not inactivate the enzyme but may react with LPO substrates at a rate even higher than that of LPO compound II with substrates. As a result, oxidation of substrates in the presence of nitrite may occur faster. When nitrite concentration exceeds 2.5−5 mmol dm$^{-3}$, partial enzyme inactivation is observed. On the other hand, the rate constants for the reaction of HRP compounds I and II with nitrite at neutral pH are much lower than those for typical HRP substrates and, in most cases, nitrite does not compete with the latter in the catalytic process.

REFERENCES


