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Exponential Amplification by Redox Cross-Catalysis and Unmasking of Doubly Protected Molecular Probes

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The strength of autocatalytic reactions lies in their ability to provide a powerful means of molecular amplification, which can be very useful for improving the analytical performances of a multitude of analytical and bioanalytical methods. However, one of the major difficulties in designing an efficient autocatalytic amplification system is the requirement for reactants that are both highly reactive and chemically stable in order to avoid limitations imposed by undesirable background amplifications. In the present work, we devised a reaction network based on a redox cross-catalysis principle, in which two catalytic loops activate each other. The first loop, catalyzed by H₂O₂, involves the oxidative deprotection of a naphthylboronate ester probe into a redox-active naphthohydroquinone, which in turn catalyzes the production of H₂O₂ by redox cycling in the presence of a reducing enzyme/substrate couple. We present here a set of new molecular probes with improved reactivity and stability, resulting in particularly steep sigmoidal kinetic traces and enhanced discrimination between specific and nonspecific responses. This translates into the sensitive detection of H₂O₂ down to a few nM in less than 10 minutes or a redox cycling compound such as the 2-amino-3-chloro-1,4-naphthoquinone down to 50 pM in less than 30 minutes. The critical reason leading to these remarkably good performances is the extended stability stemming from the double masking of the naphthohydroquinone core by two boronate groups, a counterintuitive strategy if we consider the need for two equivalents of H₂O₂ for full deprotection. An in-depth study of the mechanism and dynamics of this complex reaction network is conducted in order to better understand, predict and optimize its functioning. From this investigation, the time response as well as detection limit are found highly dependent on pH, nature of buffer, and concentration of the reducing enzyme.

Introduction

In the context of chemical or biochemical analysis, it is common to rely on molecular or biomolecular amplifications to improve the analytical performances of a detection method (*i.e.*, sensitivity, limit of detection (LoD), analysis time, dynamic range of concentration, ...).^{1,2} This is even more important when the targeted molecules or biomolecules are present at low concentrations. In most cases, molecular amplifications result from catalytic reactions wherein a molecule (either the analyte acting as a catalyst or a label catalyst revealing indirectly the presence of the analyte) triggers the catalysis of a reaction capable to generate a detectable signal (which can be optical, electrochemical, thermal, gravimetric, *etc* ...). Under these conditions, the product linearly accumulates with time and the amplification factor depends only on the catalytic rate and reaction time.³ A popular example is the signal amplification resulting from the enzyme label involved in enzyme-linked immunosorbent assays.^{4, 5} However, greater molecular amplifications can be achieved by adequately combining the reactivity of different catalysts. One possibility is to cascade the reactivity of two catalysts, wherein a first one catalyses the production of a second one. In such situation, the product of the secondary catalytic reaction accumulates quadratically with time.⁶⁻¹¹ An even more powerful approach is to produce an exponential signal gain from an autocatalytic reaction, for which one of the products catalyses its own formation.¹²⁻¹⁴ A well-known example is the polymerase chain reaction, which through self-replication of a targeted DNA sequence allows the ultimate detection of a few molecules of nucleic acids in a few tens of microliters.¹⁵ While autocatalytic replication is well suited to nucleic acids,¹⁶⁻¹⁹ it is more challenging to design for

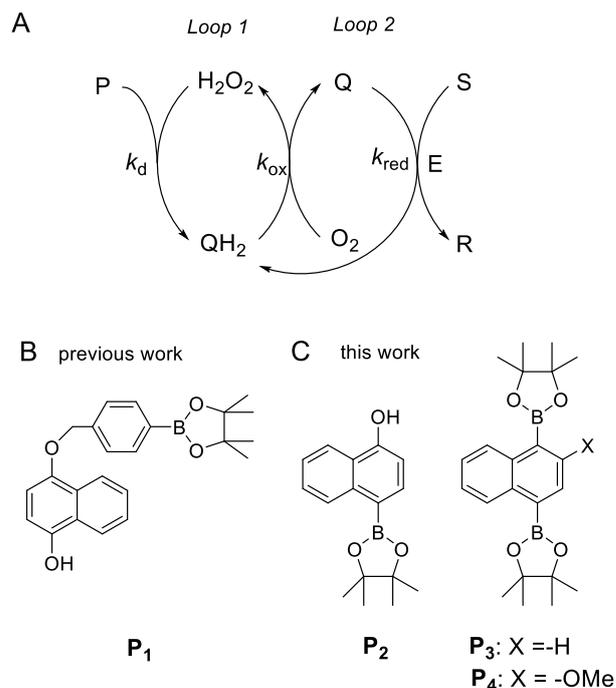
other molecules.¹³ Among the few examples of molecular self-replication strategies developed to date for the detection of small molecules is the dendritic chain reaction (DCR) first described in 2009 by Shabat and his collaborators.^{20, 21} In this concept, the analyte triggers its self-replication through the decomposition of a dendritic molecular probe containing at least two masked equivalents of the analyte. This methodology was applied to the autocatalytic detection of H₂O₂^{20, 22, 23} and fluoride ions.²⁴⁻²⁷ More recently, alternative strategies based on self-propagating thiolate-disulfide exchange,^{28, 29} methionase chain reaction,³⁰ or photoinduced autocatalysis^{31, 32} were also reported. However, all these methods suffer from undesirable background reactions, which prevent the achievement of very low analyte concentrations. The main reason of these background reactions is the intrinsic lack of stability of the starting coreactants, leading to self-initiation of the amplification cycle in the absence of analyte. Therefore, a critical issue is to design systems involving coreactants with high chemical stability and, at the same time, high reactivity and selectivity towards the trigger of the reaction. This is certainly the major challenge facing the exponential signal amplification strategies.³³ An additional drawback is the slowness of the approaches proposed to date, sometimes requiring waiting several hours for an analytical response.^{21, 23}

To address these drawbacks and contribute to the development of alternative solutions, we recently proposed a new versatile molecular-based autocatalytic system^{34, 35} in which the exponential gain is generated from cross-activation of two catalytic loops, as depicted in Scheme 1A. The first loop (*Loop 1*) is based on a H₂O₂-mediated deprotection of a boronate ester probe (P) to a hydroquinone derivative (QH₂), which, in the presence of O₂ in solution, spontaneously autoxidizes to a

quinone (Q) with the concomitant regeneration of H_2O_2 (the latter acting thus as a catalyst in *Loop 1*). The autocatalysis is then engendered from the back conversion of Q into its reduced QH_2 form in the presence of a reducing substrate S (or alternatively an enzyme/reducing substrate couple E/S), thus forming the second catalytic loop (*Loop 2*, which is also a redox cycle wherein Q/ QH_2 serves as a redox catalyst). Analytical derivations as well as numerical simulations of the kinetics of this redox cross-catalytic system allowed us to demonstrate the overstoichiometric production of H_2O_2 coupled with the exponential decrease of probe and substrate concentrations with time.^{34, 36} This autocatalytic set is conceptually different from the previously reported branched-chain or self-propagating amplification reactions in that it does not require the design of a dendritic probe capable of releasing two or more equivalents of the trigger by self-immolation. Another advantage is that it offers the possibility to trigger the reaction either by H_2O_2 or any redox cycling compound, thus providing an extended degree of freedom to couple this molecular amplification not only to the detection of H_2O_2 (directly or indirectly produced), but also to the detection of molecules exhibiting redox cycling properties (also either directly or indirectly generated).

Using the boronate ester probe **P**₁ shown in Scheme 1B, detection of submicromolar or subnanomolar concentrations of H_2O_2 or a naphthoquinone, respectively, within 1 hr was demonstrated.³⁴ Although these analytical performances are significantly better than those previously achieved by DCR,^{20, 22, 23, 37, 38} they are still limited by the lack of chemical stability of **P**₁, which in the absence of the trigger tends to slowly self-deprotect into a naphthohydroquinone. This lack of stability turns out to be intrinsic to the self-immolating spacer installed on the probe,³⁴ a recurrent issue of probes based on self-immolation and which is at the origin of the unsatisfactory analytical performances encountered so far. To access more robust and stable probes, it is thus important to design probes that no longer rely on self-immolation. This is what we address here with the new boronate-based probes shown in Scheme 1C. Instead of introducing the masking boronate ester function *via* a remote self-immolating spacer, we have chosen to place it directly on the naphthyl ring, leading either to the single-protected naphthol monoboronate ester probe **P**₂ or the doubly protected naphthyl diboronate ester probes **P**₃ and **P**₄. By reaction with H_2O_2 , these probes are expected to release a 1,4-naphthohydroquinone (1,4-NQH₂), which in the presence of O_2 spontaneously autoxidizes into 1,4-naphthoquinone (1,4-NQ) with concomitant regeneration of H_2O_2 , as previously shown with **P**₁.³⁴ Another important feature we have considered here is to design boronate probes with an improved reactivity towards H_2O_2 as well as an optimized redox cycling capability, with the underlying idea to outcompete the background reactions.³⁶

As it will be shown, the single-protected probe **P**₂ generates a remarkably efficient autocatalytic reaction (LoD of 0.1 μM H_2O_2



Scheme 1. (A) General principle of the autocatalytic reaction scheme based on cross-activation of two catalytic loops. (B) Boronate-based probe previously used in ref 34. This probe is converted by H_2O_2 in an intermediate phenolic compound that self-immolates into a quinone methide and a 1,4-naphthohydroquinone. (C) The new probes used in the present study.

in less than 10 min at pH 8.5) thanks to a fast H_2O_2 -mediated deprotection rate (k_d) (faster than **P**₁). However, the lack of chemical stability of **P**₂ pushed us to opt for a double masking strategy, as exemplified with probes **P**₃ and **P**₄, which finally enable us to obtain much more stable boronate probes while keeping a good reactivity towards H_2O_2 . With these doubly protected probes, detection of nanomolar concentrations of H_2O_2 in less than 10 min was achieved using a simple UV-visible absorbance readout, which in terms of analytical performance is competitive with the best fluorescent-based molecular probes currently available on the market for quantitative analysis of H_2O_2 by fluorescence.³⁹⁻⁴¹ Finally, we will show that these excellent analytical performances could be obtained thanks to a comprehensive kinetic analysis of the complex cross-catalytic reaction network, thus providing an access to its quantitative prediction and rationalization. In particular, we were able to explain why the autocatalysis is still possible for doubly protected probes, even in the absence of instability of the monoprotected intermediate.

Results and discussion

Characterization of the single-protected probe **P**₂

The ability to induce an autocatalytic reaction by mixing **P**₂, H_2O_2 and a reducing agent has been investigated in a Tris buffer at pH

8.5. As a reductant, we used the same enzyme/reducing substrate as previously, *i.e.* the diaphorase/NADH couple.³⁴ The diaphorase (DI) from *Bacillus stearothermophilus* (EC 1.6.99.-) is a thermostable flavin mononucleotide-dependent oxidoreductase that efficiently catalyses the obligatory two-electron reduction of naphthoquinones into naphthohydroquinones in the pH range 7 to 10.⁴² It is also insensitive to O₂ and its NADH substrate is quite stable in slightly basic aerated buffers⁴³ (showing almost no cross-reactivity with dissolved dioxygen unless either O₂ or NADH is activated⁴⁴). The time course of the autocatalytic reaction was monitored by UV-visible spectrophotometry at $\lambda = 340$ nm, a wavelength characteristic of the transformation of NADH into NAD⁺. A spectrophotometric microplate reader for high-throughput kinetic analysis was used to simultaneously monitor a set of different reaction mixtures (200 μ L working solution per microwell). During the whole reaction, the microplate was left open to the atmosphere so as to operate under an approximately constant O₂ concentration.

Kinetic traces recorded at different H₂O₂ concentrations are shown in Fig. 1A. Their particularly steep sigmoidal shape is characteristic of an efficient autocatalytic process, for which the lag phase that precedes the exponential decrease of absorbance is a function of the triggering H₂O₂ concentration. Indeed, it is observed that the lower the starting H₂O₂ concentration, the longer the lag phase. It is also important to note the S-shaped kinetic curve obtained in the absence of H₂O₂ (blank) attests to a nonspecific autocatalysis triggered by undesirable reactions (the latter initiating the production of either H₂O₂ or Q/QH₂ to eventually lead to the exponential and full consumption of NADH). The time-to-threshold values (*i.e.*, $t_{1/2}$ defined here as the time required to reach 50% of the maximal absorbance change) extracted from these kinetic traces were used to plot, on a semi-logarithmic scale, the H₂O₂ calibration curve reported in Fig. 1B. The fact that this calibration plot shows a quasilinear relationship between $t_{1/2}$ and $\log[\text{H}_2\text{O}_2]$ over nearly two order of magnitude (*i.e.*, from 0.5 to 20 μ M H₂O₂) confirms the autocatalytic nature of the reaction. The deviation from linearity observed at the lowest concentrations (with a tendency to merge progressively with the $t_{1/2}$ value determined in the absence of H₂O₂) illustrates the limitation imposed by the nonspecific response on the analytical performances, leading here to a LoD of only 0.2 μ M H₂O₂, a value which is finally not much different from that we had previously obtained with the self-immolating probe **P**₁ under similar experimental conditions.³⁴ Nevertheless, the kinetics with **P**₂ are much faster than previously (by a factor 6), which is clearly an advantage to shorten the analysis time. The influence of pH was investigated by repeating the same experiments at two slightly less basic pHs (*i.e.*, pH 8 and pH 7.5). The resulting plots in Fig. S1 show a shift in the kinetic traces to longer times along with a reduction in the slope of the sigmoid as the pH decreases, a behaviour consistent with slower autoxidation and deprotection rates at lower pHs.³⁴ Unfortunately, no gain in LoD was obtained, suggesting a relatively similar pH dependence of both probe reactivity and stability.³⁶

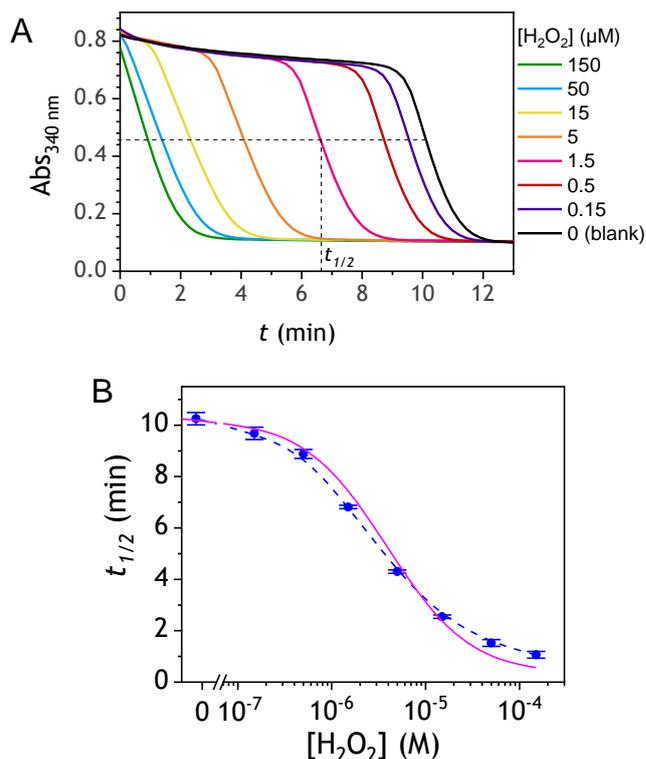


Fig. 1 (A) UV-vis kinetic traces monitored at 340 nm in a 96-wells microtiter plate during the reaction generated from the mixing of different concentrations of H₂O₂ (see the caption on the graph) to solutions containing 50 μ M **P**₂, 10 nM DI and 250 μ M NADH. All experiments were conducted in an air-saturated Tris buffer (0.1 M, pH 8.5) containing 10 μ M EDTA. Each curve is the average of triplicate experiments. (B) Semi-logarithmic calibration curve obtained from the threshold times ($t_{1/2}$) recovered from the crossing of the horizontal dashed line in A to the kinetic traces ($t_{1/2}$ is defined here as the time for which the absorbance is decreased approximately half to the maximal absorbance change). Errors bars are standard deviations from triplicates. The magenta solid line is the theoretical plot obtained by simulation (see text and ESI for details).

To understand why this new probe results in such a fast autocatalysis but also in this unexpectedly large nonspecific response, we have examined the fate of the probe alone in a Tris buffer (pH 8.5) and its reactivity towards H₂O₂. The rather fast evolution of the UV-vis spectrum of a dilute solution of **P**₂ (50 μ M) over 20 minutes (graph A in Fig. S2) indicates the relatively poor stability of **P**₂ in the aqueous buffer. Moreover, the lack of isobestic point reveals the probe degradation into several byproducts. Their structural identity was determined from analyses by HPLC and comparison of the UV-visible spectra of degraded probe with reference spectra of different byproduct candidates (see Figs. S2, S3, S4 and ESI for details). Independently of the naphthol boronic acid derivative generated from fast hydrolysis of the pinacol group, two byproducts were found, *i.e.* the 1-naphthol and 1,4-NQ (according to HPLC analysis, in a 1:3 ratio in the fully degraded probe solution). The formation of 1-naphthol is the

consequence of the slow first-order protodeboronation of P_2 , a well-established reaction for aromatic boronates in aqueous media,^{45, 46} while the formation of 1,4-NQ was attributed to the slow oxidation of C-B bond by dissolved O_2 (indeed, under argon, we observed that P_2 and its hydrolysed homologue only degrade into 1-naphthol – see Fig. S3). Since the mechanism of this slow probe autoxidation is unknown, we have at first assumed it can be assimilated to a bimolecular reaction between P_2 and O_2 , leading to H_2O_2 and 1,4-NQH₂. Even if slow, this probe autoxidation is clearly a major issue because, unlike protodeboronation which leads to a dead-end product, it triggers the catalytic degradation of P_2 *via* the production of 1,4-NQH₂ and regeneration of H_2O_2 , which in turns further deprotects P_2 until the latter is fully converted (Scheme 2). This undesirable reaction finally ends up in an important nonspecific response in the absence of H_2O_2 , which is of course detrimental to the analytical performances. Owing to the pinacol hydrolysis occurring together with the other two nonspecific competitive reactions, it was tricky to access the individual kinetics of these reactions. For the sake of simplicity, we have assumed that the 3 rate constants characterizing the probe protodeboronation (k_p), autoxidation (k_n), and H_2O_2 deprotection (k_d) are almost the same either P_2 is with or without its pinacol group. To determine these rates constants, we relied on a UV-vis absorbance measurement at 262 nm, a wavelength selective of the 1,4-NQ.³⁴ The kinetic traces in Figs. 2A and S5 were obtained either in the absence or presence of different H_2O_2 concentrations. Without H_2O_2 , the change in absorbance is characterized by a small induction period followed by a nonlinear increase until it asymptotically reaches a constant value after more than 20 min. This increase in absorbance indirectly measures the autoxidation rate of P_2 through the nonspecific catalytic formation of 1,4-NQ, from which we can estimate k_n . In the presence of H_2O_2 , the increase in absorbance starts sooner (and without delay) and faster as the H_2O_2 concentration is higher (Fig. S5). Furthermore, with the increase of $[H_2O_2]$, the maximum absorbance converges progressively towards a same maximal value, indicating complete probe transformation into 1,4-NQ. For the lowest and intermediate concentrations, the fact that the absorbance levels off at lower values suggests that a non-negligible fraction of the probe converts into another

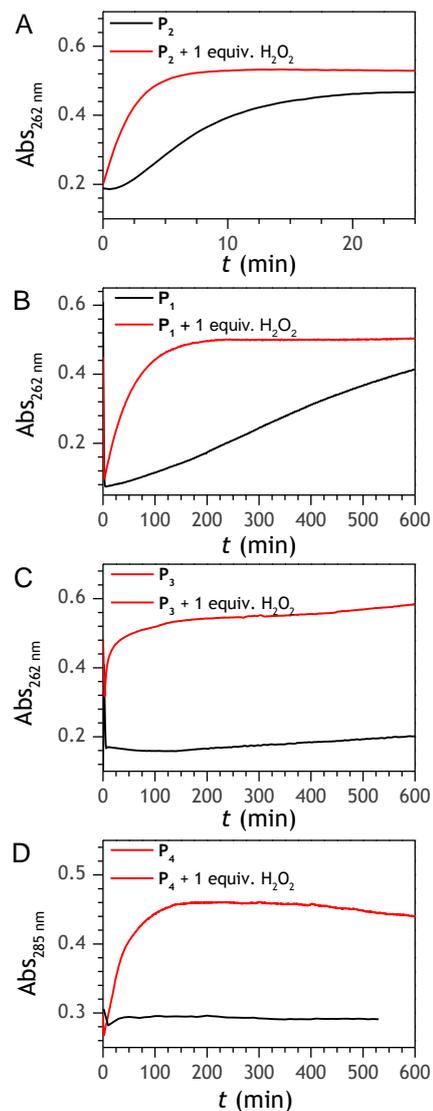
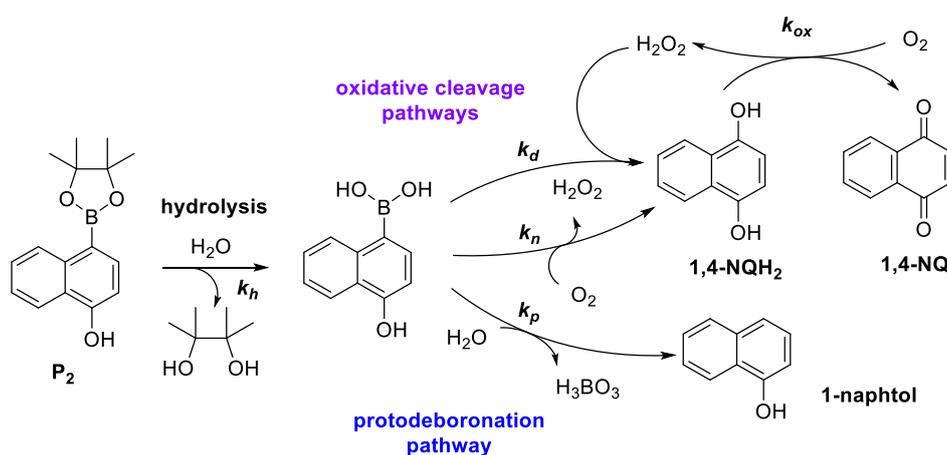


Fig. 2 UV-vis kinetic traces recorded at 262 or 285 nm (wavelengths specific of 1,4-NQ) for solutions containing 50 μ M of (A) P_1 , (B) P_2 , (C) P_3 , and (D) P_4 . The black lines correspond to the probe alone in solution, while the red lines correspond to a 1:1 mixture of probe and H_2O_2 . All experiments were performed at 25°C in an air-saturated Tris buffer (0.1 M, pH 8.5) containing 10 μ M EDTA.



Scheme 2. Reaction scheme associated to the instability of probe P_2 in aqueous solution.

compound, not detectable at this wavelength, which we attributed to 1-naphthol. The fast kinetics recorded in the presence of H_2O_2 demonstrate the fast H_2O_2 -mediated deprotection of P_2 , clearly much faster than previously reported for the self-immolative probe P_1 (the kinetic traces of which are reminded in Fig. 2B for comparison). The trade-off is that the nonspecific self-deprotection of P_2 is also faster than P_1 , therefore explaining the unsatisfactory detection limit of H_2O_2 . Assuming that the probe has similar reactivity, whether it is hydrolysed or not, and the set of reaction in Scheme 2 is adequate to account for the different pathways of P_2 transformation, one can expect to reproduce by simulation the experimental curves in Figs. 2A and S5. For such purpose, we have considered for the 1,4-NQ autoxidation the same global apparent bimolecular rate constant (k_{ox}) than the one we had previously electrochemically determined (*i.e.*, $3300 \text{ M}^{-1}\cdot\text{s}^{-1}$),³⁴ a value high enough for the autoxidation of 1,4-NQH₂ to not be rate limiting. From iterative adjustment of the three other rate constants, *i.e.* k_d , k_n and k_p , and assuming $[O_2]$ constant, it was possible to reproduce quite well the experimental kinetic traces in Fig. S5 using the following set of values: $k_d = 300 \text{ M}^{-1}\cdot\text{s}^{-1}$, $k_n = 1.2 \text{ M}^{-1}\cdot\text{s}^{-1}$, and $k_p = 4 \times 10^{-4} \text{ s}^{-1}$. The extracted k_d indicates a rather fast H_2O_2 deprotection, much faster than that previously determined for the arylboronate P_1 probe in a same Tris buffer ($\sim 5 \text{ M}^{-1}\cdot\text{s}^{-1}$).³⁴ To our knowledge this is even the highest rate constant so far reported for an aromatic boronate compound (at neutral or slightly basic pHs, H_2O_2 -mediated deprotection rate constants of arylboronates generally range from 0.07 to $4 \text{ M}^{-1}\cdot\text{s}^{-1}$ ^{45,47-51}). We attribute this to the presence of the hydroxy group in *para* position of the naphthylboronate, whose effect is to destabilize the C-B bond by inductive and/or mesomeric effects. This is supported by the much lower k_d value we separately determined for the 1-naphthylboronate in the same Tris buffer (*i.e.*, $7.5 \text{ M}^{-1}\cdot\text{s}^{-1}$). The protodeboronation rate is also significantly higher (by *ca.* one order of magnitude) than previously reported for arylboronic acids,⁴⁵ a behaviour we also attribute to the same electronic effects.

Investigation of the double-protected probe P_3

To address the instability issue of the single-protected probe P_2 , we hypothesized that double substitution of the -OH groups of 1,4-NQH₂ by boronates could make the probe more stable. Therefore, the probe P_3 was synthesized and then tested for its long-term stability in the Tris buffer. The data in Fig. 2C show the absorbance of a dilute P_3 solution (recorded at 262 nm, for an overview of the full spectrum see Fig. S6), which after a rapid decrease during the first few minutes (ensuing from the hydrolysis of pinacols), stabilizes at ~ 0.17 over several hours. This result clearly demonstrates the high chemical stability of P_3 , which once hydrolysed into diboronic acid does no longer decomposes into 1,4-NQ, therefore validating our initial hypothesis. This much higher chemical stability also correlates with a slower protodeboronation rate since the kinetic trace recorded at 290 nm (specific to the probe absorbance) is also very stable over several hours (see Fig. S7). Adding one equivalent of H_2O_2 to the solution results in a fast absorbance increase at 262 nm, rising to a maximal value in less than 5 min (red plot in Fig. 2C). It thus attests to an efficient H_2O_2 -mediated deprotection of P_3 into 1,4-NQ and so to a high responsiveness of the doubly protected probe to H_2O_2 . In order to determine the deprotection rate of P_3 , we repeated the experiment at different H_2O_2 concentrations (Fig. S8). As expected, the kinetics show a faster increase in absorbance as the H_2O_2 concentration is higher. However, unlike P_2 the maximal absorbance change scales with the stoichiometry of H_2O_2 (see ESI and Fig. S8 for details), which means that for each substoichiometric H_2O_2 concentration added to the solution of P_3 , only the corresponding substoichiometric equivalent in [1,4-NQ] can be recovered. This distinct behaviour of P_3 can be explained from the need to provide two equivalents of H_2O_2 for full deprotection of one equivalent, but also from the ability to regenerate an additional equivalent of H_2O_2 from each released molecule of 1,4-NQH₂. This also points out that the autoxidation of 1,4-NQH₂ is an efficient process for producing H_2O_2 at a yield close to 100% (a result which agrees with what was previously reported for the autoxidation of 1,4-NQ⁵²). The experimental kinetics reported in Fig. S8 should therefore fulfil the mechanism presented by reactions S5 to S9 in ESI, wherein P_3 is first converted into the monoprotected intermediate P_2 and

then into 1,4-NQ according to the same H₂O₂-mediated catalytic transformation described above for P₂. This reaction scheme is therefore characterized by two deprotection rate constants, namely k_{d,P_3} and k_{d,P_2} . The value of k_{d,P_2} can be considered the same than the one determined above for P₂, a consideration which remains also valid for all the other rate constants, the only new rate being k_{d,P_3} . We have thus numerically simulated the set of reactions S5 to S9 using the value k_{d,P_2} , k_{ox} , k_n and k_p previously recovered from P₂ and iteratively adjusted the rate constant k_{d,P_3} . The best fits of the simulated curves to the experimental data are shown in Fig. S8 (dashed plots in graph D) using $k_{d,P_3} = 35 \text{ M}^{-1}\cdot\text{s}^{-1}$. Despite all of our attempts, it was not possible to perfectly fit the data, the simulated plots showing a systematic slower increase of 1,4-NQ with time whatever k_{d,P_3} . In fact, much better fits were obtained by artificially considering the second deprotection rate fast enough (e.g., $k_{d,P_3} > 3000 \text{ M}^{-1}\cdot\text{s}^{-1}$) to no longer interfere on the kinetics (solid plots in graph D of Fig. S8). This apparent inconsistency suggests that in parallel to the H₂O₂ deprotection of the intermediate P₂, additional fast reactions contribute to an accelerated deprotection of P₂. We emitted the hypothesis that it could be due to the autoxidation of P₂ (reaction S8), which instead of a two-electron transfer reaction, as we have at first assumed, occurs through a one-electron transfer between O₂ and P₂ with the concomitant production of radical species (notably O₂[•]), which by dismutation can produce H₂O₂ for additional deprotection of both P₃ and intermediate P₂). One can then conceive that these radical species can either directly or indirectly further contribute to the accelerated deprotection of the intermediate P₂. We will come back later on this specific point. It is to note that the value of $k_{d,P_3} = 35 \text{ M}^{-1}\cdot\text{s}^{-1}$ recovered from the best fits, even if lower than k_{d,P_2} , remains high and attractive for an efficient autocatalysis.

Autocatalysis with P₃

The autocatalysis with P₃ was performed under the same conditions than P₂. The resulting kinetics curves (see Fig. 3A) characterizing the NADH consumption as a function of time, show, similarly to P₂, well-defined S-shaped kinetics traces with longer lags as the triggering H₂O₂ concentration is lower. However, a main difference with P₂ is the much slower kinetics, requiring for the lowest concentrations more than 5 hrs to reach $t_{1/2}$, while it takes less than 10 min with P₂. Such a huge difference can be understood from the slower deprotection rate of P₃ with respect to P₂, as well as from the need of two equivalents of H₂O₂ to fully deprotect each P₃ molecule. The analytical performances grasped from the semi-logarithmic calibration plot in Fig. 3C indicate a dynamic concentration range over more than two decades (i.e., from 0.1 to 20 μM H₂O₂) and a LoD of 0.2 μM H₂O₂. These analytical performances are not as good as expected, leading to a LoD similar to that of P₂, which is surprising if we consider the much higher chemical stability of P₃. In order to improve the reaction rate and at the same time the analytical performances, we reproduced the autocatalysis at a slightly more basic pH (pH 9.5). The resulting autocatalytic responses in Fig. 3B show a clear kinetic gain, reducing the $t_{1/2}$ values to a time window < 40 min. Compared

to the results obtained at pH 8.5, this corresponds to an analysis time improvement by a factor 10. Moreover, the extracted H₂O₂ calibration curve (Fig. 3C) demonstrates a better LoD of ~20 nM, which is again a 10-fold improvement compared to pH 8.5 (Fig. 3D). Clearly, the pH has a strong impact on the analytical performances of the system, since by changing the pH of one unit, both LoD and analysis time are improved 10-fold. All these results led us to investigate what could lead to such a drastic pH

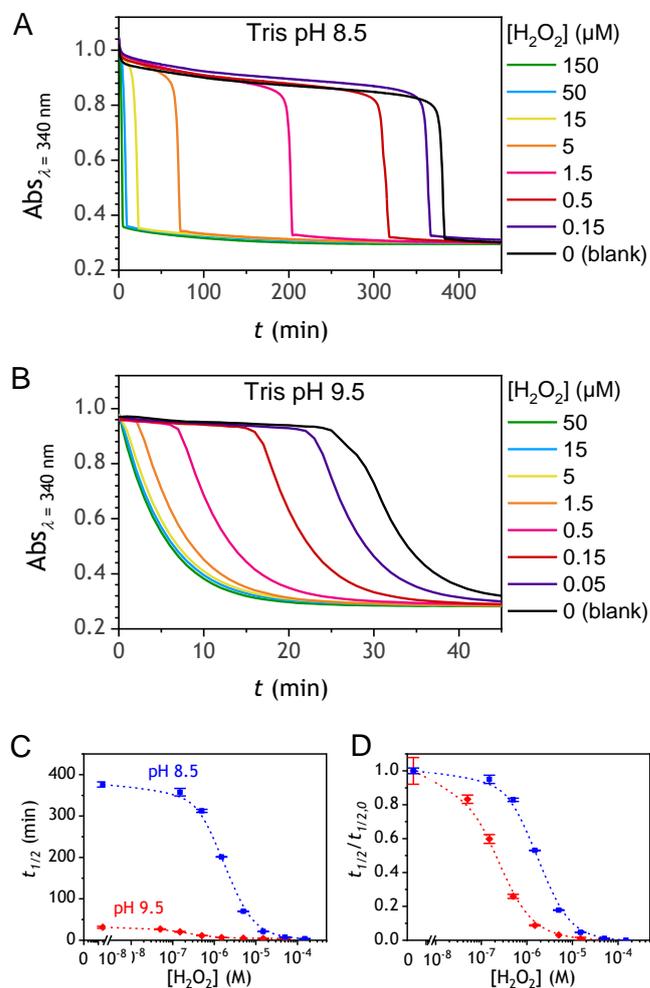


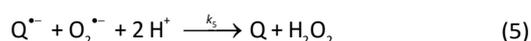
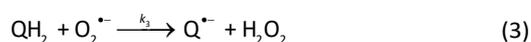
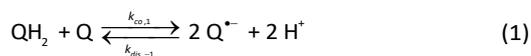
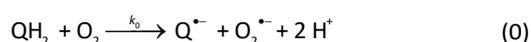
Fig. 3 (A, B) UV-vis kinetic traces monitored at 340 nm immediately after the injection of different concentrations of H₂O₂ (see the caption on the graph) to solutions containing 50 μM P₃, 10 nM DI and 250 μM NADH. The experiments were conducted in an air-saturated Tris buffer (0.1 M) of (A) pH 8.5 or (B) pH 9.5. Each curve is the average of triplicate experiments. (C) Semi-logarithmic calibration curves obtained from the plot of $t_{1/2}$ (recovered from the top and middle kinetic traces) as a function of $\log[\text{H}_2\text{O}_2]$. Red dots: pH 9.5. Blue dots: pH 8.5. Errors bars are standard deviations from triplicates. (D) Same graphs than in C but normalized to the $t_{1/2,0}$ recovered from the blank.

effect, but also how to interpret the large rate difference between P₃ and P₂. To this end, we examined the possibility to simulate the autocatalytic kinetic traces on the basis of a reasonable set of reactions for describing the cross coupling

between the H₂O₂-mediated catalytic deprotection of **P**₃ (*Loop 1*) and the enzyme-assisted redox cycling (*Loop 2*) (see thereafter).

Mechanistic and kinetics analysis of the autocatalytic process

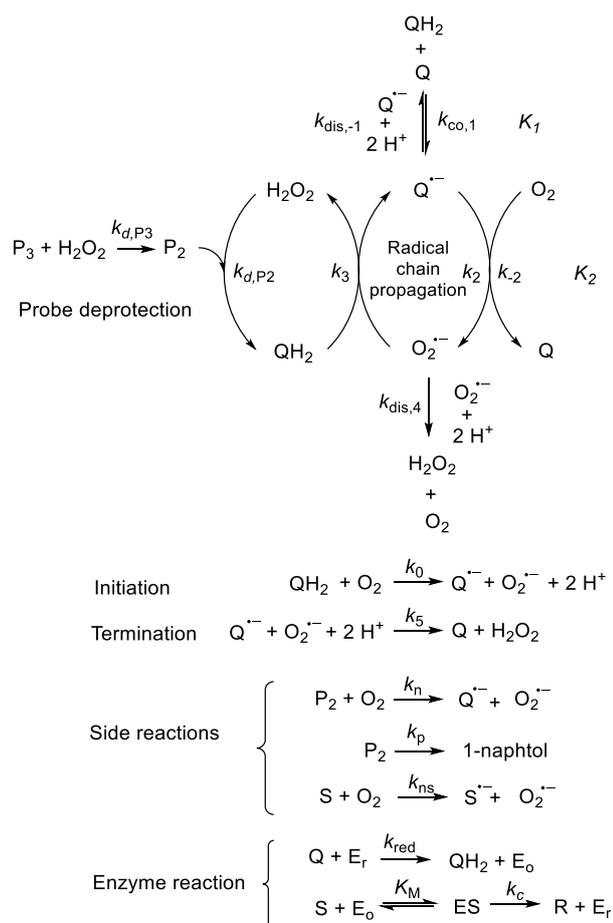
The catalysed reduction of quinones by DI is a well-established “ping-pong” mechanism,⁴² which under turnover conditions can be defined by two half enzyme reactions (and three rate constants), *i.e.* (i) a two-step Michaelis-Menten type reaction between NADH and the oxidized form of the enzyme (characterized thus by a Michaelis-Menten constant K_M and a catalytic turnover rate constant k_c) and (ii) a bimolecular reaction characterizing the obligatory two-electron transfer reduction of quinones (here the 1,4-NQ) by the reduced form of the enzyme (governed thus by a second-order rate constant k_{red}). The coupling of this enzymatic “ping-pong” mechanism to the H₂O₂-mediated deprotection of **P**₃ leads to the autocatalytic set of reactions S10 to S16 in ESI. To simulate this mechanism, we first used for *Loop 2* the values of K_M , k_c and k_{red} (*i.e.*, 60 μM, 1200 s⁻¹, 1.5 × 10⁹ M⁻¹·s⁻¹, respectively) previously determined by us for an analogue of the 1,4-NQ, *i.e.* the menadione,⁴² while the other rate constants associated to *Loop 1* were the same as those above determined (for clarity, the list of rate constants are gathered in the entry 1 of Table 1). The simulated time course of NADH consumption at different H₂O₂ concentrations are shown in Fig. S9. Their sigmoidal shapes are, as expected, in agreement with an autocatalytic reaction and also similar with those obtained experimentally (Fig. 3A). However, a main issue is that their $t_{1/2}$ do not match those obtained experimentally, differing by a factor as high as 28. This result indicates that something is missing in the proposed mechanism. We figured out that it might be due to our oversimplified description of the autoxidation of 1,4-NQ. It is well established that the autoxidation mechanism of hydroquinones is more complex than a simple bimolecular reaction. It is indeed an autoinductive radical chain mechanism, in which the semiquinone (Q^{•-}) and anion superoxide are involved as chain propagating species as follow:⁵²⁻⁵⁴



This chain reaction is initiated by reaction 0, which is however considered to be very slow under near neutral conditions due to its spin-forbidden nature.⁵⁵ However, once a small amount of Q^{•-} is produced, it is able to trigger a cascade of faster reactions, notably reactions 1, 2 and 5 (forming a set of propagation reactions), which results in an autoinductive acceleration of the oxidation of 1,4-NQ by O₂. Through this reaction network, QH₂

and O₂ are consumed by reactions 1, 3 and 2, respectively, while Q is produced in oxidation reactions 2 and 5. This reaction scheme predicts a 1:1 stoichiometry for the autoxidation of QH₂, leading thus to a 1:1 production of Q and H₂O₂. The rate acceleration of this mechanism originates from the product Q, which is the catalyst of its own formation through the comproportionation reaction 1. The majority of these reactions are pH-dependent but, for sake of simplicity, the rate constants used in this work are those at an effective pH (here pH 8.5). The rate constant values we have used are gathered in Table 1. Most of them were retrieved from the literature. A few were corrected for pH, while others were adjusted to fit the experimental data (see ESI for a detailed description of the rate constant selection).

Incorporating the above autoxidation mechanism of 1,4-NQH₂ to the coupled enzymatic reduction of 1,4-NQ and H₂O₂-mediated deprotection of **P**₃ leads to the autocatalytic set depicted in Scheme 3. Furthermore, since **P**₃ is very stable and therefore unlikely to be the source of the nonspecific triggering of autocatalysis, the alternative is to consider it originates from the slow autoxidation of NADH (characterized by the rate k_{ns}). In our model, this nonspecific reaction was assumed to be a one-electron transfer process, leading to the release of O₂^{•-} and NAD[•] (defined as S[•] in Scheme 3). Moreover, as we will see later, this reaction is apparently catalysed by DI (NADPH oxidase-like activity), which leads to a dependence of the nonspecific response on the enzyme concentration. Since there is no available information on the autoxidation kinetics of NADH in the literature (either it is catalysed or not by DI), the value of k_{ns} in our simulations was simply adjusted so that the simulated nonspecific autocatalytic response aligns with the experimental one. In this model, we also substituted the two-electron autoxidation of the intermediate **P**₂ by a one-electron transfer reaction so as to better reflect the radical chain reaction we suspect coupled to this reaction. The S-shaped kinetic curves numerically calculated using the model in Scheme 3 and the associated rate constants gathered in entry 2 of Table 1 are shown in Fig. S9, while the theoretical H₂O₂ calibration curve derived from these plots is given in Fig. 4 (green line). Compared to the previous simulations involving just a simple second order reaction between 1,4-NQH₂ and O₂, the new computed kinetics traces are now much closer to the experimental ones, a results which highlights the importance to consider the overall complexity of 1,4-NQH₂ autoxidation. Interestingly, the discrepancy we observe between simulations and experiments is clearly on the side of the lowest H₂O₂ concentrations (Fig. 4), since at the highest concentrations the theoretical plot fits the experimental data almost perfectly. We suspected that this could be related to an inhibition of the autoxidation of **P**₂ intermediate induced by the presence of the reducing enzyme/NADH substrate. As we have already mentioned, the autoxidation reaction mechanism of **P**₂ could well be of a radical nature accompanied by a chain process. If we assume that this is the case, then one can consider that such a radical chain reaction is promoted under oxidizing conditions and inhibited under reducing conditions. To evaluate such an effect on the autocatalysis *via* simulation, we disabled in our model the autoxidation reaction of the monoboronic intermediate by



Scheme 3. Overall reactions involved in the autocatalytic deprotection of P_3 in the presence of the reducing NADH/DI couple and H_2O_2 . The enzyme substrate NADH and product NAD^+ are here symbolized by S and R, and the different states of the diaphorase enzyme by E_o (oxidized), E_r (reduced) and ES (enzyme/substrate precursor complex).

setting its k_n value to zero. Thanks to this, we were able to obtain the magenta simulated plot in Fig. 4, which now predicts the experimental data much more closely. This finding tends to support the hypothesis that the slow self-deprotection of P_2 by O_2 occurs through a one-electron transfer reaction with concomitant formation of radical species (possibly $\text{O}_2^{\bullet-}$ and a naphthoxyl radical), which are then inhibited or scavenged under the reducing conditions (NADH has been shown to efficiently scavenge either HO_2^{\bullet} ⁵⁶ or phenoxyl radicals⁵⁷ by one-electron reduction). This also demonstrates that the double masking does not prevent the autocatalytic reaction from working, even when the instability of the monoprotected intermediate is switched-off (*i.e.* when $k_n = 0$). Indeed, when $k_n = 0$, the autocatalysis still occurs with a reasonable efficiency

(magenta curves in Fig. 4) because, statistically, the probability of having a double-deprotection of the probe is not null, even under the substoichiometric H_2O_2 conditions used here. In fact, this probability depends directly on the ratio between the first and the second deprotection rate constants (*i.e.*, k_{d,P_2} and k_{d,P_3}), which is more favourable when the first one is slower than the second, as it is the case here (*i.e.*, 35 vs. 300 $\text{M}^{-1}\cdot\text{s}^{-1}$, respectively).

As the magenta simulated plot in Fig. 4 still does not completely superimpose to the data, we considered that it could be due to the uncertainty in the selected values of k_3 and k_{red} (the latter being moreover recovered from a previous enzymatic study performed with menadione but not with the 1,4-NQ). To experimentally access the value of these two rate constants, we have performed redox cycling experiments (*Loop 2*) by mixing a fixed concentration of DI (10 nM) and NADH (250 μM) with different concentrations of 1,4-NQ and then monitoring the NADH absorbance decay (see graph A in Fig. S10). From the results reported in Fig. S10 and analysed in details in ESI, we were able to estimate the values of $k_3 = (5 - 8) \times 10^4 \text{ M}^{-1}\cdot\text{s}^{-1}$ and $k_{\text{red}} = (1.5 - 0.8) \times 10^8 \text{ M}^{-1}\cdot\text{s}^{-1}$ with a reasonably good degree of confidence. We thus considered these new values in our simulations and we succeeded to obtain an almost perfect agreement between the simulated calibration plot and the experimental data using $k_{\text{red}} = 10^8 \text{ M}^{-1}\cdot\text{s}^{-1}$ and $k_3 = 5 \times 10^4 \text{ M}^{-1}\cdot\text{s}^{-1}$ (blue curve in Fig. 4). This latter result definitely validates the model in Scheme 3 as well as the selected set of rate constants. With this tool in hand, it was next interesting to go back to the data with P_2 in order to see to what extent the autocatalytic kinetic traces in Fig. 1 can be predicted by the model in Scheme 3 and the associated rate constants (excluding of course from the model the first deprotection, *i.e.* k_{d,P_3} , which does not take place with P_2). The best-simulated curves are reported in Fig. S12 using the rate constants gathered in the entry 6 of Table 1. To obtain such a good fit, it was necessary to adjust the rate constant k_n to a non-null value, which is not surprising as for simulating the nonspecific autocatalytic response of P_2 , it is now necessary to consider the probe autoxidation as the main source of nonspecificity (thus controlled by k_n) and no longer the NADH autoxidation (k_{ns}). A k_n of 0.25 $\text{M}^{-1}\cdot\text{s}^{-1}$ was thus set from the alignment of the simulated nonspecific response to the experimental one. This value is significantly lower than the value of 1.2 $\text{M}^{-1}\cdot\text{s}^{-1}$ previously derived from the H_2O_2 -mediated deprotection of P_2 , therefore confirming that there is an inhibition of the autoxidation of P_2 under the reducing conditions of autocatalysis. The theoretical H_2O_2 calibration plot that was finally extracted from the simulated kinetic traces fits nicely to the experimental calibration plot in Fig. 1B. This is a strong plea for the validity of the proposed model. It also demonstrates the excellent self-consistency of our results. Armed with this knowledge, it was next interesting to identify the keys rate constants controlling the rate of autocatalysis but

Table 1. Rate constants values used for the different autocatalytic kinetic simulations

	Rate constant	Simulation entry ^a					
		1	2	3	4	5	6
Probe deprotection reactions	k_{d,p_1} ($M^{-1} s^{-1}$)	35	35	35	35	35	/
	k_{d,p_2} ($M^{-1} s^{-1}$)	300	300	300	300	300	300
Side reactions	k_n ($M^{-1} s^{-1}$)	1.2	1.2	0.0	0.0	0.0	0.25
	k_p (s^{-1})	0.0004	0.0004	0.0004	0.0004	0.0004	0.0004
	k_{ns} ($M^{-1} s^{-1}$)		10^{-4}	2.75×10^{-3}	4.1×10^{-3}	4.1×10^{-3}	4.1×10^{-3}
Enzyme reactions	K_M (μM)	60	60	60	60	60	60
	k_c (s^{-1})	1200	1200	1200	1200	1200	1200
	k_{red} ($M^{-1} s^{-1}$)	1.5×10^9	1.5×10^9	1.5×10^9	10^8	10^8	10^8
1,4-NQH ₂ autoxidation reactions	k_{ox} ($M^{-1} s^{-1}$)	3300	/	/	/	/	/
	k_0 ($M^{-1} s^{-1}$)		0.001	0.001	0.001	0.001	0.001
	$k_{co,1}, k_{dis,-1}$ ($M^{-1} s^{-1}$)		$66000; 10^8$	$66000; 10^8$	$66000; 10^8$	$6.6 \times 10^6; 10^8$	$66000; 10^8$
	k_2, k_{-2} ($M^{-1} s^{-1}$)		$4.4 \times 10^7; 2 \times 10^8$				
	k_3 ($M^{-1} s^{-1}$)		8×10^4	8×10^4	5×10^4	4.7×10^6	5×10^4
	k_4 ($M^{-1} s^{-1}$)		2×10^4				
	k_5 ($M^{-1} s^{-1}$)		2×10^9				

^a Are highlighted in blue the change made when going step-by-step from the first simulation entry to the last one.

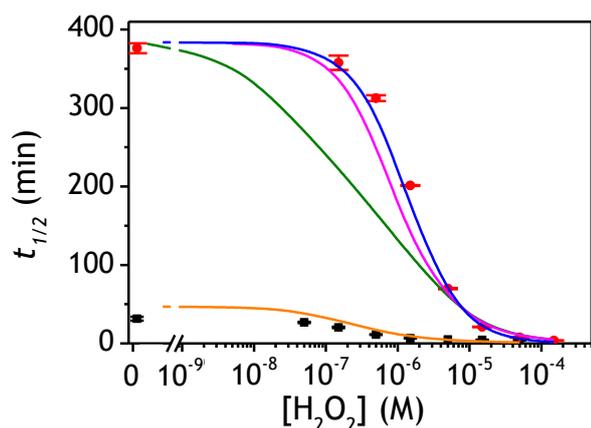


Fig. 4 Plots of the experimental and theoretical semi-logarithmic H₂O₂ calibration curves. The red and black dots are the experimental data recovered for P₃ at pH 8.5 and 9.5 respectively (same data than in Figs. 3C and D), while the solid lines are the theoretical plots predicted from numerical simulations of the autocatalytic set presented in Scheme 3. The parameters used for each simulated plots are those reported in Table 1 according to the following code color: (green) entry 2, (magenta) entry 3, (bleu) entry 4, and (orange) entry 5.

also to determine what limits the analytical performances of the method.

Rationalization of the effect of pH on the kinetics and LoD

To understand the influence of the pH on the rate of autocatalysis (as illustrated in Fig. 3), one has to start by identifying the rate constants that *a priori* are the most strongly pH dependent. On account of the two pK_as of the 1,4-NQH₂ (pK_{a,1} = 9.48 and pK_{a,2} = 11.02, see ESI), the rate constants k_0 ,

$k_{co,1}$ and k_3 are clearly much pH sensitive than the other rate constants of the model (since this is HO₂⁻ the reactive species involved in the oxidative cleavage of boronates,⁵⁸ the constants k_{d,p_1} or k_{d,p_2} are also pH dependent but to a lesser extent than k_0 , $k_{co,1}$ and k_3). Therefore, if we consider $k_{co,1}$ and k_3 (k_0 is not taken into account for the same reason as previously) are mainly responsible for the pH dependency of the autocatalysis rate, this should be verifiable through the simulation of the H₂O₂ calibration curve. From the two pK_as of the 1,4-NQH₂, we can calculate the values of $k_{co,1}$ and k_3 at pH 9.5, which are $6.6 \times 10^6 M^{-1} \cdot s^{-1}$ and $4.7 \times 10^6 M^{-1} \cdot s^{-1}$, respectively (see ESI). The theoretical H₂O₂ calibration curve (orange line in Fig. 4) determined using these new values of $k_{co,1}$ and k_3 (while keeping the other rate constants the same as previously at pH 8.5) finally agrees quite well with the experimental plot at pH 9.5 (black squares in Fig. 4). This result confirms the key role played by both $k_{co,1}$ and k_3 in the pH-rate dependence of the autocatalysis. A better fit to the data would certainly be obtained by increasing slightly the two deprotection rate constants k_{d,p_1} and k_{d,p_2} , which, as mentioned above, are also expected to rise with the pH. This kinetic analysis also explains the better detection limit obtained by increasing the pH (Fig. 3D). Indeed, if we admit that the NADH autoxidation (k_{ns}) (which determines the nonspecific response of the system and thus the LoD) varies less rapidly with the pH than the overall rate of autocatalysis (varying predominantly with the pH through $k_{co,1}$ and k_3), then we can understand why the LoD is 10-fold improved when the pH is increased from 8.5 to 9.5.

Further improving the analytical performances

To determine whether the analytical performance could be further improved and/or optimized, we performed the autocatalysis reaction in buffers of different nature adjusted at different pHs (it is to note that DI from the *Bacillus Stearothermophilus* has a bell-shaped activity in the pH range

6.5 to 10.5, with an optimum at pH 8.0⁵⁹). Among the different organic and inorganic buffers investigated (see ESI and Figs. S13 to S20), the glycine buffer at pH 10 was found superior. The analytical performances obtained with this buffer were indeed clearly improved (Fig. S18), leading to a dynamic concentration range over nearly 3 decades (*i.e.*, from ca. 0.01 to 10 μM H_2O_2) and a LoD as low as 5 nM H_2O_2 in less than 10 min reaction time. This is a very appealing performance since in terms of LoD, it is competitive with the best H_2O_2 detection kits currently commercially available (a LoD of 10 nM H_2O_2 within 30 min incubation time is reported for the popular fluorescent-based Amplex red[®] hydrogen peroxide/peroxidase assay⁴¹). Another buffer particularly attractive is the CAPS buffer at pH 9.5 (Fig. S20). In spite of a slightly longer reaction time, the data indicates the possibility to reach the nanomolar H_2O_2 concentration range in less than 50 min. It is to note that with some buffers the autocatalysis was found to be inefficient in producing well-defined or discernible autocatalytic responses. This was especially true with the borate buffer (Fig. S17). It was also observed in some buffers an anomalous shift of the nonspecific blank response to shorter times compared to the responses recorded at the lowest H_2O_2 concentrations. All these behaviours suggest that the buffer plays an active role in the autocatalysis that we have not yet been able to elucidate.

Another parameter we examined to improve the analytical performances was the concentration of DI. As shown in Fig. 5, the concentration of the reducing enzyme was found to have a considerable impact on the nonspecific autocatalytic response, dramatically accelerating the nonspecific $t_{1/2}$ value with the rise of DI concentration, an effect which suggests that the enzyme slowly catalyses the reaction between NADH and O_2 to produce $\text{O}_2^{\cdot-}$ or H_2O_2 . These results echo the residual NADH oxidase activity (< 0.01%) reported by the enzyme supplier and which would be related to the presence of traces of NADH oxidase in the purified diaphorase samples. However, another possibility that cannot be ruled out is that DI has an intrinsic residual NADH oxidase activity. Indeed, one can assume that the reaction with O_2 is not necessarily inexistent even though this flavoenzyme is known to not accept O_2 as an electron acceptor. By gradually decreasing the concentration of DI, we can see in Fig. 5 that the value of $t_{1/2}$ increases until it reaches a maximum before slightly decreasing at the lowest concentration (graph B in Fig. 5). This indicates that by lowering the concentration of DI, the nonspecific contribution associated to the residual NADH oxidase activity is reduced. This can therefore be exploited to improve the specific vs. nonspecific response, especially if the enzyme reduction rate of the naphthoquinone in *Loop 2* continues to be not rate limiting. This latter attribute can be easily verified from the S-shape of the autocatalytic response. Indeed, if the kinetic profiles are similar regardless of the enzyme concentration, it suggests that the naphthoquinone reduction is not rate limiting. On the other hand, when the rate of the naphthoquinone reduction becomes too slow, as illustrated for the lowest enzyme concentration (*i.e.* 0.1 nM in Fig. 5), the kinetic trace presents a more spread-out autocatalytic response, signing thus the slowing down of *Loop 2*. The kinetic gain also observed at the lowest enzyme

concentrations can be explained from the redox cycling (*Loop 2*) in Scheme 3, which tends to be inhibited at high DI concentration by forcing the naphthoquinone to be rapidly doubly reduced, thus bypassing the formation of seminaaphthoquinone indispensable to sustain the free radical chain reaction. This inhibitory effect is relaxed when the enzyme concentration becomes sufficiently low, which finally leads to a slightly faster autocatalytic reaction. Considering the

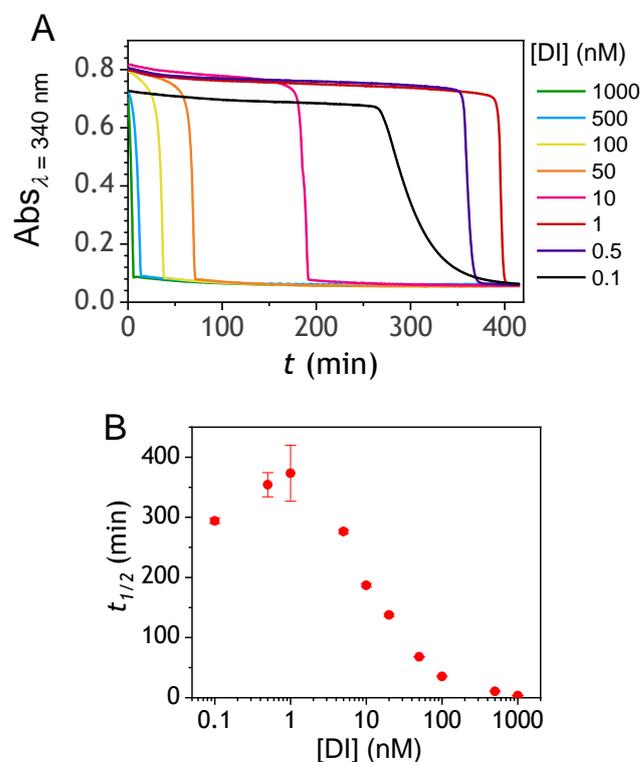


Fig. 5 (A) Influence of the reducing enzyme concentration on the nonspecific autocatalytic response. The kinetic traces at 340 nm were recorded immediately after 50 μM P_3 was mixed with 250 μM NADH and different concentrations of DI (see the legend on the graph) in an air-saturated Tris buffer (0.1 M) at pH 8.5. (B) Semi-logarithmic curve showing the influence of the DI concentration on the $t_{1/2}$ values recovered from the nonspecific autocatalytic responses.

importance of the enzyme concentration on the nonspecific response, we reproduced the autocatalysis experiments at pH 8.5 with a DI concentration 100 times lower, *i.e.* 0.1 nM. The results in Fig. 6 show a significant gain in the H_2O_2 detection limit, allowing now to detect as low as 15 nM H_2O_2 . Compared to the previous experiments performed at a higher enzyme concentration, this corresponds to a 10-fold gain in LoD (compare Figs. 3 and 6).

To illustrate the potential of the method to indirectly detect the activity of an oxidase through its H_2O_2 production, we performed the quantitative detection of glucose oxidase (GOD) under the experimental conditions used above. Indeed, in the presence of glucose, GOD catalyses the oxidation of glucose into gluconolactone with concomitant production of H_2O_2 from catalytic reduction of dissolved O_2 . The results gathered in Fig.

S23 demonstrate the feasibility of the approach, allowing indirect quantification of GOD by autocatalytic molecular amplification down to subpicomolar concentrations.

Triggering the autocatalysis with a naphthoquinone

A compelling property of the present cross-catalytic reaction scheme lies in its ability to be triggered not only by H_2O_2 but also any redox cycling compound (*i.e.*, by definition a redox-active compound able in the presence of O_2 and a reducing species to cycle and to generate reactive oxygen species). This

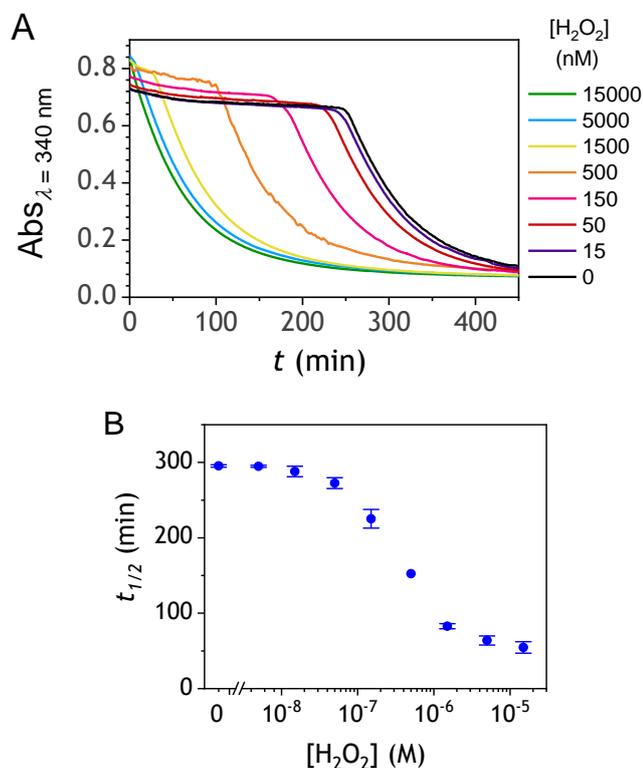


Fig. 6 (A) Autocatalytic responses obtained with 0.1 nM DI, 50 μM P_3 , 250 μM NADH and different H_2O_2 concentrations (see the legend on the graph) in an air-saturated Tris buffer (0.1 M) at pH 8.5. Each curve is the average of triplicate experiments. (B) Semi-logarithmic H_2O_2 calibration curve extracted from kinetics experiments analogous to those reported in A. Errors bars are standard deviations from triplicates.

makes this approach potentially attractive for the design of new sensitive bioassays.^{60, 61} This can be for instance through the upstream production of a redox cycling compound *via* an enzyme or an enzymatic label, which can then trigger *via* one of the above boronic probes an exponential molecular amplification. With the aim to evaluate the ability of P_3 to be triggered by low concentrations of a given redox cycling compound, we have investigated the autocatalysis in the presence of different concentrations of 2-amino-3-chloro-1,4-naphthoquinone (from 0.15 nM to 0.15 μM), a compound we had previously identified as an efficient redox cycling compound.³⁴ The results gathered in Fig. 7 demonstrates particularly good analytical performances, allowing to easily detect as low as 150 pM of 2-amino-3-chloro-1,4-NQ in less than

30 min, with a LoD estimated close to 50 pM. As for H_2O_2 , the dynamic range of concentrations is also wide, extending over nearly 3 orders of magnitude. These analytical performances are definitely much better than those obtained with H_2O_2 . This asymmetry in the analytical performances was shown to be an intrinsic property of a cross-catalytic reaction scheme.³⁶ Indeed, if the two catalytic loops have similar rates, the detection efficiency of the catalyst of *Loop 1* or *Loop 2* is theoretically the same, thus leading to an identical LoD of each trigger. On the other hand, if the rates of each loop are different, the trigger of the fastest loop becomes the most easily detectable and therefore the one that theoretically leads to the lowest LoD. This is exactly what we observe here with the trigger naphthoquinone, which, due to its involvement in a faster catalytic loop (*Loop 2*) is detected more sensitively than H_2O_2 .

Towards an improvement with the design of a diboronate probe able to release a better redox cyclor

Improvement of the rates associated to each loop of the cross-catalysis reaction is a strategy to shorten the analysis time and to potentially enhance the detection limit by better discriminating the specific from nonspecific responses.³⁶ In the present case, the background amplification stems from the instability associated to *Loop 2* (*i.e.*, from NADH autoxidation or NADH-oxidase like activity of diaphorase) since the double masking of the probe makes it so stable that no nonspecific response can arise from *Loop 1*. It is therefore particularly relevant to find a strategy that aims at improving the global rate of *Loop 2*. To do so, a relatively straightforward method is to substitute the masked 1,4-NQH₂ in P_3 by an alternative faster redox cycling naphthohydroquinone. A series of different naphthoquinones were thus screened for their redox cycling properties and then compared to the 1,4-NQ (see ESI and Fig. S21 for details). From this screening, three naphthoquinones were identified as a better redox cyclor than the 1,4-NQ, *i.e.* the 2-amino-3-chloro-1,4-NQ, 1,2-NQ and 2-methoxy-1,4-NQ. The chemical stability of these three naphthoquinones in an aqueous buffered solution was also examined and the most stable was the 2-methoxy-1,4-NQ, showing almost no change in the absorbance spectra after several hours in a Tris buffer (graph C in Fig. S22). The chemical synthesis of the corresponding diboronate probe P_4 was thus planned and successfully achieved (Scheme 1C). The long-term stability of a diluted solution of P_4 was shown to be quite good (Fig. S22 and 2D), demonstrating a chemical stability of P_4 at least as good as P_3 . Concerning the H_2O_2 -mediated deprotection of P_4 (Fig. 2D), it was found a little slower than P_3 (by a factor ~ 6). We attribute this slower rate to the methoxy group in *ortho* position, which by steric effect can slow-down the nucleophilic attack of HO_2^- . The autocatalytic kinetic traces collected with P_4 over ~ 90 min at pH 8.5 attest of a rather fast autocatalytic reaction (Fig. 8A), approximately 4-fold faster than P_3 at the same pH. This gain in analysis time is clearly the result of the better redox cycling properties of 2-MeO-1,4-NQ as compared to 1,4-NQ, compensating thus for the slower H_2O_2 deprotection rate. This gain in time also almost completely translates into a gain in LoD since the detection limits for P_3 and P_4 are 0.2 and 0.06 μM H_2O_2 , respectively (Fig. 8C). The fact that the faster kinetics with

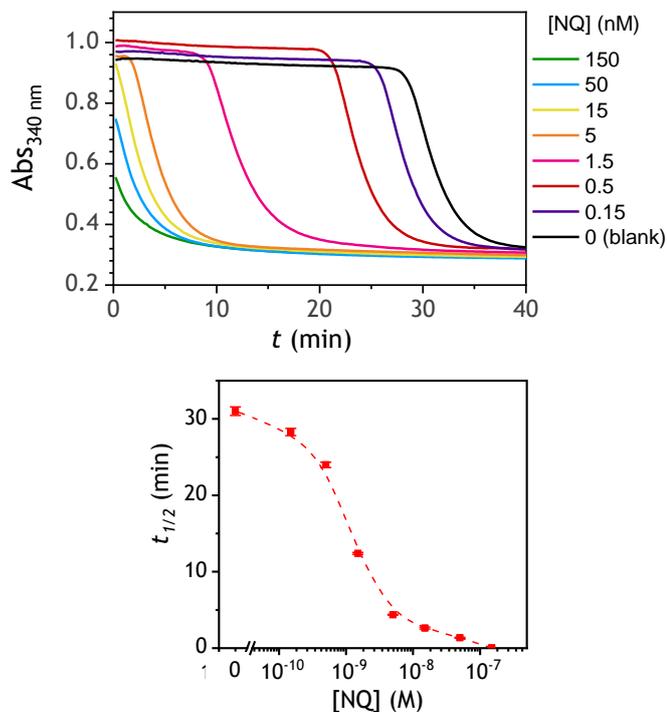


Fig. 7 (Top) UV-vis kinetic traces (average of triplicates) monitored at 340 nm immediately after the injection of different concentrations of 2-amino-3-chloro-1,4-NQ (see the caption on the graph) to solutions containing 50 μM **P**₃, 10 nM DI and 250 μM NADH in an air-saturated CAPS-EDTA buffer (0.1 M) at pH 9.5, and (bottom) the corresponding semi-logarithmic 2-amino-3-chloro-1,4-NQ calibration curve. Errors bars are standard deviations from triplicates.

P₄ results in a better LoD is not surprising because, as we had previously demonstrated from theoretical analysis of a cross-catalytic system,³⁶ reaction time and detection limit are both inversely proportional to the geometric mean of each loop's rate (this corresponds to the limiting case where detection of H_2O_2 is mainly restricted by the nonspecific response resulting from NADH autoxidation).

Conclusions

Thanks to the simple strategy consisting in the double masking of a naphthohydroquinone with boronate groups, we demonstrated the feasibility to design an efficient and versatile exponential molecular amplification system, well-suited for the sensitive detection of H_2O_2 down to a few nM in less than 10 minutes or a redox cycling compound such as the 2-amino-3-chloro-1,4-NQ down to 50 pM in less than 30 minutes. These remarkable analytical performances are the best so far reported for an autocatalytic molecular amplification strategy, amounting to a ~ 300 -fold improvement in LoD compared to H_2O_2 detection by dendritic chain reaction.²¹ Such performances are the result of the high chemical stability of the diboronate ester probes (a high stability resulting from the double protection of the probe and a non-utilization of a self-immolating protective group) as well as a particularly good

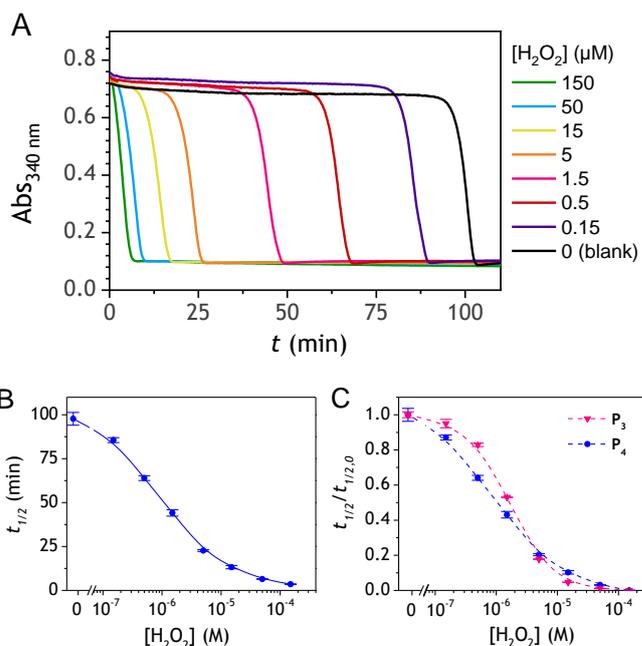


Fig. 8 (A) UV-vis kinetic traces (average of triplicates) monitored at 340 nm immediately after the injection of different concentrations of H_2O_2 (see the caption on the graph) to solutions containing 50 μM **P**₄, 10 nM DI and 250 μM NADH in an air-saturated Tris-EDTA buffer (0.1 M) at pH 8.5, and (B) the corresponding semi-logarithmic H_2O_2 calibration curve. (C) Overlay of the normalized semi-logarithmic calibration curve of H_2O_2 obtained in B to that obtained with **P**₃. Errors bars are standard deviations from triplicates.

reactivity towards H_2O_2 . Another important result of this work is that it demonstrates the usefulness of deciphering the mechanism and kinetics of complex reaction networks in order to better understand, predict and optimize their functioning. Through this mechanistic study we were able to explain how the pH can play an important role and how, by adjusting it, it is possible to improve significantly the analytical performances thanks to a better discrimination of the specific vs. nonspecific response. It also allowed us to demonstrate that the double masking does not prevent autocatalysis from working despite the fact that two equivalents of H_2O_2 are needed to completely deprotect the probe. Furthermore, as illustrated with the selection of a better redox cycle for the design of probe **P**₄, it is clear that there is room to further improve the analytical performances of the system.

The study of complex reaction networks with highly nonlinear dynamics is a rapidly expanding field which, due to its numerous implications in the field of systems chemistry, is expected to grow rapidly in the future. Further developments and exploitations in a variety of analytical or bioanalytical applications are also expected, especially considering the versatility and generic character of the present molecular amplification strategy. Finally, we believe that the approach proposed here has also potential for applications in cellular imaging or drug delivery, notably by taking advantage of probes or pro-drugs capable of selectively and autocatalytically delivering the active molecule *in vivo*.

Author Contributions

J.P. and C.R. contributed equally to this work.

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