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# Dynamics of a Key Conformational Transition in the Mechanism of Peroxiredoxin Sulfinylation

Alexandre Kriznik, Marouane Libiad, H el ene Le Cordier, Samia Boukhenouna, Michel B. Toledano, and Sophie Rahuel-Clermont\*



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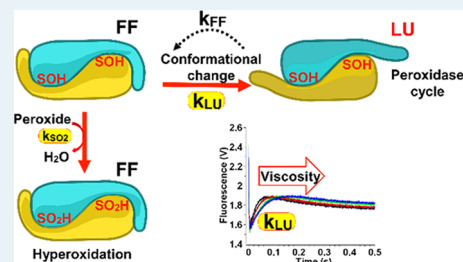
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Supporting Information

**ABSTRACT:** Peroxiredoxins from the Prx1 subfamily (Prx) are moonlighting peroxidases that operate in peroxide signaling and are regulated by sulfinylation. Prxs offer a major model of protein–thiol oxidative modification. They react with  $\text{H}_2\text{O}_2$  to form a sulfenic acid intermediate that either engages into a disulfide bond, committing the enzyme into its peroxidase cycle, or again reacts with peroxide to produce a sulfenic acid that inactivates the enzyme. Sensitivity to sulfinylation depends on the kinetics of these two competing reactions and is critically influenced by a structural transition from a fully folded (FF) to locally unfolded (LU) conformation. Analysis of the reaction of the Tsa1 *Saccharomyces cerevisiae* Prx with  $\text{H}_2\text{O}_2$  by Trp fluorescence-based rapid kinetics revealed a process linked to the FF/LU transition that is kinetically distinct from disulfide formation and suggested that sulfenate formation facilitates local unfolding. Use of mutants of distinctive sensitivities and of different peroxide substrates showed that sulfinylation sensitivity is not coupled to the resolving step kinetics but depends only on the sulfenic acid oxidation and FF-to-LU transition rate constants. In addition, stabilization of the active site FF conformation, the determinant of sulfinylation kinetics, is only moderately influenced by the Prx C-terminal tail dynamics that determine the FF  $\rightarrow$  LU kinetics. From these two parameters, the relative sensitivities of Prxs toward hyperoxidation with different substrates can be predicted, as confirmed by in vitro and in vivo patterns of sulfinylation.

**KEYWORDS:** enzyme, thiol peroxidase, sulfenic acid, sulfenic acid, redox regulation, conformational transition, rapid kinetics



## INTRODUCTION

Sulfinylation of cysteine (Cys) thiolate is increasingly recognized as important in cell regulation and signaling.<sup>1</sup> Eukaryotic peroxiredoxins (Prxs) from the Prx1 subfamily are thiol peroxidases and the first enzyme family shown to be regulated by reversible Cys sulfinylation, an hitherto thought irreversible oxidative modification that coevolved with the sulfenic acid reductase sulfiredoxin (Srx).<sup>2–5</sup> Prxs are major peroxide-reducing enzymes, reacting with substrates by attack of a peroxidatic Cys  $C_p$  that becomes sulfenylated ( $C_p$ –SOH). In the peroxidatic cycle, the  $C_p$ –SOH condenses with the enzyme resolving Cys  $C_r$  to form a disulfide bond, the reduction of which by thioredoxin (Trx) completes the cycle (Figure 1a). Alternatively, the  $C_p$ –SOH can react with a second peroxide molecule to form a sulfenic acid ( $C_p$ –SO<sub>2</sub>H), which inactivates the enzyme. Prxs inactivation by sulfinylation (also referred to as hyperoxidation) is thought as critical for allowing unimpeded  $\text{H}_2\text{O}_2$  cell signaling, as if active, they would, by their high abundance and high substrate reactivity, quench the  $\text{H}_2\text{O}_2$  signal.<sup>3,6</sup> For instance, an Srx-dependent 24 h rhythmic sulfinylation of Prx3 occurs in mammalian mitochondria, which fosters  $\text{H}_2\text{O}_2$  mitochondrial buildup and rhythmic cytosolic release, thereby contributing to establish the circadian nature of corticosteroid secretion.<sup>7</sup> As a major target of Trx, Prx sulfinylation serves to free the function of Trx

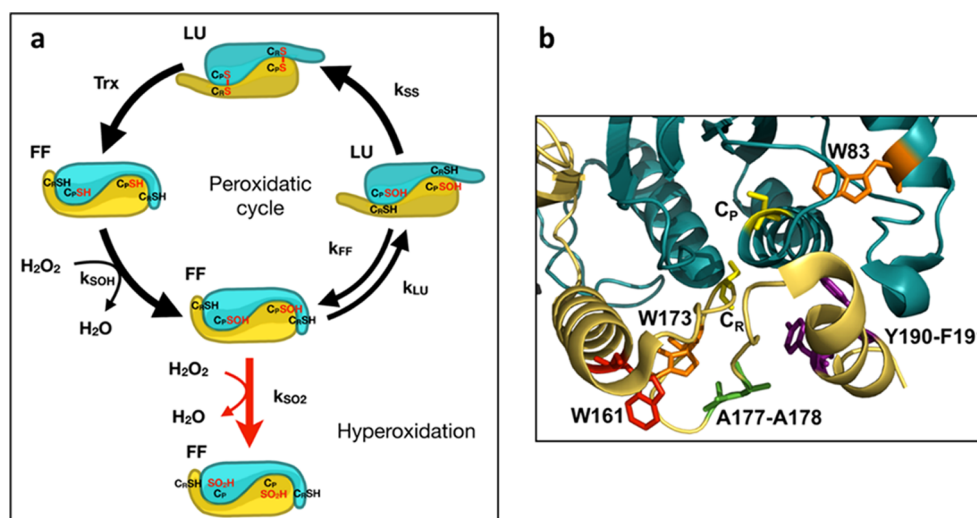
to other targets during severe oxidative stress.<sup>8</sup> Prx sulfinylation also provides a gain of function by stabilizing its oligomeric structure, thereby switching it into a molecular chaperone<sup>9</sup> or fostering protein interactions.<sup>10</sup>

Prokaryotic Prxs are resistant to hyperoxidation, whereas eukaryotic ones, especially the Prx1 subfamily, are sensitive.<sup>3</sup> Further, within sensitive Prxs, a large range of sulfinylation sensitivity levels exists, as for instance the higher sensitivity of human Prx2 relative to Prx1 and Prx3.<sup>11,12</sup> These differences may indicate differential enzyme regulation depending on the concentration of  $\text{H}_2\text{O}_2$ .<sup>13</sup> Karplus and collaborators identified structural signatures that distinguish hyperoxidation-sensitive Prxs from the resistant ones, one of which is a Tyr-Phe motif located in the C-terminal helix, and proposed a mechanistic model of how these features generate sensitivity<sup>3</sup> (Figure 1). Prxs are obligate symmetrical homodimers in which both subunits contribute to the assembly of each of the two active sites by providing either the  $C_p$  or the C-terminal tail- $C_r$  to one site (Figure 1b). In the Karplus model, reduced Prx exists

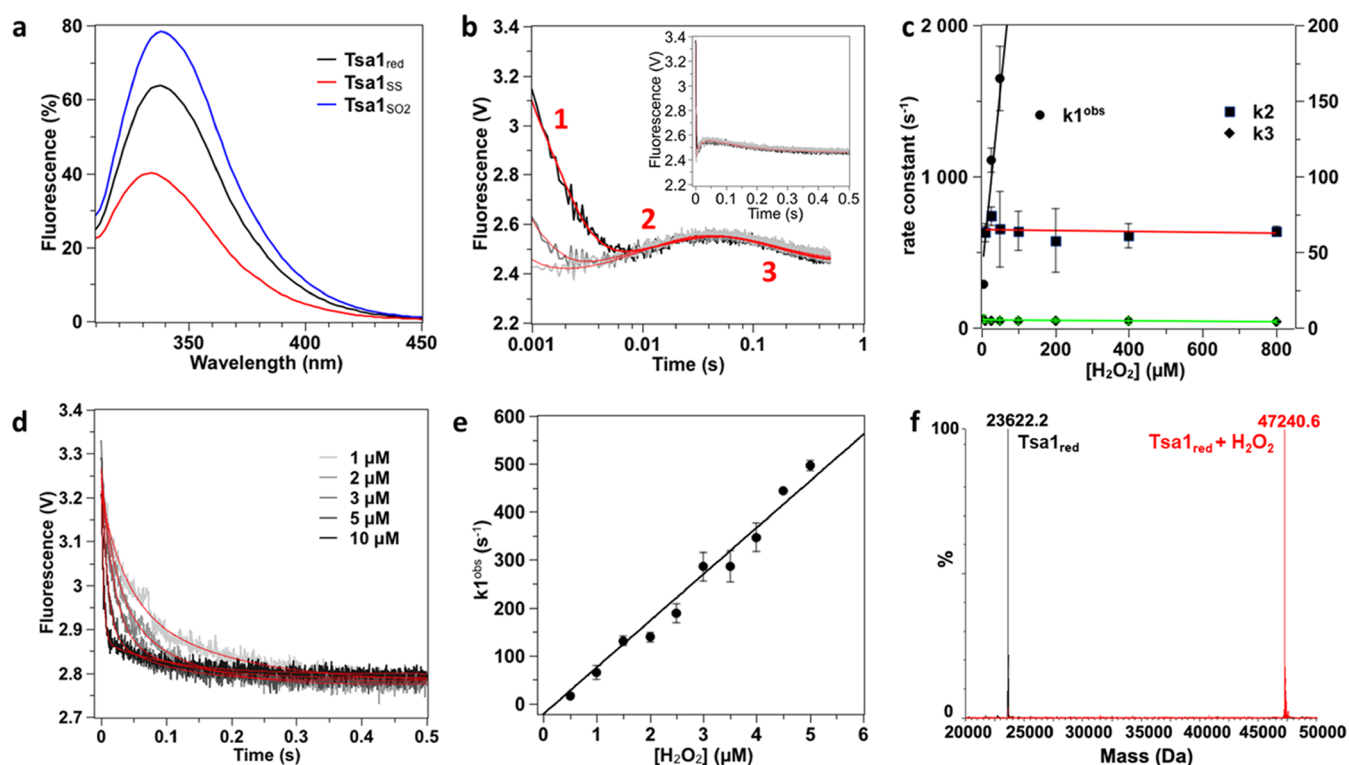
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**Figure 1.** (a) Prx1 catalytic cycle. The Prx1 peroxidase cycle and hyperoxidation mechanism are shown at the level of the Prx1 dimeric unit to highlight the conformational state [fully folded (FF) and locally unfolded (LU)] of the active site identified by the peroxidatic Cys  $C_P$  and C-terminal tail identified by resolving Cys  $C_R$ . The rate constants of peroxide reduction  $k_{SOH}$ , sulfinylation  $k_{SO_2}$ , resolution  $k_{SS}$ , and FF–LU transition  $k_{FF}$  and  $k_{LU}$  are indicated. (b) Zoom into the catalytic site of Tsa1. The subunit containing the  $C_P$  is shown in blue, and the subunit containing the  $C_R$  is shown in yellow (PDB code 3SBC). Residues  $C_P$  and  $C_R$  are shown in yellow, W83 and W173 in orange, and W161 in red. Within the C-terminal tail, residues A177 and A178 are highlighted in green and the Y190-F191 motif in purple.



**Figure 2.** Wild-type Tsa1 oxidizes  $H_2O_2$  by three-phase kinetics. (a) Trp fluorescence emission spectra of reduced (black), disulfide (red), and hyperoxidized (blue) Tsa1 ( $2 \mu M$ ) after excitation at 295 nm. (b) Pre-steady-state kinetics for the reaction of Tsa1 ( $5 \mu M$ ) with  $H_2O_2$  ( $10 \mu M$ , light gray;  $25 \mu M$ , gray; and  $50 \mu M$ , black) monitored by Trp fluorescence. The excitation wavelength is set at 295 nm, and the signal is collected above 320 nm. The time courses are shown in log time scale to highlight the three phases and fitted against a three-exponential equation (red lines). In the inset, time courses are shown in linear  $X$ -scale. Each curve is the average of six runs. (c) Second-order plots and linear fits of the observed rate constants  $k_1^{obs}$  (circles, black line),  $k_2$  (squares, red line), and  $k_3$  (diamond, green line) vs  $H_2O_2$  concentration. Very similar results were obtained using the untagged native Tsa1, showing that the N-terminal His tag has no impact on Tsa1 catalysis. (d) Pre-steady-state kinetics for the reaction of Tsa1 ( $0.5 \mu M$ ) with low  $H_2O_2$  concentrations (from  $0.5$  to  $5 \mu M$ ) monitored by Trp fluorescence. Only phases 1 and 3 are observable in these conditions. Each trace is the average of six runs and is fitted according to a biphasic equation (red line). (e) Precise determination of  $k_1$  by a second-order plot and linear fit of the fast phase rate constants  $k_1^{obs}$  measured in (d). (f) Superimposed deconvoluted mass spectra of the Tsa1 redox species before and after 5 s reaction with  $10 \mu M H_2O_2$  followed by acid quenching.

Table 1. Steady-State, Pre-Steady-State, and Hyperoxidation Kinetics Parameters of Wild-Type and Mutant Tsa1 with H<sub>2</sub>O<sub>2</sub>

Tsa1	H <sub>2</sub> O <sub>2</sub>					
	Tsa1 <sup>Y190G F191G</sup>	Tsa1 <sup>A177S A178D</sup>	wild-type		Tsa1 <sup>W161F</sup>	
$k_{\text{steady-state}}^{\text{apparent}}$ (s <sup>-1</sup> )	2.4 ± 0.1	2.6 ± 0.1	global fit		2.8 ± 0.5	
$C_{\text{hyp1\%}}$ (μM)	X	2620 ± 220	730 ± 100		440 ± 90	
$k_1$ (M <sup>-1</sup> s <sup>-1</sup> )	1.7 × 10 <sup>7a</sup>	(4 ± 1.4) × 10 <sup>7</sup>	(9.7 ± 0.4) × 10 <sup>7</sup>	$k_{\text{SOH}}$	(1.0 ± 0.01) × 10 <sup>8</sup>	(6.7 ± 0.4) × 10 <sup>7</sup>
$k_2$ (s <sup>-1</sup> )		216 ± 41	64 ± 5	$k_{\text{LU}}$	65 ± 1	37 ± 1
$k_3$ (s <sup>-1</sup> )	3 ± 0.1	5.3 ± 0.6	4.6 ± 0.3	$k_{\text{SS}}$	5.8 ± 0.1	7.0 ± 0.6
$k_4$ (M <sup>-1</sup> s <sup>-1</sup> ) <sup>b</sup>	0	(1.0 ± 0.4) × 10 <sup>3</sup>	(2.1 ± 0.2) × 10 <sup>3</sup>	$k_{\text{SO}_2}$	(2.9 ± 0.1) × 10 <sup>3</sup>	(2.9 ± 0.6) × 10 <sup>3</sup>

<sup>a</sup>Up to 50 μM. <sup>b</sup>Measured on C171A mutants. Data are reported as the mean value obtained from two independent experiments performed on distinct protein productions ± standard deviation (s.d.).

in a fully folded (FF) conformation that stabilizes an active site, endowing C<sub>P</sub> with extraordinary H<sub>2</sub>O<sub>2</sub> reactivity. A conformational transition from the FF to a locally unfolded (LU) state, in which the helix carrying C<sub>P</sub> and the enzyme C-terminal tail partially unfold, enables C<sub>P</sub>–SOH and C<sub>R</sub> condensation into a disulfide bond. When the enzyme is in the FF state, however, the C<sub>P</sub>–SOH can further react with H<sub>2</sub>O<sub>2</sub> to generate sulfenic acid. Enzyme sulfinylation and the conformational FF–LU transition equilibrium appear positioned as two reciprocally exclusive competing events. In this model, the Tyr-Phe C-terminal helix motif, by stabilizing to the FF conformation, generates sulfinylation sensitivity<sup>3,14–16</sup> (Figure 1b).

Kinetic and mass spectrometry-based evaluation of Prx intrinsic hyperoxidation sensitivity has so far relied on the hypothesis that the FF–LU transition is in rapid equilibrium, that is, that the FF → LU and LU → FF processes occur much faster than the resolving and hyperoxidation steps.<sup>11,12</sup> A steady-state sensitivity index  $C_{\text{hyp1\%}}$ , defined as the concentration of H<sub>2</sub>O<sub>2</sub> required to oxidize 1% of the sites in one enzyme catalytic cycle, has been derived from the above mechanistic model as a function of the kinetic rate constants of the disulfide formation step,  $k_{\text{SS}}$ ; the hyperoxidation step,  $k_{\text{SO}_2}$ ; and the FF–LU equilibrium constant  $K_{\text{LU}}$ <sup>17</sup> (Figure 1a). These methods enabled the identification of other structural determinants of hyperoxidation sensitivity in the C-terminal tail, in the C<sub>P</sub> environment and in the dimer–dimer interface.<sup>18–20</sup> In addition, pre-steady-state analyses using rapid kinetic approaches allowed resolution of the Prx catalytic steps for sulfenic acid and disulfide formation<sup>13,21,22</sup> and, recently, for hyperoxidation.<sup>12</sup> However, assessing nonchemical steps within a catalytic cycle is challenging, and thus few studies have provided direct information on FF → LU and LU → FF transition kinetics,<sup>23</sup> which is critical to understand the origin of Prx sulfinylation sensitivity.

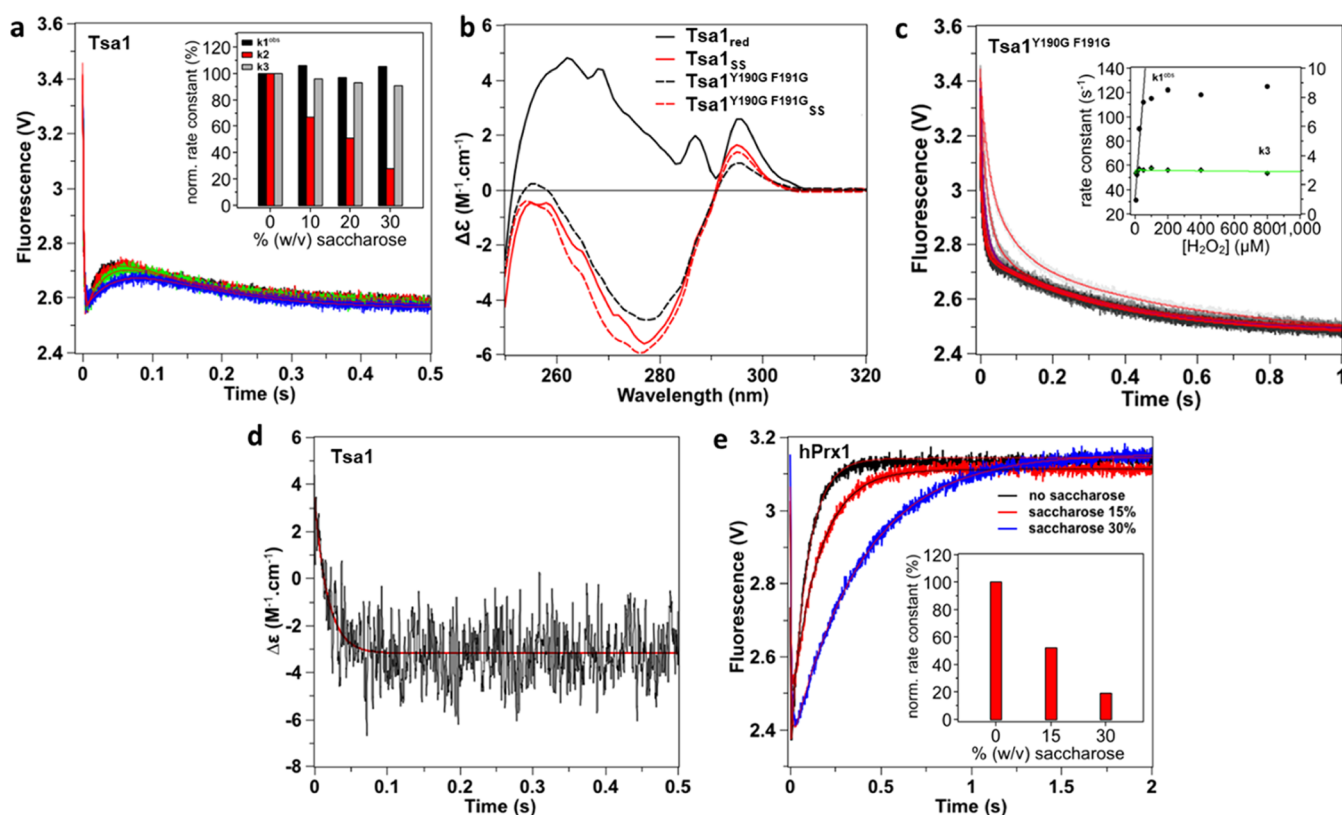
We address here the hyperoxidation mechanism of the major *Saccharomyces cerevisiae* Prx1-type enzyme Tsa1 by pre-steady-state and steady-state kinetics using Trp fluorescence, circular dichroism (CD), and in vivo analysis. The reaction of Tsa1 with peroxides monitored in single turnover using a stopped-flow apparatus showed multiphasic kinetics, of which one is kinetically distinct from the resolving step and can be assigned to a conformation change associated with the FF–LU transition, thus suggesting that the FF–LU transition does not fulfill the proposed rapid equilibrium hypothesis. Using this approach, mutants of distinctive sensitivities, and the comparison of H<sub>2</sub>O<sub>2</sub> and organic peroxide substrates, we established that hyperoxidation sensitivity is independent of the resolving step kinetics, allowing us to refine the

hyperoxidation index  $C_{\text{hyp1\%}}$ , which in fact appears to only depend on the sulfinylation and FF-to-LU transition rate constants. Our results also suggest that the molecular determinants that control each of these two parameters are distinct. We now can calculate  $C_{\text{hyp1\%}}$ , a critical parameter for in vivo modeling of redox regulation. Understanding the sulfinylation mechanism is also relevant in view of the recently discovered new sulfiredoxin sulfinylated substrates.<sup>1</sup>

## RESULTS

### Disulfide Bond Formation Is Kinetically Resolved from a Conformational Process in Tsa1 Peroxidase Cycle.

To kinetically resolve the catalytic steps, the reaction of Tsa1 with H<sub>2</sub>O<sub>2</sub> was explored by monitoring the change of enzyme Trp fluorescence over time under single turnover conditions in the absence of Trx. Tsa1 contains three Trp residues, one located close to the active site (W83), another in helix α5 preceding the C-terminal tail (W161), and the last in the C-terminal tail close to the resolving Cys (W173) (Figure 1b). The intrinsic sensitivity of Trp fluorescence to alterations of the redox state of Tsa1, with 62 and 124% emission change for the C<sub>P</sub>–C<sub>R</sub> disulfide (Tsa1<sub>SS</sub>) and hyperoxidized (Tsa1<sub>SO<sub>2</sub></sub>) forms, respectively, relative to the reduced enzyme (Tsa1<sub>red</sub>), provided a powerful reaction-monitoring probe (Figure 2a). The reaction of Tsa1 (5 μM) with H<sub>2</sub>O<sub>2</sub> generated three-exponential kinetics, fast (phase 1) and slow (phase 3) ones, both displaying a decrease in fluorescence, which flanked the remaining one (phase 2) characterized by a fluorescence increase (Figure 2b). At lower amounts of Tsa1 and H<sub>2</sub>O<sub>2</sub>, the phase 1 rate constant appeared linearly dependent on the H<sub>2</sub>O<sub>2</sub> concentration (Figure 2d,e), indicating a second-order, bimolecular process, which could thus correspond either to H<sub>2</sub>O<sub>2</sub> binding, a process expected to be reversible, or to the irreversible Cys C<sub>P</sub> attack on H<sub>2</sub>O<sub>2</sub>.<sup>22</sup> However, the low y-intercept of the  $k_1^{\text{obs}}/\text{H}_2\text{O}_2$  plot (Figure 2e), which implied an essentially irreversible process, and the absence of signal with the C<sub>P</sub> to Ser mutant Tsa1<sup>C48S</sup>, which indicated a requirement of C<sub>P</sub> (Figure S1), suggest that the kinetic phase 1 reflects the C<sub>P</sub> attack on H<sub>2</sub>O<sub>2</sub> and concomitant C<sub>P</sub>–SOH intermediate formation. The phase 1 second-order rate constant  $k_1$  of 9.7 × 10<sup>7</sup> M<sup>-1</sup> s<sup>-1</sup> (Table 1) fits previously reported values.<sup>24</sup> In contrast, phases 2 and 3 were characterized by H<sub>2</sub>O<sub>2</sub>-independent observed rate constants  $k_2$  and  $k_3$ , respectively (Figure 2c), and therefore reflect first-order, monomolecular events that do not directly involve H<sub>2</sub>O<sub>2</sub> as a reactant. As a first approach to assign the second and third phases to mechanistic steps, we analyzed by mass spectrometry the nature of redox products obtained after completion of the reaction of reduced Tsa1 with 10 μM H<sub>2</sub>O<sub>2</sub> (Figure 2f). Dimeric Tsa1<sub>SS</sub> was by far



**Figure 3.** Attribution of phase 2 to a conformational transition. (a) Effect of saccharose (0% black, 10% red, 20% green, and 30% blue) on the reaction of Tsa1 ( $5 \mu\text{M}$ ) with  $\text{H}_2\text{O}_2$  ( $10 \mu\text{M}$ ) monitored as in Figure 2, fitted against a three-exponential equation (red lines). Inset: effect of saccharose concentration on rate constants  $k_1^{\text{obs}}$ ,  $k_2$ , and  $k_3$  normalized to 0% saccharose. The stopped-flow mixer efficiency in viscous solutions was established by mixing Trp and up to 30% saccharose, which showed no artifactual effects on the dilution kinetics (Figure S2). (b) Near-UV circular dichroism spectra of  $50 \mu\text{M}$  wild-type Tsa1 (plain) and Tsa1<sup>Y190G F191G</sup> (dashed line) under the reduced (black) and disulfide (red) forms. Measurements were performed in a 1 cm cuvette in a phosphate 10 mM and NaF 100 mM buffer (pH 7) and are the average of three records. (c) Pre-steady-state kinetics for the reaction of Tsa1 ( $5 \mu\text{M}$ ) with  $\text{H}_2\text{O}_2$  (5, 10, 25, 50, 100, 200, 400, and  $800 \mu\text{M}$ , light gray to black) monitored as in Figure 2, fitted against a biexponential equation (red lines). Inset: second-order plots and linear fits of the observed rate constants for the fast phase  $k_1^{\text{obs}}$  (circles, black line) and slow phase  $k_3$  (diamond, green line). (d) Pre-steady-state kinetics for the reaction of Tsa1 ( $50 \mu\text{M}$ ) with  $\text{H}_2\text{O}_2$  ( $100 \mu\text{M}$ ) monitored by a near-UV CD signal at 270 nm. The trace is the average of 50 runs, and the first-order fit is shown in red. (e) Effect of saccharose (0% black, 15% red, and 30% blue) on the reaction of human Prx1 ( $5 \mu\text{M}$ ) with  $\text{H}_2\text{O}_2$  ( $10 \mu\text{M}$ ) monitored as in Figure 2 and fitted against a biexponential equation (red or black lines). Inset: effect of saccharose concentration on the rate constant of the increasing phase normalized to 0% saccharose.

the major species, thus indicating that no hyperoxidation occurred in these conditions. We thus hypothesized that phases 2 and 3, which are characterized by rate constants  $k_2$  of  $64 \text{ s}^{-1}$  and  $k_3$  of  $4.6 \text{ s}^{-1}$  (Table 1), reflect either a conformational change associated with the FF-to-LU transition or  $\text{C}_\text{P}$ – $\text{C}_\text{R}$  disulfide formation.

To test whether phases 2 or 3 could be attributed to a conformational transition, we repeated the stopped-flow experiment in the presence of increasing concentration of saccharose, a viscogen expected to slow down protein motions without interfering with chemical processes.<sup>25,26</sup> Conditions were chosen to ensure kinetic resolution of the three phases, that is,  $5 \mu\text{M}$  Tsa1 and  $10 \mu\text{M}$   $\text{H}_2\text{O}_2$ . The efficiency of the stopped-flow mixer in viscous solutions was verified by mixing Trp and up to 30% saccharose (Figure S2). As shown in Figure 3a, increasing saccharose from 0 to 30% significantly reduced  $k_2$  by 75%, without altering  $k_1^{\text{obs}}$  and  $k_3$  or only very slightly. Similar results were obtained with fructose or sorbitol as viscogen (Figure S3). To establish whether phase 2 corresponded to a conformational change linked to the FF-to-LU transition, we analyzed the behavior of the Y190G-F191G Tsa1 mutant (Tsa1<sup>Y190G F191G</sup>), which is expected to exist

primarily in the LU conformation due to C-terminal  $\alpha$  helix destabilization.<sup>3</sup> The conformational state of the protein was first assessed by circular dichroism (Figure 3b). The spectrum of reduced Tsa1, which exists in the FF conformation,<sup>24</sup> exhibited three positive maxima at 262, 287, and 296 nm. The spectrum of Tsa1<sub>SS</sub>, which adopts the LU conformation, exhibited a minimum at 277 nm and a positive maximum at 295 nm, thus showing much larger difference upon oxidation compared to human Prx1 (hPrx1) and human Prx2 (hPrx2).<sup>12,16</sup> Further, the reduced Tsa1<sup>Y190G F191G</sup> spectrum was similar to the Tsa1<sub>SS</sub> one and only slightly altered upon  $\text{H}_2\text{O}_2$  oxidation, which is consistent with the LU conformation, as previously observed for hPrx2.<sup>16</sup> Similar results were obtained in the far-UV CD Tsa1 spectra (Figure S4). Kinetics of Tsa1<sup>Y190G F191G</sup> oxidation by  $\text{H}_2\text{O}_2$  revealed a biphasic profile, the first phase of which has a  $\text{H}_2\text{O}_2$ -dependent rate constant  $k_1$  of  $1.7 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$  up to  $50 \mu\text{M}$ , and then, it reaches a plateau at  $\sim 120 \text{ s}^{-1}$ , likely corresponding to  $\text{C}_\text{P}$ – $\text{SOH}$  formation (Figure 3c). The second slower process was characterized by a fixed rate constant of  $3.0 \pm 0.1 \text{ s}^{-1}$  close to the  $k_3$  value measured for the wild type (Figure 3c and Table 1). However, the viscogen-sensitive phase 2 was absent here,

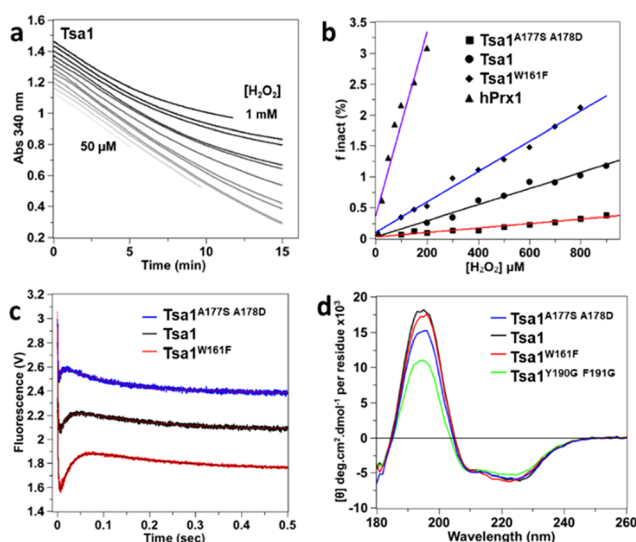
strongly suggesting that in the wild-type enzyme this phase reflects a protein structural motion associated with the FF–LU transition.

To further support this conclusion, we independently monitored the FF–LU transition using the large near-UV CD signal change observed between Tsa1 and Tsa1<sub>SS</sub> at 270 nm. The low sensitivity of the CD signal in stopped-flow mode required use of 50  $\mu\text{M}$  wild type reduced Tsa1 and 100  $\mu\text{M}$   $\text{H}_2\text{O}_2$ , which prevented visualizing the first phase of predicted very fast observed rate constant of  $10^4 \text{ s}^{-1}$ . Under these conditions, the near-UV CD-monitored reaction obeyed monophasic kinetics characterized by a rate constant of  $54 \pm 2 \text{ s}^{-1}$  close to the  $k_2$  value of  $64 \text{ s}^{-1}$  measured using Trp fluorescence, thus suggesting that the same process is monitored (Figure 3d). Overall, these data support that phase 2 reflects a conformation change linked to the FF–LU transition. Since dimeric Tsa1<sub>SS</sub> is the final product of the reaction in these conditions (Figure 2f), phase 3 must then correspond to the formation of the C<sub>P</sub>–C<sub>R</sub> disulfide bond.

To extend these results, we compared the kinetics of  $\text{H}_2\text{O}_2$  reduction of human Prx1 (hPrx1) and Tsa1 by Trp fluorescence. The hPrx1 reaction with 10  $\mu\text{M}$   $\text{H}_2\text{O}_2$  obeyed biphasic kinetics (Figure 3e) as reported,<sup>12</sup> which consists in a fast decreasing phase equivalent to Tsa1 phase 1 (sulfenic acid formation) and a second increasing phase of rate constant  $10.8 \text{ s}^{-1}$  previously attributed to C<sub>P</sub>–C<sub>R</sub> disulfide formation.<sup>12</sup> The latter rate constant value was strongly dependent on saccharose concentration, indicating that, as for Tsa1, the kinetics of this phase is at least in part controlled by a process associated with a conformation change (Figure 3e, inset). Thus, in contrast to Tsa1, the hPrx1 FF–LU associated process and disulfide bond formation are not resolved and hPrx1 phase 2 thus kinetically reflects the conformational event.

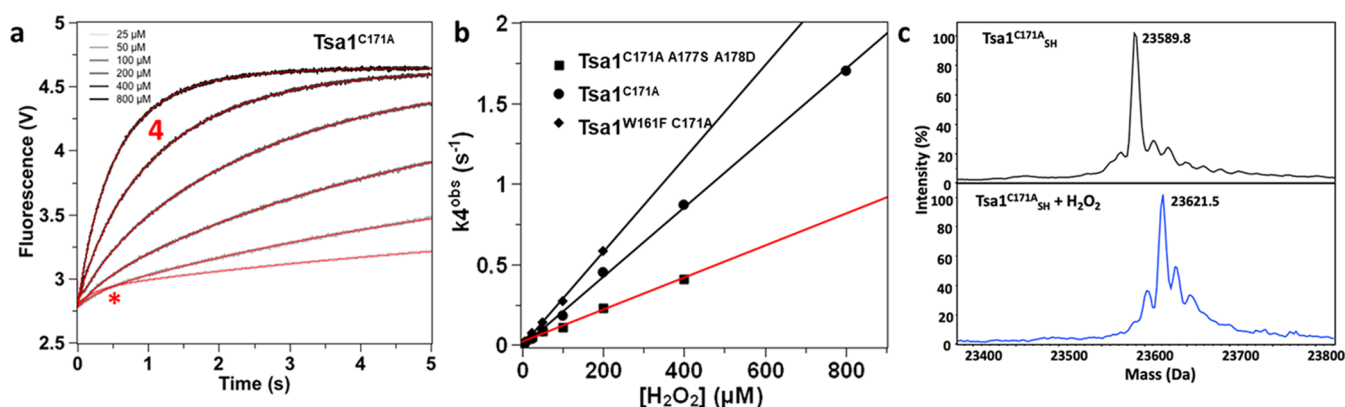
**Tsa1 Hyperoxidation Sensitivity and Conformation Change Kinetics Are Correlated.** To correlate the kinetics of the above individual events with the sensitivity to hyperoxidation, we used the method developed by Nelson,<sup>17</sup> which provides a measure of the hyperoxidation sensitivity index  $C_{\text{hyp}1\%}$  from the fraction of hyperoxidized enzyme per catalytic cycle in multiple turnover conditions, that is, in the steady state in the presence of the Trx system. Deviation from linear kinetics resulting from enzyme oxidative inactivation is monitored here by the consumption of reduced nicotinamide adenine dinucleotide phosphate (NADPH) at 340 nm in a NADPH/Tsa1/Trx/Trx reductase coupled assay (Figure 4a). As shown previously,<sup>17</sup> the deduced rate constant of inactivation normalized to the peroxidase turnover rate constant ( $f_{\text{inact}}$ ) was linearly dependent on  $\text{H}_2\text{O}_2$  concentration, yielding a slope value that reflected the sensitivity to hyperoxidation (Figure 4b) and provided the  $C_{\text{hyp}1\%}$  value, that is, the concentration of  $\text{H}_2\text{O}_2$  required to hyperoxidize 1% of sites in one cycle. For hPrx1, we measured a  $C_{\text{hyp}1\%}$  of 80  $\mu\text{M}$ , in good correlation with the reported value of 62  $\mu\text{M}$ .<sup>17</sup> Surprisingly, Tsa1 appeared 9-fold less sensitive than hPrx1 (Figure 4b), with a  $C_{\text{hyp}1\%}$  of 730  $\mu\text{M}$  (Table 1).

To identify mechanistic determinants of hyperoxidation sensitivity, we designed Tsa1 mutations located close to the C-terminal region that are expected to alter  $C_{\text{hyp}1\%}$ . The W161F mutant (Tsa1<sup>W161F</sup>) modifies the C-terminus of helix  $\alpha_5$ . The A177S-A178D mutant (Tsa1<sup>A177S A178D</sup>) replaces some known determinants of  $C_{\text{hyp}1\%}$  by the residues present at the same location in hPrx1,<sup>19</sup> and the above-mentioned Tsa1<sup>Y190G F191G</sup> adopts mostly the LU conformation (Figures 1b and 3b). All

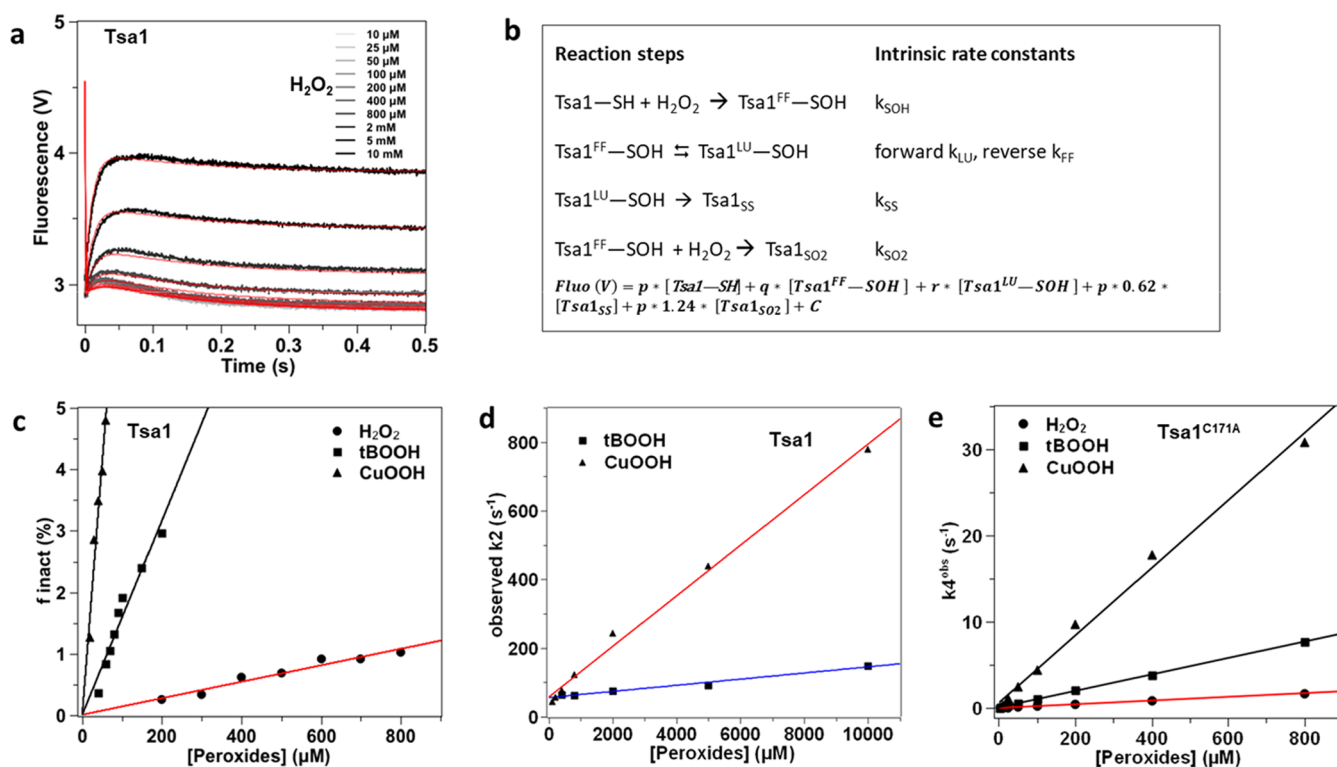


**Figure 4.** Steady-state hyperoxidation sensitivity of wild-type and mutant Tsa1 with  $\text{H}_2\text{O}_2$ . (a) Steady-state kinetics for the determination of hyperoxidation sensitivity of Tsa1 monitored by consumption of NADPH (200  $\mu\text{M}$ ) at 340 nm in the presence of thioredoxin reductase (0.25  $\mu\text{M}$ ), Trx (5  $\mu\text{M}$ ), Tsa1 (1  $\mu\text{M}$ ), and variable amounts of  $\text{H}_2\text{O}_2$  (from 50, 100, 150, 200, 300, etc. to 1 mM) in TK buffer. The time courses have been shifted on the y axis for clarity. (b) Secondary plot of the inactivated fraction  $f_{\text{inact}}$  per turnover deduced from (a) vs  $\text{H}_2\text{O}_2$  concentration. The hyperoxidation index  $C_{\text{hyp}1\%}$  is deduced from the slope of the linear fit for wild type (black circles, black line fit), mutants Tsa1<sup>W161F</sup> (black diamonds, blue line fit) and Tsa1<sup>A177S A178D</sup> (black squares, red line fit), and hPrx1 (black triangles, purple line fit). Data are the mean of two independent experiments. (c) Pre-steady-state kinetics for the reaction of Tsa1<sup>A177S A178D</sup>, Tsa1, or Tsa1<sup>W161F</sup> (5  $\mu\text{M}$ , top to bottom) with  $\text{H}_2\text{O}_2$  (10  $\mu\text{M}$ ) monitored by Trp fluorescence as in Figure 2b, fitted against a three-exponential equation (red or black line). Time courses have been shifted on the y axis for clarity. (d) Far-UV CD spectra of 5  $\mu\text{M}$  Tsa1 (black), Tsa1<sup>W161F</sup> (red), Tsa1<sup>A177S A178D</sup> (blue), and Tsa1<sup>Y190G F191G</sup> (green) under the reduced state. Measurements were performed in a 0.01 cm flat cell in phosphate (10 mM) NaF (100 mM) buffer (pH 7) and are the average of three records.

mutants had wild-type peroxidase activity, reducing  $\text{H}_2\text{O}_2$  with a steady-state rate constant of ca.  $2.4\text{--}2.8 \text{ s}^{-1}$  (Table 1). Tsa1<sup>Y190G F191G</sup> was not sensitive to hyperoxidation (Figure S5), as expected, whereas Tsa1<sup>W161F</sup> and Tsa1<sup>A177S A178D</sup> were more and less sensitive than the wild type, with  $C_{\text{hyp}1\%}$  of 440 and 2620  $\mu\text{M}$ , respectively (Figure 4b and Table 1). Under single turnover conditions, these two mutants obeyed triphasic kinetics (Figure 4c), with rate constants  $k_1$  of  $6.7 \times 10^7$  and  $4 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$  and  $k_3$  of 7.0 and 5.3  $\text{s}^{-1}$ , respectively, close to the wild type (Table 1, Figures S6, and S7). In contrast, their  $k_2$  were inversely correlated with hyperoxidation sensitivity, with values of 37 and 216  $\text{s}^{-1}$  for Tsa1<sup>W161F</sup> and Tsa1<sup>A177S A178D</sup>, respectively, relative to the wild type (64  $\text{s}^{-1}$ ). In keeping with the  $C_{\text{hyp}1\%}$  and  $k_2$  correlation, as the most hyperoxidation-sensitive enzyme, hPrx1 yielded a  $k_2$  of 10.2  $\text{s}^{-1}$ . Further, the presence of saccharose caused changes of Tsa1 mutant kinetics similar to those seen with wild type, supporting that phase 2 indeed identifies a conformational event also in both mutants (Figures S8 and S9). In addition, the far-UV CD spectra of reduced Tsa1<sup>W161F</sup> had a wild-type profile, whereas Tsa1<sup>A177S A178D</sup> displayed an intermediate signature between the FF and LU conformations, which suggested displacement of the equilibrium toward the LU



**Figure 5.** Direct observation of hyperoxidation kinetics with C171A mutants. (a) Pre-steady-state kinetics for the reaction of Tsa1<sup>C171A</sup> (5 μM) with H<sub>2</sub>O<sub>2</sub> (25, 50, 200, 400, and 800 μM as indicated) monitored by Trp fluorescence, fitted against a biexponential equation (red lines). Phase 4 is labeled. The \* indicates a small, faster phase that is not attributed. (b) Second-order plot and linear fit of the slow phase observed rate constant  $k_4^{\text{obs}}$  (deduced from panel (a) fits) against H<sub>2</sub>O<sub>2</sub> concentration, for Tsa1<sup>C171A</sup> (black circles, black fit), Tsa1<sup>W161F C171A</sup> (black diamonds, black fit), and Tsa1<sup>C171A A177S A178D</sup> (black squares, red fit). (c) Deconvoluted mass spectra of the Tsa1<sup>C171A</sup> redox species before and after 5 s reaction with 200 μM H<sub>2</sub>O<sub>2</sub>, followed by acid quenching.



**Figure 6.** Kinetic integration of all mechanistic steps. (a) Pre-steady-state kinetics for the reaction of Tsa1 (5 μM) with increasing H<sub>2</sub>O<sub>2</sub> as indicated, monitored by Trp fluorescence as in Figure 2b. The data are globally fitted (red lines) using Kintek Explorer software and the model and equation shown in (b). It must be noted that  $k_{\text{LU}}$  and  $k_{\text{SS}}$  behaved as dependent parameters in the fit, although the concordance between  $k_{\text{LU}}$  and the value deduced from CD monitoring (65 vs 54 s<sup>-1</sup>) validated the global fit results. (b) Kinetic model and equation used for Tsa1 reaction global fitting. Fluorescence intensity factors for Tsa1<sub>SS</sub> and Tsa1<sub>S02</sub> were fixed at 62 and 124% of the Tsa1-SH value ( $p$ ) (based on Figure 2a) and intensity factors for Tsa1<sup>FF</sup>-SOH ( $q$ ) and Tsa1<sup>LU</sup>-SOH ( $r$ ) were fit to 63 and 66% of the Tsa1-SH ( $p$ ) value, respectively. The  $k_{\text{SOH}}$  was fixed at the  $k_1$  value ( $9.7 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ ) measured at low Tsa1 concentration in Figure 2d,e. The parameter  $C$  was adjusted to account for the background fluorescence. The fit was not improved by fitting individual background values for each H<sub>2</sub>O<sub>2</sub> concentration. (c) Secondary plot and linear fit of the Tsa1 inactivated fraction per turnover  $f_{\text{inact}}$  with *t*BOOH (squares, deduced from Figure S12a, black fit) and CuOOH (triangle, deduced from Figure S12b, black fit), compared with H<sub>2</sub>O<sub>2</sub> (black circles, red fit from Figure 4b). The hyperoxidation index  $C_{\text{hyp1\%}}$  is deduced from the slope of the linear fits. Data are the mean of two independent experiments. (d) Second-order plots and linear fits of the observed rate constant  $k_2$  vs peroxide concentration, deduced by fit of pre-steady-state kinetics for the reaction of Tsa1 (5 μM) with *t*BOOH (squares, blue line) and CuOOH (triangles, red line) from Figure S13 against a three-exponential equation. (e) Determination of the rate constant  $k_4$  for the Tsa1<sup>C171A</sup> reaction with *t*BOOH and CuOOH. Second-order plots and linear fits of the rate constant  $k_4^{\text{obs}}$  against peroxide concentration, obtained from pre-steady-state kinetics of the reaction with *t*BOOH (squares, black fit) and CuOOH (triangle, black fit), compared to H<sub>2</sub>O<sub>2</sub> (circles, red fit, from Figure 5b).

Table 2. Calculated and Experimental Hydroperoxidation Sensitivity of Tsa1, Human Prxs, and AhpC

Prx peroxide	Tsa1 <sup>A177S A178D</sup> H <sub>2</sub> O <sub>2</sub>	Tsa1 H <sub>2</sub> O <sub>2</sub>	Tsa1 <sup>W161F</sup> H <sub>2</sub> O <sub>2</sub>	Tsa1 tBOOH	Tsa1 CuOOH
$k_{LU}$ (s <sup>-1</sup> )	216 ± 42	64 ± 5	37 ± 1	59 ± 6	57 ± 5
$k_4^a$ (M <sup>-1</sup> s <sup>-1</sup> )	10 <sup>3</sup> ± 400	(2.1 × 10 <sup>3</sup> ) ± 200	(2.9 × 10 <sup>3</sup> ) ± 70	(9.5 × 10 <sup>3</sup> ) ± 200	(39 × 10 <sup>3</sup> ) ± 500
calculated C <sub>hyp1%</sub> (μM)	2160 ± 1280	305 ± 53	128 ± 7	62 ± 8	16 ± 1
experimental C <sub>hyp1%</sub> (μM)	2620 ± 220	730 ± 100	440 ± 93	63 ± 15	11 ± 2
Prx peroxide	hPrx1 H <sub>2</sub> O <sub>2</sub>	hPrx1 ONOOH	hPrx2 H <sub>2</sub> O <sub>2</sub>	hPrx3 H <sub>2</sub> O <sub>2</sub>	AhpC H <sub>2</sub> O <sub>2</sub>
$k_{LU}$ (s <sup>-1</sup> )	10.8 ± 2	10.8 ± 2	0.2–0.6 <sup>12,13,16</sup>	20 <sup>11</sup>	>205 calculated
$k_4^a$ (M <sup>-1</sup> s <sup>-1</sup> )	1.77 × 10 <sup>3</sup> <sup>12</sup>	2.8 × 10 <sup>5</sup> <sup>12</sup>	(1.73–1.97) × 10 <sup>3</sup> <sup>12,16</sup>	2 × 10 <sup>3</sup> estimated	410 <sup>12</sup>
calculated C <sub>hyp1%</sub> (μM)	61 ± 13	0.4	1–3	100	
experimental C <sub>hyp1%</sub> (μM)	80 ± 5/50 <sup>18</sup>		5 <sup>18</sup>	127 <sup>27</sup>	>5000 <sup>18</sup>

<sup>a</sup>Measured on C171A mutants. Data are reported as the mean value obtained from two independent experiments performed on distinct protein productions ± s.d.

conformation in the reduced state (Figure 4d). The inverse correlation seen between the sensitivity to hydroperoxidation and the rate constant  $k_2$  thus suggests a direct competition between hydroperoxidation and the FF–LU conformational transition, also excluding an influence of the kinetics of C<sub>P</sub>–C<sub>R</sub> disulfide formation on the sensitivity to hydroperoxidation.

**Direct Assessment of the Kinetics of Tsa1 Hydroperoxidation.** To complete dissection of the hydroperoxidation mechanism, we sought to assess the sulfinylation rate constant (Figure 1a). To directly measure this parameter, we used the Tsa1<sup>C171A</sup> mutant, which, by lacking C<sub>R</sub>, does not form the catalytic disulfide bond and thus could provide a means of isolating the reaction of H<sub>2</sub>O<sub>2</sub> with the C<sub>P</sub>–SOH intermediate. Upon reacting Tsa1<sup>C171A</sup> with H<sub>2</sub>O<sub>2</sub> (>25 μM) under single turnover conditions, Trp fluorescence increased during 5 s with a biphasic kinetics. Here, the reaction of H<sub>2</sub>O<sub>2</sub> with C<sub>P</sub> (phase 1) was too fast to be observed, since Tsa1<sup>C171A</sup> has a wild-type reactivity toward H<sub>2</sub>O<sub>2</sub> ( $k_1 = 7.1 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ ) (Figure S10). The first phase of small amplitude had an H<sub>2</sub>O<sub>2</sub>-independent rate constant of 2–3 s<sup>-1</sup> and corresponds to a process that remains to be attributed. The second phase (phase 4) was slower, of larger but fixed amplitude, and had an H<sub>2</sub>O<sub>2</sub>-dependent rate constant  $k_4^{\text{obs}}$ , which indicated a bimolecular reaction involving H<sub>2</sub>O<sub>2</sub>, characterized by a second-order value of  $2.1 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$  (Table 1 and Figure 5a,b). Phase 4 suggested the occurrence of sulfinylation, based on its increasing fluorescence signal and on the increased fluorescence seen for sulfinylated Tsa1<sub>SO<sub>2</sub></sub> (Figure 2a). Furthermore, mass spectrometry identified Tsa1<sup>C171A</sup><sub>SO<sub>2</sub></sub> as the major redox species upon completion of a reaction with 200 μM H<sub>2</sub>O<sub>2</sub> (Figure 5c). These data thus indicate that phase 4 reflects the Tsa1<sup>C171A</sup>–SOH reaction with H<sub>2</sub>O<sub>2</sub> that leads to enzyme hydroperoxidation. The deduced rate constant of Tsa1<sup>C171A</sup> hydroperoxidation  $k_4$  of  $2.1 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$  was in agreement with those obtained for human Prx1 and Prx2,<sup>12</sup> suggesting similar kinetics of hydroperoxidation of Prx1-type Prxs from diverse origins. When combined to C171A, the W161F (Tsa1<sup>W161F C171A</sup>) and A177S-A178D (Tsa1<sup>C171A A177S A178D</sup>) Tsa1 mutants had a similar hydroperoxidation behavior to Tsa1<sup>C171A</sup>, characterized by rate constants  $k_4$  of  $2.9 \times 10^3$  and  $1.0 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$ , respectively (Figure 5b and Table 1).

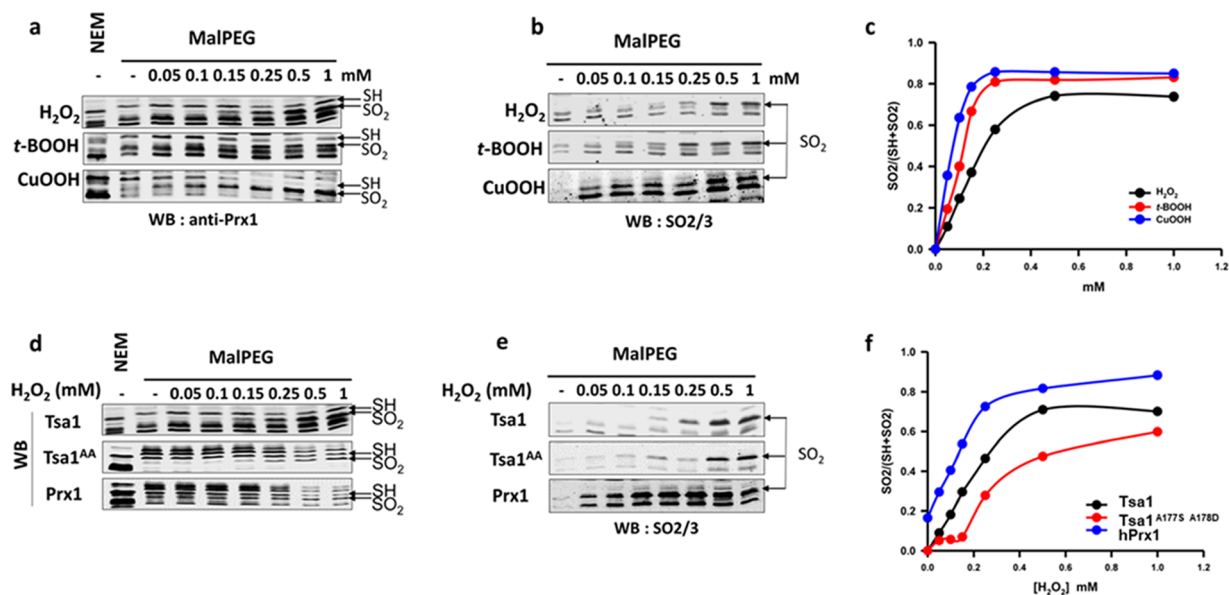
**High Peroxide Level or Reactivity Recapitulates the Full Hydroperoxidation Mechanism.** To integrate all kinetic steps within the full mechanism, we then used conditions that allow hydroperoxidation in single turnover for the wild-type

enzyme to be observed. According to the hypothesis of a direct competition between hydroperoxidation and the FF-to-LU transition, higher H<sub>2</sub>O<sub>2</sub> concentrations should overcome the kinetic barrier of the conformation step, with phase 2 becoming a composite of the fluorescence signals of the two competing reactions. Tsa1 stopped-flow kinetic series were performed with increasing concentrations of H<sub>2</sub>O<sub>2</sub>, up to 10 mM. We observed that the phase 2 amplitude increased with H<sub>2</sub>O<sub>2</sub>, suggesting that this phase must at least in part incorporate an event generating a more fluorescent species, presumably the Tsa1<sub>SO<sub>2</sub></sub> (Figure 6a). These data were globally fitted against a model (Figure 6a,b) based on the Prx catalytic cycle (Figure 1), which returned intrinsic  $k_{LU}$ ,  $k_{SS}$ , and  $k_{SO_2}$  rate constants close to the  $k_2$ ,  $k_3$ , and  $k_4$  observed values, respectively, obtained by direct fitting (Table 1). This confirmed that the hydroperoxidation rate constant  $k_4$  obtained using the C171A mutants can be used as an estimate for  $k_{SO_2}$ . Furthermore, adding an additional step in which the Tsa1–SOH in the LU form reacts with H<sub>2</sub>O<sub>2</sub> did not improve the fit, as expected. In addition, making the Tsa1–SOH LU-to-FF reaction reversible returned a very low rate constant value,  $k_{FF}$ , and did not improve the fit either. These data suggest that the rapid equilibrium hypothesis does not apply as they imply that, although intrinsically reversible, the Tsa1–SOH FF-to-LU transition nevertheless behaves as a quasi-irreversible process in the catalytic cycle.

To further assess this interpretation, simulations of the kinetics of Tsa1<sub>SO<sub>2</sub></sub> formation were performed based on the model from Figure 6b and compared to the observed values (Figure S11). Combined with the results from CD and fluorescence-based kinetics, these simulations support that for the Tsa1<sup>FF</sup>–SOH ⇌ Tsa1<sup>LU</sup>–SOH equilibrium, the forward and reverse rate constants are consistent with  $k_{LU}$  of 65 s<sup>-1</sup> and an upper  $k_{FF}$  value of 0.6 s<sup>-1</sup>, respectively. Since the  $k_{LU}$  values are of comparable order of magnitude to those of the disulfide formation and hydroperoxidation rate constants, these results support the notion that the rapid equilibrium hypothesis likely does not apply in this case.

Using alternate organic peroxide substrates with high reactivity should similarly impact hydroperoxidation sensitivity by increasing the competition against the conformation step. We found that Tsa1 was much more sensitive to hydroperoxidation by organic peroxides, with C<sub>hyp1%</sub> of 63 μM for *tert*-butyl hydroperoxide (tBOOH) and 11 μM for cumene





**Figure 7.** Hyperoxidation of wild-type and mutants Tsa1 with  $\text{H}_2\text{O}_2$  and  $t\text{BOOH}$  in *S. cerevisiae*. (a–c) Tsa1 is more reactive toward organic peroxides than with  $\text{H}_2\text{O}_2$  in vivo. Tsa1 hyperoxidation in cells exposed to  $\text{H}_2\text{O}_2$ ,  $t\text{BOOH}$ , and  $\text{CuOOH}$ , during 5 min at the indicated concentrations. Thiols were derivatized by *N*-ethylmaleimide (NEM) or mPEG, as indicated, after reduction with dithiothreitol (DTT), as described in methods. (d–f) Comparison of the  $\text{H}_2\text{O}_2$  reactivity of Prx1, Tsa1, and Tsa1<sup>A177S A178D</sup> in vivo. Thiols were derivatized by NEM or mPEG, as indicated, after reduction with DTT, as described in methods, using cell lysates of  $\Delta\text{tsa1}$  expressing human Prx1, Tsa1, or Tsa1<sup>A177S A178D</sup> and exposed to  $\text{H}_2\text{O}_2$  at the indicated concentration. (a, d) Western blot of reduced (–SH) ( $2 \times$  mPEG) and hyperoxidized (–SO<sub>2</sub>H) ( $1 \times$  mPEG) forms of Tsa1 (indicated by black arrows), revealed with an anti-Prx1 antibody. (b, e) Western blot of the Prx SO<sub>2</sub>H form using a Prx anti-SO<sub>2/3</sub> antibody and the cell lysates used in (a) and (c), respectively. (c, f) Quantification of the degree of oxidation (SO<sub>2</sub>H/SH + SO<sub>2</sub>H) vs peroxide concentration.

hydroperoxide ( $\text{CuOOH}$ ), that is, 12- and 66-fold lower, respectively, relative to  $\text{H}_2\text{O}_2$  (Figures 6c, S12a,b, and Table 2). As observed with  $\text{H}_2\text{O}_2$ , Trp fluorescence monitoring of the reduction of  $t\text{BOOH}$  by Tsa1 displayed a three-phase kinetics (Figure S13a), characterized by  $k_1$  of  $1.7 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$  that remained much higher than the subsequent steps (Figure S13a, inset). Further, the saccharose-sensitive phase 2 could be assigned to the FF–LU-linked step (Figure S14a,b). Phase 2 amplitude increased with the  $t\text{BOOH}$  concentration, likely due to the combination of the FF/LU event and the formation of Tsa1<sub>SO<sub>2</sub></sub>, as observed for the  $\text{H}_2\text{O}_2$  data set. If the two reactions were in competition, as in the proposed model, the observed rate constant for this phase should increase linearly with the peroxide concentration, with a slope corresponding to  $k_{\text{SO}_2}$ . The observed  $k_2$  indeed increased linearly with the concentration of both  $t\text{BOOH}$  and  $\text{CuOOH}$ , further supporting the direct competition mechanism (Figure 6d). Linear fit yielded  $y$ -intercept values of 59 and 57  $\text{s}^{-1}$ , respectively, corresponding to  $k_{\text{LU}}$ , unchanged relative to  $\text{H}_2\text{O}_2$ , and slopes of  $8.8 \times 10^3$  and  $7.4 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ , respectively, corresponding to  $k_{\text{SO}_2}$ . In support of this interpretation,  $t\text{BOOH}$  and  $\text{CuOOH}$  hyperoxidized Tsa1<sup>C171A</sup> with rate constants  $k_4$  of  $9.5 \times 10^3$  and  $3.9 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$  in the same range, respectively (Figure 6e and Table 2). Therefore, in addition to the enzyme intrinsic FF–LU transition kinetics, the hyperoxidation sensitivity is also a function of  $\text{C}_p$ –SOH reactivity, that is, the hyperoxidation rate constant  $k_{\text{SO}_2}$ .

**Integration of Kinetic Mechanism and Steady-State Hyperoxidation Sensitivity.** Can we predict Prx hyperoxidation sensitivity based on kinetics rate constants? The method of Nelson<sup>17</sup> measures  $\text{C}_{\text{hyp}1\%}$  based on the assumption

that the FF–LU transition behaves as a rapid equilibrium in the catalytic cycle, but our results suggest that the Tsa1–SOH FF–LU transition does not fit the rapid equilibrium hypothesis. If this is true, the fraction of hyperoxidized enzyme per catalytic cycle  $f_{\text{inact}}$  can be defined as follows

$$f_{\text{inact}} = \frac{k_{\text{SO}_2}[\text{ROOH}][\text{Prx}^{\text{FF}}\text{-SOH}]}{k_{\text{SO}_2}[\text{ROOH}][\text{Prx}^{\text{FF}}\text{-SOH}] + k_{\text{LU}}[\text{Prx}^{\text{FF}}\text{-SOH}] - k_{\text{FF}}[\text{Prx}^{\text{LU}}\text{-SOH}]}$$

with ROOH standing for the peroxide substrate,  $\text{Prx}^{\text{FF}}\text{-SOH}$  for the sulfenic acid intermediate in the FF conformation, and  $\text{Prx}^{\text{LU}}\text{-SOH}$  for the sulfenic acid intermediate in the LU conformation. Furthermore, considering the FF-to-LU transition as practically irreversible nullifies the  $k_{\text{FF}}[\text{Prx}^{\text{LU}}\text{-SOH}]$  term, simplifying the equation to

$$f_{\text{inact}} = \frac{k_{\text{SO}_2}[\text{ROOH}]}{k_{\text{SO}_2}[\text{ROOH}] + k_{\text{LU}}}$$

and

$$f_{\text{inact}} = \frac{k_{\text{SO}_2}[\text{ROOH}]}{k_{\text{LU}}} \quad (1)$$

if inactivation occurs in less than 5% of the enzyme molecules, which comply to the conditions used in this study.<sup>3</sup> Thus, as with the Nelson method,  $f_{\text{inact}}$  is expected to be linearly dependent on the peroxide concentration but with a slope equal to  $k_{\text{SO}_2}/k_{\text{LU}}$  (eq 1). A summary of the  $k_{\text{LU}}$  and  $k_4$  (used as an estimate of  $k_{\text{SO}_2}$  for enzymes possessing the  $\text{C}_R$ ) rate constants measured for Tsa1 and a comparison of the  $\text{C}_{\text{hyp}1\%}$  value calculated from these values with the experimental ones

(Figure 4b) showed a good agreement for wild-type Tsa1, Tsa1 mutants, H<sub>2</sub>O<sub>2</sub>, and organic peroxides (Table 2 and Figure S15). In the case of human Prxs, as the disulfide formation step was not kinetically resolved from the rate-limiting FF-to-LU transition (Figure 3e), we used the reported rate constants of the resolving step<sup>11,12</sup> as proxy of  $k_{LU}$ . Based on published human Prxs  $k_{SO_2}$  experimental or estimated values,<sup>11,12</sup> the hyperoxidation sensitivity index  $C_{hyp1\%}$  is again predicted with a good agreement compared to the experimental values published with H<sub>2</sub>O<sub>2</sub> (Table 2 and Figure S15). This interpretation also explains the results obtained by Randall et al.,<sup>16</sup> using the equation “direct  $C_{hyp1\%}$ ” =  $0.01 \times k_{res}/k_{SO_2}$ , with  $k_{res}$  corresponding to the observed resolving phase rate constant. These comparisons further support a mechanistic model in which the degree of hyperoxidation sensitivity is controlled by the FF-to-LU transition kinetics of the sulfenic intermediate and not by the disulfide formation event. In the case of AhpC, the reverse calculation using a  $C_{hyp1\%} > 5$  mM<sup>18</sup> and a  $k_{SO_2}$  of  $410 \text{ M}^{-1} \text{ s}^{-1}$ <sup>16</sup> gave an estimation of the FF-to-LU rate constant  $>205 \text{ s}^{-1}$ , similar to the Tsa1<sup>A177S A178D</sup> mutant.

### Relative Hyperoxidation Sensitivity of Prxs in Vivo.

To assay Prx sulfinylation in vivo, we monitored the differential migration of reduced vs sulfenylated Prx seen upon alkylation with methyl-PEG (24)-maleimide (mPEG). mPEG adds 1239.44 Da per modified residue and can only alkylate reduced, but not sulfenylated, Cys residues, thereby differentiating the reduced from the sulfenylated enzyme. We first compared the reactivity of Myc-tagged and untagged Tsa1, using a Myc-specific and a human-Prx antibody, the latter also reacting with Tsa1, and although it produces multiple nonspecific signals, it allows quantification (Figure S16a). Upon mPEG alkylation of the reduced enzyme, migration of either enzyme form was up-shifted to a molecular size corresponding to the addition of two mPEG moieties (2-mPEG), which lines up with the two Tsa1 Cys residues C<sub>P</sub> and C<sub>R</sub>. When cells were exposed to H<sub>2</sub>O<sub>2</sub>, the 2-mPEG-modified band disappeared upon increasing H<sub>2</sub>O<sub>2</sub> concentration, with appearance of a new lower-migrating band, which corresponds to Tsa1 modified by one mPEG, presumably at C<sub>R</sub>, as carrying C<sub>P</sub> in the sulfinate form, as confirmed by immunoblotting with an anti-SO<sub>2/3</sub> antibody (Figure S16b). The Tsa1 sulfinylation level can thus be determined by the relative intensity ratio of the 1-mPEG and 2-mPEG Tsa1 bands (SO<sub>2</sub>/SO<sub>2</sub> + SH, Figure S16c), irrespective of absolute levels of the enzyme. The compared sulfinylation profiles of Myc-Tsa1 and Tsa1 showed that although they displayed similar sulfinylation sensitivity, both reaching 50% saturation at 200 μM H<sub>2</sub>O<sub>2</sub>, Myc-Tsa1 sulfinylation was delayed at H<sub>2</sub>O<sub>2</sub> < 100 μM, indicating that the N-terminal Myc tag alters enzyme reactivity (Figure S16b).

We thus used untagged Tsa1 to next compare the enzyme sulfinylation profile in response to a 5 min cell exposure to H<sub>2</sub>O<sub>2</sub> and organic peroxides (Figure 7a). Sulfinylation reached 50% at the doses of 75, 125, and 200 μM, for CuOOH, tBOOH, and H<sub>2</sub>O<sub>2</sub>, respectively, consistent with in vitro data that indicated a much higher hyperoxidation sensitivity toward organic peroxides, relative to H<sub>2</sub>O<sub>2</sub>. We next compared the in vivo sulfinylation sensitivity of Tsa1<sup>A177S A178D</sup> and hPrx1 to that of Tsa1. As expected from the in vitro kinetics, Tsa1<sup>A177S A178D</sup> required much higher doses of H<sub>2</sub>O<sub>2</sub> to reach 50% sulfinylation (500 μM), relative to Tsa1 (200 μM) (Figure 7d,e). As expected too, hPrx1 was far more reactive,

reaching 50% sulfinylation at 100 μM H<sub>2</sub>O<sub>2</sub>, also displaying a non-negligible basal sulfinylation, ca. 20%, presumably caused by endogenous H<sub>2</sub>O<sub>2</sub>. Western blot of the same lysates with the anti-PrxSO<sub>2/3</sub>, although less quantitative, provided results similar to those of the mPEG procedure (Figure 7b,e). It is worth noting that full 100% enzyme sulfinylation was never reached, also plateauing at a higher value with tBOOH and CuOOH, relative to H<sub>2</sub>O<sub>2</sub>, and at a much lower value in the case of the Tsa1<sup>A177S A178D</sup> mutant, which presumably result from the assay integrating both intrinsic enzyme reactivity rates and the cellular peroxide degradation, as best exemplified with the Tsa1<sup>A177S A178D</sup> mutant. In summary, the sulfinylation parameters measured in vitro are valid in the cellular context.

## DISCUSSION

The mechanistic foundations of Prx1-type sulfinylation were established by the seminal work of Wood, Poole, and Karplus in a model that integrated three parameters as determinants of the sensitivity to hyperoxidation.<sup>3</sup> By Trp fluorescence-based rapid kinetics of the *S. cerevisiae* Prx1-type Tsa1, we have identified a conformational event linked to the FF-to-LU transition that is kinetically distinct from the recycling step. This finding establishes that hyperoxidation sensitivity is dictated by only two parameters, the sulfinylation step per se and the FF → LU rate constants (Figure 1a). Accordingly, formation of Tsa1<sub>SO<sub>2</sub></sub> and the conformational FF-to-LU transition appear as two reciprocally exclusive competing paths, with the “kinetic pause” that enables Tsa1<sub>SO<sub>2</sub></sub> formation occurring prior to the conformational transition. A slow FF-to-LU transition, as it happens in hPrx1, however becomes rate-limiting for the subsequent resolving step (Figure 3e). In addition, our data suggest that the stability of the active site FF conformation, which sets the sulfinylation kinetics, is only moderately influenced by the C-terminal tail conformation, which sets FF → LU kinetics.

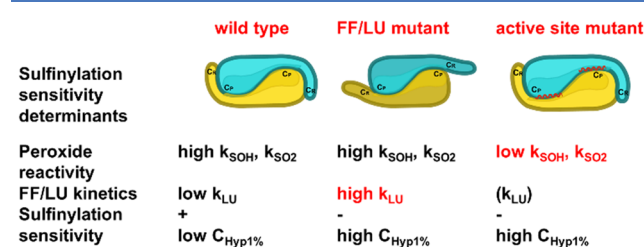
This model is supported by several lines of experimental evidence. First, using a wide range of H<sub>2</sub>O<sub>2</sub> concentrations relevant to hyperoxidation, Tsa1 reaction kinetics could be globally fitted according to the mechanism corresponding to Figure 1a. Importantly, global fitting and simulations suggest that once the C<sub>P</sub>-SOH intermediate is formed, the FF-to-LU transition is practically not reversible, as initially thought. Indeed, based on the model from Figure 6b, simulations suggest an upper  $k_{FF}$  value of  $0.6 \text{ s}^{-1}$  for the reverse Tsa1<sup>FF</sup>-SOH  $\rightleftharpoons$  Tsa1<sup>LU</sup>-SOH reaction, giving an equilibrium constant  $K_{LU}^{SOH} = k_{LU}/k_{FF} > 108$  (Figure S11). Since in the reduced state Tsa1-SH the FF conformation is favored, this therefore suggests that sulfenate formation has a profound effect on the FF/LU equilibrium in triggering local unfolding. This unexpected finding in fact fits ultrahigh-resolution structure analysis of reaction snapshots obtained with the robust bacterial PrxQ. These data suggested that the rate constant  $k_{LU}$  is higher than  $k_{FF}$ , based on the fact that the nascent sulfenate in the FF state forms in a high-energy structure, which promotes local unfolding by destabilization of the active site.<sup>28</sup> As already suggested in this study,<sup>28</sup> we propose that a slower FF-to-LU local unfolding process, as the one measured here for the sensitive Tsa1 and hPrx1, is due to an active site adjustment that accommodates sulfenate formation and movement. Second, use of our refined model to derive the steady-state hyperoxidation sensitivity,  $C_{hyp1\%}$ , predicts  $k_{LU}$  and  $k_{SO_2}$  as the only determinant kinetic steps, in

good agreement with the  $C_{\text{hyp}1\%}$  experimentally measured values (Figure S15). Third, a global fit returned a rate constant of  $\text{H}_2\text{O}_2$ -dependent hyperoxidation  $k_{\text{SO}_2}$  similar to the value measured directly using the Tsa1<sup>C171A</sup> mutant, in the  $10^3 \text{ M}^{-1} \text{ s}^{-1}$  range.

The much higher hyperoxidation rate constants of Prx, relative to other redox-sensitive proteins,<sup>29</sup> suggest that the active site in the Prx-SOH FF conformation favors hyperoxidation by activating  $\text{H}_2\text{O}_2$ , as it does for activating the initial reaction of  $\text{C}_p$  with  $\text{H}_2\text{O}_2$ .<sup>30</sup> The similarity of the  $k_{\text{SO}_2}$  for  $\text{H}_2\text{O}_2$  measured for Tsa1 and human Prxs<sup>12</sup> is consistent with the high conservation of Prxs active sites. The hyperoxidation rate constants measured with *t*BOOH and CuOOH were surprisingly much higher than those of  $\text{H}_2\text{O}_2$ . These organic peroxides may establish interactions in the vicinity of the FF active site by virtue of their aliphatic or aromatic moieties, respectively, which could favor the positioning of the peroxide function relative to the sulfenate. Conversely, the *t*BOOH  $k_{\text{SOH}}$  value was slightly lower than the very high, close to the diffusion limit,  $k_{\text{SOH}}$  value measured with  $\text{H}_2\text{O}_2$ , which could be a consequence of the organic peroxide bulkier structure relative to  $\text{H}_2\text{O}_2$ , reducing active site accessibility.

We characterized the Tsa1 A177S-A178D mutation, which substitutes these two residues in Tsa1 by those present at the same location in hPrx1 initially identified as C-terminal tail-determinants influencing  $C_{\text{hyp}1\%}$ .<sup>19</sup> We found that this mutant had wild-type values for  $k_{\text{SOH}}$  and for the amplitude of phase 1 but paradoxically a CD profile indicative of a FF-LU equilibrium shifted toward the LU conformation. Such a shifted FF-LU equilibrium would have been expected to decrease  $k_{\text{SOH}}$ , if indeed the C-terminal helix contributes to active site stability and hence to  $\text{C}_p$   $\text{H}_2\text{O}_2$  reactivity.<sup>30</sup> This result can be explained by two nonmutually exclusive hypotheses: (i) The equilibrium shifted in favor of the LU conformation is a rapid equilibrium, ensuring nonlimiting LU  $\rightarrow$  FF displacement upon  $\text{H}_2\text{O}_2$  reaction with Tsa1<sup>FF</sup>-SH. This is consistent with the high conformational exchange rate measured for PrxQ.<sup>23</sup> (ii) The Tsa1<sup>A177S A178D</sup> mutant exists in a mixed conformation, maintaining the active site in an FF conformation competent for a highly efficient  $\text{C}_p$  attack on  $\text{H}_2\text{O}_2$ <sup>30</sup> and destabilizing to some extent the C-terminal tail, as observed in some structures of AhpC.<sup>31</sup> In the Tsa1<sup>A177S A178D</sup> mutant, as in wild-type Tsa1, sulfenic acid formation would be the trigger of the active site transition to the LU state, then “pushing” the C-terminal tail toward a full LU form and shifting the FF/LU equilibrium in favor of LU, which is consistent with the conclusions obtained by Perkins et al. on AhpC.<sup>31</sup> Similar interpretation may apply for the Tsa1<sup>Y190G F191G</sup> mutant CD spectrum that displayed the full LU signature, but its kinetics paradoxically showed only a 6-fold decrease of  $k_1$  ( $1.7 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ , second-order kinetics up to  $50 \mu\text{M}$   $\text{H}_2\text{O}_2$ ) relative to the wild type. In this mutant, the absence of the buttressing effect of the C-terminal helix favors faster local active site unfolding by the formed  $\text{C}_p$ -SOH, thereby disfavoring hyperoxidation, while the C-terminal tail remaining in the LU conformation prevents phase 2 occurrence. These two examples support the notion that the unfolding of the C-terminal tail is not sufficient to induce the FF-to-LU transition at the active site, in agreement with previous studies.<sup>31</sup> We thus propose that molecular determinants of hyperoxidation sensitivity fall into two classes: those stabilizing the FF active site conformation and thus favoring

high peroxide  $\text{C}_p$  reactivity ( $k_{\text{SO}_2}$  and necessarily  $k_{\text{SOH}}$ ) and those influencing the C-terminal tail flexibility that determine  $k_{\text{LU}}$  (Figure 8).



**Figure 8.** Scheme illustrating the two proposed classes of Prx sulfinylation determinants. Mutants FF/LU affecting the C-terminal tail flexibility have high FF-to-LU kinetics, retain high peroxide reactivity, and are not sensitive to sulfinylation; mutants affecting the active site structure have poor peroxide reactivity and are predicted to be poorly sensitive to sulfinylation, irrespective of the FF-LU kinetics.

How do the in vitro Prx hypersensitivity parameters described here translate into biological contexts? To answer this question, we monitored Prx sulfinylation levels after a short, 5 min peroxide exposure. We observed a  $\text{H}_2\text{O}_2$ -dose response effect, consistent with the intracellular  $\text{H}_2\text{O}_2$  level dependence of the kinetics of sulfinylation. Under these conditions, the hyperoxidation parameters established in vitro were predictive of in vivo relative sensitivities of the enzyme sets toward hyperoxidation, when comparing both  $\text{H}_2\text{O}_2$  and organic peroxides, which have distinct  $k_{\text{SO}_2}$  and Prxs with distinct  $k_{\text{LU}}$ . The much higher sensitivity of hPrx1 relative to Tsa1, which displayed a basal level of sulfinylation when expressed in *S. cerevisiae*, is consistent with the measured lower  $C_{\text{hyp}1\%}$  of hPrx1 and might reflect the much lower intracellular peroxide levels of mammalian cells relative to yeast.<sup>32,33</sup>

In conclusion, our work provides a quantitative basis to predict in vivo relative hyperoxidation sensitivities for different Prx types, based on enzymatic parameters determined in vitro. It sets the bases for new Prxs structure-function studies and to approach the mechanism of the regulation of other proteins by reversible Cys sulfinylation, which is more common than initially thought.<sup>1</sup>

## MATERIALS AND METHODS

**Chemicals.** All chemicals were of reagent grade and were used without additional purification. Tris was from VWR (West Chester, PA). Tris, (2-carboxyethyl)phosphine hydrochloride (TCEP), *tert*-butyl hydroperoxide (*t*BOOH), cumene hydroperoxide (CuOOH), saccharose, Trp, NaF, KCl, and  $\text{MgCl}_2$  were from Merck (Darmstadt, Germany). NADPH was obtained from Roche (Basel, Switzerland), and dithiothreitol (DTT) and ammonium sulfate were from Euromedex (Souffelweyersheim, France). Hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) was from Acros Organics (Geel, Belgium). Peroxide stock concentrations were measured accurately by the peroxidase enzymatic coupled assay using Tsa1/Trx/Trx reductase/NADPH, following the total NADPH consumption at 340 nm ( $\epsilon_{340} = 6200 \text{ M}^{-1} \text{ cm}^{-1}$ ).

**Recombinant Protein Preparation.** Recombinant thio-reductin1 (Trx), Trx reductase from *Escherichia coli*, and wild-type and mutant His-tagged Tsa1 (Tsa1) from *S. cerevisiae* were produced and purified following the experimental

procedures previously described.<sup>34–36</sup> Tsa1 mutants were generated by standard polymerase chain reaction (PCR) site-directed mutagenesis and sequenced to confirm that no mutations had been introduced in the amplification reactions.

The pET28bHT-hPrx1 plasmid encoding the N-terminal His tag fusion protein of human Prx1 was obtained by cloning the *prdx1* open reading frame amplified by PCR (GC-rich system, Roche Applied Science, Basel, Switzerland) using a complementary DNA (cDNA) clone from *Homo sapiens* (RZPD, German science center for genome research, clone IRAUp969E034) as template into the pET28b(+) plasmid between the NdeI and SacI sites. The forward primer contained a NdeI restriction site, and the reverse primer contained a SacI restriction site (sequences of oligonucleotides not shown). *E. coli* C41(DE3) [ $F^-$ ompT hsdSB (rB<sup>-</sup>mB<sup>-</sup>) gal dcm (DE3)] transformants containing the pET28bHT-hPrx1 plasmid were grown by overnight culture at 37 °C for 24 h in the autoinducible ZYM-5052 medium<sup>37</sup> supplemented with kanamycin (50 mg L<sup>-1</sup>). Cells were harvested by centrifugation, resuspended in a minimal volume of buffer B (20 mM sodium phosphate, 500 mM NaCl, pH 7.5), and disrupted by sonication. hPrx1 contained in the soluble fraction was purified on a Ni-Sepharose column equilibrated with buffer B plus 50 mM imidazole, connected to an ÄKTA Avant system (GE Healthcare, France) and eluted by a 0.5 M imidazole step. At this stage, wild-type and mutant proteins were pure as checked by electrophoresis on 12.5% sodium dodecyl sulfate (SDS)-polyacrylamide gel followed by Coomassie Brilliant Blue R-250 staining and by electrospray mass spectrometry analyses. After overnight dialysis in a 20 mM sodium phosphate and 100 mM NaCl, pH 7.5, buffer, purified hPrx1 was stored at -80 °C in the presence of 10 mM DTT and was stable for several weeks under these conditions.

The plasmid pET20bTsa1 encoding the *S. cerevisiae* 2-Cys-Prx Tsa1 (referred to as native Tsa1) was obtained by cloning the Tsa1 open reading frame amplified by PCR using *S. cerevisiae* W303 genomic DNA as template into the pET20b plasmid between the NdeI and SacI sites, as described for hPrx1. The recombinant protein was produced as described for hPrx1. For native Tsa1 purification, the protein contained in the soluble fraction of the cellular extract was precipitated by ammonium sulfate at 55% saturation, followed by hydrophobic chromatography on a phenyl-Sepharose column (Amersham Biosciences) equilibrated with a 20 mM sodium phosphate and 100 mM NaCl, pH 7, buffer plus 1 M ammonium sulfate, eluted with a linear 1–0 M ammonium sulfate gradient. Final purification was achieved by anion exchange chromatography on a Q-Sepharose column equilibrated with buffer A, by an elution by a 0–1 M NaCl linear gradient. The protein was characterized and stored as described for hPrx1.

Immediately before use, the proteins were incubated with 20 mM TCEP for 20 min on ice, followed by desalting in TK buffer (50 mM Tris, 100 mM KCl, pH 7) on a PD-10 column. Protein monomer concentration was determined spectrophotometrically using a molar extinction coefficient of 29 500 M<sup>-1</sup> cm<sup>-1</sup> for Tsa1<sup>34</sup> and 18 450 M<sup>-1</sup> cm<sup>-1</sup> for hPrx1. Preparation of Tsa1<sub>SO<sub>2</sub></sub> was performed as previously described<sup>34</sup> and of Tsa1<sub>SS</sub> by addition of 1.2 equiv of H<sub>2</sub>O<sub>2</sub> to reduced Tsa1 in solution. The intrinsic fluorescence emission spectra of Tsa1 (2 μM) were recorded on a SAFAS Xenius fluorimeter at an excitation wavelength of 295 nm using a photomultiplier voltage of 450 V.

**Stopped-Flow Rapid Kinetics.** The reaction of wild-type and mutant Tsa1s with peroxide substrates was followed in single turnover conditions by monitoring intrinsic fluorescence intensity at 25 °C in buffer TK on an SX19MV-R stopped-flow apparatus (Applied Photophysics) equipped with a 5 μL cell, fitted for fluorescence measurements, with excitation wavelength set at 295 nm and emitted light collected above 320 nm using a cutoff filter. One syringe contained Tsa1 (5 μM, final concentration after mixing), and the other syringe contained the peroxide substrate. Equal volumes of each syringe were rapidly mixed to start the reaction. An average of at least six runs was recorded for each concentration of peroxide. The data set obtained at variable peroxide concentrations was fitted against multiexponential equation using Pro-Data viewer (Applied Photophysics) or SciDavis 1.2 software. For experiments in the presence of viscogen, the desired concentration (10–20 or 30%) was added to buffer TK.

**Steady-State Tsa1 Peroxidase Activity.** Tsa1 peroxidase activity was measured in TK buffer using the Trx/Trx reductase/NADPH coupled assay (1 μM Trx reductase, 200 μM NADPH, 150 μM Trx) with 100 μM H<sub>2</sub>O<sub>2</sub>, started by addition of 0.5 or 1 μM Tsa1 at 25 °C. Initial rate measurements were carried out on a UVmc2 spectrophotometer (Safas, Monaco) by following the decrease of absorbance at 340 nm due to the consumption of NADPH. A blank measurement recorded in the absence of Tsa1 was systematically deduced from the assay to account for nonspecific oxidation of Trx or Trx reductase.

**Steady-State Hyperoxidation Sensitivity.** The hyperoxidation sensitivity index  $C_{\text{hyp}1\%}$  was measured using a method adapted from Nelson et al.<sup>17</sup> Tsa1 (1 μM) peroxidase activity was measured as for the steady-state assay, in the presence of 5 μM Trx, 0.25 μM Trx reductase, 200 μM NADPH, and variable peroxide substrate in buffer TK by monitoring absorbance at 340 nm over 15 min on a UVmc2 spectrophotometer (Safas, Monaco). As Tsa1 becomes hyperoxidized over the course of the reaction (with a fraction inactivated at each turnover), the absorbance trace deviates from linear kinetics. The rate constant of inactivation  $k_{\text{inact}}$  was deduced from the slope of the kinetic trace first-order derivative and divided by the initial rate constant measured at time 0 to give the fraction of inactivation per turnover  $f_{\text{inact}}$ . Because the conditions were chosen to limit the hyperoxidized fraction to less than 5% (Figures 4b, 5b, and 6c), it increases linearly with the peroxide concentration.<sup>17</sup> The reciprocal of the slope of  $f_{\text{inact}}$  vs peroxide gives the peroxide concentration virtually required to hyperoxidize 100% of Tsa1 in one turnover. It is divided by 100 to give  $C_{\text{hyp}1\%}$ , corresponding to the peroxide concentration at which 1% of the enzyme molecules will be oxidatively inactivated per turnover.

**Circular Dichroism.** CD spectra of wild-type and mutant Tsa1s were recorded at 25 °C on a Chirascan Plus spectrometer (Applied Photophysics, U.K.). Far-UV measurements were carried out in a 0.01 cm pathlength flat quartz cell at 50 μM Tsa1 in a 10 mM sodium phosphate and 100 mM sodium fluoride, pH 7, buffer. For near-UV measurements, the same conditions were used, using a 1 cm pathlength cuvette. Scans were recorded with 1 nm steps from 260 to 180 nm (far-UV) or 320 to 250 nm (near-UV), and each experiment was averaged over three scans. The kinetics of the reaction of Tsa1 (50 μM) with H<sub>2</sub>O<sub>2</sub> (100 μM) was monitored using the CD signal at 270 nm on the same apparatus coupled with a stopped-flow module. The bandwidth was opened to 4 nm to

increase the detected light intensity. An average of 50 measurements was acquired to minimize the signal/noise ratio.

**Yeast Strains, Plasmids, and Growth Media.** The *S. cerevisiae* strains used in this study are derivatives of BY4741 (*MATa his3Δ1 leu2Δ0 met15Δ0 ura3Δ0*)<sup>38</sup> and RDKY3615 (*MATa, ura3-52, leu2Δ1, trp1Δ63, his3Δ200*) and listed in Table 3. Cells were grown at 30 °C in synthetic minimal media

**Table 3. *S. cerevisiae* Strains**

strains	genotype	reference
RDKY3615	<i>MATa, ura3-52, leu2Δ1, trp1Δ63, his3Δ200, lys2ΔBgl, hom3-10, ade2Δ1, ade8, hxt13::URA3</i>	39
BY4741	<i>MATa his3Δ1 leu2Δ0</i>	38
Δtsa1	<i>met15Δ0 ura3Δ0</i>	40
MEHY1631	<i>BY4741 tsa1Δ::kanMX4</i> <i>RDKY3615 ogg1Δ</i> <i>tsa1Δ::PRXI</i>	41

(SD) (0.67% yeast nitrogen base w/o amino acids, 2% glucose) supplemented with the appropriate amino acid or YPD (1% yeast extract, 2% peptone, and 2% glucose). The plasmids used in this study are pRS316-Myc-Tsa1<sup>4</sup> and pRS316-Tsa1 A177S-A178D that was generated by subcloning of the ORF of pET28b-Tsa1 A177S-A178D between *XbaI* and *SacI* into pRS316.

**mPEG Differential Cysteine Derivatization Procedure To Monitor Peroxiredoxin Sulfenylation in Vivo.** Yeast cells (10 mL) grown to an OD<sub>600 nm</sub> of 0.5 were exposed to H<sub>2</sub>O<sub>2</sub>, tBOOH, and CuOOH at the indicated concentration for 5 min. Trichloroacetic acid (TCA) (100%) was added to the cell culture to a final concentration of 20%. The cell culture was centrifuged at 6000g for 5 min at 4 °C. Pellets were washed with 20% TCA, and cells were lysed with glass beads in 0.2 mL of TCA (20%). Lysates were pelleted down by centrifugation at 14 000g for 15 min at 4 °C. Pellets were washed twice with acetone; dried; solubilized in 0.2 mL of a buffer containing 10 mM DTT, cOmplete mini ethylenediaminetetraacetic acid (EDTA)-free protease inhibitor cocktail (Roche) and 25 μg mL<sup>-1</sup> phenylmethylsulfonyl-fluoride, 2% SDS, and 100 mM N-(2-hydroxyethyl)piperazine-N'-ethanesulfonic acid (Hepes) (pH 7.4); and incubated at 25 °C for 30 min. Samples were precipitated by 20% TCA and centrifuged at 14 000g for 15 min at 4 °C. After two washes with acetone, the dried pellets were solubilized in 0.1 mL of a buffer containing 10 mM methyl-PEG (24)-maleimide (Thermo Fisher) or 50 mM NEM, cOmplete of mini EDTA-free protease inhibitor cocktail (Roche) and 25 μg mL<sup>-1</sup> phenylmethylsulfonylfluoride, 2% SDS, and 100 mM Hepes (pH 7.4) and incubated at 25 °C for 60 min. Protein samples were separated by SDS-polyacrylamide gel electrophoresis (PAGE) as described,<sup>42</sup> and proteins were immune-detected with an anti-Prx1 (Santa Cruz Biotechnology, ref sc-137222), anti-Myc (9E10), or Prx anti-SO<sub>2/3</sub> antibody kindly provided by S.G. Rhee.

## ■ ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acscatal.9b04471>.

Complementary kinetic experiments (Figures S1–S3, S5–S10, and S12–S14); far-UV CD spectra of wild-type Tsa1 and Tsa1<sup>Y190G F191G</sup> under the reduced and disulfide forms (Figure S4); simulation of the kinetics of formation of Tsa1<sub>SO<sub>2</sub></sub> (Figure S11); comparison of the calculated and experimental C<sub>hyp1%</sub> for wild-type and mutant Tsa1s (Figure S15); validation of the in vivo procedure to monitor Prx hyperoxidation (Figure S16) (PDF)

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All authors have given approval to the final version of the manuscript.

### Notes

The authors declare no competing financial interest.

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