



## Localized redox relays as a privileged mode of cytoplasmic hydrogen peroxide signaling



Rui D.M. Travasso<sup>a,\*</sup>, Fernando Sampaio dos Aidos<sup>a</sup>, Anahita Bayani<sup>b</sup>, Pedro Abranches<sup>c</sup>, Armindo Salvador<sup>d,e,\*\*</sup>

<sup>a</sup> Centro de Física da Universidade de Coimbra (CFisUC), Department of Physics, University of Coimbra, Coimbra, Portugal

<sup>b</sup> Department of Physics & Mathematics, School of Science & Technology, Nottingham Trent University, UK

<sup>c</sup> Department of Life Sciences, University of Coimbra, Coimbra, Portugal

<sup>d</sup> Center for Neuroscience and Cell Biology, University of Coimbra, Portugal

<sup>e</sup> CQC, Department of Chemistry, University of Coimbra, Portugal

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### ABSTRACT

Hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) is a key signaling agent. Its best characterized signaling actions in mammalian cells involve the early oxidation of thiols in cytoplasmic phosphatases, kinases and transcription factors. However, these redox targets are orders of magnitude less H<sub>2</sub>O<sub>2</sub>-reactive and abundant than cytoplasmic peroxiredoxins. How can they be oxidized in a signaling time frame? Here we investigate this question using computational reaction-diffusion models of H<sub>2</sub>O<sub>2</sub> signaling. The results show that at H<sub>2</sub>O<sub>2</sub> supply rates commensurate with mitogenic signaling a H<sub>2</sub>O<sub>2</sub> concentration gradient with a length scale of a few tenths of μm is established. Even near the supply sites H<sub>2</sub>O<sub>2</sub> concentrations are far too low to oxidize typical targets in an early mitogenic signaling time frame. Furthermore, any inhibition of the peroxiredoxin or increase in H<sub>2</sub>O<sub>2</sub> supply able to drastically increase the local H<sub>2</sub>O<sub>2</sub> concentration would collapse the concentration gradient and/or cause an extensive oxidation of the peroxiredoxins I and II, inconsistent with experimental observations. In turn, the local concentrations of peroxiredoxin sulfenate and disulfide forms exceed those of H<sub>2</sub>O<sub>2</sub> by several orders of magnitude. Redox targets reacting with these forms at rate constants much lower than that for, say, thioredoxin could be oxidized within seconds. Moreover, the spatial distribution of the concentrations of these peroxiredoxin forms allows them to reach targets within 1 μm from the H<sub>2</sub>O<sub>2</sub> sites while maintaining signaling localized. The recruitment of peroxiredoxins to specific sites such as caveolae can dramatically increase the local concentrations of the sulfenic and disulfide forms, thus further helping these species to outcompete H<sub>2</sub>O<sub>2</sub> for the oxidation of redox targets. Altogether, these results suggest that H<sub>2</sub>O<sub>2</sub> signaling is mediated by localized redox relays whereby peroxiredoxins are oxidized to sulfenate and disulfide forms at H<sub>2</sub>O<sub>2</sub> supply sites and these forms in turn oxidize the redox targets near these sites.

### 1. Introduction

Hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) is a key intermediate in many signaling pathways in mammalian cells [1]. Its best-studied signaling effects are mediated by the oxidation of thiolates in transcription factors [2], kinases [3,4] and phosphatases [5–8]. However, whereas the redox active thiolates in these targets react with H<sub>2</sub>O<sub>2</sub> with rate constants in the range of 9–164 M<sup>-1</sup> s<sup>-1</sup> [5,6,9,1], the cell cytoplasm contains very abundant peroxiredoxins I and II (PrxI, PrxII) [10,11], which react with H<sub>2</sub>O<sub>2</sub> with rate constants in the range of 10<sup>7</sup>–10<sup>8</sup> M<sup>-1</sup> s<sup>-1</sup> [12,13]. And even in the absence of these and other catalysts, such as

glutathione peroxidases and catalases, the ≈50 μM glutathione thiolate in cells would outcompete those redox targets for H<sub>2</sub>O<sub>2</sub> [14]. Explaining how those poorly reactive targets can be acted upon by H<sub>2</sub>O<sub>2</sub> despite the strong competition by other agents is one enduring open question in redox biology [15,16].

Three main types of explanations were proposed, which are not mutually exclusive [17]. First, there may be strong intracellular H<sub>2</sub>O<sub>2</sub> concentration gradients. Sufficiently high H<sub>2</sub>O<sub>2</sub> concentrations to oxidize the less reactive targets might be attained near very localized sites of H<sub>2</sub>O<sub>2</sub> production or entry into the cytoplasm despite the mean cytoplasmic concentration being very low [15]. Indeed, estimates of

\* Corresponding author at: CFisUC, Departamento de Física, Rua Larga, 3004-516 Coimbra, Portugal

\*\* Corresponding author at: Center for Neuroscience and Cell Biology, University of Coimbra, Portugal

E-mail addresses: [rui@teor.fis.uc.pt](mailto:rui@teor.fis.uc.pt) (R.D.M. Travasso), [salvador@cnc.uc.pt](mailto:salvador@cnc.uc.pt) (A. Salvador).

$\text{H}_2\text{O}_2$  mean diffusion length [15] as well as recent mathematical models [18,19] predict the occurrence of strong concentration gradients if  $\text{H}_2\text{O}_2$  sources are localized. And observations with ratiometric  $\text{H}_2\text{O}_2$  probes targeted to various cellular membranes support the existence of *microdomains* of elevated  $\text{H}_2\text{O}_2$  concentration in the cytoplasm of live cells treated with growth factors [20]. However, despite the evidence for strong cytoplasmic  $\text{H}_2\text{O}_2$  concentration gradients, the following question is still unclear. **Q1:** Are  $\text{H}_2\text{O}_2$  concentrations attained near the supply sites sufficient to oxidize the above mentioned targets?

Second, peroxiredoxins may be transiently inactivated during signaling, allowing  $\text{H}_2\text{O}_2$  to accumulate. The first such proposal was the floodgate hypothesis [21], which posits that the oxidation of the less reactive  $\text{H}_2\text{O}_2$  targets is facilitated by the oxidation of eukaryotic 2-Cys peroxiredoxins to redox-inactive sulfinic and sulfonic forms (hyperoxidation). This hypothesis is supported by observations that overexpressing 2-Cys peroxiredoxins in mammalian cells blocks peroxide activation of nuclear factor kappa-light-chain-enhancer of activated B cells (NF- $\kappa$ B) [22,23], and that treatment of cells with tumor necrosis factor causes substantial hyperoxidation of PrxII [24]. Furthermore, it could explain why eukaryotic 2-Cys peroxiredoxins are much more susceptible to hyperoxidation than their bacterial homologues [21]. This susceptibility is due to structural features that are absent in the latter peroxiredoxins but widely conserved among the former ones [21], suggesting that it is selectively favored in eukaryotes. However, hyperoxidation was undetectable during mitogenic signaling [25–27], and unnecessary for ASK1 activation in  $\text{H}_2\text{O}_2$ -induced apoptosis [28]. More recently, it was hypothesized that localized hyperoxidation of the 2-Cys peroxiredoxins sharpens  $\text{H}_2\text{O}_2$  concentration gradients, contributing to increased  $\text{H}_2\text{O}_2$  concentrations near sites of  $\text{H}_2\text{O}_2$  supply [29]. The localization of hyperoxidation could explain the difficulty in detecting it in the experiments above, but this *localized floodgate* hypothesis remains untested. Further support for the role of localized inhibition of the peroxidase activity of 2-Cys peroxiredoxins in redox signaling was provided by Woo et al. [27]. These authors showed that upon stimulation of a wide variety of cell lines by growth factor or immune receptors the peroxidase activity of PrxI associated to cell membranes is inhibited by phosphorylation. However, it is unclear if this localized inhibition can substantially affect local  $\text{H}_2\text{O}_2$  concentrations in the presence of abundant PrxII, which also associates to the platelet-derived growth factor (PDGF) receptor upon PDGF stimulation [25], and when active cytoplasmic 2-Cys peroxiredoxins can readily diffuse to the inhibition sites. Although PrxI phosphorylation promotes PDGF-dependent tyrosine phosphorylation of cellular proteins [27], this does not necessarily imply a direct inhibition of protein tyrosine phosphatases (PTP) by  $\text{H}_2\text{O}_2$ . For instance, PrxI inhibition should lead to local accumulation of sulfenic or disulfide PrxII, which in turn could oxidize a PTP, thereby inhibiting it. Overall, the following question needs to be addressed. **Q2:** Can a localized inactivation of the 2-Cys peroxiredoxins cause a sufficient elevation of local  $\text{H}_2\text{O}_2$  concentrations to directly oxidize the less reactive targets in a signaling time frame?

Third, the action of  $\text{H}_2\text{O}_2$  may be mediated by redox relays whereby peroxiredoxins and/or peroxidases act as initial  $\text{H}_2\text{O}_2$  sensors and then oxidize the end targets [15,30,31]. Such a relay was first described for the peroxiredoxin Orp1 and the transcription factor Yap1 in the yeast *Saccharomyces cerevisiae* [32], and other similar relays in yeasts were described meanwhile [33–35]. More recently, both PrxI [28] and PrxII [2,36] were shown to engage in redox relays in the cytoplasm of mammalian cells. This is an attractive hypothesis that could also explain the specificity of redox signaling. However, the following question needs to be clarified. **Q3:** Can in general the oxidized forms of cytoplasmic 2-Cys peroxiredoxins accumulate sufficiently to outcompete  $\text{H}_2\text{O}_2$  in oxidizing the less reactive targets?

Conversely to a role in relaying oxidizing equivalents to redox targets, peroxiredoxins can protect moderately reactive thiolates near sites of  $\text{H}_2\text{O}_2$  supply against oxidation. Kang et al. [37] have shown that PrxII is recruited to the vascular endothelial growth factor (VEGF)

receptor 2 (VEGFR2) upon VEGF stimulation of endothelial cells, protecting this receptor against oxidation of its Cys1199 and Cys1206 residues. More recently the same group [38] showed that in proliferating cells PrxI associates with the centrosome protecting it from  $\text{H}_2\text{O}_2$  during interphase. It is then inhibited by phosphorylation during early mitosis, thereby facilitating oxidative inactivation of centrosome-bound phosphatases. **Q4:** But, under what conditions can a local accumulation of the 2-Cys peroxiredoxins protect reactive thiols in other proteins from oxidation and what extent of protection is achievable?

Mathematical modeling has proved useful in assessing previous hypotheses and suggest new ones about the operation of thiol redox systems [39,16,40,18,19,41]. Here we draw on the present knowledge of the reactivity and cellular concentrations of 2-Cys peroxiredoxins, sulfiredoxin (Srx) and thioredoxin 1 (Trx1) [5,6,12,13,42,9,1] and apply an integrative computational approach to address the four questions raised above.

Our analyses show the following. At low to moderate localized  $\text{H}_2\text{O}_2$  supply rates commensurate to mitogenic signaling, a sharp  $\text{H}_2\text{O}_2$  concentration gradient with a length scale of  $\sim 0.3 \mu\text{m}$  is established. However, local  $\text{H}_2\text{O}_2$  concentrations remain insufficient to oxidize PTPs in a mitogenic signaling time frame (Q1). Further, only drastic inhibitions of the peroxiredoxins' peroxidase activity can have a substantial effect on the local  $\text{H}_2\text{O}_2$  concentration near supply sites (Q2), and this comes at the cost of dampening the cytoplasmic  $\text{H}_2\text{O}_2$  concentration gradient. On the other hand, the concentrations of sulfenic and disulfide forms of Prx exceed those of  $\text{H}_2\text{O}_2$  by several orders of magnitude, and also show strong gradients. Redox targets able to react with these forms at rate constants  $\sim 200 \text{M}^{-1} \text{s}^{-1}$  could be oxidized in minutes near the  $\text{H}_2\text{O}_2$  supply sites (Q3). The recruitment of peroxiredoxins to specific sites such as caveolae has a modest effect ( $\sim 20\%$  decrease) on local  $\text{H}_2\text{O}_2$  concentrations (Q4). However, it can have a dramatic effect on the local concentrations of oxidized forms of the peroxiredoxins, further helping these species outcompete  $\text{H}_2\text{O}_2$  for oxidation of redox targets. Altogether, these results suggest that sulfenate and/or disulfide forms of Prx mediate the oxidation of redox targets within  $\sim 1 \mu\text{m}$  of  $\text{H}_2\text{O}_2$  supply sites under conditions consistent with mitogenic signaling.

At higher  $\text{H}_2\text{O}_2$  supply rates, the system can show a *hyperoxidation catastrophe*, in keeping with a recent analysis [43]. At a critical  $\text{H}_2\text{O}_2$  supply rate Prx abruptly becomes almost fully hyperoxidized and the  $\text{H}_2\text{O}_2$  gradient collapses, allowing  $\text{H}_2\text{O}_2$  to penetrate deeply into the cell and oxidize some of the most reactive targets. Recovery from such a state can start only below a lower critical  $\text{H}_2\text{O}_2$  supply rate (hysteresis) and proceeds over a period of hours. The global floodgate hypothesis may thus find its place as part of a stress response in this context.

## 2. Models and methods

The reference mathematical model implemented in this work focuses on the early stages of  $\text{H}_2\text{O}_2$  signaling, and describes the processes and geometries depicted in Fig. 1. It embodies the following main simplifications. First, it neglects intracellular  $\text{H}_2\text{O}_2$  sources. It is expectable that in early signaling  $\text{H}_2\text{O}_2$  produced externally by NADPH oxidases is supplied to the cytoplasm mainly by permeation across the membrane [20]. Second, cytoplasmic  $\text{H}_2\text{O}_2$  sinks other than 2-Cys peroxiredoxins are neglected. It is estimated [16,18] that in most cells such sinks consume a small fraction of the  $\text{H}_2\text{O}_2$ , and their activity is insufficient to sustain substantial  $\text{H}_2\text{O}_2$  gradients. Furthermore, recent quantitative proteomic studies of multiple human cell lines and tissues [11] consistently show that glutathione peroxidases are one to two orders of magnitude less abundant than PrxI and PrxII. The neglect of alternative  $\text{H}_2\text{O}_2$  sinks is a conservative assumption with respect to the  $\text{H}_2\text{O}_2$  attained in the cytoplasm. Third, it neglects thioredoxin oxidation, as the redox capacity of most cells is sufficient to keep thioredoxin reduced in absence of strong oxidative stress [16,44,40,18]. Fourth, it accounts for a single 2-Cys peroxiredoxin whose concentration

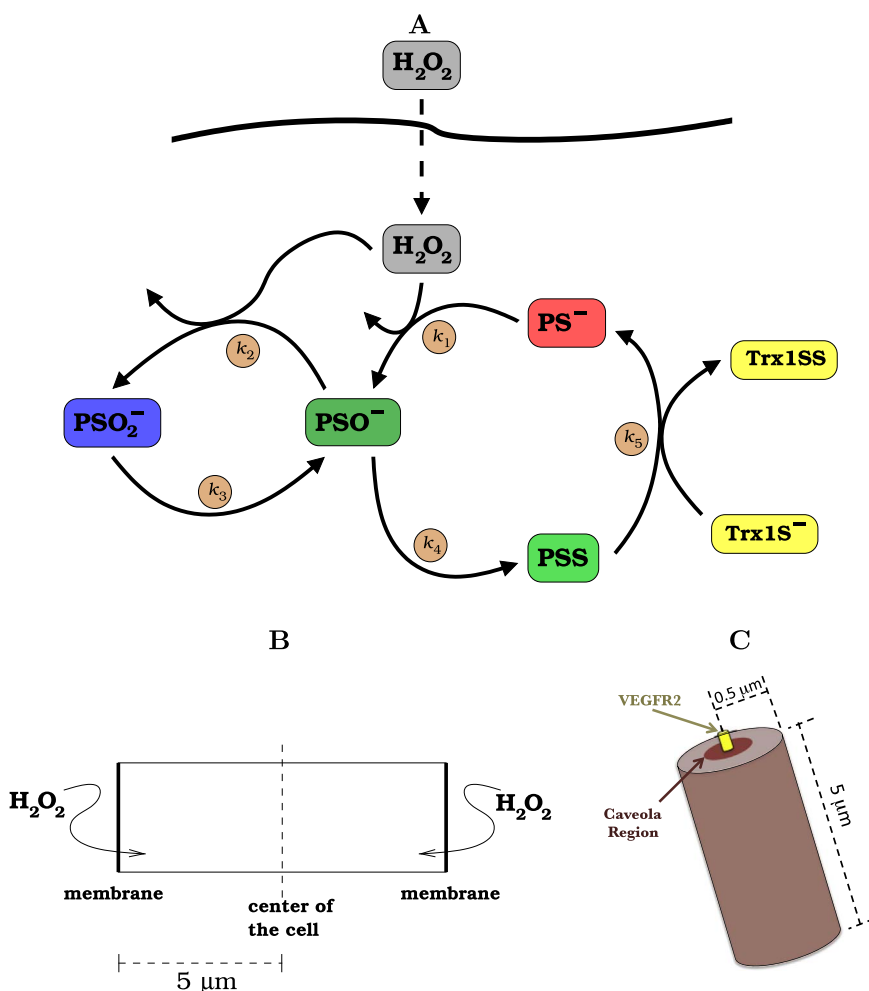
represents the sum of PrxI and PrxII, the two 2-Cys peroxiredoxins that are abundant in the cytoplasm of most mammalian cells [11]. The reference model considers that the single peroxiredoxin has the kinetic properties of PrxII. This is both because this peroxiredoxin has been more extensively characterized than PrxI, and because this is a conservative assumption with respect to most questions investigated in this work. The value we adopt for the rate constant for  $H_2O_2$  reduction by thiolate peroxiredoxin ( $k_1 = 1.0 \times 10^2 \mu M^{-1} s^{-1}$  [13]) is less conservative than that determined in ref. [45] ( $13 \mu M^{-1} s^{-1}$ ). However, even the former value has been determined at a sub-physiological temperature ( $25^\circ C$ ), and is likely still conservative. In subsequent sections we will show that the main results are robust with respect to these assumptions.

The concentrations of the various chemical species in the cytoplasm are a function of both time and space, and therefore their evolution can be obtained through the following set of reaction diffusion equations:

$$\begin{aligned} \frac{\partial c_H}{\partial t} &= D_H \nabla^2 c_H - k_1 c_{PS^-} c_H - k_2 c_{PSO^-} c_H, \\ \frac{\partial c_{PS^-}}{\partial t} &= D \nabla^2 c_{PS^-} - k_1 c_{PS^-} c_H + k_5 c_{PSS} c_{Trx}, \\ \frac{\partial c_{PSO^-}}{\partial t} &= D \nabla^2 c_{PSO^-} + k_1 c_{PS^-} c_H - k_2 c_{PSO^-} c_H + k_3 c_{PSO_2^-} - k_4 c_{PSO^-}, \\ \frac{\partial c_{PSO_2^-}}{\partial t} &= D \nabla^2 c_{PSO_2^-} + k_2 c_{PSO^-} c_H - k_3 c_{PSO_2^-}, \\ \frac{\partial c_{PSS}}{\partial t} &= D \nabla^2 c_{PSS} + k_4 c_{PSO^-} - k_5 c_{PSS} c_{Trx}, \end{aligned} \quad (1)$$

where  $c_H(\mathbf{r}, t)$ ,  $c_{PS^-}(\mathbf{r}, t)$ ,  $c_{PSO^-}(\mathbf{r}, t)$ ,  $c_{PSO_2^-}(\mathbf{r}, t)$ ,  $c_{PSS}(\mathbf{r}, t)$ , and  $c_{Trx}(\mathbf{r}, t)$  are the concentrations of  $H_2O_2$ , of the peroxiredoxin forms, and of reduced thioredoxin, respectively. The reference values of the diffusion and rate constants are as in Table 1. The diffusion constant for all peroxiredoxin forms (MW= 230 kDa) was estimated from the experimentally determined diffusion constant for Immunoglobulin G (IgG, MW=153 kDa) in the cytoplasm [46] by applying the expression

$$D_{Prx} = D_{IgG} \sqrt[3]{MW_{IgG}/MW_{Prx}},$$



**Fig. 1. Processes and geometry simulated:** **A:**  $H_2O_2$  permeates the membrane with permeability constant  $k_p$  and then oxidizes  $PS^-$  to  $PSO^-$  and  $PSO^-$  to  $PSO_2^-$  with rate constants  $k_1$  and  $k_2$  respectively.  $PSO_2^-$  is reduced back to  $PSO^-$  with rate constant  $k_3$  and  $PSO^-$  condenses with Prx's resolving Cys to yield PSS, with rate constant  $k_4$ . Thioredoxin reduces PSS to  $PS^-$ , with rate constant  $k_5$ . **B:** We consider a section of a cell described by a rectangular prism with base area  $A$ , half-length  $5 \mu m$  and a symmetry plane parallel to the top and bottom membranes.  $H_2O_2$  is assumed to permeate both membranes uniformly. This geometry models a cell in a cell layer and permits reducing the spatial problem to one dimension. Namely, along the normal to the plane of the permeant membranes. **C:** Scheme for the 3D simulation box of  $H_2O_2$  concentration near a receptor. The region of the caveolae where the various peroxiredoxin forms can bind has a radius of  $0.1 \mu m$ . The binding of 2% of the peroxiredoxin in the considered cylinder to this small region implies that the local concentration is 250-fold higher than in the bulk, considering a volume element with the depth of  $10 \text{ nm}$ . The  $H_2O_2$  is able to enter through the top surface of the cylinder. In all other surfaces we implement a zero flux boundary condition for  $H_2O_2$ . With respect to the peroxiredoxin, all boundaries of this simulation box are set to zero-flux. The various forms of peroxiredoxins bound to the membrane are laterally immobile, and unbind with a spatially uniform rate constant. For parameter values please see Table 1.

**Table 1**  
Model parameters.

Process	Rate	Parameter	Value	Ref.
$PS^- + H_2O_2 \rightarrow PSO^- + H_2O$	$k_1 c_H c_{PS^-}$	$k_1$	$1.0 \times 10^2 \mu M^{-1} s^{-1}$	[13]
$PSO^- + H_2O_2 \rightarrow PSO_2^- + H_2O$	$k_2 c_H c_{PSO^-}$	$k_2$	$0.012 \mu M^{-1} s^{-1}$	[42]
$PSO_2^- \rightarrow PSO^-$	$k_3 c_{PSO_2^-}$	$k_3$	$1.0 \times 10^{-4} s^{-1}$	[40]
$PSO^- \rightarrow PSS$	$k_4 c_{PSO^-}$	$k_4$	$1.7 s^{-1}$	[42]
$PSS + Trx1S^- \rightarrow PS^- + Trx1SS$	$k_5 c_{PSS} c_{Trx}$	$k_5$	$0.21 \mu M^{-1} s^{-1}$	[13]
Permeation of $H_2O_2$	$k_p A (c_H^0 - c_H(0, t))$	$k_p$	$7.8 \times 10^{-5} m s^{-1}$	See text
Diffusion constant of $H_2O_2$		$D_H$	$1.8 \times 10^{-9} m^2 s^{-1}$	[48]
Diffusion constant of peroxiredoxin		$D$	$5.2 \times 10^{-13} m^2 s^{-1}$	See text
Concentration of thioredoxin		$c_{Trx}$	$10 \mu M$	[11]

as per the Stokes-Einstein relationship. The estimated value is of the same order of magnitude and just slightly higher than those recently determined for untranslating polysomes [47]. The diffusion constant for  $H_2O_2$  [48] was determined for diffusion in buffer. The same authors determined a 5-fold lower diffusion constants for diffusion in hydrogels with a viscosity similar to that of the cell cytoplasm. Where pertinent we will discuss the consequences of the lower value of this diffusion constant. In Eqs. (1) we model the diffusion of the peroxiredoxin forms and of the  $H_2O_2$  according to Fick's laws [49].

Per-cell  $H_2O_2$  removal rate constants determined by Wagner et al. [50] allow to estimate the permeability constants in the range  $10^{-7} - 10^{-5} m s^{-1}$ . In order to be very conservative about intracellular  $H_2O_2$  concentrations, which in the models under appreciation are approximately proportional to the permeability constant, we adopt the higher value in Table 1.

The thioredoxin concentration is set at the constant value  $c_{Trx}(r, t) = 10 \mu M$ , which is typical for mammalian cells [11]. We will also examine the effects of lower thioredoxin concentrations.

We solve these equations in the rectangular prism domain shown in Fig. 1B. The boundary conditions differ between the two ends of the domain: the membrane side at  $x = 0 \mu m$ , and the cell center side, at  $x = 5 \mu m$ . All concentration fields are solely function of the distance to the membrane. They have zero-flux boundary conditions except for the concentration of  $H_2O_2$  at the membrane side of the domain. The amount of  $H_2O_2$  that enters the domain through the membrane per unit time is  $k_p A (c_H^0 - c_H(0, t))$ , where  $c_H^0$  is the extracellular  $H_2O_2$  concentration, and  $c_H(0, t)$  is the intracellular  $H_2O_2$  concentration adjacent to the membrane at time  $t$ . This condition is imposed as a boundary condition in the spatial derivative of  $c_H$  at the membrane:

$$\left. \frac{\partial c_H}{\partial x} \right|_{x=0} = - \frac{k_p}{D_H} (c_H^0 - c_H(0, t)). \quad (2)$$

We use these boundary conditions to solve the Eqs. (1) in the considered domain, until we obtain the stationary profiles for the different concentration fields. The equations are integrated using a finite difference scheme with  $\Delta x = 0.01 \mu m$ , on the order of magnitude of the diameter of a peroxiredoxin dimer ( $\sim 8.0$  nm, according to PdB file 4DSQ.pdb).

In order to analyze the potential protective effect of peroxiredoxin recruitment against oxidation of specific sites we consider two additional models. In the first one, we describe the binding of all peroxiredoxin species to the membrane by adding to Eqs. (1) the concentrations of the bound species. We consider that all peroxiredoxin forms have the same binding constant,  $k^+$ , to the membrane and the same unbinding constant,  $k^-$ . The ratio between the values of these constants is set such that 2% of the peroxiredoxin is bound to the membrane in the absence of  $H_2O_2$  gradients. In this setting the stationary state does not depend on the numerical values of  $k^+$  or  $k^-$ , only on their ratio. The mathematical model thus becomes:

$$\begin{aligned} \frac{\partial c_H}{\partial t} &= D_H \nabla^2 c_H - k_1 (c_{PS^-} + \delta_0 c_{PS^-}^s) c_H - k_2 (c_{PSO^-} + \delta_0 c_{PSO^-}^s) c_H \\ \frac{\partial c_{PS^-}}{\partial t} &= D \nabla^2 c_{PS^-} - k_1 c_{PS^-} c_H + k_5 c_{PSS} c_{Trx} + \delta_0 (k^- c_{PS^-}^s - k^+ c_{PS^-}) \\ \frac{\partial c_{PSO^-}}{\partial t} &= D \nabla^2 c_{PSO^-} + k_1 c_{PS^-} c_H - k_2 c_{PSO^-} c_H + k_3 c_{PSO_2^-} - k_4 c_{PSO^-} \\ &\quad + \delta_0 (k^- c_{PSO^-}^s - k^+ c_{PSO^-}) \\ \frac{\partial c_{PSO_2^-}}{\partial t} &= D \nabla^2 c_{PSO_2^-} + k_2 c_{PSO^-} c_H - k_3 c_{PSO_2^-} + \delta_0 (k^- c_{PSO_2^-}^s - k^+ c_{PSO_2^-}) \\ \frac{\partial c_{PSS}}{\partial t} &= D \nabla^2 c_{PSS} + k_4 c_{PSO^-} - k_5 c_{PSS} c_{Trx} + \delta_0 (k^- c_{PSS}^s - k^+ c_{PSS}) \quad (3) \\ \frac{dc_{PS^-}^s}{dt} &= -k_1 c_{PS^-}^s c_H(0, t) + k_5 c_{PSS}^s c_{Trx} - k^- c_{PS^-}^s + k^+ c_{PS^-}(0, t) \\ \frac{dc_{PSO^-}^s}{dt} &= k_1 c_{PS^-}^s c_H(0, t) - k_2 c_{PSO^-}^s c_H(0, t) + k_3 c_{PSO_2^-}^s - k_4 c_{PSO^-}^s - k^- c_{PSO^-}^s \\ &\quad + k^+ c_{PSO^-}(0, t) \\ \frac{dc_{PSO_2^-}^s}{dt} &= k_2 c_{PSO^-}^s c_H(0, t) - k_3 c_{PSO_2^-}^s - k^- c_{PSO_2^-}^s + k^+ c_{PSO_2^-}(0, t) \\ \frac{dc_{PSS}^s}{dt} &= k_4 c_{PSO^-}^s - k_5 c_{PSS}^s c_{Trx} - k^- c_{PSS}^s + k^+ c_{PSS}(0, t) \end{aligned}$$

where  $c_{PS^-}^s(t)$ ,  $c_{PSO^-}^s(t)$ ,  $c_{PSO_2^-}^s(t)$  and  $c_{PSS}^s(t)$  are the volume concentrations of the species bound to the membrane. We assume that all the bound peroxiredoxin species have the same reactivity towards  $H_2O_2$  and thioredoxin as the unbound counterparts. The binding of the peroxiredoxin species depends only on the concentration of the unbound species adjacent to the membrane ( $c_{PS^-}(0, t)$ ,  $c_{PSO^-}(0, t)$ ,  $c_{PSO_2^-}(0, t)$  and  $c_{PSS}(0, t)$ ). Accordingly, in Eqs. (3) the coefficient  $\delta_0$  is equal to 1 next to the membrane and equal to 0 everywhere else.

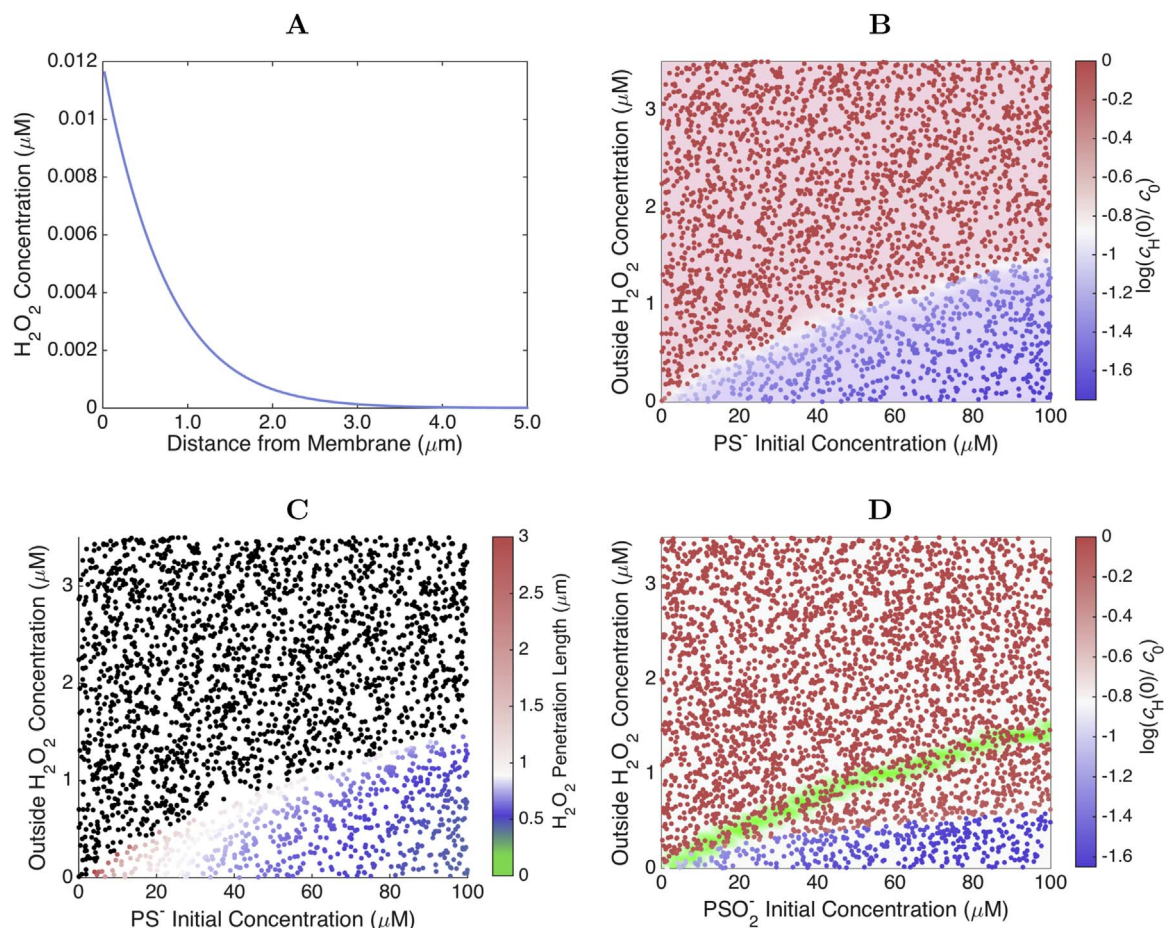
In the second model for peroxiredoxin localization the same reaction-diffusion equations are solved within a cylindrical domain with azimuthal symmetry (Fig. 1C). In this setting, we consider that the binding of the peroxiredoxin species to the membrane can only occur in a circular region representative of a caveola. In this model, the bound species are not able to diffuse within the cell membrane. This system was simulated in the  $(r, z)$  plane with a second order finite difference scheme. The increments  $\Delta x$  used were identical to those in the 1D counterparts.

### 3. Results

#### 3.1. Peroxiredoxins prevent deep penetration of $H_2O_2$ into the cell cytoplasm

We start by modeling the penetration of  $H_2O_2$  in the cell for a given fixed concentration of extracellular  $H_2O_2$ ,  $c_H^0$ . We first consider that all the peroxiredoxin is initially in reduced form. Under these conditions, for 2-Cys peroxiredoxin concentrations typical of most cells [11], the  $H_2O_2$  concentration at steady state drops steeply near the membrane (Fig. 2A). Furthermore, the steady-state concentration of  $H_2O_2$  adja-





**Fig. 2. H<sub>2</sub>O<sub>2</sub> penetration in the cell:** The H<sub>2</sub>O<sub>2</sub> can penetrate the cell depending on the amount of cytoplasmic PS<sup>-</sup> and on the extracellular H<sub>2</sub>O<sub>2</sub> concentration. **A:** Concentration of H<sub>2</sub>O<sub>2</sub> as a function of the distance from the membrane in the case of  $c_{PS^-}^0 = 50 \mu\text{M}$  and  $c_H^0 = 0.4 \mu\text{M}$ . In this situation the total amount of H<sub>2</sub>O<sub>2</sub> inside the cell is small and it is mainly located in the neighborhood of the membrane. **B:** Order of magnitude of the ratio between the H<sub>2</sub>O<sub>2</sub> concentration adjacent to the membrane and the extracellular H<sub>2</sub>O<sub>2</sub> concentration as a function of the initial cytoplasmic PS<sup>-</sup> concentration and of the extracellular H<sub>2</sub>O<sub>2</sub> concentration. Two regimes can be clearly observed: for high extracellular H<sub>2</sub>O<sub>2</sub> concentrations the H<sub>2</sub>O<sub>2</sub> oxidizes all the PS<sup>-</sup> and invades the cell, while for low extracellular H<sub>2</sub>O<sub>2</sub> concentrations the cytoplasmic PS<sup>-</sup> is able to reduce the incoming H<sub>2</sub>O<sub>2</sub>. **C:** Except for low cytoplasmic PS<sup>-</sup> concentrations, in the regime of low extracellular H<sub>2</sub>O<sub>2</sub> concentrations the depth at which the cytoplasmic H<sub>2</sub>O<sub>2</sub> concentration decays to 37% of its value near the membrane is in the range 0.5–0.7  $\mu\text{m}$ , almost independently of  $c_H^0$  and  $c_{PS^-}$ . Black dots indicate the collapse of the cytoplasmic gradient. **D:** Order of magnitude of the ratio between the H<sub>2</sub>O<sub>2</sub> concentration adjacent to the membrane and the extracellular H<sub>2</sub>O<sub>2</sub> concentration when initially all the peroxiredoxin is hyperoxidized. Each dot represents a separate simulation for the pertinent values of  $c_H^0$  and initial peroxiredoxin concentration. The green shaded area marks the transition region between both regimes in panel B, highlighting the occurrence of hysteresis. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

cent to the membrane can be almost two orders of magnitude lower than the extracellular H<sub>2</sub>O<sub>2</sub> concentration (Fig. 2B).

This decay of the H<sub>2</sub>O<sub>2</sub> concentration near the membrane is approximately exponential, similar to previous results for a simpler model [18,19]. The typical width of the membrane neighborhood where H<sub>2</sub>O<sub>2</sub> is concentrated has a weak dependence on both  $c_H^0$  and the concentration of peroxiredoxin in thiolate form (Fig. 2C), as the following approximation also shows. For low  $c_H^0$ , only a small fraction of the cell's PS<sup>-</sup> is oxidized, and therefore the steady state concentration of PS<sup>-</sup> in the cell is much higher than that of the other redox forms (Fig. 3A). Therefore, the first equation in (1) can be approximated as  $\partial_t c_H = D_H \partial_x^2 c_H - k_1 c_H c_{PS^-}$ , and  $c_{PS^-}$  is approximately equal to its initial value. The steady state for  $c_H$  is then approximately given by

$$c_H(x) \sim c_H(0) \exp(-x/\lambda), \quad (4)$$

where

$$c_H(0) \sim \frac{c_H^0}{1 + \frac{\sqrt{D_H k_1 c_{PS^-}^0}}{k_p}}, \quad (5)$$

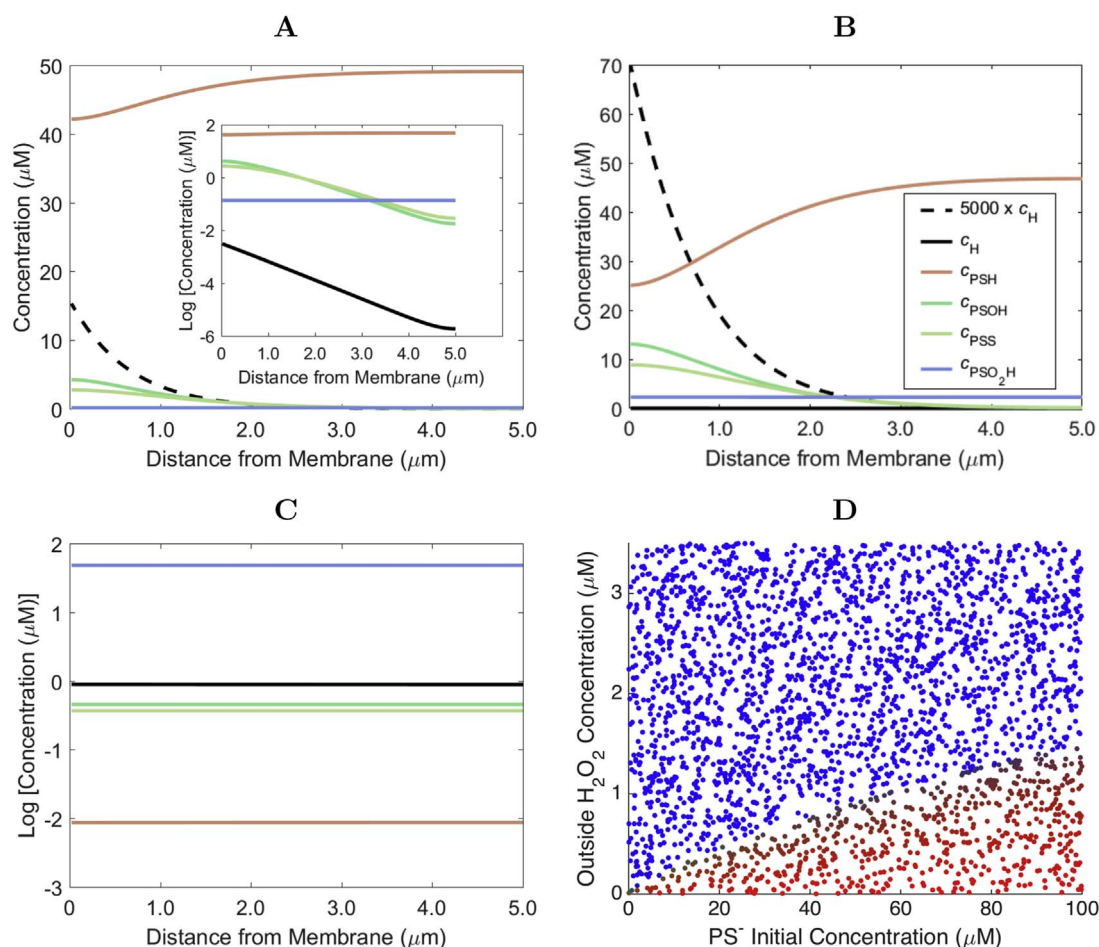
and

$$\lambda = \sqrt{\frac{D_H}{k_1 c_{PS^-}^0}} \quad (6)$$

is the length scale of the decay of the H<sub>2</sub>O<sub>2</sub> concentration. Considering a  $c_{PS^-}^0 \approx 50 \mu\text{M}$  and the parameter values in Table 1, one obtains  $\lambda \approx 0.60 \mu\text{m}$ . Considering the 5-fold lower value of  $D_H$  obtained [48] for diffusion in a hydrogel with a viscosity approaching that of the cytoplasm one obtains  $\lambda \sim 0.27 \mu\text{m}$ , which should be more representative of the situation *in vivo*. The order of magnitude of  $\lambda$  will be the same over the physiological range of peroxiredoxin concentrations, due to the shallow dependence of  $\lambda$  on  $c_{PS^-}^{-1/2}$ .

### 3.2. Cytoplasmic H<sub>2</sub>O<sub>2</sub> concentration gradients coexist with concentration gradients of sulfenic and disulfide peroxiredoxin forms

Importantly, coexisting with the cytoplasmic H<sub>2</sub>O<sub>2</sub> gradients there are also strong concentration gradients of PSO<sup>-</sup> and PSS (Fig. 3A, B). However, this is not the case of PSO<sub>2</sub><sup>-</sup>, whose concentration remains uniformly low throughout cytoplasm. The reasons for these differences among the spatial distributions of the various peroxiredoxin forms are as follows. H<sub>2</sub>O<sub>2</sub> rapidly oxidizes PS<sup>-</sup> to PSO<sup>-</sup> near the membrane, and PSO<sup>-</sup> is rapidly converted to PSS before it has time to diffuse far away. Reduction of PSS to PS<sup>-</sup> is also too fast to permit this species to diffuse



**Fig. 3. Spatial distribution of peroxiredoxin species:** **A:** Concentration profiles of  $\text{H}_2\text{O}_2$ ,  $\text{PS}^-$ ,  $\text{PSO}^-$ ,  $\text{PSS}$  and  $\text{PSO}_2$  as a function of the distance from the membrane in the case of  $c_{\text{PS}^-}(0) = 50 \mu\text{M}$  and  $c_{\text{H}}^0 = 0.1 \mu\text{M}$ . For this low  $\text{H}_2\text{O}_2$  influx rate, only a small fraction of  $\text{PS}^-$  is oxidized to  $\text{PSO}^-$  and  $\text{PSS}$ . The inset highlights the different orders of magnitude of the concentration of the various species and the strong gradients of  $c_{\text{PSO}^-}$  and  $c_{\text{PSS}}$ . The color code is as indicated in panel B. Note that in the main plot the concentration of  $\text{H}_2\text{O}_2$  is multiplied by 5000 for visualization in the same scale. **B:** Same as A but for  $c_{\text{H}}^0 = 0.4 \mu\text{M}$ . In this case  $c_{\text{PS}^-}$  is slightly depressed near the membrane, and  $c_{\text{PSO}^-}$  and  $c_{\text{PSS}}$  are elevated. Due to the very low reduction rate constant for  $\text{PSO}_2$ , the concentration of this form becomes almost constant over the cytoplasm. **C:** Same as A but for  $c_{\text{H}}^0 = 1.0 \mu\text{M}$ . For these high extracellular concentrations, the  $\text{H}_2\text{O}_2$  invades the cell and oxidizes most of the  $\text{PS}^-$  to  $\text{PSO}_2$ . **D:** Ratio between the average concentration of the various forms of peroxiredoxin and the initial concentration of  $\text{PS}^-$ . Each point is colored according to the relative fractions of peroxiredoxin in each form. Pure red and blue indicate that 100% of the peroxiredoxin is in the  $\text{PS}^-$  or  $\text{PSO}_2$  forms, respectively. Pure green indicates that 100% of the peroxiredoxin is in the  $\text{PSO}^- + \text{PSS}$  forms. Comparing this figure to Fig. 2B, we observe that when the outside  $\text{H}_2\text{O}_2$  concentration is very high, all the peroxiredoxin is hyperoxidized and the  $\text{H}_2\text{O}_2$  is able to penetrate the whole cell. For low extracellular  $\text{H}_2\text{O}_2$  concentrations, most of the peroxiredoxin is in its reduced form. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

far away from the membrane. In turn, both the formation of  $\text{PSO}_2$  and its reduction to  $\text{PSO}^-$  are much slower, allowing  $\text{PSO}_2$  to diffuse throughout the cytoplasm.

In a narrow range of  $c_{\text{H}}^0$  approaching the upper border of the blue region in Fig. 2B, the concentration of  $\text{PS}^-$  close to the membrane drops strongly (Fig. 3B) and the approximation of constant  $c_{\text{PS}^-}$  breaks down. Due to the lower  $\text{PS}^-$  concentration, both the  $\text{H}_2\text{O}_2$  concentration near the membrane (Fig. 2B) and the penetration length (Fig. 2C) increase in this range of  $c_{\text{H}}^0$ . However, the increase in the  $\text{H}_2\text{O}_2$  concentration near the membrane in this range of  $c_{\text{H}}^0$  is relatively modest.

### 3.3. Cytoplasmic $\text{H}_2\text{O}_2$ gradients collapse at high $\text{H}_2\text{O}_2$ supplies

As characterized more extensively elsewhere for a model neglecting concentration gradients [43], in cells where a high peroxiredoxin reduction capacity coexists with limited alternative  $\text{H}_2\text{O}_2$  sinks the system shows bi-stability and hysteresis. This dynamic characteristic manifests itself as a run-away hyperoxidation of nearly all the peroxiredoxin and a drastic increase in cytoplasmic  $\text{H}_2\text{O}_2$ , once the extracellular  $\text{H}_2\text{O}_2$  concentration exceeds a critical value ( $c_{\text{H}}^{0*}$ ). Once this new steady state is established, recovery starts only after  $c_{\text{H}}^0$

decreases below another lower critical value ( $c_{\text{H}}^{0**} < c_{\text{H}}^{0*}$ ) and takes several hours. The reaction-diffusion model considered in the present work shows this behavior as well. When the extracellular  $\text{H}_2\text{O}_2$  concentration reaches a critical value ( $c_{\text{H}}^{0*}$ ), the peroxiredoxin accumulates in hyperoxidized form (Fig. 3D). As a consequence, the cytoplasmic concentration gradients of  $\text{H}_2\text{O}_2$ ,  $\text{PS}^-$ ,  $\text{PSO}^-$  and  $\text{PSS}$  collapse (Fig. 3C), and  $\text{H}_2\text{O}_2$  concentrations become much higher throughout all the cytoplasm (Fig. 2B). The transition between these regimes is quite abrupt, and  $c_{\text{H}}^{0*}$  increases approximately as the square root of the initial concentration of peroxiredoxin in the cell (Fig. 2B).

Remarkably, if one starts with a system where the peroxiredoxin is initially hyperoxidized, the critical values of  $c_{\text{H}}^0$  ( $c_{\text{H}}^{0**}$ ) below which the peroxiredoxins are again able to sustain a  $\text{H}_2\text{O}_2$  gradient are considerably lower than  $c_{\text{H}}^{0*}$  (Fig. 2D). The system thus presents hysteresis.

Note that the actual values of  $c_{\text{H}}^{0*}$  and  $c_{\text{H}}^{0**}$  depend strongly on the permeability of the membrane and on the activity of alternative  $\text{H}_2\text{O}_2$  sinks. Here we are neglecting these sinks and considering a high permeability constant (see Section 2). Therefore, we expect that in most cells these critical concentrations are higher than depicted in Figs. 2 and 3.



### 3.4. The concentration of $H_2O_2$ near the membrane is insufficient to oxidize PTPs

Known rate constants for direct oxidation of PTPs by  $H_2O_2$  are  $k_{ox} \leq 164 M^{-1} s^{-1}$ , which raises the question of whether  $H_2O_2$  concentrations sufficient to directly oxidize PTPs are attainable in the cytoplasm. In order to frame this question, we first examine how the maximal cytoplasmic  $H_2O_2$  concentrations relate to the extracellular concentrations, peroxiredoxin concentrations and cytoplasmic gradients to highlight a fundamental trade-off between signal localization and the maximal cytoplasmic  $H_2O_2$  concentration. The latter concentration occurs adjacent to the membrane. Under conditions where only a small fraction of the peroxiredoxin is oxidized it is approximated by Eq. (5). The product  $k_{sink} = k_1 c_{PS^-}$  in this equation represents the pseudo-first-order rate constant for  $H_2O_2$  consumption in the cytoplasm, and therefore the following conclusions apply irrespective of the main cytoplasmic process contributing to  $k_{sink}$ . Considering a generic eukaryotic cell with  $10 \mu m$  width, for signaling to be localized the length scale of the cytoplasmic  $H_2O_2$  should be  $< 1 \mu m$ . As per Eq. (6) this implies  $k_{sink} > 310 s^{-1}$ , conservatively considering  $D_H = 3.1 \times 10^{-10} m^2 s^{-1}$  as for  $H_2O_2$  diffusion in a hydrogel [48]. Because even for the most permeable cells  $k_p < 10^{-5} m s^{-1}$ , Eq. (5) implies  $c_H(0)/c_H^0 < 1/30$ . Therefore, in most cells the maximal cytoplasmic  $H_2O_2$  concentrations are one to two orders of magnitude lower than the extracellular ones under conditions that warrant localized signaling. Furthermore, because  $\sqrt{D_H k_1 c_{PS^-}}/k_p \gg 1$ , for fixed  $D_H$  and  $k_p$  Eqs. (4) and (5) imply that  $c_H(0) \propto \lambda$ . Hence, any decrease in  $k_{sink}$  will proportionately increase the length scale of the gradient. Moreover, the approximate proportionality of  $c_H(0)$  to  $k_{sink}^{-1/2}$  implies that only very drastic inhibitions of cytoplasmic  $H_2O_2$  clearance can have a substantial impact on the  $H_2O_2$  concentrations near supply sites. For instance, in order increase  $c_H(0)$  2-fold Prx would have to be at least 75% inhibited.

Considering the  $c_H(0)/c_H^0$  ratio estimated above, a  $c_H^0 = 0.5 \mu M$  would take  $\sim 5.6$  days to oxidize 63% of a very reactive PTPs residing at the cytoplasmic face of the membrane. Therefore, the results above indicate that in the regime where steep cytoplasmic gradients are maintained,  $H_2O_2$  concentrations are far too low to directly oxidize PTPs in a mitogenic signaling time frame. Indeed, numerical results considering more generic conditions (Fig. 4A) fully substantiate this view: in this regime the oxidation of the PTPs could take weeks to years even at the points of highest  $H_2O_2$  concentration. At higher  $c_H^0$  values above  $c_H^{0*}$  the  $H_2O_2$  invades the cytoplasm, as per the previous section,

and the time for oxidation decreases. [These results remain qualitatively the same if the higher condensation rate constant estimated for peroxiredoxin I [43] is considered (Fig. S1A).] However, although peroxisome-sequestered catalase and remaining glutathione thiolate cannot sustain a cytoplasmic  $H_2O_2$  gradient where NADPH depletion prevents the action of peroxiredoxins and peroxidases, they can still substantially delay the oxidation of PTPs by  $H_2O_2$  [14].

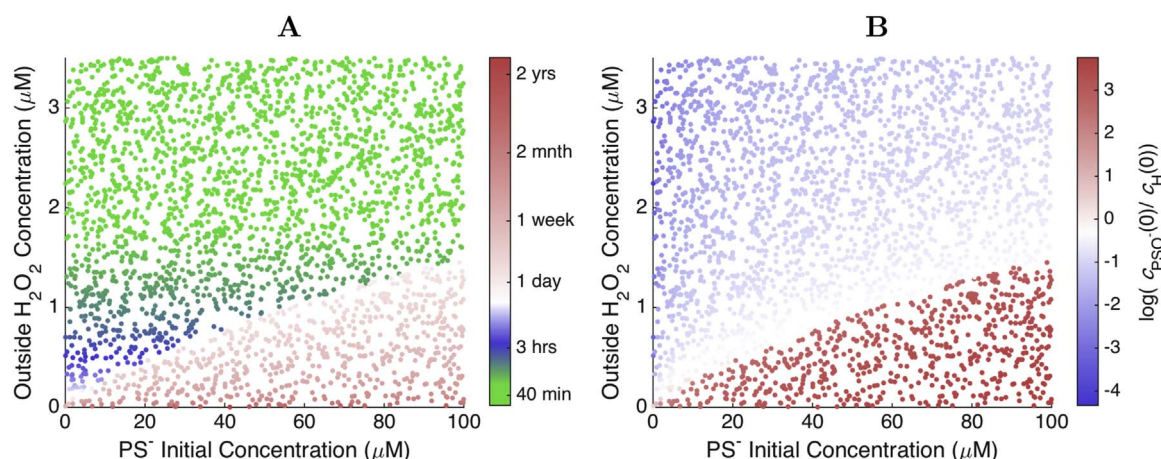
Although the results above pertain to a scenario where  $H_2O_2$  permeates uniformly across all the membrane, the maximal  $H_2O_2$  concentrations remain insufficient to quickly oxidize PTPs even in the extreme scenario where all the permeation is localized in a specific channel with a  $50 nm$  diameter (Fig. S1B,C). Therefore, PTPs cannot be directly oxidized by  $H_2O_2$  under conditions that avoid extensive peroxiredoxin hyperoxidation and keep the cytoplasm protected against excessive  $H_2O_2$ .

The hysteretic behavior described above is abolished if cells have (a) lower capacity to reduce PSS and/or (b) more-active alternative  $H_2O_2$  sinks. In case (a) high extracytoplasmic  $H_2O_2$  concentrations cause the cytoplasmic 2-Cys peroxiredoxins to accumulate in disulfide form instead (Fig. S2A), which is also accompanied by the collapse of the  $H_2O_2$  gradients (Fig. S2B). In case (b) peroxiredoxin oxidation increases more gradually with the  $H_2O_2$  supply rate, and the alternative  $H_2O_2$  sinks may eventually be able to sustain a cytoplasmic  $H_2O_2$  concentration gradient. However, cytoplasmic  $H_2O_2$  concentrations will be even lower and oxidation of PTP by  $H_2O_2$  even slower in this case.

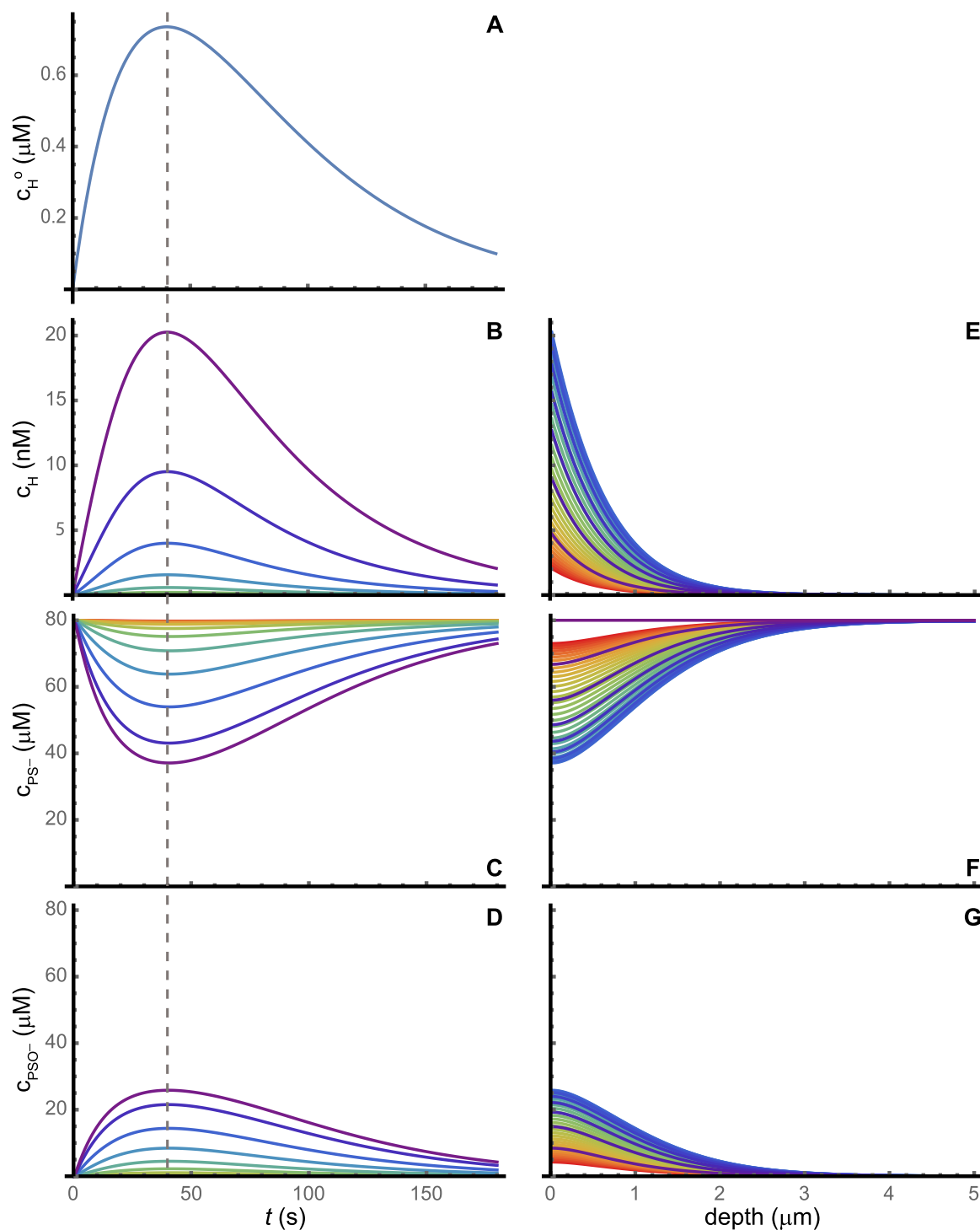
### 3.5. Peroxiredoxins show good signaling properties for moderate $H_2O_2$ pulses

Interestingly, under all conditions where strong intracellular concentration gradients of  $H_2O_2$  are sustained the concentrations of  $PSO^-$  and PSS exceed those of  $H_2O_2$  by several orders of magnitude (Fig. 4B). As a consequence, these oxidized peroxiredoxin forms can oxidize PTPs and other redox targets much faster than  $H_2O_2$  if their reactivity with these targets is comparable to that of  $H_2O_2$ .

All the results presented so far pertain to situations where the extracellular  $H_2O_2$  concentration is sustained long enough for the system to approach a steady state. However,  $H_2O_2$  production is very dynamic in the seconds and minutes subsequent to growth factor binding to receptors [25,27,20]. Therefore, in order to further assess the signaling potentialities of  $PSO^-$  and PSS vs.  $H_2O_2$  in this signaling context we simulated pulses of extracellular  $H_2O_2$  concentration. The



**Fig. 4.  $H_2O_2$  capacity for signaling:** **A:** Minimum time scale for direct oxidation of PTPs by  $H_2O_2$ . The inverse of the pseudo-first-order rate constant for the direct oxidation of the active form of a PTP by  $H_2O_2$  with  $k_{ox}=164 M^{-1} s^{-1}$  near the membrane is color coded as indicated. Values give the time needed to oxidize 63% of the target. Note the extremely slow oxidation under conditions that permit strong  $H_2O_2$  gradients. Especially in the high hyperoxidation regime, times for oxidation are underestimated due to the neglect of alternative  $H_2O_2$  sinks. **B:** Ratio between average intracellular concentrations of  $PSO^-$  and  $H_2O_2$ . Note the very high ratios ( $> 1000$ ) under conditions that permit strong  $H_2O_2$  gradients. Results for the ratios between average intracellular concentrations of PSS and  $H_2O_2$  are qualitatively similar. (The reader is referred to the web version of this article for a color version of this figure.).



**Fig. 5. Response to extracellular  $\text{H}_2\text{O}_2$  pulse.** **A:** Time course of the pulse. **B, C, D:** Time course of cytoplasmic concentrations of  $\text{H}_2\text{O}_2$ ,  $\text{PS}^-$ , and  $\text{PSO}^-$ , respectively, at depths 0 (violet) to 5  $\mu\text{m}$  (red) from the membrane (0.5  $\mu\text{m}$  steps). **E, F, G:** Spatial profiles of cytoplasmic concentrations of  $\text{H}_2\text{O}_2$ ,  $\text{PS}^-$ , and  $\text{PSO}^-$ , respectively, at times 0 (violet) to 180 s (red) from the membrane (5 s steps). The violet lines in panels E–G reflect the initial concentration change over depth. The time courses and spatial profiles for PSS are very similar to those for  $\text{PSO}^-$ . Note the different scales for extracellular  $\text{H}_2\text{O}_2$ , cytoplasmic  $\text{H}_2\text{O}_2$  and peroxiredoxin concentrations. The dashed gray line marks the time at which the cytoplasmic  $\text{H}_2\text{O}_2$  concentration reaches its maximum. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

pulse shown in Fig. 5A attains a maximum extracellular  $\text{H}_2\text{O}_2$  concentration of  $\approx 0.75 \mu\text{M}$ . The brief pulse causes little ( $< 0.05\%$ ) hyperoxidation, but substantial peroxiredoxin oxidation (Fig. 5C,F). The results highlight two desirable properties of these peroxiredoxin forms as signaling intermediates. First, and as observed for the steady-state results, they attain much higher cytoplasmic concentrations than  $\text{H}_2\text{O}_2$  (compare Fig. 5B,D). Second, they propagate the signal deeper

into the cell than  $\text{H}_2\text{O}_2$  (compare Fig. 5E,G). Namely, whereas the concentration of  $\text{H}_2\text{O}_2$  decays near-exponentially with depth, high  $\text{PSO}^-$  and  $\text{PSS}$  concentrations are attained even 0.5  $\mu\text{m}$  from the membrane (Fig. 5D,G), allowing the oxidation of potential targets even at that depth.

These good signaling properties deteriorate for stronger pulses. A pulse attaining a maximum extracellular  $\text{H}_2\text{O}_2$  concentration of 5.9  $\mu\text{M}$  already



causes  $\approx 35\%$  hyperoxidation (Fig. S3E), and substantial  $\text{H}_2\text{O}_2$  penetration in the cytoplasm (Fig. S3F). The concentrations of  $\text{PSO}^-$  and  $\text{PSS}$  start to decrease earlier than for the weaker pulse, and still in the ascending phase of the pulse, due to gradual hyperoxidation (Fig. S3D). The concentration gradients of these species are temporarily flattened out. Furthermore, the recovery of the peroxiredoxin redox status is strongly delayed, due to the slow reduction of  $\text{PSO}_2^-$  (Fig. S3C,D,E).

A pulse attaining a maximum extracellular  $\text{H}_2\text{O}_2$  concentration of  $12\ \mu\text{M}$  triggers the hyperoxidation “catastrophe” described above (Fig. S4). Virtually all peroxiredoxin is hyperoxidized within 1 min (Fig. S4E), all the cytoplasmic gradients collapse (Fig. S4F–H). Although maximal  $\text{PSO}^-$  and  $\text{PSS}$  concentrations similar to the previous pulse are attained at  $\approx 10\ \text{s}$  after the onset of the pulse, the concentrations of these species fall precipitously thereafter and nearly vanish after  $\approx 1\ \text{min}$  (Fig. S4D). Recovery of the peroxiredoxin redox state after the pulse occurs in a time scale of hours.

### 3.6. Peroxiredoxin recruitment causes modest local $\text{H}_2\text{O}_2$ depletion

A small fraction (up to a few percent) of PrxI and PrxII is able to associate to the cellular membrane [51,25,27]. Moreover, PrxII is recruited to VEGFR2 upon VEGF stimulation of endothelial cells, protecting this receptor against oxidative inactivation by  $\text{H}_2\text{O}_2$  [37]. The most straightforward explanation for this protection is that the colocalization of PrxII locally depletes  $\text{H}_2\text{O}_2$ . In order to assess the extent of local  $\text{H}_2\text{O}_2$  depletion that is achievable in this way, we model two different scenarios. In the first scenario we consider that 2% of all the peroxiredoxin is reversibly and uniformly bound to the membrane, and that all redox states of peroxiredoxin show similar binding and unbinding rate constants. This system is described by Eqs. (3). Similarly to the system analyzed in the previous sections, the  $\text{H}_2\text{O}_2$  concentration either drops exponentially near the membrane, or invades the cell, depending on the value of  $c_{\text{H}}^0$  and on the total concentration of peroxiredoxin. The binding of this small fraction of peroxiredoxin to the membrane does not alter the critical values  $c_{\text{H}}^{0*}$  (compare Figs. 6A and 3B). However, it decreases the concentration of  $\text{H}_2\text{O}_2$  adjacent to the membrane under the conditions where the peroxiredoxin is able to sustain a strong cytoplasmic  $\text{H}_2\text{O}_2$  concentration gradient (*i.e.*, where  $c_{\text{H}}^0 < c_{\text{H}}^{0*}$ ). This decrease is in the range of 10–25%, and is the stronger the higher the initial  $c_{\text{PS}^-}$  and the lower  $c_{\text{H}}^0$  are (Fig. 6A).

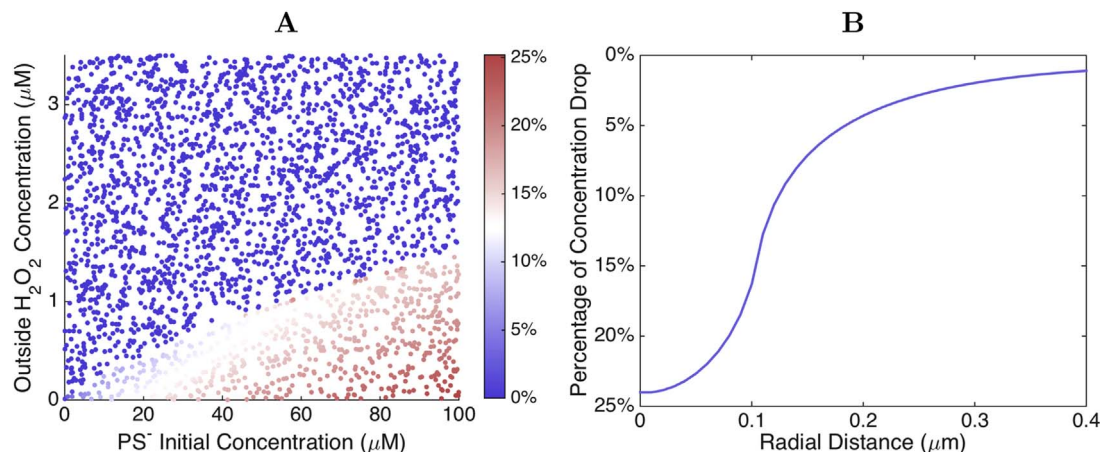
In the second scenario 2% of all peroxiredoxin is bound specifically to one caveola. We considered a cylindrical 3D simulation box as depicted in Fig. 1C. We ran the simulation in the regime of interest, *i.e.*,

for high initial concentration of  $\text{PS}^-$  ( $c_{\text{PS}^-}(\mathbf{r}, 0) = 80\ \mu\text{M}$ ) and relatively low  $c_{\text{H}}^0$  ( $0.40\ \mu\text{M}$ ). The 1D results for the previous scenario indicate that these parameter values yield a moderate protection of the receptor by the bound peroxiredoxins, with little  $\text{PSO}_2^-$  accumulation. In the current 3D scenario, the concentration of  $\text{PSO}_2^-$  is also negligible, and a steady state is achieved by 3 s of simulation time. In this simulation, the steady state depends strongly on the magnitude of the binding and unbinding constants. We chose values of  $k^+ = 1.0 \times 10^4\ \text{s}^{-1}$  and  $k^- = 39.2\ \text{s}^{-1}$ , which guarantee the binding of 2% of the peroxiredoxin to the membrane.

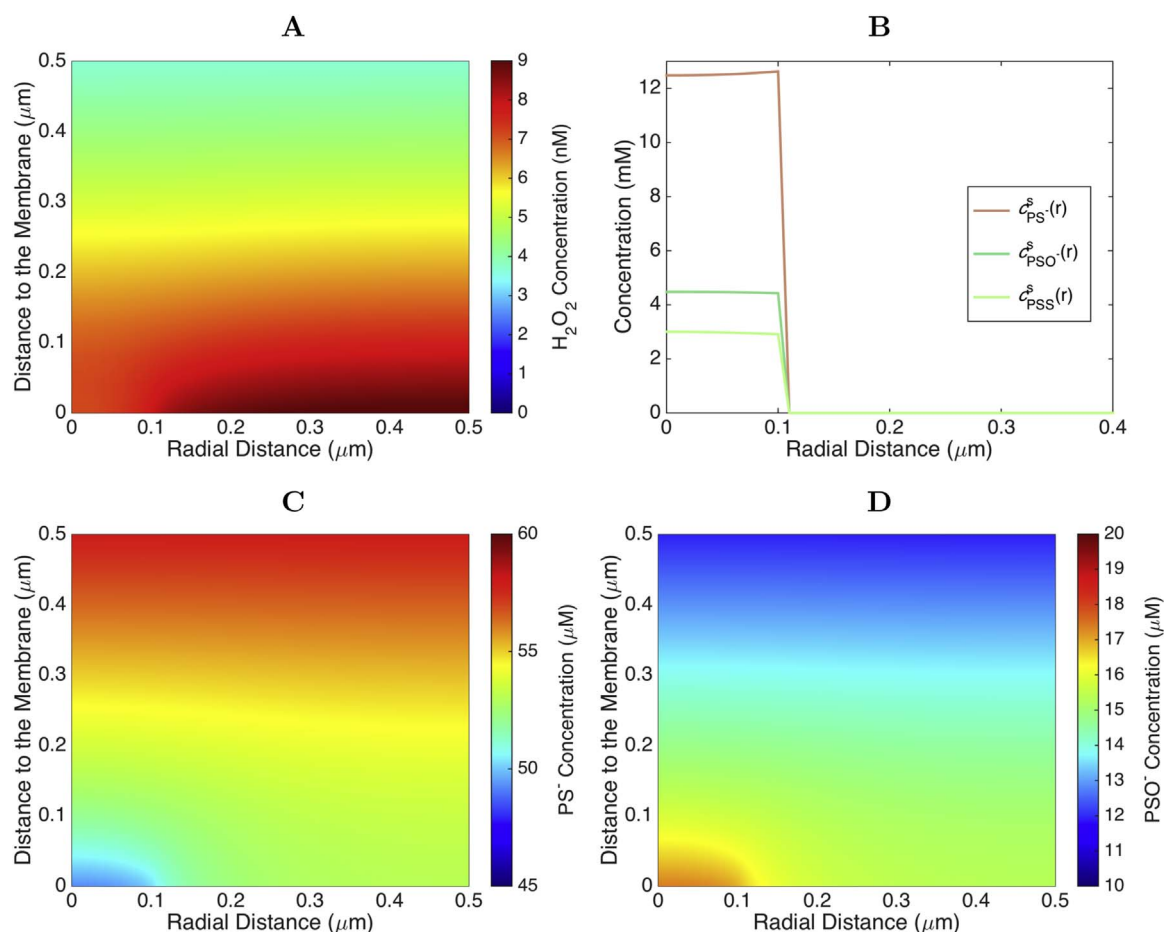
Under these conditions  $\text{H}_2\text{O}_2$  is appreciably depleted near the caveola (Figs. 6B and 7A). However, even this very strong concentration of peroxiredoxin at a caveola decreases the local  $\text{H}_2\text{O}_2$  concentration adjacent to the membrane by no more than  $\approx 25\%$ , similar to the 1D simulation. This happens despite the local concentration of  $\text{PS}^-$  at the caveolae ( $\approx 10\ \text{mM}$ , Fig. 7B) strongly exceeding the concentration at the membrane in the previous scenario, which should apparently lead to a stronger protection of the receptor. However, as the  $\text{PS}^-$  at the caveola is oxidized by incoming  $\text{H}_2\text{O}_2$  the quick binding of new  $\text{PS}^-$  molecules depletes them in the cytoplasm around the caveola (Fig. 7C), owing to their relatively slow diffusion. In turn, this allows more  $\text{H}_2\text{O}_2$  to diffuse from neighboring regions in the cytoplasm, thereby attenuating the effectiveness of protection. Remarkably, in the plane of the membrane not only the bound  $\text{PS}^-$  but also the bound  $\text{PSS}$  and  $\text{PSO}^-$  are strongly concentrated at the caveola region; that is, within  $0.10\ \mu\text{m}$  from the receptor (Fig. 7B). Furthermore, the concentrations of unbound  $\text{PSS}$  and  $\text{PSO}^-$  are also elevated in the cytoplasm near the caveola (Fig. 7D). Therefore, the recruitment of peroxiredoxins to caveolae can strongly favor the peroxiredoxin-mediated oxidation of redox targets over their direct oxidation by  $\text{H}_2\text{O}_2$ .

## 4. Discussion

The results above address several key questions about  $\text{H}_2\text{O}_2$  signaling and the participation of cytoplasmic 2-Cys peroxiredoxins in this process. Consistent with previous estimates [15,40], models [18,19] and experimental observations [20], we found that typical concentrations of 2-Cys peroxiredoxins are sufficient to sustain strong  $\text{H}_2\text{O}_2$  concentration gradients in the cytoplasm of mammalian cells. As a result, in the absence of oxidative stresses able to extensively oxidize these peroxiredoxins most of the  $\text{H}_2\text{O}_2$  in the cell cytoplasm is contained within a few tenths of  $\mu\text{m}$  of the supply sites (Fig. 2A,C). The existence of such steep  $\text{H}_2\text{O}_2$  concentration gradients could explain how effective antioxidant protection can be achieved despite the high



**Fig. 6. Effects of peroxiredoxin recruitment: (1) Local  $\text{H}_2\text{O}_2$  depletion.** **A:** Decrease in  $\text{H}_2\text{O}_2$  concentration near the membrane if 2% of the peroxiredoxin is uniformly distributed at the membrane. **B:** Decrease in  $\text{H}_2\text{O}_2$  concentration near caveolae if 2% of the peroxiredoxin is concentrated at one caveola. For the explored range of conditions local  $\text{H}_2\text{O}_2$  concentrations can decrease by up to 25%. (The reader is referred to the web version of this article for a color version of this figure.).



**Fig. 7. Effects of peroxiredoxin recruitment (2): Distribution of  $\text{H}_2\text{O}_2$  and oxidized peroxiredoxin for the case where 2% of the peroxiredoxin is bound at one caveola.** **A:** Distribution of  $\text{H}_2\text{O}_2$  close to the caveola. **B:** Distribution of the various peroxiredoxin forms at the membrane. **C:** Distribution of non-bound  $\text{PS}^-$  close to the caveola. We observe that in the caveola vicinity there is a minimum in the  $\text{PS}^-$  concentration. **D:** Distribution of non-bound  $\text{PSO}^-$  close to the caveola. The distribution of  $\text{PSS}^-$  is qualitatively similar. The concentrations of these peroxiredoxin forms is maximal next to the caveola. (The reader is referred to the web version of this article for a color version of this figure.).

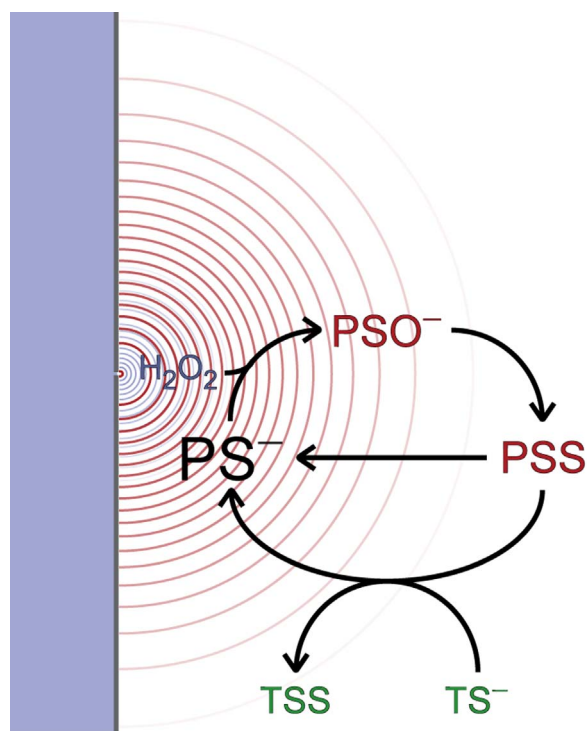
local  $\text{H}_2\text{O}_2$  production rates involved in redox signaling [15]. However, the  $\text{H}_2\text{O}_2$  concentrations attained even at the apex of these gradients are by and large insufficient to substantially oxidize PTPs in a mitogenic signaling time frame. Conservatively, we considered very high values for the  $\text{H}_2\text{O}_2$  permeability constant and for the PTP oxidation rate constant, assumed diffusion constants for peroxiredoxin consistent with the mobility of its decameric form in a crowded cellular environment, and neglected alternative cytoplasmic  $\text{H}_2\text{O}_2$  sinks. But even under these conservative assumptions oxidation of the PTPs would occur in a time scale of weeks (Fig. 4A) or longer. In an extreme scenario where all the  $\text{H}_2\text{O}_2$  would permeate through a single channel with 50 nm diameter the local  $\text{H}_2\text{O}_2$  concentrations would oxidize PTPs in a time scale of several hours (Fig. S1B,C), still too slow for mitogenic signaling. This outcome ensues from the fundamental trade-off demonstrated in Section 3.4. Namely, given physiologically plausible  $\text{H}_2\text{O}_2$  membrane permeabilities and diffusion constants, any agent consuming  $\text{H}_2\text{O}_2$  fast enough to localize it within  $\approx 1 \mu\text{m}$  of a permeant membrane also generate a  $>30$ -fold trans-membrane gradient.

At moderately high  $\text{H}_2\text{O}_2$  supply rates, peroxiredoxins can become substantially oxidized near the membrane, thus allowing for somewhat higher local  $\text{H}_2\text{O}_2$  concentrations (Fig. 3B) at the cost of shallower cytoplasmic concentration gradients (Fig. 2C). However, the effect is modest (Fig. 2B,C) and restricted to a narrow range of  $\text{H}_2\text{O}_2$  supply rates. These results also imply that the local inhibition of peroxiredoxins' peroxidatic activity – be it by hyperoxidation [21], phosphorylation [27] or non-covalent [40] – can only significantly increase local  $\text{H}_2\text{O}_2$  concentrations if it substantially decreases the total cytoplasmic

peroxidatic activity. Otherwise, the local inhibitory effect is efficiently counteracted by the diffusion of active peroxiredoxin from the bulk. However, whereas PrxI is inhibited by phosphorylation but not very susceptible to hyperoxidation, PrxII is susceptible to hyperoxidation but not inhibited by phosphorylation [27]. Because the concentrations of these two peroxiredoxins in the cytoplasm of mammalian cells are broadly similar [11], neither of these inhibitory mechanisms in isolation can dramatically increase local  $\text{H}_2\text{O}_2$  concentrations. Indeed, it follows from Eq. (5) that a 50% decrease in total peroxidase activity will increase the  $\text{H}_2\text{O}_2$  concentration near the membrane by at most  $1 - \sqrt{0.5} = 29\%$ . And as per Eq. (4) this will come at the cost of extending the length scale of  $\text{H}_2\text{O}_2$  gradient by 29% as well.

At  $\text{H}_2\text{O}_2$  supply rates slightly higher than those discussed in the previous paragraph, the peroxiredoxins become fully oxidized and  $\text{H}_2\text{O}_2$  can then invade all the cytoplasm. But even in this situation where the interior of the cytoplasm would no longer be protected the oxidation of PTPs would be slow (Fig. 4A). A simple calculation shows that in order to oxidize 50% of the most reactive PTPs characterized to date ( $k_{ox} = 164 \text{ M}^{-1} \text{ s}^{-1}$  [6]) in 10 min a  $\text{H}_2\text{O}_2$  concentration of  $\log(2)/(164 \text{ M}^{-1} \text{ s}^{-1} \times 600 \text{ s}) = 7 \mu\text{M}$  would be required. The oxidation of other PTPs in the same time frame would require  $\text{H}_2\text{O}_2$  concentrations in the order of 100  $\mu\text{M}$ . However, sustained 7  $\mu\text{M}$  extracellular  $\text{H}_2\text{O}_2$  is sufficient to make Jurkat T cells apoptose [52]. It is thus very unlikely that such high concentrations are attained anywhere in the cytoplasm.

The sulfenic and/or disulfide forms of 2-Cys peroxiredoxins oxidize redox targets *in vitro* and *in vivo* forming redox relays [34,35,28,2,36].



**Fig. 8. H<sub>2</sub>O<sub>2</sub> signaling through localized redox relays.** At H<sub>2</sub>O<sub>2</sub> (blue) supply rates commensurate with mitogenic signaling typical 2-Cys peroxiredoxins in the cytoplasm of mammalian cells sustain steep gradients but keep H<sub>2</sub>O<sub>2</sub> concentrations very low throughout. This outcome is determined by the high cytoplasmic concentrations and high reactivity of these proteins. But these factors together with the limited rates at which the sulfenic and disulfenic forms of these proteins (red) are reduced also determine that the concentrations of these forms strongly exceed those of H<sub>2</sub>O<sub>2</sub>. The oxidized peroxiredoxin species also show steep concentration gradients over the cytoplasm, here represented by level curves in red lines. Therefore, the oxidation of redox targets (green) mediated by them is also restricted to a vicinity of the H<sub>2</sub>O<sub>2</sub> supply sites. But unlike the H<sub>2</sub>O<sub>2</sub> concentration gradients (level curves in blue), those of the peroxiredoxin species plateau near the H<sub>2</sub>O<sub>2</sub> supply sites, allowing the redox signal to penetrate slightly deeper into the cytoplasm. The competitive advantage of the peroxiredoxin sulfenic or disulfide forms over H<sub>2</sub>O<sub>2</sub> in oxidizing targets at specific sites (*e.g.*, membranes, caveolae or endosomes) can be strongly amplified by localizing even a small fraction of the cytoplasmic peroxiredoxins to those sites. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Remarkably, the computational analyses presented here highlight three important advantages of peroxiredoxin-mediated oxidation of redox targets (*i.e.*, redox relays) over direct oxidation of the targets by H<sub>2</sub>O<sub>2</sub>. First, the concentrations of PSO<sup>−</sup> and PSS exceed those of H<sub>2</sub>O<sub>2</sub> by several orders of magnitude under conditions where H<sub>2</sub>O<sub>2</sub> is contained near its supply sites (Fig. 4B). Therefore, even if these peroxiredoxin forms are just as reactive with the redox targets as H<sub>2</sub>O<sub>2</sub> they can oxidize them much faster. However, the case of thioredoxin ( $k_5 = 2.1 \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$  [13]) illustrates that PSS can oxidize some targets at very high rate constants. Second, there are also important concentration gradients of these peroxiredoxin forms, thus retaining the capacity for localized signaling. Third, unlike the H<sub>2</sub>O<sub>2</sub> gradients, those of PSO<sup>−</sup> and PSS plateau at the membrane (Figs. 3A,B and 5G). This further favors the oxidation of the targets and allows the redox signal to propagate deeper into the cytoplasm. The latter may be important in some signaling contexts where the targets are located in intracellular membranes [53,54]. The penetration of these peroxiredoxin forms in the cytoplasm is less problematic from the point of view of antioxidant protection than that of H<sub>2</sub>O<sub>2</sub> because their interactions are expected to be specific. These considerations suggest a “localized redox relay” model of redox signaling (Fig. 8). This model is consistent with the observations by Klomsiri et al. [55] of localized formation of

reversibly oxidized protein forms near H<sub>2</sub>O<sub>2</sub> sources in LPA-stimulated PC3 and SKOV3 cells.

The localized oxidation of the Prx could regulate targets in at least the following four alternative ways. First, PSO<sup>−</sup> and PSS can directly oxidize some targets [33,34,28,35,2]. Second, because these Prx forms can be glutathionylated (PSSG) [56,57], PSSG might in turn glutathionylate redox targets either directly or mediated by glutaredoxin, as suggested in ref. [14]. S-glutathionylation is a well-established regulatory post-translational modification for many proteins, and this mode of operation would mirror similar proposals with glutathione peroxidases as peroxide sensors and glutaredoxins as key players [58–61]. However, the presently available information about the kinetics of PSSG deglutathionylation does not permit to determine if this species is also localized. Third, the transfer of oxidizing equivalents to targets might also be mediated by Trx [62]. Fourth, a conformational change induced by Prx oxidation might either enable or hinder a non-covalent interaction with a target [63]. Kinetic experiments *in vitro* will be instrumental for testing if PSO<sup>−</sup>, PSS and/or PSSG can outcompete H<sub>2</sub>O<sub>2</sub> for actuation of redox targets.

The results above highlight that despite small and very diffusible, H<sub>2</sub>O<sub>2</sub> cannot diffuse far away from production sites due to the very high activity of cytoplasmic peroxiredoxins, and the redox signal is actually carried to distal sites by proteins.

Although under conditions of mitogenic signaling hyperoxidation has just a minor effect on local H<sub>2</sub>O<sub>2</sub> concentrations and weakens the cytoplasmic concentration gradient, at higher H<sub>2</sub>O<sub>2</sub> the opening of the hyperoxidation “floodgate” could be part of a stress response. The analysis in ref. [43] predicts that in cells with abundant peroxiredoxin reduction capacity and limited alternative H<sub>2</sub>O<sub>2</sub> sinks the peroxiredoxin/thioredoxin/sulfiredoxin system shows bi-stability and hysteresis. The simulations in Figs. 2 and 3 show that the predicted bi-stability and hysteresis persist when concentration gradients are taken into account. Furthermore, they show that the onset of the high-hyperoxidation state is accompanied by the collapse of the cytoplasmic H<sub>2</sub>O<sub>2</sub> and peroxiredoxin concentration gradients, and by very low PSO<sup>−</sup> and PSS concentrations despite the high H<sub>2</sub>O<sub>2</sub> concentrations. Under these particular conditions, signal transmission through redox relays is interrupted and H<sub>2</sub>O<sub>2</sub> may become able to oxidize some intermediate-reactivity targets deep into the cell. This predicted behavior is reminiscent of experimental observations for *Schizosaccharomyces pombe*, where exposure to high H<sub>2</sub>O<sub>2</sub> concentrations caused strong hyperoxidation of the Tpx1 peroxiredoxin, interruption of redox-relay signaling to the Pap1 transcription factor and accumulation sufficient cytoplasmic H<sub>2</sub>O<sub>2</sub> to trigger the stress response mediated by mitogen-activated protein kinase Sty1 [33].

A small fraction of the cytoplasmic 2-Cys peroxiredoxins can be recruited to the cellular membrane [51,25,37] or to the centrosome [38], where they can protect sensitive proteins against oxidation by H<sub>2</sub>O<sub>2</sub> [37,38]. Our simulations of the effects of localizing 2% of the cytoplasmic peroxiredoxin (Figs. 6 and 7) show a modest (~25%) decrease in local H<sub>2</sub>O<sub>2</sub> concentrations despite the very high (up to 10 mM) local concentration of peroxiredoxin considered. Moreover, the effect is quantitatively similar irrespective of the peroxiredoxin binding uniformly to the membrane or just at caveolae. This is because the fast binding of PS<sup>−</sup> to the membrane in replacement of oxidized molecules depletes PS<sup>−</sup> from the neighboring cytoplasm, facilitating the inflow of H<sub>2</sub>O<sub>2</sub> into this region.

On the other hand, the recruitment of peroxiredoxin to specific sites can originate very high local concentrations of PSO<sup>−</sup> and PSS. This effect can be further amplified if the H<sub>2</sub>O<sub>2</sub> supply is also localized at the same sites, for instance through permeation by aquaporins [64,65]. Oxidation by PSO<sup>−</sup> and PSS of targets also recruited to the same sites could thus be very fast. Likewise, a local inactivation of PrxI as described in ref. [27] although having a modest effect on local H<sub>2</sub>O<sub>2</sub> concentrations could effectively block signal transmission through PrxI-mediated redox relays. Therefore, the possibility that the recruit-



ment of peroxiredoxins to specific sites plays a more important role in facilitating and regulating the operation of localized redox relays than on modulating local H<sub>2</sub>O<sub>2</sub> concentrations deserves further consideration. The observed protective effect of PrxII against VEGFR2 oxidation by H<sub>2</sub>O<sub>2</sub> can as well be explained in this framework: the selective binding of oxidized PrxII, but not PrxIIS<sup>-</sup>, to VEGFR2 might protect the sensitive thiols in the latter protein by blocking access to H<sub>2</sub>O<sub>2</sub> and/or by inducing a conformational change that hindered its reactivity.

Although the geometry addressed in this work applies most properly to situations where H<sub>2</sub>O<sub>2</sub> permeates the apical and basolateral membranes of a cell in a cell layer, the qualitative results are valid for other geometries. Namely, for H<sub>2</sub>O<sub>2</sub> release from endosomal vesicles or other intracellular membranes. Likewise, the key features determining the occurrence and functional advantages of localized redox relays over direct oxidation of targets by H<sub>2</sub>O<sub>2</sub> are common to most eukaryotic cells known to date.

### Author contributions

AS, FSA, RDMT designed analyses; AB, FSA, PA, RDMT performed analyses; AS, RDMT wrote the paper.

### Competing financial interests

The authors declare no competing financial interests.

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### Appendix A. Supplementary data

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.redox.2017.01.003>.

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