Novel Redox-Sensing Modules: Accessory Protein- and Nucleic Acid-Mediated Signaling

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Abstract

Significance: Organisms have evolved both enzymatic and nonenzymatic pathways to prevent oxidative damage to essential macromolecules, including proteins and nucleic acids. Pathways modulated by different protein-based sensory and regulatory modules ensure a rapid and appropriate response. **Recent Advances:** In contrast to classical two-component systems that possess internal sensory and regulatory modules, an accessory protein-dependent redox-signaling system has been recently characterized in bacteria. This system senses extracellular iron-mediated oxidative stress signals *via* an extracellularly located protein (HbpS). *In vivo* and *in vitro* studies allowed the elucidation of molecular mechanisms governing this system. Moreover, recent studies show that nucleic acids may also participate in redox-signaling during antioxidative stress response. **Critical Issues:** Research for novel redox-signaling systems is often focused on known types of sensory and regulatory modules. It is also often considered that the oxidative attack of macromolecules, leading to modification and degradation processes, is the final step during oxidative stress. However, recent studies have demonstrated that oxidatively modified macromolecules can be intermediary states in the process of redox-signaling. **Future Directions:** Analyses of adjacent regions of genes encoding for known sensory and regulatory modules can identify potential accessory modules that may increase the complexity of sensing systems. Despite the fact that the involvement of DNA-mediated signaling in the modulation of one bacterial regulator protein has been analyzed in detail, further studies are necessary to identify additional regulators. Given the role of DNA in oxidative-stress response, it is tempting to hypothesize that RNA modules may also mediate redox-signaling. *Antioxid. Redox Signal.* 16, 668–677.

Introduction

THE ABILITY OF ORGANISMS to respond to oxidative-stress events is strictly coupled to the available repertoire of macromolecules that can sense different redox states in the external environment. The main role of redox-sensing macromolecules is the protection of cell components against the toxicity of reactive oxygen species (ROS), including hydroxyl radicals ($\bullet\text{OH}$) or singlet oxygen ($^1\text{O}_2$), which are produced not only under oxidative-stress conditions but also during normal physiological processes, such as respiration or biosynthesis of a wide range of organic molecules (*i.e.*, sugars, amino acids, or fatty acids) (28, 32). A relevant source for the generation of highly reactive $\bullet\text{OH}$ within cells is the interaction of ferrous iron with hydrogen peroxide (H_2O_2) *via* the so-called Fenton reaction (22). The reaction of cellular cuprous ions with H_2O_2 results in the generation of singlet oxygen ($^1\text{O}_2$) (24). Additionally, $^1\text{O}_2$ can be produced either through

photodynamic effects, mediated by endogenous photosensitizers (*i.e.*, cytochromes, flavin, heme, NAD(P)H, and porphyrins) or as a side product of myeloperoxidase reactions (14). One exogenous source of ROS is the exposure of living organisms to ionizing (γ) and to ultraviolet (UV) radiation. The generation of $\bullet\text{OH}$ has also been suggested to be involved in the action of bactericidal antibiotics during the killing of bacteria and $\bullet\text{OH}$ is also used in the degradation of natural existing biopolymers (cellulose, chitin, or xylan) by microorganisms (5, 36). Moreover, ROS are generated during phagocytosis of a pathogenic bacterium by a host cell (1, 43).

ROS provoke oxidative damage of biomolecules, including nucleic acids, proteins, and lipids. For instance, $\bullet\text{OH}$ efficiently reacts with the four nucleobases, whereas $^1\text{O}_2$ is only able to oxidize guanine bases (15, 16). During the formation of 8-hydroxyguanosine (8-OHG) by $\bullet\text{OH}$, but not $^1\text{O}_2$, a guanine radical cation ($\text{G}^{\bullet+}$, so-called electron hole) is formed with a subsequent release of an electron and proton (14, 16).

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Moreover, the presence of 8-OHG is considered to be a biomarker of oxidative stress (12, 13, 16, 37). It is noteworthy that $G^{\bullet+}$ can migrate long distances (hundred of Å) along DNA by a reversible hopping process before being irreversibly trapped by a reaction with H_2O and O_2 (14, 34). Additionally, guanine is the main target of oxidation in RNA and DNA, especially at multiple guanine sequences, that is, GG or GGG. Among nucleobases it has the lowest ionization potential that causes the migrating $G^{\bullet+}$ to briefly pause at guanine multiplets, facilitating a trapping reaction (26, 34). The formation as well as hopping and trapping of $G^{\bullet+}$ can be considered to be a kind of signal-sensing and transfer mechanism, and hence as a DNA-mediated signaling process. Over recent years researchers have indeed reported the existence of such signaling mechanisms. These results will be presented and discussed in the frame of this review.

Many oxidative base lesions in DNA are mutagenic. These can provoke structural alterations (including transversions: $G \rightarrow T$ or $A \rightarrow C$) or overall conformational changes, which might have an effect on transcription and/or replication; ultimately, this can lead to chromosome deletions with lethal effects (18, 32). To counteract these deleterious effects organisms have evolved enzyme-based repair systems that comprise base excision repair (BER) and nucleotide excision repair (NER). During BER, a single lesion is removed *via* a glycosylase, while during NER a lesion-containing oligonucleotide is removed. Both DNA repair pathways are initiated by damage recognition followed by incision/excision steps, gap filling accompanied by ligation (53, 63). While these enzymatic-based processes may be rapid, they still require a finite period of time between the generation of the mutation and competition of the repair. Thus, the transfer of $G^{\bullet+}$ may provide a supporting mechanism to localize and mitigate the effects of oxidative attack to guanosine nucleotides. Taken together, these processes provide the cell with an efficient and variable DNA damage response mechanism under conditions of oxidative stress.

The main consequence of oxidative modification of messenger RNA (mRNA) is reduced protein production. Studies with human RNAs led to the proposal that RNA is more vulnerable to oxidative damage than DNA and that RNA oxidation is not random but highly selective (55). Further studies have shown that ribosomal RNAs (rRNAs) can be also oxidized *via* a bound redox-active metal ion and that the higher order structure of rRNA may contribute to its greater metal ion-binding ability (31). To our knowledge, RNA oxidation processes in bacteria have not yet been reported. Nonetheless, it is likely that such processes exist, in particular those in which regulatory RNA elements play a role (30, 67). Interestingly, a bacterial RNA element—designated to be a metal-sensing riboswitch—was characterized *in vivo* and *in vitro* and shown to be involved in the regulation of gene expression in a metal-dependent fashion (21).

Proteins with coordinated metal ions are also modified during oxidative stress. Metal ion (*e.g.*, iron)-mediated generation of hydroxyl radicals occurs mostly at metal centers of proteins as $\bullet OH$ can travel only 4 nm within its short half-life ($\sim 10^{-9}$ s), often inducing site-specific damage (38). With a redox potential ranging from -300 to $+700$ mV, iron is a versatile prosthetic component that can be incorporated into proteins either as a mono- or as a binuclear species, or in a more complex form as part of iron-sulfur clusters or heme groups (2). These redox properties provide iron with high

sensitivity, and the ability to sense the redox status of the cell. This in turn provokes an appropriate cellular response. Thus, iron-based production of $\bullet OH$ can also be considered to be beneficial for cells, as these radicals function as redox-signaling molecules that lead to a coordinated response under oxidative-stress conditions.

Proteins exposed to $\bullet OH$ undergo a variety of oxidative modifications, including nitration of aromatic amino acid residues, hydroxylation of aromatic groups and aliphatic amino acid side-chains, sulfoxidation of methionine residues, and conversion of some amino acid residues to carbonyl derivatives (57). For example, oxidation of tyrosines leads to the formation of dityrosine cross-linking products, causing irreversible oxidative damage and often leading to a loss of function due to aggregation and degradation. Indeed, the presence of dityrosines has previously been used as a marker for whole-body oxidative stress (20). Cysteines are additional targets for protein oxidation, as they react easily with H_2O_2 and other free radicals. Reversible oxidation of cysteines are highly complex chemical reactions and can lead to multiple sulfur oxidation states, including thiols, sulfenic and sulfinic acids, thiyl radicals, and disulfide S-oxides (33).

In common with all living organisms, bacteria have evolved nonenzymatic and enzymatic pathways to prevent damage to essential macromolecules during redox-stress events (3). For example, single transcription regulators (*e.g.*, OxyR or FurS) use cysteine thiol groups for signal sensing. Upon signal sensing a conformational change occurs within the regulator, leading to activation or de-repression of transcription of genes involved in antioxidative stress response (17, 49). SoxR is an example of a redox-sensitive transcriptional regulator that utilizes iron-sulfur $[2Fe-2S]$ clusters. Upon oxidative stress, the cluster is oxidized, resulting in the SoxR-dependent transcriptional activation of *soxS*. This encodes for a further transcriptional activator, SoxS, which regulates the expression of genes involved in antioxidant production or oxidative damage repair (51).

The majority of protein-based sensory systems in bacteria are the so-called two-component systems (TCSs) consisting of a sensor kinase (SK) and a response regulator (RR) that controls cellular response by a change in gene expression. One of the best studied SKs involved in redox sensing is FixL from *Rhizobium* sp. FixL possesses a Per-Arnt-Sim domain that contains a heme prosthetic group. Reversible binding of oxygen to the heme group regulates its kinase activity, and thereby the phosphorylation state of its cognate RR FixJ (27). Allosteric changes of FixL upon oxygen binding lead to deactivation of kinase activity (52). Oxygen binding may induce a flattening of the heme group, promoting the formation of a hydrogen bond to the liganded oxygen (8).

In the first part of the presented review, we describe novel studies performed on nucleic acid-mediated processes involved in redox signaling. Under these conditions specific DNA or RNA modules can be considered to be a type of signaling module. Our main aim in this first section is to provide the readers with current knowledge on novel signaling paradigms and avenues with promising research perspectives. In the second part, we focus further on a novel protein-based redox-sensing system, namely, the three component system HbpS-SenS-SenR, in which the accessory protein module HbpS directly switches on/off the corresponding signal cascade.

Nucleic Acids and Redox-Signaling

DNA-mediated electron transfer

Although DNA is not primarily an electron-transfer module, the ordered π -electron system of common DNA bases in duplex B-form DNA provides a pathway for the transfer of excess charges once generated on extended and well-defined stacks of base pairs (7, 23, 58). This concept of charge mobility is an important consideration with respect to environmentally induced DNA damage. Such excess charges can be generated in defined DNA sequences by exposure to free radicals or radiation (either ionizing [7] or nonionizing [UV]) and a number of different studies [reviewed by refs. (25, 26)] demonstrated that long-range charge transfer occurs. Experiments have been performed showing that guanosine (G) undergoes oxidative modifications (to 8-oxoG) upon oxidative stress (12, 56). Studies have also shown that oxidation of guanosines results in a guanine radical cation ($G^{\bullet+}$), which induces an electron migration (hole transport) from a distant guanine through the base pairs of DNA (Fig. 1). Holes (electron vacancies) induced by oxidants are transferred across DNA ~ 50 Å by hopping between CG base pairs and can oxidize remote G-sites, particularly sites comprising sequences of multiple GC base pairs (26). Such a mechanism would imply that upon oxidative attack of an essential DNA element, hole transport would result in oxidative damage to a less essential element. In some bacteria several genes possess G:C-rich sequences outside of the encoding area (19). These G:C-rich domains would act as sinks for the damage, and function as a driving force for hole transport. As a consequence, mutagenic damage would be concentrated in noncoding regions of the genome (Fig. 1).

DNA-mediated redox-signaling

A common feature of protein-based redox-signaling is the modulation of the transcriptional activity of the effector DNA-binding protein *via* a redox sensory protein module. In the last few years, a novel type of signaling, namely, DNA-mediated redox-signaling pathway, has been identified (4, 25). Barton and co-workers elucidated a mechanism in which DNA functions as redox sensor and mediates the oxidation of a regulatory protein, and hence the cellular response to oxidative stress. *In vivo* and *in vitro* analyses showed that ROS can rapidly migrate to regions in the chromosome that are rich in

FIG. 1. Schematic view of charge transport to DNA. Oxidative attack (by $\bullet\text{OH}$) of a guanosine (in the coding DNA region) results in a guanine radical cation ($G^{\bullet+}$), which is subsequently transported (charge transfer) in to DNA regions containing guanine multiplets (GGG). This illustration was adapted from ref. (26).

guanine multiplets (GG or GGG). Such regions are abundant near the SoxR operator within the *soxS* promoter. According to the above model, guanine radicals, arising from guanine multiplet regions, lead to oxidation of the SoxR protein, which subsequently causes the activation of SoxR-mediated transcription of *soxS* (Fig. 2). The mechanism of DNA charge transfer provides this regulator with a long-distance redox-sensing mechanism (up to 27 nm).

RNA-based processes in redox-signaling

The main consequence of oxidative modification of mRNA is reduced protein synthesis, due to the oxidative degradation of mRNA (55). Recently, the discovery of RNA elements within mRNAs, riboswitches, led us to hypothesize a role for RNA in the sensing of redox events. In general, riboswitches directly sense and bind cellular metabolites. This binding induces a conformational change within the RNA element or directly participates in a chemical reaction that modulates riboswitch activity. These events affect transcriptional initiation and termination, as well as translation (30, 67). Interestingly, these metabolites include the redox-active compound riboflavin (from which both flavin adenine dinucleotide and flavin mononucleotide [FMN] are derived). The FMN-mediated riboswitch is exclusively located upstream of genes involved in riboflavin biosynthesis and transport (66). FMN can accept electrons and exists in three different states, all possessing different chemical properties (oxidized, singly reduced, and doubly reduced) (40). Recently, structural analyses of FMN riboswitches (with and without bound FMN) from *Fusobacterium nucleatum* have shown that FMN-free RNA element may adopt a bound-like structure, suggesting that structural rearrangements occurring upon binding are restricted to a few of the loop regions that form the binding pocket of the riboswitch (54, 64). Studies on the FMN riboswitch from *Bacillus subtilis* demonstrated that there are different parameters, that is, kinetic of RNA polymerase, concentration of FMN, and thermodynamic equilibrium that together govern the highly complex mode of action of this riboswitch *in vivo* (65). However, these studies do not provide direct evidence for the redox state of FMN. We believe that under certain oxidative-stress conditions FMN may adopt different redox states, which would result in further conformational changes of the riboswitch, fine-tuning regulatory response.

Interestingly, *in vivo* and *in vitro* studies allowed the identification and characterization of a metal-sensing riboswitch in *B. subtilis* that interacts with Mg^{2+} and is involved in Mg^{2+} homeostasis (21). These studies revealed that Mg^{2+} can induce a compacted tertiary architecture for the so-called M-box RNAs that ultimately facilitate the accessibility of nucleotides involved in genetic control. Dann *et al.* suggest that additional RNA-based metal sensors exist and that these may have a high relevance during metal ion-mediated signaling (21). In this context it is noteworthy to mention that a human rRNA has been shown to be oxidized *via* a bound redox-active metal ion and that the higher order structure of rRNA may contribute to its greater metal ion-binding ability (31). Therefore, it will be challenging to search within bacterial genomes for riboswitches that are specific for redox-active metal ions, that is, iron or copper, and to investigate the genetic response in these organisms upon exposure to

FIG. 2. A model showing DNA-mediated activation of SoxR. Oxidative stress generates guanine radical cations ($G^{\bullet+}$) that are transferred into the DNA-binding sites of SoxR. These radicals oxidize SoxR that in turn activates the transcription of *soxS*. This illustration was adapted from ref. (25).

oxidative stress. We assume in general that redox-active ligands (*i.e.*, ions or metabolites) from riboswitches may sense the redox state of the cell environment and subsequently undergo conformational or electrostatic changes that ultimately result in changes of the riboswitch structure, eliciting the appropriate response in terms of inhibition or activation of gene expression (Fig. 3).

Over the last decades researches were able to *in vitro* synthesize small RNA molecules, called RNA aptamers, *via* the SELEX (systematic evolution of ligands by exponential enrichment) procedure (62). These aptamers show high specificity for macromolecules (*i.e.*, proteins, metabolites, or redox cofactors). For instance, Burgstaller *et al.* obtained an RNA aptamer that displayed high affinity to FMN (11). Binding studies revealed that the aptamer exclusively recognizes the flavin portion of the cofactor with a K_d of ~ 500 nM. Nevertheless, a detailed analysis of redox-dependent modulation of these aptamers has yet to be performed. Interestingly, Lauhon and Szostak synthesized an anti-NMN⁺ (nicotinamide mononucleotide) aptamer that was shown to be able to discriminate between NMN⁺ and NMNH in solution by ~ 15 -fold. Their studies additionally show that there are specific parts in the aptamer that are responsible for the binding activity and stability of the aptamer–ligand interaction (39).

Despite the fact that naturally occurring redox riboswitches and *in vitro* synthesized aptamer–redox cofactor exist (61), an open question remains: are these cofactors used as a prosthetic group for RNA-mediated redox signaling?

RNA-binding redox sensor proteins

Most systems identified to date rely on DNA-binding proteins to control differential gene expression upon redox stress. However, increasing information is available on pro-

teins that interact with RNA to regulate cellular response on a translational level, thereby exerting additional levels of redox regulation. Careful regulation of mRNA half-lives is a fundamental mechanism that allows cells to quickly respond to changing environmental conditions. Examples are the iron responsive proteins (IRP1 and IRP2), which function under low iron conditions as RNA-binding proteins to modulate the translation of mRNAs. These proteins are regulated by iron-mediated oxidative stress in which the reactivity of H_2O_2 with iron directly connects oxidative stress and intracellular iron metabolism (50). In IRP1 there is a further link between the presence of iron and metabolism. Under high iron conditions IRP1 contains an intact 4Fe-4S cluster and functions as aconitase within the citric acid cycle. Under low iron conditions the protein incorporates a 3Fe-4S cluster and functions as an RNA-binding translational regulator. Iron responsive elements (IREs)-like sequences are present at the untranslated regions of open reading frames that are involved in either iron or oxygen metabolism of the cell (6, 29). IRP2 is also redox regulated under low iron conditions. Two cysteine residues of the RNA-binding cleft form a disulfide bridge and close off the terminal-loop binding pocket preventing insertion of the IRE terminal loop (69). However, in contrast to IRP1 it does not contain an iron–sulfur cluster under high iron conditions but is polyubiquitinated before proteasomal degradation (41).

Accessory Proteins-Mediated Redox Signaling

TCSs and accessory proteins

Sensing of environmental signals in bacteria is often coupled to TCSs (48). Recently, the existence of accessory proteins modulating the activity of certain TCSs has been reported (42,

FIG. 3. Schematic view of a ligand-binding riboswitch RNA. Under reducing conditions (red., bottom) the reduced form (hexagon, black) of the ligand interacts with the ligand-binding domain of the riboswitch, leading to a structure that sequesters the ribosomal binding site (RBS). Consequently, the 30S subunit of the bacterial ribosome cannot interact with RBS, blocking the initiation of translation. Under oxidizing conditions (oxid., top) the oxidized form (quadrangle, gray) of the ligand cannot interact with the ligand-binding domain that remains unoccupied in a conformation that favors the accessibility of RBS for the ribosome. This allows the expression of the downstream coding sequence.

48, 59). For instance, the activity of the TCS YycGF from *B. subtilis* seems to be modulated by two proteins, YycH and YycI, which are tethered to the periplasmic side of the membrane via a single N-terminal helix. Theoretical calculations combined with mutagenesis studies indicate that the SK YycG forms a complex with YycH and YycI. It is proposed that subtle alterations in the structure of the transmembrane complex influence YycG kinase activity (59). Because the

predicted interface between YycH and YycI is distinct from their interface with YycG the implication of the proposed model is that the two proteins tether the cellular YycG pool in a large complex (59). However, the exact mechanism of interaction and signaling between YycH and YycI with YycGF remains unclear. Szurmant and co-workers hypothesize that YycH and YycI are involved in sensing the redox state of the electron transport over the membrane and link this to the YycGF system (60).

The HbpS-SenS-SenR system: HbpS as an accessory module

The soil bacterium and cellulose degrader *Streptomyces reticuli* possesses a TCS composed of SenS-SenR (10, 47). SenS is an SK that transfers the phosphate group to its cognate RR, SenR. Dependent upon its phosphorylation state the RR SenR regulates the transcription of the *senS-senR* operon, the gene *hbpS* encoding for a heme-binding protein (HbpS), the *furS-cpeB* operon encoding for the redox regulator (FurS), and the mycelia-associated catalase-peroxidase CpeB (9, 44, 68). Physiological studies have demonstrated that CpeB protects *S. reticuli* against redox-cycling compounds (35), indicating that CpeB plays an important role in minimizing reactions caused by ROS. The production of two additional proteins that have features of redox-active proteins (xylanase-like and tyrosinase-like) has also been shown to be under the control of SenS-SenR (10). Thus, we have proposed SenS-SenR to be a redox-sensing system involved in antioxidative stress response. Indeed, sensitivity and viability assays showed that an *S. reticuli senS-senR* disruption mutant is more sensitive to redox-active compounds, including iron ions, heme, H₂O₂, or plumbagin (47).

We have characterized the *hbpS* gene that is located between the operons *furS-cpeB* and *senS-senR* (Fig. 4A). Immunological studies combined with mutagenesis have demonstrated that HbpS is located extracellularly (46). Interestingly, comparative physiological studies revealed that the presence of HbpS increases the synthesis of the catalase-peroxidase CpeB and provides *S. reticuli* with resistance against redox-cycling compounds. Therefore, we have proposed that HbpS interacts with extracellular, membrane-associated, or membrane-embedded proteins involved in a signal transduction cascade, regulating *cpeB* transcription and resistance to redox stress. As the membrane-embedded SK SenS together with the RR SenR regulates *cpeB* transcription, an interaction between HbpS and SenS has been postulated. Further *in vivo* and *in vitro* analyses showed that in fact HbpS specifically interacts with SenS, and subsequently modulates its autophosphorylation: while HbpS alone inhibits the autophosphorylation of SenS, in the additional presence of iron ions or heme and redox stress considerably enhances SenS autokinase activity (Fig. 4B) (10, 44).

Domain and sequence alignments of HbpS yielded a number of putative homologs encoded within ecologically relevant bacteria [i.e., *Streptomyces coelicolor* A3(2), *Streptomyces kasugaensis*, *Arthrobacter aureus*, *Pseudomonas fluorescens*, *Pseudomonas putida*, and *Sphingomonas aromaticivorans*] and medically relevant bacteria (i.e., *Vibrio cholera*, *Yersinia enterocolitica*, *Leifsonia xyli*, *Acinetobacter baumannii*, *Photobacterium luminescens*, and *Bordetella avium*). The majority of the corresponding *hbpS*-like genes are clustered with genes encoding

FIG. 4. Position of genes and modulation of SenS by HbpS. Relative location and transcriptional orientation of *furS*, *cpeB*, *hbpS*, *senS*, and *senR* on the *Streptomyces reticuli* genome (A). Regulation of the activity of SenS by HbpS (B). HbpS alone represses SenS autophosphorylation (left). In the presence of heme alone (middle) or in combination of H₂O₂ (right), HbpS enhances (~3-fold or ~12-fold, respectively) the level of SenS autophosphorylation.

for SKs and RRs (48). Interestingly, the genes of the accessory proteins YycH and YycI as well as those of the TCS YycGF are also clustered within their corresponding bacterial genomes (60).

HbpS binds and degrades heme

Domain analysis using the Pfam Protein Families Database revealed that HbpS displays homologies with small stretches within the N-terminal regions of heme-binding proteins belonging to the NapC/NirT cytochrome c-family, suggesting that HbpS might interact with heme. The existence of an HbpS-heme complex was colorimetrically demonstrated within the extracellular proteins from *S. reticuli*. In addition, a typical heme spectrum (Soret peak between 409 and 412 nm) was recorded using purified HbpS proteins (44, 46). As HbpS does not possess any known heme-binding signature (*e.g.*,

C-X-X-C-H), it was suggested to be a novel heme-binding protein.

Electron density analysis from the three-dimensional structure of HbpS elucidated from crystals growing in the presence of heme revealed the presence of bound iron ions at the protein surface, potentially as a result of heme degradation. Subsequent spectroscopic and biochemical studies confirmed that HbpS degrades heme through a nonenzymatic and H₂O₂-dependent mechanism known as coupled oxidation (44).

HbpS has an octomeric assembly

Size exclusion chromatography combined with static light scattering analyses allowed the identification of HbpS as an octamer (44). Moreover, HbpS was successfully crystallized either in the presence or in the absence of heme. Both crystal structures (1.6 and 2.3 Å, respectively) shared this octomeric assembly (Fig. 5A). The extensive interactions between N-terminal residues within the HbpS octamer are suggested to control the oligomerization process. A detailed analysis allowed the identification of two N-terminal residues (Ser26 and His28) as essential for octomeric assembly. Ser26 from one subunit interacts *via* hydrogen bonds with His28 from an immediately adjacent subunit, and vice versa (Fig. 5B). The octomeric assembly was additionally demonstrated in solution using site-directed spin labeling combined with pulse electron paramagnetic resonance (SDSL EPR) spectroscopy (35). Further protein-protein interaction studies combined with phosphorylation assays revealed that the octomeric assembly in HbpS is required for efficient interaction with SenS, and the modulation of SenS autophosphorylation (10, 48).

Molecular switches governing the HbpS-mediated cascade

HbpS inhibits SenS autophosphorylation under nonstressing conditions. However, in the presence of heme or iron ions and redox-stress this activity is highly enhanced. Using fluorescence resonance energy transfer (FRET) a decrease of the distance between Trp90 and Ser139 (from 21 to 16.2 Å) under iron-based oxidative-stressing conditions was recorded (45), indicating that HbpS undergoes a conformational change upon exposure to oxidative stress.

FIG. 5. Assembly of HbpS.

(A) The three-dimensional crystal structure of HbpS reveals an octomeric assembly (protein data bank: 3FPV). (B) Amino acids involved in molecular interactions supporting oligomerization are shown. Ser26 (green or red, respectively) from one subunit interacts with the His28 (red or green, respectively) from the neighboring subunit. (To see this illustration in color the reader is referred to the web version of this article at www.liebertonline.com/ars).

FIG. 6. Schematic view of the mode of action of the redox-sensing system HbpS-SenS-SenR. Under iron-mediated oxidative-stressing conditions (*left*) the oxidized and structure-changed form of HbpS (*red*) activates the autophosphorylation of the sensor kinase SenS, leading SenS ~ P [1]. SenS ~ P phosphorylates the response regulator SenR, leading SenR ~ P [2]. This activates the transcription of *hbpS* and *senS-senR*, and de-represses the transcription of *furS-cpeB* [3]. Consequently, FurS, CpeB, HbpS, SenS, and SenR are produced [4] and participate in the antioxidative stress response. While SenS, CpeB, and HbpS are transported into the membrane (SenS) or extracellular space (CpeB and HbpS), FurS and SenR remain cytoplasmic. The oxidized and structure-changed form of HbpS undergoes a time-dependent degradation during oxidative stress [5]. Once CpeB associates with the mycelia, it degrades H₂O₂ providing *S. reticuli* with a nonoxidized environment and blocking the oxidation of HbpS [6] (*right*). Freshly synthesized and unmodified HbpS (*mixed color, turquoise, and green*) interacts with SenS and inhibits its autophosphorylation [7]. SenS in turn dephosphorylates SenR ~ P, leading SenR [8]. This, together with FurS, represses the transcription of *furS-cpeB* and de-activates the transcription of *hbpS* and *senS-senR* [9]. (To see this illustration in color the reader is referred to the web version of this article at www.liebertonline.com/ars).

The FRET data are in line with those obtained by SDSL EPR, which also show an iron-based oxidative stress-dependent distance decrease between Ser139 and Trp90 within the HbpS octamer (35). Moreover, analyses of secondary structures by circular dichroism spectroscopy in the presence of increasing levels of iron-mediated oxidative stress showed a reduction in α -helical structures (from 42% to 20%) combined with an increase of β -sheets (from 15% to 25%) after addition of higher concentrations of iron ions (45).

Biochemical analyses revealed that iron-catalyzed oxidative stress led to the formation of sodium dodecyl sulfate (SDS)-resistant cross-linked forms of HbpS. The formation of intra- or inter-protein cross-linked derivatives can be induced by several different mechanisms: formation of dityrosines, direct interaction of two carbon-centered radicals, or interactions of the carbonyl groups of oxidized proteins with the primary amino groups of lysine residues in the same or a different protein (57). Carbonyl groups have been identified in iron-stressed HbpS proteins using specific antibodies. Fur-

thermore, the formation of dityrosines in HbpS has also been demonstrated by fluorescence spectroscopy. In combination with mutational analyses it was shown that Tyr77-Tyr77 bridges between neighboring subunits within the octamer are formed. Additional *in vivo* analyses showed that upon iron-mediated oxidative stress the levels of *cpeB* as well as of *hbpS* transcripts are highly elevated in an HbpS-SenS-SenR-dependent manner. Thus, oxidative modifications accompanied by structural and conformational changes in HbpS are responsible for the up-regulation of the activity of the HbpS-SenS-SenR system (Fig. 6).

In vitro and *in vivo* studies demonstrated that oxidatively modified HbpS is degraded, either autonomously or enzymatically through action of naturally occurring extracellular *Streptomyces* proteases (35, 45). Recently obtained results showed that the mycelia-associated catalase-peroxidase CpeB protects HbpS *in vivo* and *in vitro* from iron-mediated oxidative modifications. In this manner, a freshly synthesized HbpS remains unmodified and inhibits SenS autophosphorylation,

providing negative feedback for the HbpS-SenS-SenR signaling cascade (9, 35, 45) (Fig. 6).

Concluding Remarks and Perspectives

Over the last years studies have shown that oxidative attack on macromolecules, including proteins as well as nucleic acids, results in specific damage events. A number of systems have been described by which living organisms can sense and react to an increase in oxidative damage, which may be as a result of environmental changes, or through the presence of competing organisms. Some of these systems have evolved to efficiently recognize this oxidative damage, using the modified molecules as mediators in signaling cascades that control the expression of genes involved in antioxidative stress responses. DNA-based protection can be generated by the transport of charges across the DNA, facilitating the transport of a mutagen (guanine radical) from more (coding) to less (noncoding) important DNA regions. Studies on the redox regulator SoxR have also demonstrated that guanine radicals generated *via* guanine multiplets, near the SoxR-binding site, can oxidize SoxR, modulating its activity. Thus, analyses of DNA-binding sites of other redox transcriptional regulators in order to identify neighboring guanine multiplets may provide supporting data for a general mechanism of DNA-based signaling. Interestingly, guanine multiplets (GG and GGG) are to be found near to the DNA-binding site of the iron-dependent redox regulator FurS from *S. reticuli* (49). This transcriptional repressor undergoes a H₂O₂-mediated intramolecular disulfide bond formation that leads to conformational changes, resulting in a loss of FurS-binding to its own DNA operator sequence within the regulatory region of the target operon, *furS-cpeB*. In future, studies will be pursued in order to establish the extent by which DNA-mediated signaling co-regulates the activity of FurS under oxidative stress. It should be noted that the *furS-cpeB* operon is also under the control of the redox-sensing system HbpS-SenS-SenR that mediates the response to extracellular iron-based oxidative stress.

It will be also challenging to analyze whether a comparable RNA-based signaling exists. Such a process might provide a redox-based mechanism to modulate the activity of RNA-binding regulators (*e.g.*, IRPs). In support of this a regulatory function of RNA has been demonstrated for small RNAs and riboswitches can regulate gene expression, at transcriptional or translational levels, in a ligand-dependent manner. As redox metabolites and metal-ion-containing riboswitches have been identified, it will be interesting to explore the ability of such RNA elements to sense redox stress-based signals that would result in changes of the oxidation state of the ligand. Such changes may subsequently be able to induce conformational changes of the riboswitch, resulting in a fine-tuning of regulatory response.

Protein-based sensory and regulatory systems are well known. Most of these systems possess a sensory and regulatory module. An exception is the HbpS-SenS-SenR system in which the extracellular protein HbpS acts as an accessory module, interacting with the membrane-embedded SK SenS. The octomeric HbpS senses extracellular iron-mediated oxidative stress signals, which lead to oxidative modifications accompanied by conformational and structural changes of HbpS, activating SenS. The activity of the SK SenS leads to the expression of antioxidative genes (*i.e.*, those coding for the mycelium-

associated catalase-peroxidase CpeB) *via* the RR SenR. The CpeB enzyme has been shown to protect HbpS from oxidation; and unmodified HbpS inhibits SenS autophosphorylation, resulting in the down-regulation of the HbpS-SenS-SenR signaling cascade (Fig. 6). Given the presence of homologous proteins in the genomes of other organisms, it is highly likely that other accessory-based TCSs sensing systems are present. However, further work is necessary to elucidate the exact nature of the signals that are detected.

Despite the fact that the molecular mechanisms involved in HbpS-based activation/inhibition of SenS are understood, further aspects merit study. As HbpS does not possess any known heme-binding signature, it was postulated to be a novel type of heme-binding protein. Therefore, the elucidation of heme-binding site(s) within the octomeric assembly of HbpS using structural biology will deepen the understanding of the heme-mediated regulation of HbpS. Moreover, it will give further insights into the heme-degradation processes and the ability of HbpS to transport and coordinate the released iron. The coordination of extracellular iron ions or sources of iron is highly relevant, as these processes, together with other HbpS-independent ones, may ensure that under iron-rich conditions these potentially damaging ions are maintained in the extracellular space and are not transported in to the cytoplasm.

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Abbreviations Used

8-OHG = 8-hydroxyguanosine
 BER = base excision repair
 EPR = electron paramagnetic resonance
 FMN = flavin mononucleotide
 FRET = fluorescence resonance energy transfer
 H₂O₂ = hydrogen peroxide
 IRE = iron responsive element
 IRP = iron responsive protein
 mRNA = messenger RNA
 NER = nucleotide excision repair
 RBS = ribosomal binding site
 ROS = reactive oxygen species
 RR = response regulator
 rRNA = ribosomal RNA
 SDSL = site-directed spin labeling
 SK = sensor kinase
 TCS = two-component system
 UV = ultraviolet