

The functional differences between paralogous regulators define the control of the general stress response in *Sphingopyxis granuli* TFA

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Summary

Sphingopyxis granuli TFA is a contaminant degrading alphaproteobacterium that responds to adverse conditions by inducing the general stress response (GSR), an adaptive response that controls the transcription of a variety of genes to overcome adverse conditions. The core GSR regulators (the response regulator PhyR, the anti- σ factor NepR and the σ factor EcfG) are duplicated in TFA, being PhyR1 and PhyR2, NepR1 and NepR2 and EcfG1 and EcfG2. Based on multiple genetic, phenotypical and biochemical evidences including *in vitro* transcription assays, we have assigned distinct functional features to each paralogue and assessed their contribution to the GSR regulation, dictating its timing and the intensity. We show that different stress signals are differentially integrated into the GSR by PhyR1 and PhyR2, therefore producing different levels of GSR activation. We demonstrate *in vitro* that both NepR1 and NepR2 bind EcfG1 and EcfG2, although NepR1 produces a more stable interaction than NepR2. Conversely, NepR2 interacts with phosphorylated PhyR1 and PhyR2 more efficiently than NepR1. We propose an integrative model where NepR2 would play a dual negative role: it would directly inhibit the σ factors upon activation of the GSR and it would modulate the GSR activity indirectly by titrating the PhyR regulators.

Introduction

Microbial survivability in natural habitats is usually threatened by fluctuations in the environmental conditions. In order to adapt to these stressful situations, bacteria react by adjusting their transcriptional profile, triggering either specific or global responses, depending on the extent of the transcriptional remodelling. Frequent mechanisms used to control these responses upon exposure to a stimulus are one- or two-component systems, as well as alternative σ factors (Starón *et al.*, 2009). One relevant example of a bacterial global response that is regulated by alternative σ factors is the general stress response (GSR), which is a protective broad response that generates cross-protection against a number of unrelated stresses (Starón and Mascher, 2010). In *Bacillus subtilis* and related Gram-positive bacteria, the GSR is controlled by σ^B (Pané-Farré *et al.*, 2017), whereas this response is regulated by σ^S in many of the proteobacterial representatives of the Gram-negative species (Hengge, 2010; Battesti *et al.*, 2011). However, Alphaproteobacteria lack a σ^S orthologue (Starón and Mascher, 2010). In this case, the GSR is regulated by a unique mechanism that combines two-component signalling and transcriptional activation by an extracytoplasmic function σ factor (ECF) (Francez-Charlot *et al.*, 2015), which are the most diverse and abundant alternative σ factors (Starón *et al.*, 2009).

In the last decade, the GSR regulatory pathway has been described for a number of alphaproteobacterial representatives (Gourion *et al.*, 2009; Bastiat *et al.*, 2010; Herrou *et al.*, 2012; Jans *et al.*, 2013; Kim *et al.*, 2013; Fiebig *et al.*, 2015; Francez-Charlot *et al.*, 2016; Gottschlich *et al.*, 2018; Ledermann *et al.*, 2018; Lori *et al.*, 2018; Gottschlich *et al.*, 2019). The central regulatory elements (the ECF EcfG, its cognate anti- σ factor NepR and the response regulator PhyR) and the mechanistic principles of the signal transduction (Francez-Charlot *et al.*, 2009; Herrou *et al.*, 2010; Campagne *et al.*, 2012; Campagne *et al.*, 2014) are conserved in most members of this phylogenetic group (Fiebig *et al.*, 2015). In the absence of stress, EcfG is sequestered by NepR, preventing the transcription of the GSR regulon (Campagne *et al.*, 2012; Herrou *et al.*, 2015).

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Besides, PhyR would remain in its inactive conformation. When a stress appears, it would be sensed by one or more GSR-specific HRXXN histidine kinases, which would phosphorylate PhyR turning it into its active form. In this conformation, PhyR exposes a σ -like domain that is able to interact with NepR more efficiently than its cognate EcfG σ factor, promoting a partner switch (Gourion *et al.*, 2008; Francez-Charlot *et al.*, 2009; Campagne *et al.*, 2012; Herrou *et al.*, 2015). This would release EcfG from inhibition, hence activating the transcription of the GSR regulon. Nevertheless, a number of species-specific variations in the signalling circuit may appear (Fiebig *et al.*, 2015). Such diversity includes the presence of paralogues of some of the core regulators (Bastiat *et al.*, 2010; Staroń and Mascher, 2010; Jans *et al.*, 2013; Fiebig *et al.*, 2015; Francez-Charlot *et al.*, 2015; Francez-Charlot *et al.*, 2016), accessory elements involved in the phospho-signalling (Kaczmarczyk *et al.*, 2014; Gottschlich *et al.*, 2018; Lori *et al.*, 2018) or further control at the level of protein stability (Kim *et al.*, 2013). Involvement of paralogous regulators is the most common addition to the canonical regulatory pathway. In most cases, the different paralogues display specific functions in the control of the GSR, although with a certain level of redundancy in some instances. For example, in *Sinorhizobium meliloti*, two PhyR homologues (RsiB1 and RsiB2) regulate the GSR to similar extents (Bastiat *et al.*, 2010) in response to high temperature and stationary phase. On the other hand, in the same species, the NepR-like anti- σ factors RsiA1 and RsiA2 seemed to control different aspects of the regulation, since the deletion of *rsiA2* led to derepression of the response, whereas *rsiA1* mutation resulted in lethality (Bastiat *et al.*, 2010). The most accentuated known example of GSR regulator multiplicity is found in *Methylobacterium extorquens*, in which up to six EcfG paralogues are involved in the control of the response, with EcfG1 and EcfG2 playing a major role in the stress resistance (Francez-Charlot *et al.*, 2016). Furthermore, a main NepR protein seems to play a canonical anti- σ role, inhibiting two EcfG paralogues (EcfG1 and EcfG5 to a certain extent) and being amenable to PhyR sequestration, whereas an additional NepR copy (MexAM1_META2p0735) is unable to interact with any of the EcfG paralogues.

Sphingopyxis granuli TFA is an alphaproteobacterium that has been deeply characterized regarding its ability to use the organic solvent tetralin as carbon and energy source, both at the biochemical and genetic level (reviewed in Floriano *et al.*, 2019). Also, since the annotation of its genome and after confirmation by functional characterization (García-Romero *et al.*, 2016), it has

been defined as the first facultative anaerobe within the *Sphingopyxis* genus due to its capability to respire nitrate anaerobically, and its global regulatory response to this condition has been described (González-Flores *et al.*, 2019; González-Flores *et al.*, 2020). Recently, the GSR regulators encoded in TFA were identified (de Dios *et al.*, 2020). This strain encodes two paralogues of each of the regulators of the central GSR pathway, distributed in two genomic loci: one bearing *nepR1* and *phyR1*, and other genomic location containing *nepR2* and *ecfG1* in a bicistronic operon, *ecfG2* and *phyR2*. The individual roles of EcfG1 and EcfG2 in the regulation have been investigated (de Dios *et al.*, 2020), being EcfG2 the main GSR activator, as it confers stress resistance by itself and is able to control the expression of the whole GSR regulon. On the other hand, EcfG1 seems to play an accessory role, since its expression is EcfG2-dependent and it is only able to fully activate the transcription of part of the GSR target genes.

In this work we have further characterized the GSR regulatory pathway in TFA by combining *in vivo* and *in vitro* approaches. We show a functional differentiation between NepR1 and NepR2 in the control of the response and a different specificity in the stress signalling by PhyR1 and PhyR2. Finally, after reproducing the regulatory system *in vitro*, we propose an integrative model in which the PhyR regulators would produce different levels of activation of the GSR according to the stress that triggers it. We demonstrate that functional differences between paralogous GSR regulators allow an intrinsic feedback regulation in this pathway. Also, for the first time we provide direct evidence of the dual role of the NepR2 anti- σ factor in the regulation: it would directly inhibit the EcfG σ factors and it would negatively modulate the GSR activity indirectly, by titrating the PhyR regulators and releasing NepR1 to further inhibit EcfG1 and EcfG2, thus preventing an overactivation of the response.

Results

NepR1 and NepR2 play specific roles in the regulation of the GSR

Previous analysis of the TFA genome annotation revealed that the elements involved in the core GSR signalling pathway appear duplicated (de Dios *et al.*, 2020). EcfG1 and EcfG2 are the σ factors that drive the transcription of the GSR regulon, with EcfG2 having the leading role in the activation (de Dios *et al.*, 2020). Upstream in the signalling cascade, the NepR1 and NepR2 paralogues would act as anti- σ factors, inhibiting the GSR in the absence of stress. In the genome, *nepR1* is transcribed in a monocistronic operon, presenting up to two

suboptimal GSR target promoters upstream of its coding region (Sup. Fig. S1A). This is coherent with a subtle increase in transcription under GSR-inducing conditions, according to differential RNA-seq (dRNA-seq) data and RT-qPCR (Sup. Fig. S1B). In contrast, *nepR2* is transcribed as the first gene in the *nepR2ecfG1* operon in a GSR-dependent manner, presenting a canonical GSR target promoter upstream (Sup. Fig. S1A) (de Dios *et al.*, 2020). This causes a strong upregulation of *nepR2* transcription under GSR-inducing conditions, as shown by previous dRNA-seq data (de Dios *et al.*, 2020) and RT-qPCR measurements (Sup. Fig. S1B). Due to their inhibitory function, their absence would theoretically lead to a derepression of the response under non-stress conditions.

In order to address their role in the regulation, the construction of the different *nepR* deletion mutants was attempted. In other Alphaproteobacteria (Bastiat *et al.*, 2010; Lourenço *et al.*, 2011), the deletion of a *nepR* homologue that is co-transcribed together with an EcfG coding gene in an autoregulated operon resulted in lethality. This has been argued to be due to an uncontrolled transcriptional activity of the respective EcfG orthologue on its own promoter in the absence of NepR, which may lead to a deleterious overactivation of the GSR. In agreement with this, *nepR1* could be deleted in TFA, contrarily to *nepR2*. Nevertheless, a deletion mutant in the whole *nepR2ecfG1* operon could be constructed. To address the cause of the *nepR2* essentiality, *in trans* complementation experiments were performed. In these assays, the viability of the Δ *nepR2ecfG1* mutant was assessed after transformation with a plasmid bearing *ecfG1* without the promoter region, preceded by its own promoter or by a GSR-insensitive promoter. As shown in Sup. Fig. S2, the plasmid bearing *ecfG1* under its own promoter was the only one unable to be stabilized in the mutant. In contrast, an inducible expression of *ecfG1* (from the IPTG-inducible P_{trc} promoter) producing a functional amount of EcfG1 (de Dios *et al.*, 2020) that is not subjected to positive feedback on its own promoter would not affect viability. This indicates that, in the absence of NepR2, EcfG1 would persistently initiate transcription from its own promoter to eventually produce more of this σ factor, entering a positive feedback loop without any counter-regulation to stop it. Therefore, this highlights the essentiality of NepR2 to control the autoinduction of *ecfG1*.

To distinguish the specific role of each NepR paralogue in the GSR regulation, a *nepR2::lacZ* reporter [which has been previously used to assess the GSR activity in TFA (de Dios *et al.*, 2020)] was integrated into the chromosome of the Δ *nepR1* and the Δ *nepR2ecfG1* mutants, as well as in the Δ *nepR1* Δ *nepR2ecfG1* triple mutant. Next, their β -galactosidase activity was measured

in exponential (GSR repressed) and stationary phase (GSR active) and compared to those of the wild type and the Δ *ecfG1* single mutant (the time points of activity measurement are specified in Sup. Fig. S3 and activity measurements throughout the whole growth curves are shown in Sup. Fig. S4). According to the results shown in Fig. 1, the Δ *ecfG1* mutant showed a slightly lower level of GSR activity in stationary phase (as previously reported in de Dios *et al.*, 2020), whilst the Δ *nepR1* mutant presented a derepressed GSR in exponential phase compared to the wild type, with a slight increase in stationary phase. The Δ *nepR2ecfG1* performed similarly to the wild type and the Δ *ecfG1* mutant in the repression of the GSR under exponential growth. However, in stationary phase, this mutant nearly doubled the activity of the wild type strain. In the case of the Δ *nepR1* Δ *nepR2ecfG1* mutant, in which a constitutively active EcfG2 would be alone to activate the response, a strong derepression was observed in exponential phase, presenting approximately a 40-fold increase in activity compared to the wild type TFA in exponential phase, which continued to increase in stationary phase to even higher levels. The levels of activity reached by the triple mutant indicate that, even under stress conditions, the maximum levels of GSR expression are not reached by the wild type strain, suggesting that a proportion of the anti- σ factors remain active under stationary phase. Altogether, these results suggest that NepR1 and NepR2 have specific roles in GSR regulation, with NepR1 playing a main role in the global repression of the GSR in TFA in the absence of stress, and hence in its initial activation, and with NepR2 modulating the intensity of the response once it is active.

NepR1 and NepR2 show different binding affinities for EcfG1 and EcfG2

Structural studies describing the molecular aspects of the partner switching mechanism that mediate the GSR activation in Alphaproteobacteria revealed that the EcfG inhibition by NepR occurs by a direct protein–protein interaction (Campagne *et al.*, 2012). Since TFA encodes two paralogues of each of these proteins, one possible model would be that each of the EcfG proteins were specifically titrated by one of the NepR anti- σ factors. To explore this option, combinatorial mutants were constructed (namely, Δ *nepR1* Δ *ecfG1*, Δ *nepR1* Δ *ecfG2* double mutants and Δ *nepR1* Δ *ecfG1* Δ *ecfG2* triple mutant) and their ability to resist to heavy metals and osmotic stress were tested. As a result (Sup. Fig. S5A) only those strains lacking *ecfG2* showed an increased sensitivity compared to the wild type. Contrarily, when β -galactosidase activity from the *nepR2::lacZ* fusion was

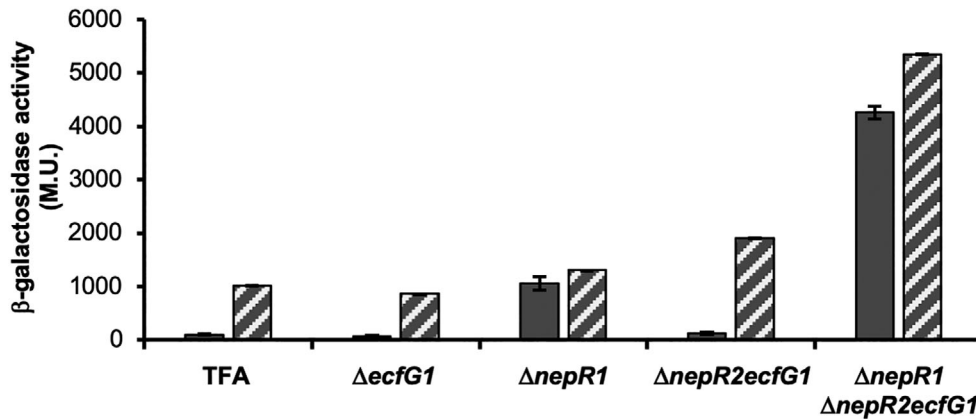


Fig. 1. β-galactosidase activity from the *nepR2::lacZ* translational fusion in different Δ*nepR* mutant backgrounds compared to the wild type and the Δ*ecfG1* single mutant. The activity was measured in exponential (whole bars) and stationary phase (striped bars).

measured in those backgrounds, it reached higher levels in the Δ*nepR1*Δ*ecfG2* mutant compared to those of the Δ*nepR1*Δ*ecfG1* (even beyond those of the wild type TFA) as shown in Sup. Fig. S5B. These results imply that each EcfG paralogue is not specifically titrated by one NepR protein. Rather, they would suggest a more complex interplay at the NepR–EcfG interface, which may be defined by the protein–protein affinities between each of the σ-anti-σ pairs and the relative abundance of these regulators in the cell. In order to characterize the four possible NepR–EcfG interactions (NepR1 with EcfG1 or EcfG2 and NepR2 with EcfG1 or EcfG2) and their effect on the transcriptional output of the response, each of these regulators were purified. After that, they were used in different combinations in an *in vitro* transcription (IVT) setup together with the native core RNA polymerase (RNAP) purified from TFA and using the *P_{nepR2}* promoter as template (de Dios *et al.*, 2020). After fixing a common concentration for each EcfG paralogue below RNAP saturation levels (de Dios *et al.*, 2020), either NepR1 or NepR2 were added to the reactions in increasing molecular proportion with respect to them (Fig. 2A). As a result, NepR1 was able to titrate either EcfG protein nearly in a 1:1 proportion, achieving a complete inhibition of transcription. In contrast, a 10:1 molecular excess of NepR2 with respect to either EcfG1 or EcfG2 could not reach similar levels of inhibition to those of NepR1, indicating a weaker interaction between NepR2 and the EcfG σ factors compared to that of NepR1. To further address this interplay, the NepR–EcfG protein–protein interactions were quantified by surface plasmon resonance. For these experiments either NepR1 or NepR2 were immobilized on CM5 chip and either EcfG1 or EcfG2 were injected as analytes under a continuous flow. Kinetic analysis of the interactions gave dissociation constants (K_D) of 2.0 ± 0.8 nM and 14.2 ± 2.0 nM for NepR1–EcfG1 and NepR1–EcfG2 respectively, and 38.8 ± 3.2 nM and

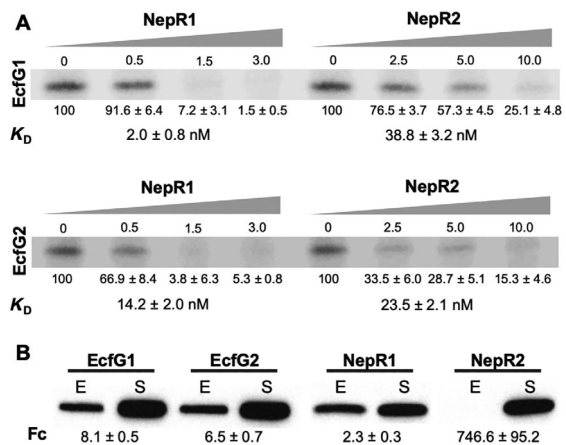


Fig. 2. *In vitro* transcription levels defined by the interaction between the different EcfG–NepR pairs encoded in TFA and protein quantification of the different regulators.

A. IVT results using either EcfG1 or EcfG2 as σ factor and increasing concentrations of either NepR1 or NepR2. Transcription quantifications are referred to those obtained in the absence of anti-σ factor. The dissociation constant (K_D) measured for each EcfG–NepR pair using surface plasmon resonance is indicated underneath each combination.

B. Immunodetection of EcfG1, EcfG2, NepR1 and NepR2 tagged in their C-terminal end with a 3xFLAG epitope. Samples were collected in exponential (E) and stationary phase (S). Protein accumulation fold-change (Fc) is indicated underneath.

23.5 ± 2.1 nM for NepR2–EcfG1 and NepR2–EcfG2 respectively (Fig. 2A). These results agree with those obtained with the IVT system, showing a correlation between lower K_D values and stronger repression of gene transcription. Thus, a stronger interaction between those EcfG–NepR pairs including NepR1 would be responsible for a more efficient repression of transcription compared to those pairs including NepR2.

Apart from the affinity between the different NepR–EcfG pairs, the relative amounts of each of the elements involved in an interaction also determine its output. To

have an impression of the evolution of the *in vivo* protein accumulation of each of the NepR and EcfG regulators, 3xFLAG-tagged versions of each of them were constructed in a wild type background. Their accumulation was assessed by Western blot in exponential phase (in which the GSR would be off due to NepR inhibition) and in stationary phase (in which the GSR is active because of prevention of the NepR–EcfG interaction). As a result, a general increase in the accumulation of the four regulators was observed in stationary phase, with the most drastic change being that of NepR2 (Fig. 2B). These results are coherent with those obtained in the IVT assays, since NepR2 would be needed in bigger amounts than NepR1 in order to perform an efficient inhibition.

The GSR is specifically activated by PhyR1, PhyR2 or both of them depending on the stress

The role of PhyR response regulators consists in derepressing the GSR upon receiving the stress signal in the shape of phosphorylation by sequestering NepR proteins, thus acting as indirect activators of the GSR regulon. In other alphaproteobacterial species, *phyR* mutants behave similarly to *ecfG* mutants regarding their stress resistance, displaying an increased sensitivity compared to the parental wild type strain. This is due to the inability of these strains to prevent EcfG titration by NepR.

In order to address the role of each PhyR paralogue encoded in TFA in the GSR signalling, deletion mutants were constructed in each *phyR* gene, as well as a double mutant. Subsequently, the resulting mutant strains were challenged to resist a variety of stresses compared to the wild type strain and a $\Delta ecfG1\Delta ecfG2$ double mutant, which is totally impaired in the GSR activation. The results revealed that an increased sensitivity to heavy metals (copper) was only observed in those $\Delta phyR$ mutant backgrounds lacking *phyR2* (Fig. 3A). On the other hand, an increased sensitivity to oxidative stress was obtained only in the absence of *phyR1* (Fig. 3B). Regarding the resistance to desiccation, all $\Delta phyR$ mutant strains were affected compared to the wild type, with a milder sensitivity observed for the $\Delta phyR2$ mutant (Fig. 3C). In contrast, only the $\Delta phyR1\Delta phyR2$ double mutant resulted more affected than the wild type under osmotic stress conditions (Fig. 3A). Altogether, this suggests that PhyR1 and PhyR2 are activated specifically depending on the stress that triggers the GSR signalling. A reasonable explanation would be that the specificity came from upstream, being subsequently channelled through PhyR1 or PhyR2 or both of them. This possibility is further explored in the discussion of this work.

PhyR1 and PhyR2 produce different levels of activation of the GSR

The results presented previously conveyed the idea that each of the PhyR regulators encoded in TFA performed distinctive roles in the GSR activation. To evaluate their ability to activate the response, the *nepR2::lacZ* reporter was introduced in each of the $\Delta phyR$ mutant backgrounds and their β -galactosidase activity was measured in exponential and stationary phase compared to that of the wild type (Fig. 4, extended figure showing activities throughout the whole growth curves is shown in Sup. Fig. S6). As expected, the $\Delta phyR1\Delta phyR2$ mutant showed a similar level of activity to that of the $\Delta ecfG1\Delta ecfG2$ mutant. The $\Delta phyR1$ single mutant showed a marked decrease in the activity, mainly observed in stationary phase, whereas the $\Delta phyR2$ mutant produced slightly lower levels of activity than the wild type. These results indicate that PhyR1 is able to produce a stronger activation of the GSR than PhyR2, at least in stationary phase induced by carbon starvation.

After comparing β -galactosidase activity from the *nepR2::lacZ* fusion in the different $\Delta phyR$ mutants to those of the $\Delta ecfG$ mutants (de Dios *et al.*, 2020), similarities in both expression patterns were observed (i.e. the expression phenotype of the $\Delta phyR1$ mutant resembled that of an $\Delta ecfG2$ mutant, and the phenotype of the $\Delta phyR2$ mutant resembled that of the $\Delta ecfG1$). This raised the question of whether there would be a specific signalling from PhyR1 toward EcfG2 and from PhyR2 toward EcfG1. Nevertheless, a $\Delta phyR1\Delta ecfG1$ double mutant, in which the only signalling stream possible would be from PhyR2 to EcfG2, showed a similar expression to that observed in the $\Delta phyR1$ single mutant (Sup. Fig. S7). This suggests that PhyR2, as well as PhyR1, are able to communicate stress to EcfG2 (also probably to EcfG1 in the extent in which this σ factor participates in the activation), opening the possibility of a signal convergence via the NepR anti- σ factors.

PhyR1 and PhyR2 are able to interact more efficiently with NepR2 than with NepR1

As demonstrated for other alphaproteobacterial species, the only stream that the GSR signalling pathway follows is the PhyR–NepR–EcfG cascade, with no accessory regulation occurring between the PhyR and EcfG regulators known so far. Therefore, the only possibility that PhyR1 or PhyR2 may have to activate the transcription would be the direct interaction with either NepR1 or NepR2 in a 1:1 theoretical proportion.

In order to determine the ability of PhyR1 and PhyR2 to activate the GSR, they were purified and added to the previously set up IVT system. All possible PhyR–NepR–EcfG combinations were assayed, using a

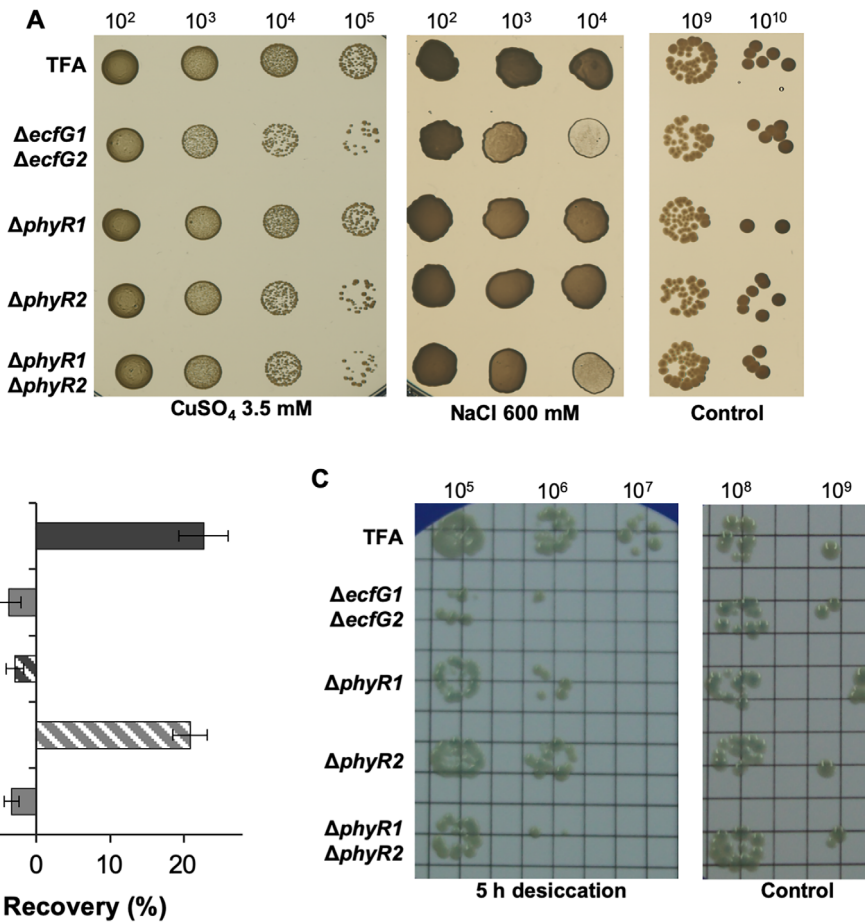


Fig. 3. Stress resistance phenotypes of the $\Delta phyR1$ and $\Delta phyR2$ single mutants and the $\Delta phyR1\Delta phyR2$ double mutant compared to the wild type TFA and the $\Delta ecfG1\Delta ecfG2$ double mutant (stress-sensitive control). The phenotypes tested were (A) resistance to $CuSO_4$ 3.5 mM and NaCl 600 mM, (B) exposure to desiccation during 5 h and (C) recovery of the growth after the addition of H_2O_2 10 mM.

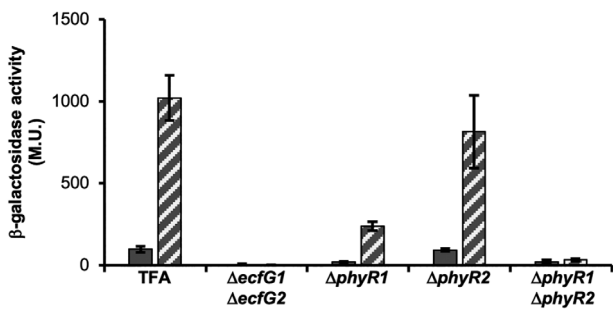


Fig. 4. β -galactosidase activity from the *nepR2::lacZ* translational fusion in different $\Delta phyR$ mutant backgrounds compared to the wild type and the $\Delta ecfG1\Delta ecfG2$ double mutant (negative control). The activity was measured in exponential (whole bars) and stationary phase (striped bars).

molecular NepR-EcfG proportion that would inhibit transcription (see Fig. 2A), such as 1.5:1 for the NepR1-EcfG pairs and 10:1 for the NepR2-EcfG pairs. For the PhyR proteins, two different PhyR-NepR ratios were used

depending on the NepR anti- σ factor present in the reaction: 1:1 and 2:1 with respect to NepR1 and 0.5:1 and 1:1 with respect to NepR2. To simulate an active or inactive status of the GSR, defined by the phosphorylation state of the PhyR proteins, the universal phosphor-donor acetyl phosphate (or a mock treatment) was added to the reactions accordingly. The results (Fig. 5) show that only phosphorylated PhyR1 and PhyR2 were able to stimulate transcription using either EcfG1 or EcfG2. Therefore, when acetyl phosphate was not added, transcription levels remained insensitive to the presence of either PhyR1 or PhyR2. Regarding the anti- σ factor used in each case, whereas both active PhyR1 and PhyR2 could relieve the inhibition exerted by NepR2 to different extents, only PhyR1 was able to activate transcription *in vitro* to detectable levels in the presence of NepR1 in the conditions tested (6.1-fold for PhyR1 versus 1.3-fold for PhyR2 using EcfG1 as σ factor with respect to the absence of PhyR; 1.2-fold for PhyR1 and no transcription

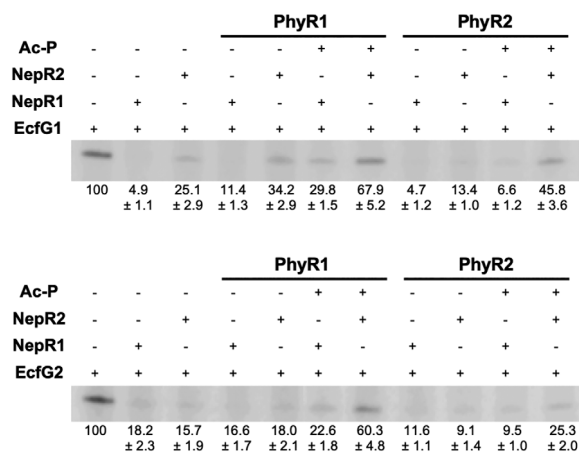


Fig. 5. *In vitro* reconstruction of the GSR using 0.2 μ M of either EcfG1 (upper part) or EcfG2 (lower part) as σ factor in an *in vitro* transcription system. The molecular proportions among all proteins added to each reaction (EcfG:NepR:PhyR) were 1:1.5:1.5 and 1:1.5:3 when using NepR1 as anti- σ factor and 1:10:5 and 1:10:10 when using NepR2. Presence or absence of the different elements is indicated with '+' or '-' signs respectively. Molecular proportion of PhyR proteins with respect to the EcfG proteins and, in brackets, with respect to the NepR protein used in each case are indicated above the lanes. When required, 15 mM acetyl phosphate (AcP) was added to obtain phosphorylated versions of the PhyR proteins. For control reactions using PhyR1 or PhyR2 in the absence of AcP, the highest PhyR concentration was added depending on the NepR protein used. Transcription quantifications are referred to those obtained in the absence of NepR and PhyR proteins.

stimulation by PhyR2 when adding EcfG2). This is coherent with the β -galactosidase activity results obtained using the *nepR2::lacZ* reporter, (814.6 M.U. in the Δ *phyR2* mutant versus 237.8 M.U. in the Δ *phyR1*, as shown in Fig. 4) thus confirming the greater potential of PhyR1 to trigger the GSR compared to PhyR2.

An intriguing observation from this data is the higher transcription levels obtained when using NepR2 in the presence of any of the phosphorylated PhyR proteins than when using NepR1. Furthermore, even when NepR2 was in sufficient amounts to titrate all EcfG and active PhyR proteins present in the reaction (mixtures with proportion 1:10:5, with a more marked effect using EcfG1), transcription was stimulated, thus indicating that NepR2 has a greater potential to interact with the PhyR proteins than with the EcfG σ factors. Contrarily, in the presence of NepR1, PhyR1 was only able to produce a modest activation of the transcription and PhyR2 could barely stimulate transcription only when using EcfG1 as σ factor. This would mean that NepR1 has a stronger affinity for the EcfG σ factors than for the PhyR regulators.

Discussion

Shingopyxis granulii TFA is an alphaproteobacterium that encodes two paralogues of each of the central

regulators of the GSR. Paralogy in the regulatory elements of this pathway is usual among the Alphaproteobacteria. Although the signalling flow is usually straightforward to assess due to the configuration of the regulatory cascade [e.g. convergence from various PhyR and NepR paralogues to one EcfG σ factor (Bastiat *et al.*, 2010) or divergence from one PhyR-NepR stream to a number of EcfG representatives that act in series or in parallel (Lourenço *et al.*, 2011; Francez-Charlot *et al.*, 2016; Gottschlich *et al.*, 2019)], establishing functional differences between *a priori* redundant regulators may be challenging. In the case of TFA, the interplay between EcfG1 and EcfG2 in the activation of the GSR regulon had already been addressed (de Dios *et al.*, 2020), depicting a model in which EcfG2 would be the master activator and EcfG1 would play an accessory role upon activation of the response, most likely as an amplifier of part of the regulon. In this work, we elucidate the signalling flow from the PhyR regulators to the EcfG σ factors via the NepR anti- σ factors based on multiple genetic, phenotypical and biochemical evidences, highlighting the specific functional differences between paralogous elements.

In vitro experiments addressing the interaction between the NepR1 and NepR2 anti- σ factors and the EcfG1 and EcfG2 σ factors clearly indicate that, although both proteins bind EcfG1 and EcfG2, NepR1 interacts more efficiently with EcfG1 and EcfG2 than NepR2. This is a remarkable difference with previously described NepR-EcfG interactions, such as those of *C. crescentus* (Lourenço *et al.*, 2011) and *S. melonis* (Kaczmarczyk *et al.*, 2011). In the first case, the main NepR element does not interact with the secondary EcfG paralogue. In TFA, the IVT assays and interaction quantifications show that NepR1 efficiently binds both EcfG σ factors, ruling out that possibility. In the case of *S. melonis*, a secondary NepR protein (also termed NepR2) is unable to be co-expressed with EcfG1 in a bacterial two-hybrid system, which has suggested an inefficient interaction between them (Gottschlich *et al.*, 2019). In TFA, IVT assays show that NepR2, in amounts sufficiently high (10:1 molecular excess), is able to inhibit around 75% of the transcription driven by either EcfG1 or EcfG2. This hints that the interactions between NepR2 and both EcfG1 and EcfG2 in TFA would occur mainly upon GSR activation, when the *nepR2* transcription has already been induced and the respective protein product is present in sufficiently high cellular concentrations. On the other hand, before GSR activation, inhibition by NepR2 would be less prominent due to its negligible amounts (Fig. 2B), yet essential (Sup. Fig. S2), compared to the inhibition exerted by NepR1. The *in vitro* differences between NepR1 and NepR2 are coherent with the *in vivo* expression measurements obtained with the *nepR2::lacZ* reporter in the

nepR mutant backgrounds, assigning to NepR1 the role of controlling the initial activation of the GSR upon stress exposure. Later on, NepR2 would act once the response is active by modulating its final intensity. Prior to this work, this role as feedback modulator had been only hypothesised for other additional NepR orthologues (Francez-Charlot *et al.*, 2016; Gottschlich *et al.*, 2019) without further experimental development. Based on our results, the possible mechanistic insight of this regulation is discussed below.

When various NepR paralogues are present, they may exhibit functional differences, such as those NepR pairs characterized in *M. extorquens* and *S. melonis*. In these species, the main NepR element binds either the main EcfG σ factor or PhyR, depending on the phosphorylation state of the response regulator. Oppositely, the secondary NepR paralogue (MexAM1_META2p0735 and NepR2 respectively) interacts with PhyR, but it seems unable to form a stable complex with any of the EcfG paralogues encoded in these species (Francez-Charlot *et al.*, 2016; Gottschlich *et al.*, 2019). This regulatory interplay would imply a model in which, once the activation of the GSR is triggered by the PhyR-dependent sequestration of the main NepR, the production of a paralogous NepR would titrate PhyR in a negative feedback loop so that a proportion of the primary NepR is available to inhibit the σ factor activity. The balance in the amounts of NepR bound either to EcfG or to PhyR would determine the levels of GSR activity. The IVT results obtained with the TFA regulators, together with the protein amounts of the two NepR anti- σ factors before and after triggering the response, provide direct evidence for the first time to support an indirect negative feedback regulation. Also, NepR1 binds EcfG1 and EcfG2 more efficiently than NepR2, whereas the latter is able to interact with PhyR1 and PhyR2 (in their phosphorylated state) more efficiently than NepR1. Hence, the GSR would be modulated by a two-level negative feedback loop in TFA, with NepR2 playing a dual role: (i) directly inhibiting the EcfG1 and EcfG2 activity (mainly under GSR-inducing conditions and to a lesser extent in the absence of stress), and (ii) indirectly inhibiting the GSR activity by titrating the active PhyR proteins (and thus releasing NepR1 to inhibit EcfG1 and EcfG2) to prevent the over-activation of the system.

Biochemical studies on the NepR-PhyR interaction (Luebke *et al.*, 2018) revealed that its specificity is determined by the NepR intrinsically disordered N-terminal region, termed FR1, particularly in the residues adjacent to the helix α 1. This region also participates in the PhyR activation by enhancing its phosphorylation (Kaczmarczyk *et al.*, 2014; Herrou *et al.*, 2015; Luebke *et al.*, 2018). Also, the FR1 fragment shows a strong divergence even comparing NepR paralogues encoded

within the same strain, such as the TFA NepR1-NepR2 pair and the *S. melonis* NepR-NepR2 pair (Sup. Fig. S8). In agreement with Luebke *et al.* (2018), this region, especially in the fragment right next to the α 1 helix, was the most divergent between main and additional NepR paralogues, which may suggest different specificities for the respective EcfG and PhyR proteins. These observations might explain the distinct interplay between NepR1 and NepR2 and the rest of the regulators in this pathway, hinting at a modulatory role of NepR2 beyond the usual σ -anti- σ titration.

Ascending further upstream in the GSR cascade, we tackled the characterization of the two PhyR proteins encoded in TFA. In other Alphaproteobacteria with two PhyR paralogues [e.g. RsiB1 and RsiB2 from *S. melliloti* (Bastiat *et al.*, 2010)], both elements appear to exert a similar control on the GSR, since their mutation led to similar phenotypes. However, in TFA both PhyR1 and PhyR2 seem to play different functional roles as judged by the stress resistance assays testing the single and double Δ *phyR* mutants. These experiments indicate a specificity in the signalling depending on the stress that triggers the response. Nevertheless, given the nature of these regulators and their role in the signalling, it seems clear that they do not participate in the specific sensing themselves. Instead, there would be other elements above the PhyR level, such as the four putative HRXXN histidine kinases predicted in the TFA genome (SGRAN_1165, SGRAN_1773, SGRAN_2544 and SGRAN_3483) or any other phosphor-transfer element yet unknown, the ones differentiating among signals and/or transducing them selectively to either PhyR1, PhyR2 or both of them. The role of each PhyR regulator in this pathway was addressed by measuring the activity of the *nepR2::lacZ* reporter under carbon starvation, a condition that seems to trigger the signalling through both PhyR elements (Fig. 4), although to different extents. The differences in GSR activation *in vivo* and the ability of each PhyR protein to stimulate transcription *in vitro* indicate that PhyR1 is able to produce a stronger activation of the GSR compared to PhyR2. This would imply that PhyR1 and PhyR2 have different binding affinities for NepR1 and NepR2, eventually affecting the proportion of active EcfG σ factors, and thus, the intensity of the response. Taken together, the stress specificity showed by PhyR1 and PhyR2 and their different abilities to bind NepR1 and NepR2 would suggest a mechanism to modulate the intensity of the GSR output accordingly to the stress that triggered it.

Interestingly, an observation taken from our results is that EcfG2 increases its protein amounts in stationary phase (Fig. 2), concomitantly with an increase in GSR activity in the absence of anti- σ repression in stationary phase with respect to exponential phase

($\Delta nepR1\Delta nepR2ecfG1$ mutant shown in Fig. 1). This contrasts with previous reports indicating a constitutive expression of EcfG2 when measuring a *lacZ* fusion to the beginning of its coding region including the promoter (de Dios *et al.*, 2020). Taking them together, these data open the possibility for the first time to documenting a post-transcriptional control of a GSR regulator in Alphaproteobacteria.

Taking together all the results obtained throughout this work, a step-wise GSR regulatory model proposed for TFA would be as depicted in Fig. 6. When some kind of stress appears either in the environment or in the cytoplasm, it would be sensed by one or more of the predicted HRXXN histidine kinases, causing an

autophosphorylation in their conserved His residue. The signal would be transduced in a specific manner, either directly or indirectly, to PhyR1 and/or PhyR2, which would receive the phosphoryl group in an Asp residue. The phosphorylation would trigger a conformational change to expose their σ -like domains. This would lead to the sequestration of NepR1 in a different proportion, depending on whether the signalling occurred through PhyR1 and/or PhyR2. NepR1 titration would release EcfG2 and the basal amount of EcfG1 from inhibition, thus activating the GSR regulon. As part of that regulon, the expression of the *nepR2ecfG1* operon would be induced, increasing EcfG1 and, more importantly, NepR2 levels. In a negative feedback loop, NepR2 would inhibit

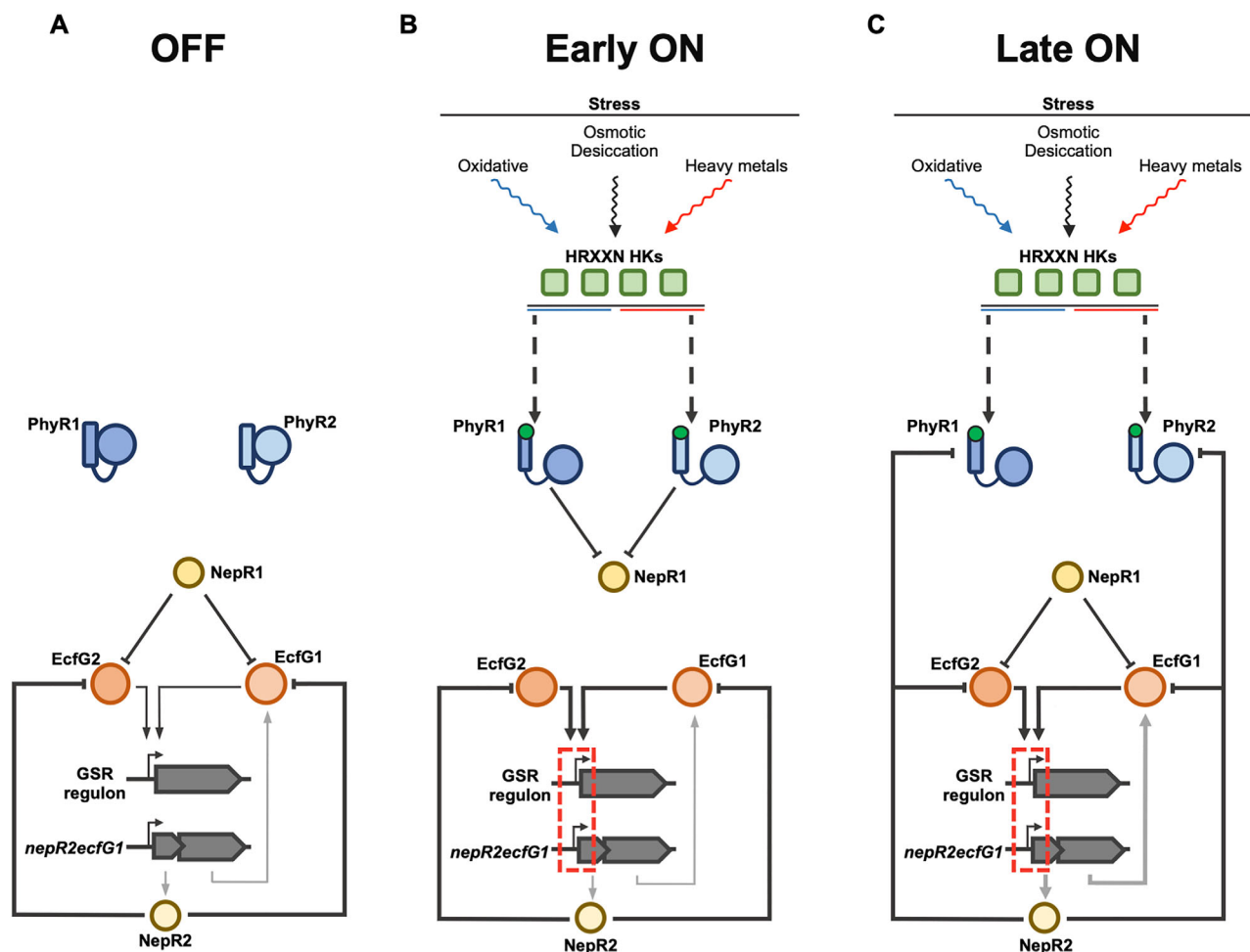


Fig. 6. Step-wise representation of the regulatory model for the GSR signalling pathway in *S. granuli* TFA, indicating the interplay among the regulators and/or their state in the absence of stress (A), at the onset of the stress signalling (B) and once the GSR is fully active (C). Green squares represent the four histidine kinases annotated in the TFA genome, PhyR regulators are represented in dark (PhyR1) and light blue (PhyR2), NepR anti- σ factors are represented in dark (NepR1) and light yellow (NepR2), EcfG σ factors are represented in light (EcfG1) and dark orange (EcfG2), genes are represented in grey. Wavy arrows indicate stress sensing (black lines indicate signalling through PhyR1 and PhyR2, blue lines indicate signalling through PhyR2); dashed arrows indicate phosphorylation of the PhyR regulators (either directly or through intermediate elements); a green circle represents the phosphorylation of PhyR1 and PhyR2; black arrows indicate a regulatory relationship by direct interaction (triangular arrowheads indicate a positive effect, flat arrowheads indicate a negative effect); grey arrows represent transcription and translation.

EcfG1 and EcfG2 in a direct manner by protein–protein interaction. Also, NepR2 would bind PhyR1 and/or PhyR2 with higher affinity than NepR1, titrating them away from the latter. After its release, NepR1 would be again available for directly inhibiting EcfG1 and EcfG2 together with the remaining NepR2. The effect of NepR2 at the EcfG and PhyR levels, together with its high accumulation, would ensure autoregulated levels of GSR by a negative feedback loop to prevent over-activation or to quickly switch GSR off.

Experimental procedures

Media and growth conditions

Escherichia coli and *Sphingopyxis granuli* strains were routinely grown in LB rich medium (Sambrook *et al.*, 1989) at 37°C or MML mineral medium (Andújar *et al.*, 2000) at 30°C respectively. When indicated, *S. granuli* strains were grown in minimal medium (Dorn *et al.*, 1974) supplemented with β -hydroxybutyrate (BHB) as a carbon source in concentrations 8 or 40 mM, depending on the experimental conditions. When appropriate, solid and liquid media were supplemented with kanamycin (25 mg L⁻¹ for *E. coli*, 20 mg L⁻¹ for *S. granuli*), ampicillin (100 mg L⁻¹ for *E. coli*, 5 mg L⁻¹ for *S. granuli*), streptomycin (50 mg L⁻¹ for routine selection, 200 mg L⁻¹ for selection of co-integrates of pMPO1412-derivative plasmids) or X-gal (25 mg L⁻¹).

Plasmids, strains and oligonucleotides

Bacterial strains, plasmids and oligonucleotides used in this work are indicated in Sup. Table S1.

For the generation of mutant strains with scar-less chromosomal modifications (deletions/insertions), the *SceI* double-strand break mediated double recombination procedure was followed as previously described (González-Flores *et al.*, 2019; de Dios *et al.*, 2020). Briefly, a pMPO1412-derivative plasmid containing upstream and downstream 1 kb flanking regions of the fragment to be deleted or the position where the insertion will be placed was transformed in the respective *S. granuli* parental strain and its recombination into the chromosome was selected in the presence of kanamycin. Double-check of recombinant candidates or co-integrates was performed by growing them in the presence of streptomycin (200 mg L⁻¹). Subsequently, plasmid pSWI (Martínez-García and de Lorenzo, 2011), bearing the *SceI* open reading frame, was transformed into the co-integrate strain to force a second recombination event. Candidates bearing the desired modifications were checked by PCR. For the construction of strains with multiple modifications, this procedure was performed serially

with the different pMPO1412-derivative plasmids. Deletion mutants constructed with this strategy were MPO865 (with pMPO1416), MPO866 (with pMPO1414), MPO867 (with pMPO1415), MPO868 (with pMPO1414 and pMPO1415), MPO889 (with pMPO1416 using MPO860 as parental strain), MPO898 (with pMPO1428) and MPO899 (with pMPO1428 using MPO865 as parental strain). Strains bearing 3xFLAG-tagged genes constructed following this protocol were MPO906 (with pMPO1453), MPO907 (with pMPO1454), MPO908 (with pMPO1457) and MPO909 (with pMPO1458).

Strains bearing the *nepR2::lacZ* reporter inserted in the chromosome (MPO871, MPO872, MPO873, MPO874, MPO890, MPO900 and MPO902) were constructed by transforming the respective parental strain with plasmid pMPO1408 and selecting its chromosomal integration by a single recombination event.

For construction of pMPO1412 derivatives, the respective upstream and downstream flanking regions were amplified using *S. granuli* TFA genomic DNA as template and were subsequently assembled together by overlapping PCR. Oligonucleotide pairs used in each case were phyR1 del1 *SacI*-phyR1 del2 and phyR1 del3-phyR1 del4 *Bam*HI for pMPO1414; phyR2 del1 *Bam*HI-phyR2 del2 and phyR2 del3-phyR2 del4 *Eco*RI for pMPO1415; nepR1 del1 *SacI*-nepR1 del2 and nepR1 del3-nepR1 del4 *Bam*HI for pMPO1416; ecfG1 FLAG-1 *Bam*HI-ecfG1 FLAG-2 and ecfG1 FLAG-3-ecfG1 FLAG-4 *SacI* for pMPO1453; ecfG2 FLAG-1 *Bam*HI-ecfG2 FLAG-2 and ecfG2 FLAG-3-ecfG2 FLAG-4 *Eco*RI for pMPO1454; nepR1-FLAG-1-nepR1-FLAG-2 and nepR1-FLAG-3-nepR1-FLAG-4 for pMPO1457; nepR2-FLAG-1-nepR2-FLAG-2 and nepR2-FLAG-3-nepR2-FLAG-4 for pMPO1458. For construction of pMPO1414, pMPO1415, pMPO1416, pMPO1453 and pMPO1454, the assembled PCR fragments were digested with the appropriate restriction enzymes (included in the name of the respective oligonucleotides) and ligated into pMPO1412 digested with the same enzymes. In the case of pMPO1457, pMPO1458, pMPO1459 and pMPO1460, the assembled PCR fragments were directly cloned into pMPO1412 cut with *Sma*I.

The pMPO1412 derivative pMPO1428, used for the deletion of the *nepR2ecfG1* operon, was constructed based on the previously constructed pMPO1407 and pMPO1413 (de Dios *et al.*, 2020). pMPO1407 was digested with *Xho*I, blunted with *Klenow* and subsequently cut with *Acc*65I. The resulting 1 kb fragment was ligated into pMPO1413 digested with *Stu*I and *Acc*65I.

pTXB1 and pTYB21 derivatives for protein overproduction were constructed based on the guidelines provided with the IMPACT kit (New England Biolabs). The coding sequences of *nepR1*, *nepR2*, *phyR1* and *phyR2* were amplified by PCR from *S. granuli* TFA genomic DNA using oligonucleotide pairs ORF-nepR1 fw-ORF-

nepR1 rv BamHI, ORF-nepR2 fw-ORF-nepR2 rv BamHI, ORF-phyR1 fw NdeI-ORF-phyR1 rv and ORF-phyR2 fw NdeI-ORF-phyR2 rv respectively. *nepR1* and *nepR2* fragments were digested with BamHI and ligated into pTYB21 cut with SapI, blunted with Klenow and digested with BamHI, resulting in plasmids pMPO1434 and pMPO1435 respectively. *phyR1* and *phyR2* fragments were digested with NdeI and ligated into pTXB1 cut with SapI, blunted with Klenow and digested with NdeI, resulting in plasmids pMPO1436 and pMPO1437 respectively.

Stress phenotypic assays

Stress resistance assays were performed as in de Dios *et al.* (2020). Briefly, to test the resistance to osmotic stress and copper, 10 µl spots of serial dilutions of late-exponential phase cultures were placed on solid MML rich medium plates supplemented with NaCl 0.6 M or CuSO₄ 3.5 mM and incubated for 5 days at 30°C. For desiccation assays, 5 µl spots of serial dilutions of late-exponential phase cultures were placed on 0.45 µm pore size filters (Sartorius Stedim Biotech GmbH) and they were left to air-dry in a laminar flow cabin for 5 h (5 min in the control assay). Then, filters were placed on MML rich medium plates supplemented with bromophenol blue 0.002% and incubated for 5 days at 30°C. In the case of recovery from oxidative shock, late-exponential phase cultures were diluted to an OD₆₀₀ of 0.1 in MML medium. When an OD₆₀₀ 0.5 was reached, H₂O₂ was added to the medium in a final concentration of 10 mM. Recovery from the treatment is represented by a percentage of the OD₆₀₀ reached by treated cultures after 5 h of growth compared to non-treated cultures. At least three independent replicates of each experiment were performed, and most representative examples are shown.

GSR activation assays and expression measurements

Saturated preinocula were diluted to an OD₆₀₀ of 0.05 in minimal medium supplemented with BHB 40 mM and incubated at 30°C in an orbital shaker for 16 h. Then, 20 ml of minimal medium with BHB 8 mM were inoculated at OD₆₀₀ 0.1. β-galactosidase activity (Miller, 1972) from the *nepR2::lacZ* reporter was measured after 10 and 58 h of growth, representing exponential and stationary phase respectively (Sup. Fig. 3). Averages of three independent replicates are represented.

Protein overexpression and purification

Spingopyxis granuli TFA core RNAP, EcfG1 and EcfG2 were purified as previously published in de Dios *et al.* (2020).

NepR1, NepR2, PhyR1 and PhyR2 proteins were overexpressed and purified using the IMPACT kit (New England Biolabs) following the manufacturer's instructions and equal procedures for the four of them. Briefly, pMPO1434, pMPO1435, pMPO1436 and pMPO1437 (for overexpression of *nepR1*, *nepR2*, *phyR1* and *phyR2* respectively) were transformed into *E. coli* ER2566 host strain. Saturated pre-inocula of each plasmid-bearing strain were diluted to an OD₆₀₀ of 0.1 in different total volumes of LB medium, depending on the gene to be overexpressed (2 L for *nepR1*, 1 L for *nepR2*, 4 L for *phyR1* and 1 L for *phyR2*), and incubated at 37°C in an orbital shaker until reaching OD₆₀₀ 0.7. Then, cultures were chilled on ice and subsequently induced with IPTG 0.5 mM and incubated overnight in a shaker at 16°C. After harvesting the cultures and assessing the induction by SDS-PAGE, cell pellets were resuspended in binding buffer (Tris-HCl 20 mM pH 8, NaCl 0.5 M), lysed by sonication and clarified by centrifugation. Once the chitin resin was packed in a purification column and washed with binding buffer, the respective clarified lysates were loaded on the column and left to flow through the resin by gravity at a low flow rate. Afterwards, the column was flushed with 100 ml of binding buffer prior to the induction of the on-column protein cleavage. To release the target protein, the resin was incubated with TEDG buffer (Tris-HCl 50 mM pH 8, glycerol 10%, Triton X-100 0.01%, EDTA 0.1 mM, NaCl 50 mM) supplemented with DTT 50 mM at 18°C for 40 h approximately. The eluate content in the target protein was assessed by SDS-PAGE. Then, DTT concentration in the buffer was reduced by dialysis against TEDG buffer with DTT 0.1 mM at 4°C overnight using a 3 kDa pore size dialysis cassette (Thermo Fisher Scientific). Finally, purity and concentration of the protein mixtures were evaluated by densitometry compared to different dilutions of BSA using a Typhoon scanner and the ImageLab software. For long-term storage, protein mixtures were aliquoted and frozen at -80°C.

In vitro transcription

Multi-round IVT reactions were performed as in Porrúa *et al.* (2009) with modifications from de Dios *et al.* (2020). Briefly, reactions were run in a final volume of 22.5 µl in IVT buffer (Tris-HCl 10 mM pH 8, NaCl 50 mM, MgCl₂ 5 mM, KCl 100 mM, BSA 0.2 mg ml⁻¹, DTT 2 µM) at 30°C. A mixture containing the appropriate combination of the different GSR regulators, either supplemented or not with acetyl phosphate 15 mM depending on the experiment, was preincubated at 30°C for 5 min. In this mixture, to ensure that any transcriptional activation would be due to disruption of the EcfG-NepR interaction by PhyR, the right amount of each PhyR protein (with or

without acetyl phosphate) was added first in a tube chilled on ice followed by a volume bearing the appropriate EcfG-NepR pair also pre-incubated on ice. 0.2 μM of the respective EcfG σ factor was set as reference to establish molecular proportions with the rest of the regulators present in the reaction. After that, the core RNAP mix was added to the reaction and it was incubated for 5 min. Subsequently, 0.5 μg of plasmid pMPO1440 (pTE103 derivative plasmid bearing the P_{nepR2} promoter from the -35 box to the $+1$, both sites included) (de Dios *et al.*, 2020) were added as circular template. 10 min later, a mix of ATP, GTP, CTP (final concentration of 0.4 mM), UTP (0.07 mM) and [α - ^{32}P]-UTP (0.33 mM, Perkin Elmer) was added to start the reaction. After 10 min, reaction re-initiation was prevented by adding heparin to a final concentration of 0.1 mg ml^{-1} , and 10 min later reactions were arrested by adding 5 μl of stop/loading buffer (0.5% formamide, 20 mM EDTA, 0.05% bromophenol blue, 0.05% xylene cyanol). Samples were boiled for 3 min and run in a 4% polyacrylamide-urea denaturing gel in TBE buffer at room temperature. Gels were dried and exposed in a phosphoscreen and results were visualized in an Amersham Typhoon scanner and analyzed using the ImageQuant software (both provided by GE Healthcare Bio-Sciences AB). Quantifications refer to the median intensity of each band normalized against the levels of transcription obtained by each EcfG protein alone, in the absence of PhyR and NepR. Fig. 2A and Fig. 5 show representative assays of this experiment and quantifications are the average of three independent replicates.

Protein immunodetection (Western blot)

Samples were obtained from cultures in exponential and stationary phase as explained above for gene expression assays. For each sample, 1 OD_{600} unit was harvested by centrifugation and the cell pellet was resuspended in 25 μl bidistilled water. Whole-cell protein content was measured using the RC DC Protein Assay kit (Bio-Rad) and the remaining sample was mixed with loading buffer 2 \times , boiled for 5 min and centrifuged. The equivalent volume to 10 μg of protein was run in a Stain-Free FastCast 12.5% polyacrylamide gel (Bio-Rad) and transferred to a nitrocellulose membrane using the Trans-Blot Turbo semi-dry system (Bio-Rad) following the manufacturer's instructions. The membrane was washed with TTBS buffer and blocked with 5% skimmed milk powder in TTBS buffer (blocking solution). Subsequently, the membrane was incubated overnight with a 1:2000 dilution of mouse monoclonal anti-FLAG antibody (Sigma-Aldrich) in blocking solution at 4°C with mild shaking. Then, the membrane was washed with TTBS, incubated for 2 h with a 1:10 000 dilution of anti-mouse secondary

antibody (Sigma-Aldrich) in blocking solution at room temperature with mild shaking and washed again with TTBS. Finally, the membrane was developed with the Immuno-Star AP Chemiluminescence kit (Bio-Rad) and the signal was detected with a ChemiDoc image system (Bio-Rad) and analyzed with the ImageLab software (Bio-Rad). Representative experiments from three independent replicates are shown in Fig. 2B. Quantifications refer to the average fold-change in stationary phase compared to exponential phase of three independent replicates.

Surface plasmon resonance

EcfG1 and EcfG2 interaction kinetics with respect to immobilized NepR1 or NepR2 were measured using a BIAcore X100 device (GE Healthcare Life Sciences). Assays were performed at 30°C in TEDG buffer. NepR1 (12.1 RU) or NepR2 (36.4 RU) were immobilized on the surface of a CM5 chip using 10 mM acetate buffer pH 4.0 or 5 mM malate buffer pH 5.5 respectively, at 30°C with a contact time of 300 s, following the manufacturer's instructions. Serial two-fold dilutions of EcfG1 and EcfG2 in TEDG buffer were injected into the system at a flow rate of 20 $\mu\text{l min}^{-1}$ in concentrations ranging from 60 to 0.469 nM. Analyte contact time was enough to reach interaction equilibrium and dissociation time was 300 s. After each interaction cycle, the chip was regenerated by injection of 10 mM glycine-HCl buffer pH 2.0. Data were fitted to a 1:1 interaction model using the evaluation software provided by the manufacturer (GE Healthcare Life Sciences). Reliability of the results was assessed according to U -value < 15 and $\chi^2 < 5\%R_{\text{max}}$. Interaction affinity was defined by the dissociation constant (K_D) obtained for each NepR-EcfG pair. At least three independent replicates were assayed for each pair.

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Author Contributions

R.D. performed the experiments, R.D., F.R.-R. and E.S. designed the experimental strategy and analyzed the results. F.R.-R. and E.S. supervised the work. R.D. and F.R.-R. wrote the manuscript considering the revisions of all the authors.

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Supporting Information

Additional Supporting Information may be found in the online version of this article at the publisher's web-site:

Supplementary Table S1. Strains, plasmids and oligonucleotides used in this work and mentioned both in the main text as well as in the Supplemental Material (included in Excel document).

Appendix S1: Supporting Information.