Staphylococcus aureus Uses the Bacilliredoxin (BrxAB)/Bacillithiol Disulfide Reductase (YpdA) Redox Pathway to Defend Against Oxidative Stress Under Infections

Nico Linzner1, Vu Van Loi1, Verena Nadin Fritsch1, Quach Ngoc Tung1, Saskia Stenzel1, Markus Wirtz2, Rüdiger Hell2, Chris J. Hamilton3, Karsten Tedin4, Marcus Fulde4 and Haike Antelmann*1

1 Institute for Biology – Microbiology, Freie Universität Berlin, Berlin, Germany, 2 Plant Molecular Biology, Centre for Organismal Studies Heidelberg, Heidelberg University, Heidelberg, Germany, 3 School of Pharmacy, University of East Anglia, Norwich, United Kingdom, 4 Institute of Microbiology and Epizootics, Centre for Infection Medicine, Freie Universität Berlin, Berlin, Germany

Staphylococcus aureus is a major human pathogen and has to cope with reactive oxygen and chlorine species (ROS, RCS) during infections. The low molecular weight thiol bacillithiol (BSH) is an important defense mechanism of S. aureus for detoxification of ROS and HOCl stress to maintain the reduced state of the cytoplasm. Under HOCl stress, BSH forms mixed disulfides with proteins, termed as S-bacillithiolations, which are reduced by bacilliredoxins (BrxA and BrxB). The NADPH-dependent flavin disulfide reductase YpdA is phylogenetically associated with the BSH synthesis and BrxA/B enzymes and was recently suggested to function as BSSB reductase (Mikheyeva et al., 2019). Here, we investigated the role of the complete bacilliredoxin BrxAB/BSH/YpdA pathway in S. aureus COL under oxidative stress and macrophage infection conditions in vivo and in biochemical assays in vitro. Using HPLC thiol metabolomics, a strongly enhanced BSSB level and a decreased BSH/BSSB ratio were measured in the S. aureus COL ypda deletion mutant under control and NaOCl stress. Monitoring the oxidation degree (OxD) of the Brx-roGFP2 biosensor revealed that YpdA is required for regeneration of the reduced BSH redox potential ($E_{BSH}$) upon recovery from oxidative stress. In addition, the ypda mutant was impaired in $H_2O_2$ detoxification as measured with the novel $H_2O_2$-specific Tpx-roGFP2 biosensor. Phenotype analyses further showed that BrxA and YpdA are required for survival under NaOCl and $H_2O_2$ stress in vitro and inside murine J-774A.1 macrophages in infection assays in vivo. Using NADPH-coupled electron transfer assays provide evidence for the function of YpdA in BSSB reduction, which depends on the conserved Cys14 residue. YpdA acts together with BrxA and BSH in de-bacillithiolation of S-bacillithiolated GapDH. In conclusion, our results point to a major role of the BrxA/BSH/YpdA pathway in BSH redox homeostasis in S. aureus during recovery from oxidative stress and under infections.

Keywords: Staphylococcus aureus, oxidative stress, bacillithiol, bacilliredoxin, bacillithiol disulfide reductase, YpdA, roGFP2
INTRODUCTION

Staphylococcus aureus is an important human pathogen, which can cause many diseases, ranging from local soft-tissue and wound infections to life-threatening systemic and chronic infections, such as endocarditis, septicemia, bacteremia, pneumonia or osteomyelitis (Archer, 1998; Lowy, 1998; Boucher and Corey, 2008). Due to the prevalence of methicillin-resistant S. aureus isolates, which are often resistant to multiple antibiotics, treatment options are limited to combat S. aureus infections (Livermore, 2000). Therefore, the “European Center of Disease Prevention and Control” has classified S. aureus as one of six ESKAPE pathogens which are the leading causes of nosocomial infections worldwide (Pendleton et al., 2013). During infections, activated macrophages and neutrophils produce reactive oxygen and chlorine species (ROS, RCS) in large quantities, including H$_2$O$_2$ and HOCl with the aim to kill invading pathogens (Winterbourn and Kettle, 2013; Hillion and Antelmann, 2015; Beavers and Skaar, 2016; Winterbourn et al., 2016).

Low molecular weight thiols play important roles in the defense against ROS and HOCl in bacterial pathogens and are required for survival, host colonization, and pathogenicity (Loi et al., 2015; Tung et al., 2018). Gram-negative bacteria produce GSH as major LMW thiol, which is absent in most Gram-positive bacteria (Fahy, 2013). Instead, many firmicutes utilize BSH as alternative LMW thiol (Figure 1A), which is essential for virulence of S. aureus in macrophage infection assays (Newton et al., 2012; Pöther et al., 2013; Posada et al., 2014; Chandrangsu et al., 2018). A recent study identified a BSH derivative with an N-methylated cysteine as N-methyl-BSH in anaerobic phototrophic Chlorobiaceae, suggesting that BSH derivatives are more widely distributed and not restricted to Gram-positive firmicutes (Hiras et al., 2018). In B. subtilis and S. aureus, the glycolytic glyceraldehyde-3-phosphate dehydrogenase (GapDH) and the antioxidant enzymes (Chi et al., 2011, 2013; Loi et al., 2015; Imber et al., 2018a,c). In addition, BSH participates in post-translational thiol-modifications under HOCl stress by formation of BSH mixed protein disulfides, termed as protein S-bacillilihthiolations (Chi et al., 2011, 2013; Imber et al., 2018a,c).

Protein S-bacillilihthiolation functions in thiol-protection and redox regulation of redox-sensing regulators, metabolic enzymes and antioxidant enzymes (Chi et al., 2011, 2013; Loi et al., 2015; Imber et al., 2018a,b,c). In S. aureus, the glycolytic glyceraldehyde-3-phosphate dehydrogenase (GapDH) and the aldehyde dehydrogenase AldA were identified as most abundant S-bacillilihthiolated proteins that are inactivated under HOCl stress (Imber et al., 2018a,b). In B. subtilis, the methionine synthase MetE and the OhrR repressor are S-bacillilihthiolated under HOCl stress leading to methionine auxotrophy and derepression of the OhrR-controlled ohrA peroxiredoxin gene, respectively (Fuangthong et al., 2001; Lee et al., 2007; Chi et al., 2011).

Reduction of S-bacillilihthiolated OhrR, MetE, and GapDH proteins is catalyzed by the bacilliredoxins (BrxA/B) in B. subtilis and S. aureus in vitro (Gaballa et al., 2014; Chandrangsu et al., 2018). BrxA (YphP) and BrxB (YqiW) are paralogous thioredoxin-fold proteins of the UPF0403 family with an unusual CGC active site that are conserved in BSH-producing firmicutes (Supplementary Figure S1). Upon de-bacillilihthiolation, the BSH moiety is transferred to the Brx active site, resulting in BrxA-SSB formation (Figure 1B). However, the Brx associated thiol-disulfide reductase involved in regeneration of Brx activity is not known. In GSH-producing bacteria, Grx catalyze the reduction of S-glutathionylated proteins, which requires GSH for regeneration of Grx, resulting in GSSG formation (Lillig et al., 2008; Allen and Miyel, 2012). The regeneration of GSH is catalyzed by the flavoenzyme Gor, which belongs to the pyridine nucleotide disulfide reductases and recycles GSSG on expense of NADPH (Argyrou and Blanchard, 2004; Deponte, 2013).

Phylogenomic profiling of protein interaction networks using EMBL STRING search has suggested the flavoenzyme YpdA (SACOL1520) as putative NADPH-dependent BSSB reductase (Supplementary Figure S1), since YpdA co-occurs together with BrxA/B and the BSH biosynthesis enzymes (BshA/B/C) only in BSH-producing bacteria, such as B. subtilis and S. aureus (Supplementary Figure S2; Gaballa et al., 2010). While our work was in progress, a recent study provides first evidence for the function of YpdA as putative BSSB reductase in S. aureus in vivo since an increased BSSB level and a decreased BSH/BSSB ratio was measured in the ΔypdA mutant under control and H$_2$O$_2$ stress conditions (Mikheyeva et al., 2019). YpdA overproduction was shown to increase the BSH level and contributes to MetE and the OhrR repressor are S-bacillilihthiolated under HOCl stress leading to methionine auxotrophy and derepression of the OhrR-controlled ohrA peroxiredoxin gene, respectively (Fuangthong et al., 2001; Lee et al., 2007; Chi et al., 2011).

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**Abbreviations:** BSH, bacillilihthiol; BSSB, bacillilihthiol disulfide; BrxA/B, bacilliredoxin A (YphP)/bacilliredoxin B (YqiW); CFUs, colony forming units; DTT, dithiothreitol; E$_{BSSB}$, bacillilihthiol redox potential; GapDH, glyceraldehyde 3-phosphate dehydrogenase; GSH, glutathione; GSSG, glutathione disulfide; Gor, glutathione disulfide reductase; Grx, glutaredoxins; HOCl, hypochlorous acid; LMW, low molecular weight; Mtr, mycothiol disulfide reductase; NaOCl, sodium hypochlorite; OD$_{500}$, optical density at 500 nm; rdw, raw dry weight; RCS, reactive chlorine species; ROS, reactive oxygen species; YpdA, bacillilihthiol disulfide reductase.

**FIGURE 1** Structure of the LMW thiol bacillilihthiol (BSH) (A) and mechanism of the bacillilihthiol (Brx)/BSH/YpdA de-bacillilihthiolation pathway (B). (A) Bacillilihthiol is composed of glucosamine, malate, and cysteine. (B) Under HOCl stress, BSH leads to S-bacillilihthiolation of proteins which are reduced by bacilliredoxins (BrxA/B), resulting in the transfer of BSH to the Brx active site (Brx-SSB). BSH functions in Brx-SSB reduction to restore Brx activity, leading to BSSB formation. The BSSB reductase YpdA (SACOL1520) regenerates BSH on expense of NADPH.
oxidative stress resistance, fitness, and virulence of *S. aureus* (Mikheyeva et al., 2019). However, biochemical evidence for the function of YpdA as BSSB reductase and the association of YpdA to the BrxA/B enzymes have not been demonstrated in *B. subtilis* or *S. aureus*.

In this work, we aimed to investigate the role of the complete BrxAB/BSH/YpdA pathway in *S. aureus* in vivo and in vitro. We used phenotype and biochemical analyses, HPLC metabolomics and redox biosensor measurements to study the physiological role of the Brx/BSH/YpdA redox pathway in *S. aureus* under oxidative stress and macrophage infection assays. Our data point to important roles of both BrxA and YpdA in the oxidative stress defense for regeneration of reduced E_BSH and de-bacillithiolation upon recovery from oxidative stress. Biochemical assays further provide evidence for the function of YpdA as BSSB reductase and de-bacillithiolation.

**MATERIALS AND METHODS**

**Bacterial Strains, Growth, and Survival Assays**

Bacterial strains, plasmids and primers used in this study are listed in Supplementary Tables S1, S2, S3. For cloning and genetic manipulation, *Escherichia coli* was cultivated in LB medium. For stress experiments, *S. aureus* COL wild type and mutant strains were cultivated in LB, RPMI, or Belitsky minimal medium and exposed to the different compounds during the exponential growth as described previously (Loi et al., 2017, 2018b). 

**Cloning, Expression, and Purification of His-Tagged Brx-roGFP2, Tpx-roGFP2, GapDH, BrxA, YpdA, and YpdAC14A Proteins in E. coli**

Construction of plasmids pET11b-brx-roGFP2 for expression of the Brx-roGFP2 biosensor was described previously (Loi et al., 2017). The pET11b-derived plasmids for overexpression of the His-tagged GapDH and BrxA (SACOL1321) proteins were generated previously (Imber et al., 2018a). The plasmid pET11b-brx-roGFP2 was used as a template for construction of the Tpx-roGFP2 biosensor to replace brx by the *tpx* gene of *S. aureus*. The *tpx* gene (SACOL1762) was PCR-amplified from chromosomal DNA of *S. aureus* COL using primers pET-tpx-for-Nhel and pET-tpx-rev-Spal (Supplementary Table S3), digested with Nhel and BamHI and cloned into plasmid pET11b-brx-roGFP2 to generate pET11b-tpx-roGFP2. To construct plasmids pET11b-ypdA or pET11b-ypdAC14A, the *ypdA* gene (SACOL1520) was PCR-amplified from chromosomal DNA of *S. aureus* COL with pET-ypdA-for-Ndel or pET-ypdAC14A-for-Ndel as forward primers and pET-ypdA-rev-BamHI as reverse primer (Supplementary Table S3), digested with Ndel and BamHI and inserted into plasmid pET11b (Novagen). For expression of His-tagged proteins (GapDH, BrxA, YpdA, YpdAC14A, Tpx-roGFP2), *E. coli* BL21(DE3) pLysS carrying plasmids pET11b-gap, pET11b-brxA, pET11b-ypdA, pET11b-ypdAC14A and pET11b-tpx-roGFP2 was cultivated in 1 l LB medium until an OD600 of 0.8 followed by addition of 1 mM IPTG (isopropyl-β-D-thiogalactopyranoside) for 16 h at 25°C. His6-tagged GapDH, BrxA, YpdA, YpdAC14A, and Tpx-roGFP2 proteins were purified using His Trap™ HP Ni-NTA columns (5 ml; GE Healthcare, Chalfont St. Giles, United Kingdom) and the ÄKTA purifier liquid chromatography system (GE Healthcare) as described (Loi et al., 2018b).

**Construction of S. aureus COL ΔypdA, ΔbrxA and ΔbrxA ΔypdA Clean Deletion Mutants and Complemented Mutant Strains**

*Staphylococcus aureus* COL ΔypdA (SACOL1520), ΔbrxA (SACOL1464), and ΔbrxB (SACOL1558) single deletion mutants as well as the ΔbrxAΔbrxB double and ΔbrxAΔypdA triple mutants were constructed using pMAD as described (Arnaud et al., 2004; Loi et al., 2018b). Briefly, the 500 bp up- and downstream regions of *ypdA*, *brxA*, and *brxB* were amplified using gene-specific primers (Supplementary Table S3), fused by overlap extension PCR and ligated into the Bgl II and Sall sites of plasmid pMAD. The *pMAD* constructs were electroporated into *S. aureus* RN4220 and further transduced into *S. aureus* COL using phage 81 (Rosenblum and Tyrone, 1964). The clean marker-less deletions of *ypdA*, *brxA*, or *brxB* were selected after plasmid excision as described (Loi et al., 2018b). All mutants were clean deletions of internal gene regions with no genetic changes in the up- and downstream encoding genes. The deletions of the internal gene regions were verified by PCR and DNA sequencing. The ΔbrxA and ΔbrxAΔypdA double and triple mutants were obtained by transduction and excision of pMAD-ΔbrxB into the ΔbrxA mutant, leading to the ΔbrxAΔbrxB deletion of plasmid pMAD-ΔypdA into the ΔbrxAΔbrxB mutant, resulting in the ΔbrxAΔbrxBΔypdA knockout. For construction of *ypdA*, *brxA*, and *brxB* complemented strains, the xylose-inducible ectopic *E. coli*/S. aureus shuttle vector pRB473 was applied (Brückner et al., 1993). Primers pRB-ypdA, pRB-brxA, and pRB-brxB (Supplementary Table S3) were used for amplification of the genes, which were cloned into pRB473 after digestion with *Bam*HI and *Kpn*I to generate plasmids pRB473-ypdA, pRB473-brxA, and pRB473-brxB, respectively. The pRB473 constructs were confirmed by PCR and DNA sequencing and transduced into the ΔypdA and ΔbrxAΔbrxB deletion mutants as described (Loi et al., 2017).

**Construction of Tpx-roGFP2 and Brx-roGFP2 Biosensor Fusions in S. aureus COL**

The *tpx-roGFP2* fusion was amplified from plasmid pET11b-tpx-roGFP2 with primers pRB-tpx-roGFP2-for-BamHI and pRB-tpx-roGFP2-rev-Sacl and digested with *Bam*HI and *Sac*I (Supplementary Table S3). The PCR product was cloned into
fluorescence emission was measured at 510 nm after excitation respectively. The Brx-roGFP2 and Tpx-roGFP2 biosensor treated with 15 mM DTT and 20 mM cumene hydroperoxide, oxidized control samples of Tpx-roGFP2 expression strains were described previously (Loi et al., 2017). The fully reduced and 2
and after injection of the oxidants H₂O₂ and NaOCl, the biosensor oxidation degree (OxD) along the growth curves of 2, transferred to Belitsky minimal medium and treated with 2 mM NaOCl for 60 and 90 min. Cytoplasmic proteins were prepared and subjected to non-reducing BSH-specific Western blot analysis using the polyclonal rabbit anti-BSH antiserum as described previously (Chi et al., 2013). The de-bacillithiolation reactions with purified GapDH-SSB and the BSSB, cysteine and cystine) were extracted from the S. aureus cells, labeled with monobromobimane and measured by HPLC thiol metabolomics as described (Chi et al., 2013).

Northern Blot Experiments

Northern blot analyses were performed using RNA isolated from S. aureus COL before and after exposure to 0.5 mM methylglyoxal, 0.75 mM formaldehyde, 1 mM NaOCl, 10 mM H₂O₂, 2 mM diamide, and 45 µM methylhydroquinone as described (Wetzstein et al., 1992). Hybridizations were conducted using digoxigenin-labeled antisense RNA probes for ypdA, brxA, and brxB that were synthesized in vitro using T7 RNA polymerase and primers ypdA-NB-for/rev, brxA-NB-for/rev, or brxB-NB-for/rev (Supplementary Table S3) as in previous studies (Tam le et al., 2006).

HPLC Thiol Metabolomics for Quantification of LMW Thiols and Disulfides

For preparation of thiol metabolomics samples, S. aureus COL WT, ΔypdA and ΔbrxA/ΔbrxB mutants as well as the ypdA complemented strains were grown in RPMI medium to an OD₅₀₀ of 0.9 and exposed to 2 mM NaOCl stress for 30 min. The intracellular amounts of reduced and oxidized LMW thiols and disulfides (BSH, BSSB, cysteine and cystine) were extracted from the S. aureus cells, labeled with monobromobimane and measured by HPLC thiol metabolomics as described (Chi et al., 2013).

Western Blot Analysis

Staphylococcus aureus strains were grown in LB until an OD₅₀₀ of 2, transferred to Belitsky minimal medium and treated with 100 µM NaOCl for 60 and 90 min. Cytoplasmic proteins were prepared and subjected to non-reducing BSH-specific Western blot analysis using the polyclonal rabbit anti-BSH antiserum as described previously (Chi et al., 2013). The de-bacillithiolation reactions with purified GapDH-SSB and the BSH, BSSB, cysteine and cystine) were extracted from the S. aureus cells, labeled with monobromobimane and measured by HPLC thiol metabolomics as described (Chi et al., 2013). The de-bacillithiolation reactions with purified GapDH-SSB and the BSH, BSSB, cysteine and cystine) were extracted from the S. aureus cells, labeled with monobromobimane and measured by HPLC thiol metabolomics as described (Chi et al., 2013).

Biochemical Assays for NADPH-Dependent BSSB Reduction by YpdA and De-Bacillithioliation of GapDH-SSB Using the BrxA/BSH/YpdA Electron Pathway in vitro

Before the activity assays, the purified BrxA, YpdA, and YpdAC14A proteins were incubated with 40 µM GSSG, 40 µM GSH, or 40 µM coenzyme A disulfide and 500 µM NADPH in 20 mM Tris, 1.25 mM EDTA, pH 8.0. NADPH consumption of YpdA and YpdAC14A was measured immediately after the start of the reaction as absorbance change at 340 nm using the Clariostar microplate reader. The NADPH-dependent BrxA/BSH/YpdA electron pathway was reconstituted in vitro for de-bacillithioliation of GapDH-SSB. About 60 µM of purified GapDH was S-bacillithiolated with 600 µM BSH in the presence of 6 mM H₂O₂ for 5 min. Excess of BSH and H₂O₂ were removed with Micro Biospin 6 columns, which were equilibrated with 20 mM Tris, 1.25 mM EDTA, pH 8.0. Before starting the de-bacillithiolation assay using the BrxA/BSH/YpdA electron pathway, 2.5 µM GapDH-SSB was incubated with 12.5 µM BrxA, 40 µM BSH, and 500 µM NADPH in 20 mM Tris, 1.25 mM EDTA, pH 8.0 at room temperature for 30 min. Next, 12.5 µM YpdA or YpdAC14A proteins were added to the reaction mix at 30°C for 8 min and NADPH consumption was measured at 340 nm. The biochemical activity assays were performed in four replicate experiments.

Infection Assays With Murine Macrophage Cell Line J-774A.1

The murine cell line J774A.1 was cultivated in Iscove’s modified Dulbecco MEM medium (Biochrom) with 10% heat inactivated
fetal bovine serum (FBS) and used for *S. aureus* infection assays as described (Loi et al., 2018b). Macrophages were infected with *S. aureus* cells at a multiplicity of infection (MOI) of 1:25. One hour after infection, the cell culture medium was replaced and 150 μg/ml gentamycin was added for 1 h to kill extracellular bacteria and to stop the uptake of *S. aureus*. The *S. aureus* cells were harvested at 2, 4, and 24 h post infection. To determine the percentage of surviving *S. aureus* cells, infected macrophages were lysed with 0.1% Triton X-100 and the supernatant of internalized bacteria was plated on brain heart infusion (BHI) agar plates. The CFUs were counted after incubation for 24–36 h at 37°C (Loi et al., 2018b).

**Statistical Analyses**

Statistical analysis of growth and survival assays was performed using the Student’s unpaired two-tailed *t*-test by the graph prism software. The statistics of the J-774.1 macrophage infection assays was calculated using the one-way ANOVA and Tukey’s multiple comparisons *post hoc* test by the graph prism software. The results of the statistical tests are included in the figure legends.

**RESULTS**

**Transcription of ypdA, brxA, and brxB Is Induced Under Disulfide Stress by Diamide and NaOCl in S. aureus COL**

The bacilliredoxins BrxA (SACOL1464) and BrxB (SACOL1558) of *S. aureus* share an unusual CGC active site and are highly conserved in BSH-producing firmicutes (Supplementary Figure S1; Gaballa et al., 2014). The pyridine nucleotide disulfide oxidoreductase YpdA (SACOL1520) belongs to the FAD/NAD(P)-binding domain superfamily (IPR036188) and was annotated as putative BSSB reductase due to its phylogenetic co-occurrence with the BSH biosynthesis enzymes and BrxA/B in BSH-producing firmicutes (Supplementary Figure S2; Gaballa et al., 2010). We used Northern blot analysis to investigate whether transcription of *brxA*, *brxB*, and *ypdA* is co-regulated and up-regulated under thiol-specific stress conditions, such as 0.5 mM methylglyoxal, 0.75 mM formaldehyde, 1 mM NaOCl, 10 mM H$_2$O$_2$, 2 mM diamide and 45 μM methlyhydroquinone (Figure 2). The *brxA* gene is co-transcribed with SACOL1465-66-67 in a 2 kb operon and *brxB* is located in the 1.6 kb SACOL1557-brxB-SACOL1559 operon. The genes co-transcribed together with *brxA* and *brxB* encode proteins of unknown functions. The Northern blot results revealed significant basal transcription of the *brxA*, *brxB*, and *ypdA* genes and operons in the control, and strong induction under disulfide stress provoked by NaOCl and diamide. Of note, the *brxB* operon was stronger induced under disulfide stress compared to the *brxA* operon (Figure 2). No up-regulation of the *brxA*, *brxB*, and *ypdA* specific mRNAs was detected upon H$_2$O$_2$, aldehyde and quinone stress. The co-regulation of BrxA/B and YpdA under disulfide stress suggests that they act in the same pathway to regenerate S-bacillithiolated proteins under NaOCl stress upon recovery from oxidative stress.

**The BSSB Level Is Significantly Increased and the BSH/BSSB Ratio Is Decreased in the *S. aureus* ΔypdA Mutant**

To investigate the physiological role of BrxA/B and YpdA under oxidative stress and in BSH redox homeostasis, we constructed ΔbrxAΔ and ΔypdA deletion mutants. Using HPLC thiol metabolomics, the intracellular levels of BSH and BSSB were determined in the ΔbrxAΔ and ΔypdA mutants under control and NaOCl stress after monobromobimane derivatisation of LMW thios and disulfides. In the *S. aureus* COL wild type, a BSH level of 1.6–1.9 μmol/g rdw was determined, which was not significantly different in the ΔypdA and ΔbrxAΔ mutants (Figure 3A). Exposure of *S. aureus* to 2 mM NaOCl stress caused a five to sixfold decreased intracellular BSH level in the wild type, ΔypdA and ΔbrxAΔ mutants (Figure 3A). The level of BSSB in similar control and NaOCl-treated cells of the wild type and ΔbrxAΔ mutant (~0.05 μmol/g rdw) (Figure 3B). Most interestingly, the ΔypdA mutant showed a significantly twofold increased BSSB level under control and NaOCl stress compared to the wild type (Figure 3B), confirming previous data (Mikheyeva et al., 2019). Thus, the BSH/BSSB ratio is ~2–3-fold decreased in the ΔypdA mutant under control and NaOCl relative to the parent (Figure 3C). The increased BSSB levels and the decreased BSH/BSSB redox ratio in the ΔypdA mutant could be restored to wild type levels in the ypdA complemented strain. In addition, a significantly 1.5-fold increased cysteine level was measured in the ΔypdA mutant under NaOCl stress, but no changes in the level of cystine (Supplementary Figures S3A–C). The cystine levels could be also restored to wild type level in the ypdA complemented
strain. These results indicate that YpdA is important to maintain the reduced level of BSH under control and NaOCl stress, supporting previous results (Mikheyeva et al., 2019), while the bacilliredoxins BrxA/B are dispensible for the cellular BSH/BSSB redox balance during the growth and under oxidative stress in S. aureus.

The S. aureus ΔypdA Mutant Is Impaired to Regenerate the Reduced BSH Redox Potential and to Detoxify H₂O₂ Under Oxidative Stress

Next, we applied the Brx-roGFP2 biosensor to monitor the changes of its OxD in S. aureus COL wild type, the ΔypdA and ΔbrxAB mutants during the growth and under oxidative stress (Loi et al., 2017). Using the Nernst equation the OxD values were used to calculate the changes in the BSH redox potential (E_{BSH}) in wild type and mutant strains (see section “Materials and Methods” for details). Measurements of the Brx-roGFP2 OxD in LB medium along the growth did not reveal notable differences in the basal level of E_{BSH} between wild type, ΔypdA and ΔbrxAB mutant strains (Supplementary Figures S4A,B, S5A,B and Supplementary Table S4). The basal level of E_{BSH} varied from −282 to −286 mV in the wild type and from −286 to −299 mV in the ΔypdA and ΔbrxAB mutants in different growth phases (Supplementary Figures S5A,B and Supplementary Table S4). Thus, we monitored the biosensor OxD and calculated the E_{BSH} changes in ΔypdA and ΔbrxAB mutants after exposure to sub-lethal doses of 100 μM NaOCl and 100 mM H₂O₂ to identify functions for BrxA or YpdA under oxidative stress. The Brx-roGFP2 biosensor was strongly oxidized under NaOCl and H₂O₂ stress in the wild type, the ΔypdA and ΔbrxAB mutants (Figures 4A–D). The calculated E_{BSH} increased upon NaOCl stress from −286 to −254 mV in the wild type, from −285 to −247 mV in the ΔypdA mutant and from −288 to −259 mV in the ΔbrxAB mutant (Supplementary Figures S5C,D and Supplementary Table S5). This indicates a stronger increase of E_{BSH} by NaOCl stress in the ΔypdA mutant compared to the wild type. Regeneration of the reduced basal level E_{BSH} occurred already after 2 h reaching values of −269 mV in the wild type and −274 mV in the ΔbrxAB mutant (Figure 4A, Supplementary Figure S5D, and Supplementary Table S5). However, the ΔypdA mutant was significantly impaired to recover the reduced state of E_{BSH} was reproducible with both oxidants, H₂O₂ and NaOCl (Figures 4A,C, Supplementary Figures S5C,E, and Supplementary Table S6). While recovery of reduced E_{BSH} after H₂O₂ stress was fast in the wild type and ΔbrxAB mutant reaching E_{BSH} values of −280 and −283 mV already after 60 min, the ΔypdA mutant was still oxidized after 2 h with high E_{BSH} values of −264 mV (Supplementary Figures S5E,F and Supplementary Table S6). These Brx-roGFP2 measurements document the important role of YpdA to reduce BSSB and to regenerate the reduced E_{BSH} during the recovery phase of cells from oxidative stress.

We further hypothesized that the ΔypdA mutant is defective in H₂O₂ detoxification due to its increased BSSB levels. To analyse the kinetics of H₂O₂ detoxification in the ΔypdA mutant, we constructed a genetically encoded H₂O₂-specific Tpx-roGFP2 biosensor. First, we verified that Tpx-roGFP2 showed the same ratiometric changes of the excitation spectrum in the fully reduced and oxidized state in vitro and in vivo as previously measured for Brx-roGFP2 (Supplementary Figures S6A,B). Tpx-roGFP2 was shown to respond strongly to low levels of 0.5–1 μM H₂O₂ in vitro and was fully oxidized with 100 mM H₂O₂ inside S. aureus COL wild type cells indicating the utility of the probe to measure H₂O₂ detoxification kinetics in S. aureus (Supplementary Figures S6C,D). Measurements of Tpx-roGFP2 oxidation along the growth in LB medium...
revealed a similar high OxD of \(~0.5–0.6\) in the wild type, \(\Delta brxA\) and \(\Delta ypdA\) mutant strains (Supplementary Figures S4C,D). The absence of BrxA/B or YpdA did not affect the biosensor OxD under non-stress conditions, which further provides evidence for roles under oxidative stress. Thus, we monitored the \(H_2O_2\) response of Tpx-roGFP2 and the kinetics of \(H_2O_2\) detoxification in the \(\Delta ypdA\) and \(\Delta brxA\) mutants. Interestingly, Tpx-roGFP2 showed a similar response to 100 mM \(H_2O_2\) in all strains, but the \(\Delta ypdA\) mutant was significantly impaired in \(H_2O_2\) detoxification compared to the wild type (Figures 4E,F). These results clearly confirmed that the \(\Delta ypdA\) mutant is defective to recover from oxidative stress due to its higher BSSB level resulting in an oxidized \(E_{BSH}\) as revealed using Brx-roGFP2 and thiol-metabolomics studies.

**S-Bacillithiolation of GapDH Is Not Affected in \(\Delta ypdA\) and \(\Delta brxA\) Mutants or in ypdA, brxA, and brxB Complemented Strains**

In \(S. aureus\), the glyceraldehyde-3 phosphate dehydrogenase GapDH was previously identified as most abundant S-bacillithiolated protein under NaOCl stress that is visible as major band in BSH-specific non-reducing Western blots (Imber et al., 2018a). Since GapDH activity could be recovered with purified BrxA in vitro previously (Imber et al., 2018a), we analyzed the pattern of GapDH S-bacillithiolation in the \(\Delta brxA\) and \(\Delta ypdA\) mutants as well as in ypdA, brxA and brxB complemented strains in vivo. However, the amount of S-bacillithiolated GapDH was similar after
100 μM NaOCl stress between wild type, ΔbrxAB and ΔypdA mutants and complemented strains (Figures 5A,B). This indicates that the absence of the BrxAB/YpdA pathway does not affect the level of S-bacillithiolation of GapDH under NaOCl stress.

The Bacilliredoxins BrxA/B and the Putative BSSB Reductase YpdA Are Important for Growth and Survival Under Oxidative Stress and Macrophage Infections

Next, we analyzed the physiological role of the BrxA/B/YpdA pathway for growth and survival of S. aureus under H₂O₂ and NaOCl stress. The growth of the ΔypdA and ΔbrxAB mutants in RPMI medium without stress exposure was comparable to the wild type (Figures 6A,C). Interestingly, both ΔbrxAB and ΔypdA mutants displayed a small, but statistically significant growth delay after exposure to sub-lethal amounts of 1.5 mM NaOCl compared to the wild type, while no growth delay was observed with sub-lethal 10 mM H₂O₂ (Figures 6A,C, 7A,B). This might indicate that BrxAB and YpdA function in the same pathway as already suggested by phylogenomic profiling using STRING search (Supplementary Figure S2). Determination of viable counts revealed significantly ~2-fold decreased survival rates of both ΔbrxAB and ΔypdA mutants after exposure to lethal doses of 3.5 mM NaOCl and 40 mM H₂O₂ relative to the wild type (Figures 6E,G, 7C,D). These oxidant sensitive growth and survival phenotypes of the ΔbrxAB and ΔypdA mutants could be restored back to wild type levels by complementation with brxA and ypdA, respectively (Figures 6B,D,F,G, 7C,D). However, complementation of the ΔbrxAB mutant with brxB did not restore the growth and viability of the wild type under NaOCl stress (Figures 6E,G), although xylose-inducible brxB expression of plasmid pRB473-brxB could be verified in Northern blots (Supplementary Figure S7). Moreover, the ΔbrxABΔypdA triple mutant displayed the same sensitivity as the ΔbrxAB mutant to 40 mM H₂O₂ and 3 mM NaOCl indicating that BrxA and YpdA function in the same pathway for reduction of S-bacillithiolated proteins (Figures 7D and Supplementary Figure S8C).

To investigate the function of the BrxA/B/YpdA pathway under infection-relevant conditions, we measured the intracellular survival of the ΔbrxAB and ΔypdA mutants in phagocytosis assays inside murine macrophages of the cell line J-774A.1, as previously (Loi et al., 2018b). The viable counts (CFUs) of internalized S. aureus cells were determined at 2, 4, and 24 h post infection of the macrophages. The number of surviving cells decreased to 21.3% at 24 h post infection for the S. aureus COL wild type, but more strongly to 11.4 and 10.2% for the ΔypdA and ΔbrxAB mutants (Figures 8A,C). Thus, the number of viable counts was significantly ~2-fold lower for both ΔbrxAB and ΔypdA mutants at 24 h post infection compared to the wild type. These sensitive phenotypes of the ΔypdA and ΔbrxAB mutants under macrophage infections could be restored to 80% of wild type levels after complementation with plasmid-encoded ypdA or brxA, respectively (Figures 8B,D). However, complementation with brxB did not restore the survival defect of the ΔbrxAB mutant, pointing again to the major role of BrxA in this pathway.

![FIGURE 5](image-url) Protein S-bacillithiolation of GapDH is not affected in the ΔypdA and ΔbrxAB mutants (A) or in the ypdA, brxA, and brxB complemented strains (B) as revealed by non-reducing BSH Western blots. The prominent GapDH-SSB band is visible in the cell extracts of NaOCl-treated S. aureus cells using non-reducing BSH Western blots. Other bands visible under control and stress conditions are proteins cross-reactive with the polyclonal rabbit anti-BSH antibodies. The amount of GapDH-SSB is similar in the WT, ΔypdA and ΔbrxAB mutants (A) as well as in the ypdA, brxA, and brxB complemented strains (B). The SDS PAGE loading control is shown at the bottom for comparison.
FIGURE 6 | The *S. aureus* ΔypdA and ΔbrxAB mutants are more sensitive under NaOCl stress. (A–E) Growth curves of *S. aureus* COL WT, ΔypdA and ΔbrxAB mutants as well as ypdA, brxA, and brxB complemented strains in RPMI medium after exposure to 1.5 mM NaOCl stress at an OD<sub>500</sub> of 0.5. (F,G) Survival rates were determined as CFUs for *S. aureus* COL WT, ΔypdA and ΔbrxAB mutants as well as ypdA, brxA, and brxB complemented strains at 2, 3, and 4 h after treatment with 3.5 mM NaOCl. Survival of the untreated control was set to 100%. Mean values and SD of 3–4 biological replicates are presented. The statistics was calculated using a Student’s unpaired two-tailed t-test by the graph prism software. Symbols are: ns p > 0.05, * p ≤ 0.05, and ** p ≤ 0.01.

Taken together, our results revealed that the bacilliredoxin BrxA and the putative BSSB reductase YpdA are required for improved survival of *S. aureus* inside macrophages to resist the oxidative burst. Our data suggest that BrxA and YpdA act together in the BrxA/BSH/YpdA pathway to regenerate S-bacillithiolated proteins and to restore the BSH redox potential upon recovery from oxidative stress during infections.

**The Flavin Disulfide Reductase YpdA Functions in BSSB Reduction and De-Bacillithiolation of GapDH-SSB in the BrxA/BSH/YpdA Electron Transfer Assay in vitro**

Next, we aimed to analyze the catalytic activity of purified YpdA in a NADPH-coupled assay with BSSB as substrate in vitro, since biochemical evidence for the function of YpdA as BSSB reductase activity in vitro is still missing (Mikheyeva et al., 2019). The His-tagged YpdA protein was purified as yellow colored enzyme and the UV-visible spectrum revealed the presence of the FAD co-factor indicated by the two absorbance peaks at 375 and 450 nm (Supplementary Figure S9). Incubation of YpdA protein with BSSB resulted in significant and fast consumption of NADPH as measured by a rapid absorbance decrease at 340 nm (Figure 9A). Only little NADPH consumption was measured with YpdA alone in the absence of the BSSB substrate supporting previous finding that YpdA consumes NADPH alone (Mikheyeva et al., 2019). However, in our assays, BSSB significantly enhanced NADPH consumption by YpdA compared to the control reaction without BSSB. No increased NADPH consumption was measured with coenzyme A disulphide (CoAS<sub>2</sub>) or GSSG as substrate indicating the specificity of YpdA for BSSB (Figure 9A). In addition, we investigated the role of the conserved Cys14 of YpdA for the BSSB reductase activity in the NADPH-coupled assay.
NADPH-consumption of YpdAC14A upon BSSB reduction was much slower and similar to the control reaction of YpdA and YpdAC14A without BSSB (Figure 9B). Our in vivo data support that YpdA and BrxA act together in the BrxA/BSH/YpdA de-bacillithiolation pathway. Thus, we analyzed NADPH-consumption by the BrxA/BSH/YpdA electron pathway in de-bacillithiolation of GapDH-SSB in vitro. The de-bacillithiolation assays revealed fast NADPH consumption in the complete BrxA/BSH/YpdA coupled assays (Figure 9C). NADPH consumption by YpdA was slower in the absence of BrxA and might be caused by residual BSSB in the BSH samples. The control reaction of GapDH-SSB with BrxA did not consume NADPH and only little NADPH consumption was measured with BrxA, BSH and the YpdAC14A mutant protein in de-bacillithiolation of GapDH-SSB (Figure 9D).

In addition, BSH-specific non-reducing Western blots were used to investigate if BrxA and the complete BrxA/BSH/YpdA pathway catalyze de-bacillithiolation of GapDH-SSB (Figure 9E). The BSH-blots showed that BrxA is sufficient for de-bacillithiolation of GapDH-SSB, since all reactions of GapDH-SSB with BrxA lead to complete de-bacillithiolation with and without YpdA or YpdAC14A plus NADPH. However, the reactions of GapDH-SSB with YpdA/NADPH alone did not lead to reduction of GapDH-SSB, indicating the main role of BrxA in de-bacillithiolation while YpdA functions in regeneration of BSH in the BrxA/BSH/YpdA/NADPH redox cycle.

In conclusion, our biochemical assays revealed that YpdA functions as BSSB reductase in an NADPH coupled assay. Cys14 of YpdA is important for the BSSB reductase activity in vitro. Thus, YpdA facilitates together with BrxA the reduction of S-bacillithiolated GapDH in the BrxA/BSH/YpdA redox pathway upon recovery from oxidative stress.

DISCUSSION

The putative disulfide reductase YpdA was previously shown to be phylogenetically associated with the BSH biosynthesis enzymes and bacilliredoxins (Supplementary Figure S2), providing evidence for a functional Brx/BSH/YpdA pathway in BSH-producing bacteria (Gaballa et al., 2010). Recent work confirmed the importance of YpdA for the BSH/BSSB redox balance and survival under oxidative stress and neutrophil infections in S. aureus in vivo (Mikheyeva et al., 2019). Here, we have studied the role of the bacilliredoxins BrxA/B and the BSSB reductase YpdA in the defense of S. aureus against oxidative stress in vivo and their biochemical function in the de-bacillithiolation pathway in vitro. Transcription of brxA, brxB and ypdA is strongly upregulated under disulfide stress, provoked by diamide and NaOCl. About two to fourfold increased transcription of ypdA, brxA, and brxB was previously found under H2O2, diamide and NaOCl stress, by the antimicrobial surface coating.
FIGURE 8 | YpdA and BrxA/B promote the intracellular survival of S. aureus inside murine macrophages during infections. (A,B) The survival of S. aureus WT, \( \Delta ypdA \) and \( \Delta brxA/\Delta brxB \) mutants and complemented strains was analyzed 2, 4 and 24 h post infection (p.i.) of the murine macrophage cell line J-774A.1 by CFU counting. The percentages in survival of the \( \Delta ypdA \) and \( \Delta brxA/\Delta brxB \) mutants and complemented strains were calculated after 4 and 24 h in relation to the 2 h time point, which was set to 100%. (C,D) The average percentage in survival was calculated for \( \Delta ypdA \) and \( \Delta brxA/\Delta brxB \) mutants (C) and complemented strain (D) in relation to the WT and WT with empty plasmid pRB473, which were set to 100%. Error bars represent the SEM and the statistics were calculated using one-way ANOVA and Tukey’s multiple comparisons post hoc test using the graph prism software (\( p = 0.0050 \) for WT/\( \Delta ypdA \), \( p = 0.0022 \) for WT/\( \Delta brxA/\Delta brxB \) and \( p = 0.026 \) for WT pRB473/\( \Delta brxA/\Delta brxB \)). Symbols: ns \( p > 0.05 \); * \( p \leq 0.05 \), and ** \( p \leq 0.01 \).

composed of Ag\(^{+}\) and Ru\(^{+}\) (AGXX\(^{\circ}\)) and after exposure to azurophilic granule proteins in S. aureus (Palazzolo-Ballance et al., 2008; Posada et al., 2014; Mader et al., 2016; Loi et al., 2018a,b; Mikheyeva et al., 2019). The elevated transcription of \( brxA, brxB, \) and \( ypdA \) under disulfide stress correlated with the up-regulation of the \( bshA, bshB, \) and \( bshC \) genes for BSH biosynthesis in S. aureus and B. subtilis (Chi et al., 2011; Nicolas et al., 2012; Loi et al., 2018a,b). The \( bshA, bshB, \) and \( bshC \) genes and operons are under control of the disulfide stress-specific Spx regulator in B. subtilis, which controls a large regulon for thiol-redox homeostasis (Gaballa et al., 2013). Thus, genes for BSH biosynthesis and the BrxA/B/YpdA pathway might be also regulated by Spx in S. aureus.

The co-regulation of BrxA/B and YpdA under disulfide stress points to their function in the same pathway in S. aureus. HOCl, diamide and AGXX\(^{\circ}\) were shown to cause a strong disulfide stress response in the transcriptome and protein S-bacillithiolation in the proteome of S. aureus (Imber et al., 2018a; Loi et al., 2018a,b). Thus, the BrxA/B and YpdA redox enzymes are up-regulated under conditions of protein S-bacillithiolations, connecting their functions to the de-bacillithiolation pathway. We could show here that NaOCl stress leads to five to sixfold depletion of the cellular pool of reduced BSH in the S. aureus COL wild type, which was not accompanied by an enhanced BSSB level. In the previous study, 20 mM H\(_2\)O\(_2\) resulted in twofold reduction of BSH and threefold increase of BSSB in the S. aureus wild type (Mikheyeva et al., 2019). Most probably, the increased BSSB level under NaOCl stress was used for protein S-bacillithiolation in our study (Imber et al., 2018a), while sub-lethal 20 mM H\(_2\)O\(_2\) might not lead to an increase in S-bacillithiolation in the previous study (Mikheyeva et al., 2019).

The BSH/BSSB redox ratio of S. aureus wild type cells was determined as \( \sim 35:1 \) under control conditions and decreased threefold to 10:1 under NaOCl. Of note, this basal BSH/BSSB ratio in S. aureus COL was higher compared to the basal BSH/BSSB ratio of \( \sim 17:1 \) as determined previously in the bshC repaired SH1000 strain (Mikheyeva et al., 2019). In E. coli, the GSH/GSSG redox ratio was determined in the range between 30:1 and 100:1 (Hwang et al., 1995; Van Laer et al., 2013), which is similar as measured for the basal BSH/BSSB ratio in S. aureus COL. The differences in the BSH/BSSB ratios might be related to different S. aureus strain
support the important role of YpdA as BSSB reductase in mutant cells during the recovery phase. These results clearly indicate the function of the conserved Cys14 as active site of YpdA for BSSB reduction. (C) NADPH consumption of YpdA was measured in the coupled de-bacillithiolation assay for reduction of GapDH-SSB. While fast NADPH consumption was measured upon de-bacillithiolation of GapDH-SSB with purified YpdA (C), the reaction was much slower with the YpdAC14A mutant (D). The coupled assays were conducted with 2.5 µM Gap-SSB, 12.5 µM BrxA, 40 µM BSH, 500 µM NADPH in 20 mM Tris, 1.25 mM EDTA, pH 8.0. After 30-min incubation, 12.5 µM YpdA or YpdAC14A proteins were added to the reaction mix and NADPH consumption was monitored at 340 nm as a function of time. Mean values and SEM of four independent experiments are shown. (E) The de-bacillithiolation of GapDH-SSB is catalyzed by BrxA or the complete BrxA/BSH/YpdA pathway, but not by YpdA alone as shown by non-reducing BSH-specific Western blots. The loading controls are shown below as Coomassie-stained SDS-PAGE with the same de-bacillithiolation reactions of GapDH-SSB as in the BSH-blot.

FIGURE 9 | YpdA functions as BSSB reductase in the NADPH-coupled BrxA/BSH/YpdA electron pathway for de-bacillithiolation of GapDH-SSB in vitro. (A) Purified YpdA is able to reduce BSSB back to BSH with electrons from NADPH as measured by the absorbance change at 340 nm. Only little NADPH consumption was measured with YpdA alone in the absence of BSSB, with coenzyme A disulfide (CoA-S2) or glutathione disulfide (GSSG) (A) and in the YpdAC14A mutant (B) indicating the function of the conserved Cys14 as active site of YpdA for BSSB reduction. (C) NADPH consumption of YpdA was measured in the coupled de-bacillithiolation assay for reduction of GapDH-SSB. While fast NADPH consumption was measured upon de-bacillithiolation of GapDH-SSB with purified YpdA (C), the reaction was much slower with the YpdAC14A mutant (D). The coupled assays were conducted with 2.5 µM Gap-SSB, 12.5 µM BrxA, 40 µM BSH, 500 µM NADPH in 20 mM Tris, 1.25 mM EDTA, pH 8.0. After 30-min incubation, 12.5 µM YpdA or YpdAC14A proteins were added to the reaction mix and NADPH consumption was monitored at 340 nm as a function of time. Mean values and SEM of four independent experiments are shown. (E) The de-bacillithiolation of GapDH-SSB is catalyzed by BrxA or the complete BrxA/BSH/YpdA pathway, but not by YpdA alone as shown by non-reducing BSH-specific Western blots. The loading controls are shown below as Coomassie-stained SDS-PAGE with the same de-bacillithiolation reactions of GapDH-SSB as in the BSH-blot.

backgrounds or growth conditions. Nevertheless, NaOCl and H₂O₂ decreased the BSH/BSSB ratio in our and the previous study (Mikheyeva et al., 2019). In the S. aureus ΔbrxAB mutant, we also measured a threefold decrease of the BSH/BSSB ratio from control conditions (38:1) to NaOCl (12:1). However, the ΔypdA mutant showed a twofold enhanced BSSB level in control and NaOCl-treated cells, leading to a significantly decreased BSH/BSSB ratio under control (17:1) and NaOCl stress (5:1). These results support previous results of the bshC mutant, showing a decreased BSH/BSSB ratio under control (6:1) to NaOCl (2:1) (Mikheyeva et al., 2019), although both ratios were again much lower as in our study. Taken together, our data indicate that BrxA are dispensable for the BSH redox homeostasis, while YpdA is essential for BSSB reduction to maintain the reduced pool of BSH and a high BSH/BSSB ratio in S. aureus.

Brx-roGFP2 biosensor measurements provide further support that YpdA is the candidate BSSB reductase. The ΔypdA mutant was significantly impaired to restore reduced E₅₅₀ required for detoxification of ROS.

These in vivo data were further corroborated by biochemical activity assays of YpdA for BSSB reduction in a NADPH-coupled assay. While little NADPH consumption was measured in the presence of YpdA alone, BSSB significantly enhanced NADPH consumption, supporting the crucial role of YpdA as BSSB reductase in vitro. Further electron transfer assays revealed that YpdA functions together with BrxA and BSH in reduction of GapDH-SSB in vitro. Previous de-bacillithiolation assays have revealed regeneration of GapDH activity by BrxA in vitro (Imber et al., 2018a). Here, we confirmed that BrxA activity is sufficient for complete de-bacillithiolation of GapDH-SSB in vitro, while YpdA alone had no effect on the GapDH-SSB reduction. Thus, BrxA catalyzes reduction of S-bacillithiolated proteins and YpdA is involved in BSH regeneration in the complete BrxA/BSH/YpdA redox cycle.

The BSSB reductase activity of YpdA was shown to be dependent on the conserved Cys14, which is located in the glycine-rich Rossmann-fold NAD(P)H binding domain (GGGPCCG) (Bragg et al., 1997; Mikheyeva et al., 2019). Cys14 might be S-bacillithiolated by BSSB and reduced by electron transfer from NADPH via the FAD co-factor. Cys14 was previously identified as oxidized under NaOCl stress in the S. aureus redox proteome using the OxICAT method, further supporting its role as active site Cys and its S-bacillithiolation.
during the BrxA/BSH/YpdA catalytic cycle (Imber et al., 2018a). The catalytic mechanism of BSSB reduction via Cys14 of YpdA is an interesting subject of future studies.

Previous phenotype results of the ΔypdA mutant revealed that YpdA is important for survival of S. aureus in infection assays with human neutrophils (Mikheyeva et al., 2019). Our phenotype analyses further showed protective functions of the complete BrxA/BSH/YpdA redox pathway for growth and survival of S. aureus under oxidative stress in vitro and in macrophage infections in vivo. The ΔypdA and ΔbrxA/ΔYpdA mutants were significantly impaired in growth and survival after exposure to sub-lethal and lethal doses of NaOCl and displayed survival defects under lethal H₂O₂. Moreover, the H₂O₂ and NaOCl-sensitivity and the defect to recover reduced EBSH in the ΔbrxAΔbrxBΔypdA triple mutant was comparable with that of the ΔypdA mutant (Figure 7D and Supplementary Figure S8). These results clearly indicate that BrxA/B and YpdA function in the same de-bacillithiolation pathway, which is an important defense mechanism of S. aureus against oxidative stress.

Based on previous bacilliredoxin activity assays in vitro, both BrxA and BrxB should use a monothiol mechanism to reduce S-bacillithiolated client proteins, such as OhrR, GapDH and MetE in B. subtilis and S. aureus (Gaballa et al., 2014; Imber et al., 2018a). Most di-thiol Grx of E. coli (Grx1, Grx2, and Grx3) use the monothiol mechanism for de-glutathionylation of proteins (Lillig et al., 2008; Allen and Mieyal, 2012; Loi et al., 2015). In the monothiol mechanism, the nucleophilic thiolate of the Brx CGC motif attacks the S-bacillithiolated protein, resulting in reduction of the protein substrate and Brx-SSB formation. Brx-SSB is then recycled by BSH, leading to increased BSSB formation. YpdA reduces BSSB back to BSH with electrons from NADPH (Figure 1B). The oxidation-sensitive phenotypes of ΔypdA and ΔbrxA/ΔYpdA mutants could be complemented by plasmid-encoded ypdA and brxA, but not brxB, respectively. These results provide evidence for the function of the BrxA/BSH/YpdA de-bacillithiolation pathway using the monothiol-Brx mechanism in S. aureus.

Similar phenotypes were found for mutants lacking related redox enzymes of the GSH and mycothiol pathways in other bacteria. In E. coli, strains lacking the Gor and Grx are more sensitive under diamide and cumene hydroperoxide stress (Alonso-Moraga et al., 1987; Vlamis-Gardikas et al., 2002; Lillig et al., 2008). In Mycobacterium smegmatis, the mycoredoxin-1 mutant displayed an oxidative stress-sensitive phenotype (Van Laer et al., 2012). In Corynebacterium glutamicum, deficiency of the Mtr resulted in an oxidized mycothiol redox potential (Tung et al., 2019), and Mtr overexpression contributed to improved oxidative stress resistance (Si et al., 2016). Taken together, our results revealed that not only BSH, but also BrxA and YpdA are required for virulence and promote survival in infection assays inside murine macrophages.

In several human pathogens, such as Streptococcus pneumoniae, Listeria monocytogenes, Salmonella Typhimurium, and Pseudomonas aeruginosa, LMW thiols or the Gor are required for virulence, colonization and to resist host-derived oxidative or nitrosative stress (Potter et al., 2012; Song et al., 2013; Reniere et al., 2015; Tung et al., 2018; Wongsonroj et al., 2018). S. aureus BSH deficient mutants showed decreased survival in murine macrophages and in human whole blood infections (Pöther et al., 2013; Posada et al., 2014). The virulence mechanisms might be related to a lack of BSH regeneration and decreased recovery of inactivated S-bacillithiolated proteins inside macrophages. Future studies should elucidate the targets for S-bacillithiolations that are reduced by the BrxA/BSH/YpdA pathway inside macrophages, increasing survival, metabolism or persistence under infections.

In summary, our results showed the importance of the BrxA/BSH/YpdA redox pathway to resist oxidative stress and macrophage infection in S. aureus. Through measurements of the BSH/BSSB redox ratio and EBSH, we provide evidence that the NADPH-dependent disulfide reductase YpdA regenerates BSH and restores reduced EBSH upon recovery from oxidative stress in S. aureus. Finally, biochemical evidence for YpdA as BSSB reductase and for the role of BrxA/BSH/YpdA pathway in de-bacillithiolation was provided in vitro. The detailed biochemical mechanism of YpdA and the cross-talk of the Trx and Brx systems in de-bacillithiolation under oxidative stress and infections are subject of our future studies.

**AUTHOR CONTRIBUTIONS**

HA and NL designed the experiments of this study. NL, VVL, VNF, QNT and SS constructed the mutants, performed the experiments and analyzed the data of this manuscript. MW and RH performed the HPLC thiol metabolomics analyses and analyzed the data. KT and MF contributed with the infection assays to this work. CH synthesized BSH and BSSB for the biochemical assays of the manuscript. NL and HA wrote the manuscript. All authors contributed with corrections of the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.01355/full#supplementary-material
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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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