

Role and Regulation of Poly-3-Hydroxybutyrate in Nitrogen Fixation in *Azorhizobium caulinodans*

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An *Azorhizobium caulinodans phaC* mutant (OPS0865) unable to make poly-3-hydroxybutyrate (PHB), grows poorly on many carbon sources and cannot fix nitrogen in laboratory culture. However, when inoculated onto its host plant, *Sesbania rostrata*, the *phaC* mutant consistently fixed nitrogen. Upon re-isolation from *S. rostrata* root nodules, a suppressor strain (OPS0921) was isolated that has significantly improved growth on a variety of carbon sources and also fixes nitrogen in laboratory culture. The suppressor retains the original mutation and is unable to synthesize PHB. Genome sequencing revealed a suppressor transition mutation, G to A (position 357,354), 13 bases upstream of the ATG start codon of *phaR* in its putative ribosome binding site (RBS). PhaR is the global regulator of PHB synthesis but also has other roles in regulation within the cell. In comparison with the wild type, translation from the *phaR* native RBS is increased approximately sixfold in the *phaC* mutant background, suggesting that the level of PhaR is controlled by PHB. Translation from the *phaR* mutated RBS (RBS*) of the suppressor mutant strain (OPS0921) is locked at a low basal rate and unaffected by the *phaC* mutation, suggesting that RBS* renders the level of PhaR insensitive to regulation by PHB. In the original *phaC* mutant (OPS0865), the lack of nitrogen fixation and poor growth on many carbon sources is likely to be due to increased levels of PhaR causing dysregulation of its complex regulon, because PHB formation, per se, is not required for effective nitrogen fixation in *A. caulinodans*.

Keywords: *Azorhizobium caulinodans*, bacteria-plant symbiosis, carbon sink, PhaR, PHB, poly-3-hydroxybutyrate, reductant storage, *Rhizobium*-legume symbiosis, *Sesbania rostrata*

Biological nitrogen fixation by diazotrophic bacteria enables them to convert atmospheric dinitrogen into ammonia using the enzyme nitrogenase (Oldroyd et al. 2011). *Azorhizobium caulinodans*, a species of the order *Rhizobiales*, is able to fix nitrogen under free-living conditions, as well as during symbiosis with its host plant, *Sesbania rostrata* (Dreyfus et al. 1983, 1988;

Elmerich et al. 1982). This is unusual for nitrogen-fixing rhizobia, because most can only fix nitrogen when in symbiosis with their host plant (Terpolilli et al. 2016).

Given that the nitrogenase enzyme is permanently inactivated by oxygen, a low-oxygen environment is essential for successful nitrogen fixation to occur (Rutten and Poole 2019). This presents a problem, because rhizobia, including *A. caulinodans*, are typically obligate aerobic organisms using the tricarboxylic acid (TCA) cycle as their primary means of energy generation. Under low-oxygen conditions, there may be insufficient oxygen to act as a terminal acceptor for electron transport to balance reductant formation. One of the main ways to remove an excess of reductant is channeling reductant into the generation of carbon storage molecules such as lipids or poly-3-hydroxybutyrate (PHB) (Terpolilli et al. 2016).

PHB is a carbon storage molecule synthesized by most rhizobia (Trainer and Charles 2006) using molecules of acetyl CoA in a multistage reaction, the polymerization step of which is catalyzed by PHB synthase (PhaC) (Schubert et al. 1988; Sharma and Dhingra 2016) (Fig. 1). In addition to its role as a reductant sink, the carbon storage provided by PHB has been shown to support nitrogen fixation during periods of carbon starvation and it has been suggested that it has a role in initial nodule formation (Gerson et al. 1978; Wang et al. 2007). Many species of rhizobia, including *A. caulinodans*, *Bradyrhizobium japonicum*, *Rhizobium etli*, and *R. tropici*, accumulate PHB both under free-living conditions (Stam et al. 1986; Tombolini and Nuti 1989) and within the nitrogen-fixing nodule form of rhizobia known as bacteroids (Karr et al. 1984; Ndoeye et al. 1994).

Mutagenesis studies performed in a number of species to determine the importance of PHB in nitrogen fixation have produced a mixture of results. In *R. etli*, *phaC* mutants fix nitrogen at higher levels than the wild type (Cevallos et al. 1996), whereas reduced symbiotic efficiency was seen in *Sinorhizobium meliloti* in symbiosis with *Medicago truncatula* but not with *M. sativa* (Wang et al. 2007). No change was seen in the fixation rates of *phaC* mutants of *R. leguminosarum* in nodules on *Pisum sativum* or *Phaseolus vulgaris* (Lodwig et al. 2005).

Formation of PHB is regulated by the interaction of PhaR, an autoregulating negative regulator (Maehara et al. 2002), and phasin proteins (PhaP). Phasins are present in both prokaryotes and archaea (Cai et al. 2015; Jang et al. 2017; Kim et al. 2017), where they are sequestered to the surface of polyhydroxyalkanoate granules such as PHB, reducing the apparent concentration within the cytoplasm despite continuous synthesis. When the granules reach a critical size, PhaP displaces PhaR from the surface due to the greater binding affinity to PHB of PhaP relative to PhaR. This results in a buildup of cytoplasmic PhaR, which then binds to the promoters of *phaR*,

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phaP, and *phaC*, preventing their transcription and halting production of further PHB (Kuchta et al. 2007; Pötter et al. 2002).

Previous knockout mutations of *phaC* showed it to be essential for nitrogen fixation and effective growth in *A. caulinodans*, suggesting that PHB synthesis is required for fixation in this species (Mandon et al. 1998). During symbiosis, this mutant strain was found to primarily produce small, nonfixing nodules containing no bacteria; however, it was noted at the time that some plants also contained “a few large nitrogen-fixing nodules”, which were theorized to be revertants (Mandon et al. 1998). In this work, we confirm that an *A. caulinodans phaC* mutant cannot fix nitrogen in the laboratory but consistently forms suppressor strains in planta that recover N₂ fixation. PHB synthesis, per se, is unnecessary for effective fixation by *A. caulinodans* in either free-living or symbiotic conditions. Instead, *phaR* dysregulation prevents N₂ fixation and reduces growth of a *phaC* mutant.

RESULTS

Phenotypic characterization of *phaC* mutants.

An *A. caulinodans phaC* mutant strain (OPS0865) was isolated via single integration using the plasmid pK19mob. Under free-living conditions, OPS0865 did not fix nitrogen (Fig. 2A). Additionally, at 21% oxygen, the *phaC* mutant had reduced growth on acetate, arabinose, and pyruvate, though growth on succinate was unaffected (Fig. 3A). This growth defect was even more severe at 1% oxygen (Fig. 3B). Growth on succinate had the shortest generation time and was equivalent in the mutant and wild type at 21% oxygen (approximately 3.2 ± 0.1 h). However, when grown on succinate at 1% oxygen, the *phaC* mutant had a generation time of 19.3 ± 1.5 h, compared with 7.5 ± 1.2 h for wild-type ORS571. Thus, mutation of *phaC* disrupts growth of *A. caulinodans* and this disruption is exacerbated by oxygen limitation. Several attempts to complement the mutation with a plasmid-borne copy of *phaC* failed. Therefore, to determine whether the mutation in *phaC* causes the observed phenotype, we isolated 12 revertants, where the integrated pK19mob in *phaC* was deleted, by repeated growth on media

lacking antibiotics. Revertants were screened for restored sensitivity to neomycin and the presence of a full-length *phaC* gene using primers oxp3394 and oxp3395. All 12 revertants were restored to wild-type growth on tryptone-yeast (TY) media, and 1 such revertant strain (OPS2077) was retained for further analysis. Wild-type levels of growth on all carbon sources were

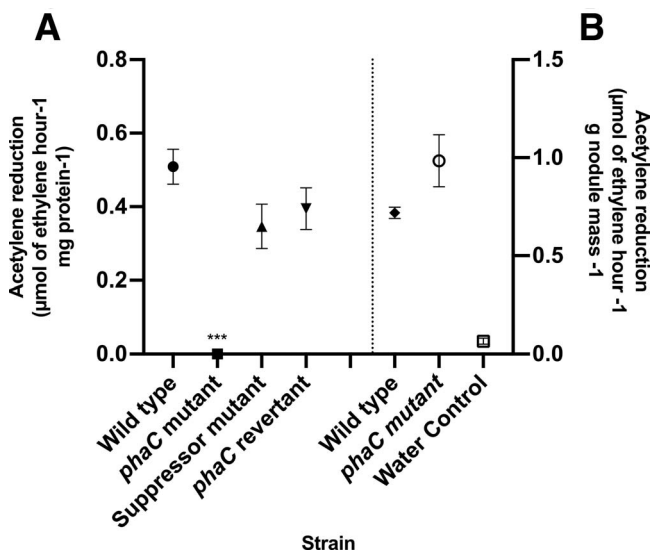


Fig. 2. Acetylene reduction activity (ARA) under free-living and symbiotic nitrogen-fixing conditions for wild-type ORS571, the *phaC* mutant (OPS0865), suppressor mutant (OPS0921), and *phaC* revertant (OPS2077) strains of *Azorhizobium caulinodans*. **A**, Mean ARA of free-living bacteria in micromoles of ethylene per hour per milligram of protein. The three strains ORS571, OPS0921, and OPS2077 are significantly different from strain OPS0865 ($P < 0.005$), Mann-Whitney test; $n = 6$. **B**, Mean ARA during symbiosis at 28 days postinoculation of *Sesbania rostrata* in micromoles of ethylene per hour per gram of nodule. Water control = plants inoculated with water alone. No significant difference was observed between the wild type and the *phaC* mutant ($P = 0.4$), two-tailed Mann-Whitney test. Error bars show standard error; $n = 3$.

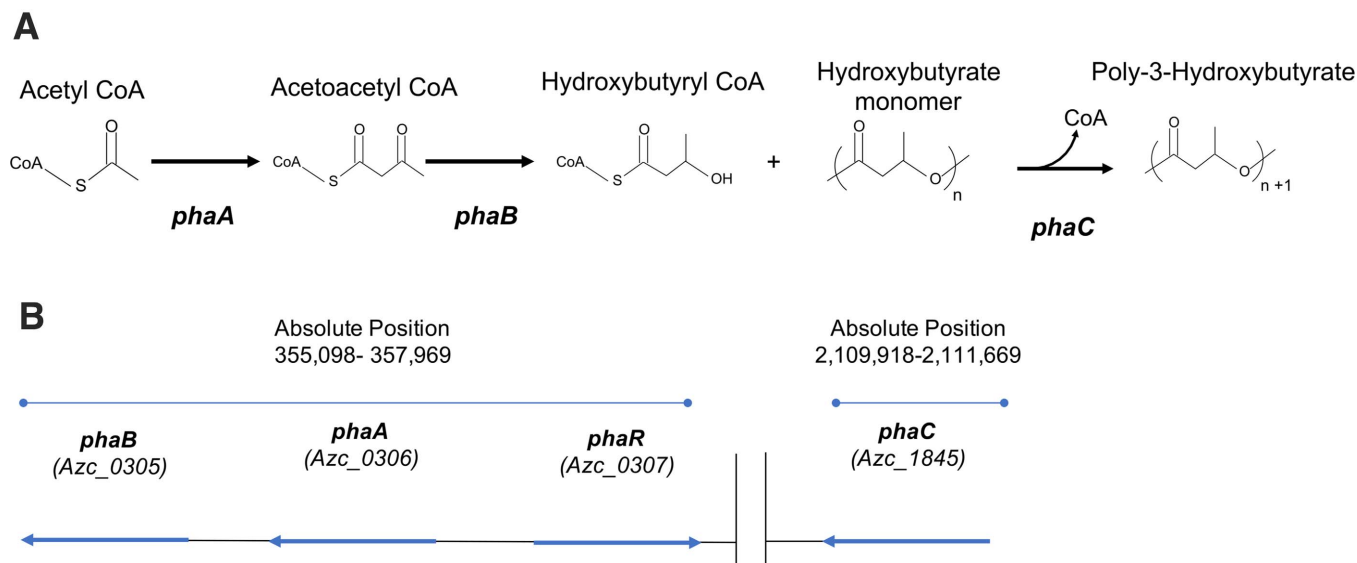


Fig. 1. Poly-3-hydroxybutyrate (PHB) synthesis in *Azorhizobium caulinodans* and the genes encoding the proteins involved. **A**, Enzymic reactions in PHB synthesis. Condensation of two acetyl CoA molecules by PhaA to acetoacetyl CoA, followed by reduction to hydroxybutyryl CoA by PhaB. Monomeric hydroxybutyrate is linked to hydroxybutyryl CoA by PhaC to form the polymer, releasing CoA. This polymer is extended by PhaC-catalyzed reactions with additional hydroxybutyryl CoA. **B**, Genome architecture of genes encoding the proteins involved in PHB synthesis. Gene numbers and absolute genome positions for the two regions are indicated. PhaA is an acetyl CoA acetyl-transferase, PhaB is an acetoacetyl CoA reductase, PhaC is a PHB synthase, and PhaR is a regulator.

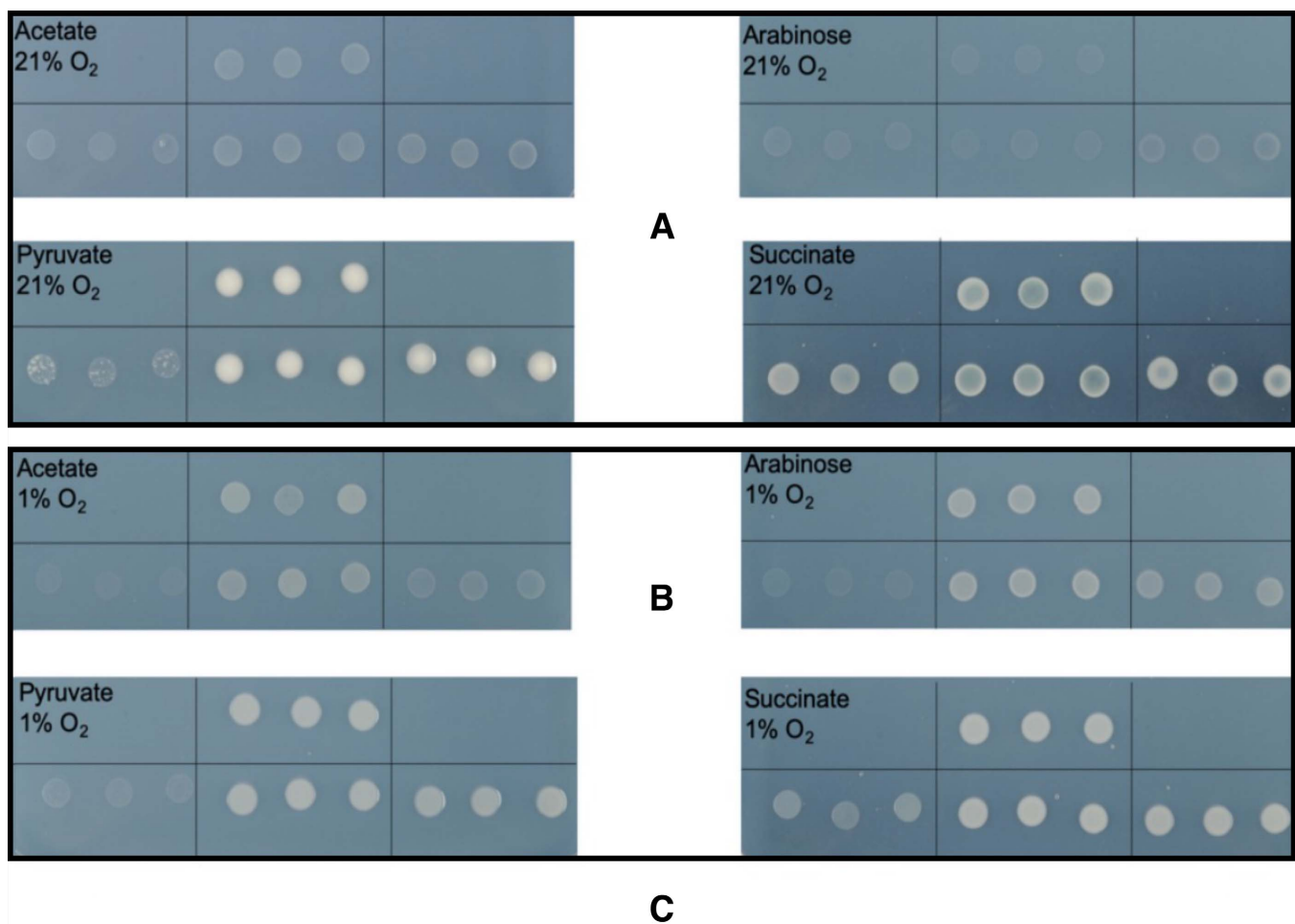
restored in the revertant strain (Fig. 3) along with wild-type rates of free-living nitrogen fixation (Fig. 2A). These observations are in line with previous reports (Mandon et al. 1998) that PHB is required for effective growth on a variety of carbon sources and alters N₂ fixation in *A. caulinodans*.

To our surprise, when the *phaC* mutant was inoculated onto *Sesbania rostrata* plants, we observed wild-type symbiotic nitrogen fixation rates on all plants measured; furthermore, all nodules on the plants were large and had a pink, wild-type appearance (Fig. 2B). Our initial suspicion was that these fixation rates were due to a spontaneous reversion mutant being favored by the plant, resulting in the loss of the *phaC* mutation and causing the wild-type appearance of the nodules. To test whether a such a mutation had occurred, bacteria were isolated from six separate nodules from plants inoculated with the *phaC* mutant. These colonies were initially isolated on TY without added antibiotics and individual purified colonies were then screened for neomycin resistance and the presence of the pK19mob plasmid, using PCR detection with a chromosomal

primer (oxp1043) and a pK19mob-specific primer (pK19B), amplifying outwards from the plasmid backbone. All purified colonies retained neomycin resistance, while PCR amplification gave a 1.5-kb fragment in both the original *phaC* mutant strain (OPS0865) and in the nodule-derived isolates but gave no band in the wild type. Glycerol stocks were made of three PCR-mapped nodule isolates (OPS0921, OPS0922, and OPS0923). Wild-type levels of free-living nitrogen fixation were observed in free-living cultures of the nodule-isolated strain OPS0921, whereas the original *phaC* mutant retained a Fix⁻ phenotype (Fig. 2A). Thus, passage through the plant restored nitrogen fixation to the *phaC* mutant strain. Because these nodule isolated strains (OPS0921, OPS0922, and OPS0923) retained the original *phaC* mutation, we subsequently refer to them as suppressor strains.

Quantification of PHB and growth of *phaC* mutants.

To check for the presence of PHB, we made transmission electron micrographs of *S. rostrata* nodules. White, electron-



Sole carbon source O ₂ concentration	Wild type (ORS571)	
<i>phaC</i> mutant (OPS0865)	<i>phaC</i> reversion mutant (OPS2077)	<i>phaC</i> suppressor mutant (OPS0921)

Fig. 3. Growth of the wild type (ORS571), *phaC* mutant (OPS0865), *phaC* revertant (OPS2077), and suppressor mutant (OPS0921) strains of *Azorhizobium caulinodans* at 28°C on acetate, arabinose, pyruvate, or succinate as a sole carbon source. **A**, At 21% oxygen; **B**, at 1% oxygen; and **C**, relative position of bacterial strains. Bacteria (5 µl at an optical density at 600 nm of approximately 0.1) were drop plated onto solid media with sole carbon source shown; *n* = 3.

transparent PHB granules were present in the wild type (Fig. 4A) but neither in the *phaC* mutant (OPS0865) nor the putative *phaC* suppressor mutant (OPS0921) (Fig. 4).

Because the *phaC* mutant and the putative suppressor mutant OPS0921 had such a clear difference in their ability to fix nitrogen, their growth was assessed at both 21 and 1% oxygen to identify any differences (Fig. 3). The *phaC* mutant showed the biggest reduction in growth relative to the wild type, with the putative suppressor (OPS0921) producing an intermediate growth phenotype. Of particular note was the poor growth of the *phaC* mutant on acetate and pyruvate as sole carbon sources at both oxygen tensions.

Given the altered growth of the putative suppressor strain (OPS0921), the retention of neomycin resistance and pK19mob insertion, together with the lack of PHB synthesis, it was apparent that, in addition to the original *phaC* mutation, there must be a suppressor mutation present in this strain. To identify the mutation, whole-genome sequencing was undertaken utilizing the sequencing services of MicrobesNG. The three isolated putative suppressor strains (OPS0921, OPS0922, and OPS0923) were sequenced and compared with both the *phaC* mutant and the wild type, with the assumption that any mutation shared by the putative suppressor strains (OPS0921, 0922 and 0923) but absent from the original *phaC* mutant and ORS571 may cause the suppression. As expected, the draft genomes of all putative suppressors (OPS0921, 0922, and 0923) retained the pK19mob plasmid backbone. However, there is a single-nucleotide polymorphism (SNP) present in all three putative suppressor mutants (OPS0921, OPS0922, and OPS0923) and absent from both the wild type and the original *phaC* mutant. This is a G-to-A transition (position 357,354), 13 bases upstream of the *phaR* ATG start codon. The location of the SNP is likely to be within the ribosome binding site (RBS) of the *phaR* gene and, thus, may affect translation of *phaR*. Because the three putative suppressor strains were otherwise genetically identical, further investigations were carried out on the strain OPS0921 alone, hereafter referred to as the suppressor mutant. Hereafter, this SNP will be referred to as 0921 SNP and this mutated RBS as RBS*.

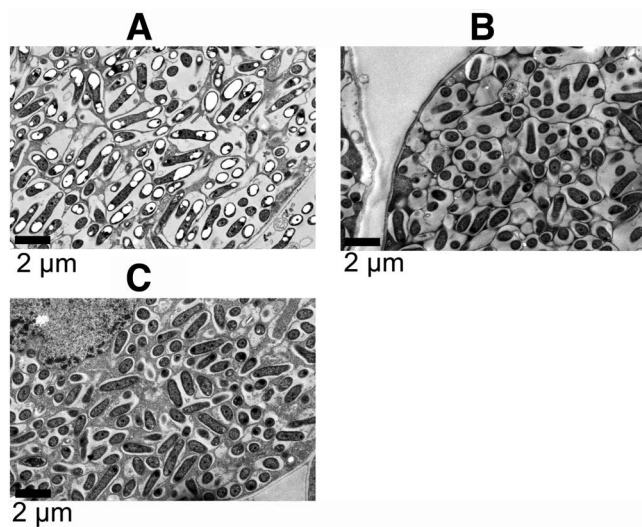


Fig. 4. Transmission electron micrographs showing bacteroids within sections from nodules of *Sesbania rostrata* at 28 days postinoculation. Poly-3-hydroxybutyrate (PHB) granules are white inclusions within bacteroids **A**, Wild type (ORS571) in which PHB granules are clearly visible; **B**, *phaC* mutant (OPS0865), no PHB visible; and **C**, *phaC* suppressor mutant (OPS0921), no PHB visible. Scale bars are shown.

Translation and transcription of *phaR* in different strain backgrounds.

Translation of *phaR* was measured using translational fusions to green fluorescent protein (GFP). The native *phaR* RBS fusion produced sixfold more GFP in the *phaC* mutant than in that of the wild type (Fig. 5A). This increase in GFP translation (and by extension, PhaR protein) was likely to be a response to the lack of PHB in the *phaC* mutant strain. Thus, translation of *phaR* is reduced by the presence of PHB and, consequently, in the absence of PHB, the level of PhaR increases. In contrast, the *phaR* RBS* translational fusion gave a consistently low GFP signal in both the *phaC* and wild-type ORS571 backgrounds (Fig. 5A), suggesting that RBS* is insensitive to the presence of PHB in the bacterial cell. Thus, the RBS* 0921 SNP mutation has locked translation of *phaR* at a low level independent of PHB.

Transcription of *phaR* and *phaA* was measured in the wild type and the *phaC* mutant by quantitative reverse-transcription PCR (qRT-PCR) (using *gyrA* as a control housekeeping gene). There was no significant difference between the two strains in the transcription of either *phaR* or *phaA* (Fig. 5B), consistent with the effect of the suppressor mutations being at the translational level.

DISCUSSION

The *phaC* mutant has a reduced growth rate on acetate, arabinose, and pyruvate at atmospheric oxygen levels but not on succinate (Fig. 3A). This was not unexpected, given the high levels of acetyl-CoA generated by growth on these three carbon sources, and is in line with previous PHB mutants in *R. etli* and *R. leguminosarum* bv. *viciae* 3841 (Cevallos et al. 1996; Ludwig et al. 2005). The observed growth defect was far greater at low oxygen (1%), with reduced growth even on succinate. This limited growth is likely to result from flux imbalance, with acetate and pyruvate generating acetyl-CoA directly, which feeds into the TCA cycle. This generates 4 mol of reductant for every 1 mol of pyruvate utilized. Succinate, by contrast, can generate small amounts of reductant via malic enzyme, or malate dehydrogenase activity, and can be converted to sugar via gluconeogenesis. This sugar can be converted to storage molecules or catabolized and fed into the TCA cycle at a more controlled rate. As a result, the *phaC* mutants grown on succinate displayed the smallest growth defect. The wild-type strain and the revertant strain (OPS2077) do not display any growth impairment on acetate or pyruvate because they can use PHB as an acetyl-CoA and reductant sink. Thus, under a low oxygen concentration, synthesis of PHB helps to maintain redox balance in *A. caulinodans*. Although growth of *A. caulinodans* strains on arabinose also showed the importance of PHB at 1% oxygen, growth was unexpectedly but consistently better at 1 than 21%.

Although the *phaC* mutant did not fix nitrogen in free-living conditions (Fig. 2A), *phaC* suppressor mutants (e.g., OPS0921), capable of nitrogen fixation, were consistently isolated from root nodules of *S. rostrata* (Fig. 2B). Transmission electron microscopy (TEM) micrographs (Fig. 4) confirmed that the suppressor strains lacked PHB; thus, its absence does not prevent nitrogen fixation in *A. caulinodans*. Whole-genome sequencing showed that the suppressor strains have a mutation in the putative ribosome binding site of *phaR*. GFP translational fusions to *phaR* demonstrated that the wild-type RBS had a sixfold increase in translation in the *phaC* mutant compared with the wild type, while the mutated RBS from the suppressor strains (RBS*) had a uniformly low translation rate in both wild-type and *phaC* backgrounds (Fig. 5). PhaR levels are usually regulated by PHB levels within the cell (Maehara et al. 2002). The location of the SNP within the RBS of the major regulatory gene *phaR* in the

suppressor strain suggests lack of PHB results in the dysregulation of PhaR. In *B. japonicum*, such a mutation would presumably result in the downregulation of more than 200 genes, including the key regulator *fixK2* (Quelas et al. 2016). This downregulation, combined with the lack of the major reductant sink PHB, would explain the extremely poor growth rate of the *phaC* mutant strain (OPS0865). The partial rescue of growth in the suppressor strain (OPS0921) appears to have been achieved by reducing translation of *phaR* to basal levels (Figs. 3 and 5A). This appears to prevent some of the effects of PhaR dysregulation; however, growth rates were not restored to wild-type levels, probably due to the suppressor strain lacking PHB as a reductant sink. The dysregulation of PhaR levels may also explain why attempts to rescue the initial *phaC* mutant strain with plasmid-borne copies of the *phaC* gene were unsuccessful, because multiple copies of *phaC* may also result in PhaR dysregulation as a result of excess PHB production, though this remains speculation.

It was shown by qRT-PCR that there was no significant difference in transcription of either *phaR* or *phaA* in either the wild type or the *phaC* mutant (Fig. 5B). The lack of a significant change in *phaA* transcription in the *phaC* mutant strain is surprising given the sixfold increase in translation in this strain of the repressor PhaR. We would have expected to observe a much greater reduction in *phaA* transcription, and this suggests that any reduction in expression caused directly by PhaR may be modest. The numerous genes encoding regulatory proteins whose expression is controlled by PhaR (e.g., *fixK2*) could serve to amplify the effect of this modest reduction (Quelas et al. 2016), contributing to the reduced growth seen in the *phaC* mutant strain compared with the suppressor strain (Fig. 3).

Because a reduced level of the global PhaR regulator restored wild-type nitrogen fixation in suppressor strains of *phaC* (Fig. 2A), this suggests that the loss of fixation was due to an excess of PhaR rather than the lack of a PHB as a reductant

sink. This brings *A. caulinodans* in line with most other rhizobial species, which are able to fix nitrogen without PHB synthesis (Cevallos et al. 1996; Lodwig et al. 2005; Quelas et al. 2013) by making use of other reductant sinks such as lipid or, potentially, glycogen (Lodwig et al. 2005; Wang et al. 2007).

Overall, it is clear that, while PHB is not strictly necessary for nitrogen fixation in *A. caulinodans*, it has an important role both as a redox sink and for maintaining the level of PhaR in the cell. Disruption of either of these roles can result in a severe growth defect, particularly when grown on carbon sources that result in high levels of acetyl-CoA.

MATERIALS AND METHODS

Bacterial strains, media, and bacterial growth conditions.

All bacterial strains, plasmids, and primers used in this study are listed in Table 1. *A. caulinodans* strains were routinely grown on solid media at 37°C on TY (Beringer 1974) or universal minimal salts (UMS) with 1× stock vitamins (thiamine hydrochloride at 1 g liter⁻¹, D-pantothenic acid calcium salt at 2 g liter⁻¹, and Biotin at 100 mg liter⁻¹); 300 μM nicotinic acid; either 10 mM glutamate or 10 mM ammonium chloride (as a nitrogen source); and 20 mM succinate, 6 mM acetate, 20 mM arabinose, or 30 mM pyruvate (as a carbon source). UMS is a modified version of ammonium mineral salt medium (Poole et al. 1994), with alterations previously published (Pini et al. 2017). Experiments using *A. caulinodans* liquid cultures were grown at 28°C (this was the maximum temperature that could be maintained by the oxygen cabinet which was used for acetylene reduction activity [ARA] experiments with free-living bacteria). Antibiotics in *A. caulinodans* were used at the following concentrations, unless stated otherwise; ampicillin (100 μg/ml), gentamycin (25 μg/ml) neomycin (80 μg/ml), and nitrofurantoin (5 μg/ml). *Escherichia coli* strains were grown at 37°C in Luria-Bertani medium using gentamycin (10 μg/ml) and kanamycin (20 μg/ml).

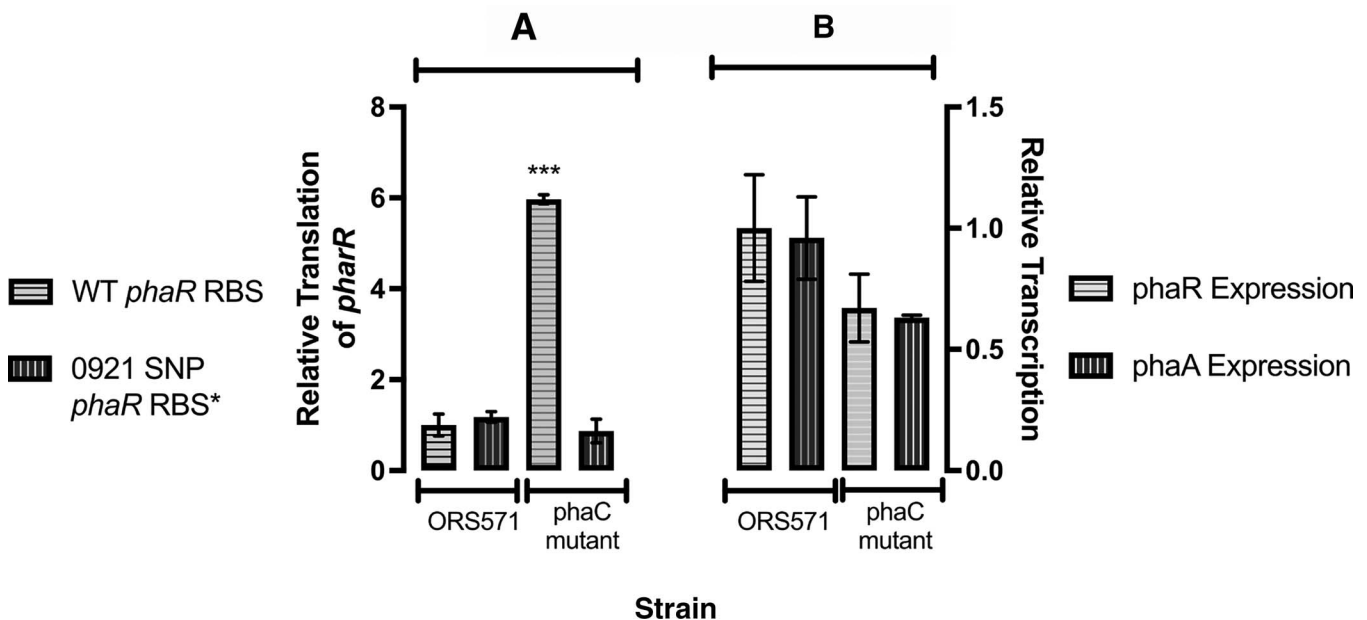


Fig. 5. Translation and transcription of *phaR* and transcription of *phaA* in the wild type (WT) (ORS571) and *phaC* mutant (OPS0865) backgrounds of free-living *Azorhizobium caulinodans*. **A**, Relative translation from *phaR* ribosome binding site (RBS) (WT) and *phaR* RBS* (SNP 0921) measured by fluorescence of a green fluorescent protein (GFP) translational reporter. Data are relative to GFP measured from *phaR* RBS (WT) in WT (ORS571) background. Error bars are standard error. Tukey's multiple comparison test indicated a significant difference between RBS and RBS* in *phaC* mutant (OPS0865) backgrounds; asterisks (***) indicate $P < 0.0001$, degrees of freedom = 89. No significant difference is seen between any combination of the other three samples; $n = 8$. **B**, Relative transcription measured by quantitative reverse-transcription PCR of *phaA* and *phaR* in WT (ORS571) and *phaC* mutant (OPS0865) strains, compared with housekeeping gene *gyrA*. Error bars are standard error. No significant difference was seen between expression of these genes in either strain; $n = 3$.

Table 1. Strains, plasmids, and primer sequences used in this study

Strains, plasmids, and primers	Relevant characteristics and primer sequence ^a	Source
Strain		
<i>Azorhizobium caulinodans</i> ORS571	Wild type, ampicillin resistant	Dreyfus et al. 1983
OPS0865	<i>phaC</i> mutant generated by integration of pOPS0391 into ORS571, ampicillin resistant, neomycin resistant	This study
OPS0921	Suppressor mutant isolated from nitrogen-fixing root nodules of <i>Sesbania rostrata</i> inoculated with OPS0865, ampicillin resistant, neomycin resistant	This study
OPS0922	Suppressor mutant isolated from nitrogen-fixing root nodules of <i>S. rostrata</i> inoculated with OPS0865, ampicillin resistant, neomycin resistant	This study
OPS0923	Suppressor mutant isolated from nitrogen-fixing root nodules of <i>S. rostrata</i> inoculated with OPS0865, ampicillin resistant, neomycin resistant	This study
OPS1426	pOPS1029 conjugated into ORS571, ampicillin resistant, gentamycin resistant	This study
OPS1427	pOPS1030 conjugated into ORS571, ampicillin resistant, gentamycin resistant	This study
OPS1432	pOPS1029 conjugated into OPS0865, ampicillin resistant, gentamycin resistant	This study
OPS1433	pOPS1030 conjugated into OPS0865, ampicillin resistant, gentamycin resistant	This study
OPS1438	pOPS1029 conjugated into OPS0921, ampicillin resistant, gentamycin resistant	This study
OPS1439	pOPS1030 conjugated into OPS0921, ampicillin resistant, gentamycin resistant	This study
OPS2077	OPS0865 reversion mutant which has lost the integrated <i>pk19mob</i> via multiple sub culturing on TY, confirmed via PCR across the <i>phaC</i> (PHB synthase) gene with <i>oxp3394</i> + <i>3395</i> (1,900 bp in wild type)	This study
<i>Escherichia coli</i> DH5 α	F- <i>deoR endA1 recA1 relA1 gyrA96 hsdR17</i> (rk-, mk+) <i>supE44 thi-1 -phoA Δ(lacZYA-argF)</i> U169 Φ 80 <i>lacZ</i> ΔM15 λ .	Bioline
<i>E. coli</i> 803	<i>hsdS</i> (rk-, mk-, gal-, met-) in this work contained pRK2013 helper plasmid	Wood 1996
Plasmid		
pK19mob	Mobilizable <i>E. coli</i> cloning vector, kanamycin and neomycin resistant	Schäfer et al. 1994
pOPS0391	1.1 kb of ORS571 <i>phaC</i> generated by PCR with <i>oxp1218/1219</i> InFusion cloned into <i>HindIII</i> -linearized pK19mob, kanamycin and neomycin resistant	This study
pOPS1029	<i>phaC-phaR</i> intergenic region + 90 bp of <i>phaR</i> (generated by PCR with <i>oxp2419/2420</i> from wild type ORS571 genomic DNA (RBS) InFusion cloned into pRU1097 linearized by PCR with <i>oxp2421/oxp2422</i> , gentamycin resistant	This study
pOPS1030	<i>phaC-phaR</i> intergenic region + 90 bp of <i>phaR</i> (generated by PCR with <i>oxp2419/2420</i> from suppressor mutant OPS0921 genomic DNA (RBS*) InFusion cloned into pRU1097 linearized by PCR with <i>oxp2421/oxp2422</i> , gentamycin resistant	This study
pRK2013	ColEI replicon with RK2 <i>tra</i> genes; helper plasmid used for mobilizing plasmids, kanamycin resistant	Karunakaran et al. 2005
pRU1097	p318/p319 PCR product (<i>gfpmut3.1</i>) from pBJA27 cloned in pOT2 as <i>SpeI-SacI</i> promoter-less GFP module, gentamycin resistant	Karunakaran et al. 2005
Primer		
p318	GAGAGAGAAGCTAGTGGAGGAAGAAAAATGCG-TAAAGGAGAAGAAGCTTTTCA	Karunakaran et al. 2005
P319	CTCTCGAGCTCATTGTATAGTTCATCCATGC	Karunakaran et al. 2005
M13 forward	CACGACGTTGTAACGCA; sequencing primer for insert cloned in pK19mob	Bahar et al. 1998
M13 reverse	GGATAACAATTTACACAGG; sequencing primer for insert cloned in pK19mob	Bahar et al. 1998
oxp1043	TTTTTGGTACCATGGAGGCGTTCGCCAGAA; forward mapping primer for <i>phaC</i> gene	This study
oxp1218	TGATTACGCCAAGCTATAAGCGGAGTTGGGCGG; forward PCR primer to amplify internal <i>phaC</i> fragment	This study
oxp1219	GCAGGCATGCAAGCTGTTCTTCGACGCCCTGAAGC; reverse PCR primer to amplify internal <i>phaC</i> fragment	This study
oxp2419	GTGGAGGAAGAAAAATGGGCTTCTCTCCTGGGG; forward PCR primer to amplify <i>phaC-phaR</i> intergenic region + 90-bp <i>phaR</i>	This study

(Continued on next page)

^a Primer sequences are presented 5' to 3'. PHB = poly-3-hydroxybutyrate, RBS = ribosome binding site, GFP = green fluorescent protein, and qRT-PCR = quantitative reverse-transcription PCR.

Table 1. (Continued from previous page)

Strains, plasmids, and primers	Relevant characteristics and primer sequence ^a	Source
oxp2420	TTCTTCTCCTTTACGTTTCGAGGGTCACATAGGTACTG; reverse PCR primer to amplify <i>phaC-phaR</i> intergenic region + 90-bp <i>phaR</i>	This study
oxp2421	CGTAAAGGAGAAGAACTTTTCACTGG; forward inverse PCR primer to linearize pRU1097	This study
oxp2422	TTTTTCTTCTCCACTAGTCTCTCT; reverse inverse PCR primer to linearize pRU1097	This study
oxp2475	GAACGGCTACGAGTGGAAC; forward qRT-PCR primer for <i>gyrA</i>	This study
oxp2476	GGAGAAGGTCTGCACCATG; reverse qRT-PCR primer for <i>gyrA</i>	This study
oxp2971	CCATGTTGCGGCGAACC; forward qRT-PCR primer for <i>phaR</i>	This study
oxp2972	CGCTTCTCGAGATGTCCAT; reverse qRT-PCR primer for <i>phaR</i>	This study
oxp2976	CCATGTGGTAGCCGTTGAAG; reverse qRT-PCR primer for <i>phaA</i>	This study
oxp3105	ATGAGCCAGTCCACCCATG; forward qRT-PCR primer for <i>phaA</i>	This study
oxp3394	TGCCGAATTCGGATCCGGAGAAGCTGACGGCGC-GAAA; forward PCR mapping primer <i>phaC</i>	This study
oxp3395	TCAACAGGAGTCCAAGAGCGGATTAAGGCAGGTGACAGC; reverse PCR mapping primer <i>phaC</i>	This study
pk19A	ATCAGATCTTGATCCCTGC; PCR mapping primer for pK19mob	Karunakaran et al. 2010

Testing bacterial growth on different carbon sources.

Single colonies of *A. caulinodans* strains were grown overnight in TY at 37°C, then diluted to an optical density at 600 nm (OD₆₀₀) of approximately 0.1. Three biological replicates of each strain were pipetted as 5 µl-droplets onto UMS agar plates supplemented with vitamins, nicotinic acid, ammonium chloride, and either acetate, arabinose, pyruvate, or succinate as the sole carbon source. Plates were incubated at 21 or 1% oxygen at 28°C for 60 h before being photographed.

Growth and nodulation of *S. rostrata*.

Seed were sterilized for 10 min in concentrated sulfuric acid to break seed dormancy, then washed in 400 ml of sterile MilliQ water. Seed were germinated for 3 days on 2% water agar containing a double layer of autoclaved filter paper soaked in MilliQ water. Germinated seed were planted individually in 1-liter pots containing autoclaved vermiculite and 300 ml of nitrogen-free rooting solution (4 mM Na₂HPO₄, 3.7 mM K₂PO₄, 1 mM CaCl₂, 800 µM MgSO₄, 100 µM KCl, 35 µM H₃BO₃, 10 µM Fe EDTA, 9 µM MnCl₂, 0.8 µM ZnCl₂, 0.5 µM Na₂MoO₄, and 0.3 µM CuSO₄). Inoculum was prepared from a TY slope (containing ampicillin or neomycin), grown for 3 days at 37°C by resuspending the bacteria in 3 ml of TY. Inoculum (1 ml, diluted 1/100) was applied to each germinated seed. Four germinating seeds were inoculated with each bacterial strain. Plants were watered with sterile water every 2 days and grown at 22°C for 28 days in a controlled-environment room with a photoperiod of 16 h of light and 8 h of darkness.

Generation of a *phaC* mutant.

A. caulinodans mutants were generated using integration through homologous recombination (single cross-over) into the genome of plasmid DNA cloned in the mobilizable vector pK19mob (Schäfer et al. 1994). Plasmid pOPS0391 was constructed by InFusion (Takara Bio) cloning 1.1 kb of ORS571 *phaC* gene amplified with primers oxp1218 and oxp1219 into the *Hind*III site of pK19mob. Construction of pOPS0391 was confirmed by sequencing with primers M13 forward and M13 reverse (Eurofins Genomics). Conjugation of pOPS0391 into ORS571 was performed using a donor strain of *E. coli*

(pOPS0391) and *E. coli* strain 803 (pRK2013). The conjugation mix was allowed to grow overnight on TY agar at 37°C, then counter-selected using neomycin (240 µg/ml) and nitrofurantoin. The mutation was confirmed by PCR mapping with primers oxp1043 and pK19A and sequencing the amplified fragment (Eurofins Genomics).

Reversion of *phaC* mutation.

A single colony of the *phaC* mutant was grown for 16 h in 5 ml of TY liquid culture with no antibiotics, before being subcultured into fresh TY and grown for a further 16 h. On the following day, 20 µl of culture was plated onto TY ampicillin and incubated for 2 days at 37°C. Colonies were patched onto TY neomycin and TY ampicillin agar plates. Neomycin-sensitive cultures were screened for the presence of *phaC* by PCR using oxp3394 and oxp3395. An isolate which has lost the pK19mob insertion in the chromosome and has a full-length *phaC* gene was retained as the reversion strain (OPS2077).

Assessing translation with GFP reporters.

GFP translational fusions to *phaR* gene of the wild type (RBS) and the suppressor mutant (RBS*) strains were made by PCR amplification with oxp2419 and oxp2420 (amplifying the *phaA-phaR* intergenic region from ORS571 and OPS0921, together with 90 bp of *phaR* gene). The products were cloned into pRU1097 (linearized by inverse PCR with oxp2421 and oxp2422) by InFusion HD cloning (Takara Bio). These constructs were conjugated into the *A. caulinodans* wild type (ORS571), *phaC* mutant (OPS0865), and suppressor mutant (OPS0921) backgrounds. Cultures were grown overnight in 10 ml of UMS containing 10 mM glutamate as the added nitrogen source. Cultures were grown to an OD₆₀₀ of approximately 0.4 before 200-µl aliquots were measured in triplicate for GFP fluorescence (excitation at 485 nm, absorbance at 520 nm) and OD₆₀₀ in a 96-well plate using a FLUOstar Omega spectrophotometric plate reader.

Assessing transcription by qRT-PCR.

RNA was extracted from 10-ml *A. caulinodans* TY cultures using the Direct-zol miniprep kit (Zymo). RNA was DNase treated using a Turbo DNA-free kit (Invitrogen). RNA

concentration was assessed via a Qubit RNA HS Assay Kit (Thermo Fisher Scientific) and converted to cDNA using Super-script II (Invitrogen). Housekeeping gene *gyrA* was used as a reference. qRT-PCR was performed using a SensiMix SYBR No-ROX Kit (Biolone) according to the manufacturer's instructions on a StepOnePlus qRT-PCR machine (Applied Biosystems).

Bioinformatic analysis.

Genomes of the *phaC* mutant and the three putative suppressor strains (OPS0921, OPS0922, and OPS0923) were sequenced by MicrobesNG (Birmingham, U.K.) service using Illumina Miseq 300PE technology. Reads from the *phaC* mutant and putative suppressor mutants (OPS0921, OPS0922, and OPS0923) were aligned against the chromosome of *A. caulinodans* ORS571 (NCBI: txid438753) using Bowtie2 v2.1.0 in "very-sensitive" parameters in end-to-end mode (Langmead and Salzberg 2012). The output of the alignment was transformed using SAMtools v1.4 (Li et al. 2009) and SNP detected with VarScan v2.4.3 (Koboldt et al. 2013).

Free-living *A. caulinodans* ARA.

Cultures were prepared by streaking single colonies of bacteria on 10-ml UMS slopes (0.6% agarose) in 28-ml universal tubes, supplemented with succinate, vitamins, nicotinic acid, and 10 mM NH₄Cl. After 2 days of incubation at 37°C, cultures were washed from the slopes three times with phosphate-buffered saline to remove residual nitrogen and resuspended in 2 ml of nitrogen-free UMS media at OD_{600nm} = 0.4 in universals. Cultures were placed in a controlled-atmosphere cabinet adjusted to 3% O₂ by flushing with N₂ gas, left for 1 h with shaking at 180 rpm, and sealed with a rubber top. To initiate fixation, 10% of the air (3.35 ml) from the headspaces of universals was removed with a syringe and replaced with C₂H₂ (2.8 ml). Cultures were then returned to incubation at 37°C and 180 rpm for 4 h. The production of C₂H₄ from C₂H₂ was measured at 4 h and again at 6 h postinitiation using gas chromatography mass spectrometry (Clarus 480 gas chromatograph; Perkin Elmer) and the rates of nitrogenase activity were subsequently calculated between these two time points. The protein concentrations of the cell lysates were quantified using a bicinchoninic acid protein assay (Sigma-Aldrich). Six biological replicates were included for each strain measured. Fixation rates are given in micromoles of ethylene per hour per milligram of protein.

In planta *A. caulinodans* ARA.

ARA was measured according to (Trinick et al. 1976), except 2% acetylene was used and incubation was for 2 h at room temperature before measurement on a gas chromatogram (Perkin Elmer). Fixation rates are given in micromoles of ethylene per hour per gram of nodule.

TEM of sectioned nodules.

The procedure used was modified from that of Beck et al. (2008) in the following manner: a JEOL JEM1400 transmission electron microscope was used, samples were not hexane treated, and pyroxylin-coated grids were used to collect sections cut on a Leica UCT ultramicrotome.

Data and materials availability.

All data and materials generated are available from P. S. Poole upon request.

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