An anti-antisigma factor in the response of the bacterium *Myxococcus xanthus* to blue light

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Cells of the Gram-negative bacterium *Myxococcus xanthus* respond to blue light by producing carotenoids, pigments that play a protective role against the oxidative effects of light. Blue light triggers a network of regulatory actions that lead to the transcriptional activation of the structural genes for carotenoid synthesis. The product of *carF*, similar to a family of proteins of unknown function called Kua, is an early regulator of this process. Previous genetic data indicate that CarF participates in the light-dependent inactivation of the antisigma factor CarR. In the dark, CarR sequesters the ECF-sigma factor CarQ to the membrane, thereby preventing the activation of the structural genes for carotenoid synthesis. Using a bacterial two-hybrid system, we show here that both CarF and CarQ physically interact with CarR. These results, together with the finding that CarF is located at the membrane, support the hypothesis that CarF acts as an anti-antisigma factor. Comparison of CarF with other Kua proteins shows a remarkable conservation of a number of histidine residues. The effects on CarF function of several histidine to alanine substitutions and of the truncation of specific CarF domains are also reported here.

INTRODUCTION

Carotenoids are widely distributed, pigmented compounds that are produced by both eukaryotes and prokaryotes. These compounds protect cells from photo-oxidative damage by scavenging harmful agents, such as singlet and triplet molecular species. In the non-photosynthetic, Gram-negative bacterium *Myxococcus xanthus*, the synthesis of carotenoids is known to be dependent either on illumination by blue light or on exposure to copper. Much is known on the activation mechanism triggered by the light stimulus, which is brought about by a complex interplay of regulatory proteins (Elías-Arnanz et al., 2008); all but one of these proteins also participate in the carotenogenic response to copper (Fig. 1; see below).

In *M. xanthus*, the structural genes for carotenoid synthesis are placed at two unlinked loci. One corresponds to the single gene *crtI* and the other to an operon, *carB*, where the rest of the structural genes are grouped (Fontes et al., 1993; Botella et al., 1995). The precise mechanism for the light activation of the *crtI* promoter (*P*<sub>I</sub>) differs from that of the *carB* promoter (*P*<sub>B</sub>). However, the earlier steps that lead to activation of *P*<sub>I</sub> as well as *P*<sub>B</sub> both depend on the actions of the same set of three proteins: CarQ, CarR and CarF. Two of these, a pair of sigma–antisigma factors, are the products of genes *carQ* and *carR*. They are cotranscribed, together with *carS*, in the *carQRS* operon, unlinked to other *car* genes and also controlled by a light-inducible promoter (*P*<sub>ORS</sub>). Protein CarQ, a member of the ECF family of σ<sup>70</sup> factors, is required for transcription of *P*<sub>I</sub> and *P*<sub>ORS</sub>. The activity of those two promoters is very low in the dark because CarQ is sequestered by the antisigma factor CarR, a transmembrane protein. Upon illumination, CarR is somehow inactivated, and CarQ is freed from the membrane to activate its cognate promoters (Fontes et al., 1993; Hodgson, 1993; McGowan et al., 1993; Gorham et al., 1996; Martínez-Argudo et al., 1998; Browning et al., 2003). Activation of *P*<sub>ORS</sub> by light also requires the action of some non-specific, DNA architectural elements. One of these is IhfA, a histone-like protein, and the other is a heteromultimeric complex formed by protein CarD, bearing a DNA-binding domain that resembles eukaryotic high-mobility group A proteins, and CarG, a novel zinc-binding protein (Nicolás et al., 1994, 1996; Padmanabhan et al., 2001; Moreno et al., 2001; Galbis-Martínez et al., 2004; Peñalver-Mellado et al., 2006). The *P*<sub>B</sub> promoter is switched off in the dark by the binding to its operator of the negative regulator CarA, produced independently of light from an adjacent operon. Light-induced activation of *P*<sub>B</sub> involves disrupting the CarA-operator complex (López-Rubio et al., 2004). This is achieved by CarS, the product of
the third gene of the light-inducible carQRS operon. Protein CarS does not bind DNA, but physically interacts with CarA to dismantle the CarA-DNA complex (Ruiz-Vázquez et al., 1993; Whitworth & Hodgson, 2001; Cervantes & Murillo, 2002; López-Rubio et al., 2002; Pérez-Marin et al., 2004; Navarro-Avilés et al., 2007).

The third protein required early for the activation of both ctrl and the carB operon is the product of the unlinked carF gene. Strong genetic evidence indicates that CarF is the earliest acting of all known Car elements of the light-driven regulatory cascade, being absolutely necessary for the light-dependent inactivation of CarR (Fontes et al., 2003). Contrary to all other known Car regulatory proteins, CarF does not take part in the carotenogenic response to copper (Moraleda-Muñoz et al., 2005). CarF shows sequence similarity to members of a family of proteins named Kua. When first described, this family only included eukaryotic proteins (Fontes et al., 2003), but some prokaryotic members of the Kua family have now been identified (see below). At present, none of the Kua proteins has a known function.

In the present work, we have investigated the mode of action of protein CarF, with a focus on the subcellular localization of CarF and on whether the functional interaction with CarR uncovered by genetic data is mediated by a physical association between the two proteins. The results reported here strongly argue for CarF acting as a membrane-bound anti-antisigma factor and serve to determine a series of protein domains and histidine residues that are essential for normal CarF activity.

**METHODS**

**Bacterial strains, yeast strains and growth conditions.** The following *Escherichia coli* strains were used: DH5α (Hanahan, 1983) for cloning procedures, BL21 (DE3) for protein overexpression, and JM109 (Yanisch-Perron et al., 1985) and BTH101 (adenylate-cyclase deficient) for a bacterial two-hybrid system (BACTH; Karimova et al., 1998, 2000). The host *Saccharomyces cerevisiae* strains for the yeast two-hybrid systems were EGY48 (for the LexA-based system; Golemis & Serebriskii, 2001) and cdc25H (for the Cytotrap system; Stratagene). Yeast growth media and conditions were as described elsewhere (Ausubel et al., 1988). *M. xanthus* strain DK1050 (Ruiz-Vázquez & Murillo, 1984) was the wild-type strain used in this study. The *M. xanthus* carF2 deletion mutant MR992 (Fontes et al., 2003) was also used. The rich castonite-tris (CTT) medium and the exact culture conditions for growth of *M. xanthus* in the dark and in the light have been described previously (Bretscher & Kaiser, 1978; Fontes et al., 1993).

**Nucleic acid manipulations.** Standard protocols were followed for DNA manipulation (Sambrook & Russell, 2001). For DNA amplification (PCR) three different systems were used, depending on the aim of the experiment. For amplification of short DNA fragments, *Pfu* high-fidelity DNA polymerase (Promega) was employed. For diagnostic PCR (i.e. in the analysis of strains carrying deleted versions of carF), the Amersham Ready To Go PCR kit was used. For inverse PCR (i.e. in the generation of the Ser86–Ser139 internal deletion of carF), the Roche Long Template High-Fidelity PCR system was used. Reaction mixtures were supplemented with 5% (v/v) DMSO to overcome PCR difficulties due to the high G+C content of the template DNA. Each PCR-derived clone was verified by sequencing.

**Site-directed mutagenesis.** The technique of PCR overlap extension (Ho et al., 1989) was used for the site-directed substitution of CarF His164 and His195 residues by alanine (for details see Fontes et al., 2003). The mutagenic primers used were His164Fw (5’-GGCGACACCAACTGTTTGCGGACATG-3’) and His164Rev (5’-CATGTGGCCGACACTTTCGGAACCTGTTGTTGGC-3’) for the H164A substitution; and His195Fw (5’-CACACCACGACCATTACGC-GCCCAAGCCGTACAAC-3’) and His195Rev (5’-GCGA-TGTCGCGTGTTGGTGGAATTCGCGTGTTG-3’) for the H195A substitution (nucleotides that differ from the wild-type sequence are underlined).

The products from the overlap extension, done with primers that amplify the complete carF ORF, were cloned separately into the *M. xanthus* carF2 deletion mutant strain MR992 (Fontes et al., 2003). As the plasmid could not replicate in *M. xanthus*, KmR colonies arise from an
integrative, homologous recombination event between the \textit{M. xanthus} DNA in the pMAR979-derived plasmid and the homologous chromosomal region in the recipient strain. The correct integration of every construction in the genome was tested by PCR. To obtain \textit{carF} alleles that lacked the N- or the C-terminal region, the corresponding PCR fragments were inserted into the EcoRI site of pMAR979. To obtain a CarF version lacking the region between amino acids 86 and 139, we used inverse PCR to amplify all the region flanking the fragment to be deleted, using a plasmid with 7.7 kb of the wild-type \textit{carF} region as template. The plasmids were introduced into the MR992 \textit{carF} deletion mutant or into the wild-type strain as described above. Every plasmid was verified by sequencing.

\textbf{CarF overexpression and purification.} The \textit{carF} ORF was amplified by PCR from \textit{M. xanthus} genomic DNA and cloned into the NdeI and BamHI sites of pET11b to obtain plasmid pET11b-\textit{carF} (Studier et al., 1990). Cells transformed with pET11b-\textit{carF} were grown in 50 ml LB/ampicillin/glucose at 37 °C to an OD_{600} of 0.6–1.0, harvested by centrifugation, washed and inoculated into 11 of the same medium without glucose. After growing to an OD_{600} of 0.6–0.8, protein expression was induced with 0.4 mM IPTG overnight at 25 °C. SDS-PAGE analysis of cell extracts showed that CarF was expressed mainly as an insoluble protein. Cells from 1 l induced culture were pelleted, lysed by mechanical treatment with alumina at 4 °C and resuspended in 30 ml buffer A (50 mM Tris, 5 %, v/v, glycerol, 5 mM \textit{β}-mercaptoethanol, pH 7.5) containing 300 mM NaCl, 1 mM PMSF and 1 mM benzamidine. Resuspended cells were subjected to sonication under ice-cold conditions and the alumina was eliminated by low-speed centrifugation (500 g, 5 min, 4 °C). The supernatant was then centrifuged (12,000 \textit{g}, 15 min, 4 °C) to isolate inclusion bodies and the pellet was solubilized in denaturing buffer (8 M urea, 300 mM NaCl, 77 mM Na_{2}HPO_{4}, 22.6 mM NaH_{2}PO_{4}, pH 7.6) and insoluble material was eliminated by centrifugation (12,000 \textit{g}, 15 min, 4 °C). The supernatant was used for CarF purification by TALON (Clontech) metal affinity resin under denaturing conditions, followed by imidazole elution (150 mM). Plasmid pET11b is designed for the overexpression of native proteins, but the abundance of histidine residues in CarF enables CarF purification by using a nickel resin.

\textbf{Obtention of anti-CarF antibodies.} The denatured CarF protein was gel-purified and employed to raise a rabbit polyclonal antisera. Western blotting confirmed that the unpurified antisera recognized the pure CarF protein and the protein expressed in \textit{E. coli} cells carrying pET11b-\textit{carF}. No signal was detected when a preimmune antisera was used against the same samples (data not shown).

\textbf{Isolation and solubilization of membranes from \textit{E. coli}.} The pellet from a 250 ml culture of \textit{E. coli} transformed with pET11b-\textit{carF} was treated as described above for CarF purification. The resulting supernatant was then centrifuged at 90,000 \textit{g} for 45 min at 4 °C. The supernatant was retained, as this contained the cytoplasmic and periplasmic proteins (soluble fraction). The pellet (containing membranes) was washed with cold buffer A and resuspended in 4 ml of the same buffer with 1 % Triton X-100, 1 mM benzamidine and 1 % protease inhibitor mixture (Sigma). The mixture was incubated for 1 h at 4 °C with shaking and centrifuged at 90,000 \textit{g} as before, to separate the solubilized membrane (supernatant) from the non-solubilized membrane (pellet). The protein concentration of each fraction was determined using the Bio-Rad protein assay kit, as specified by the manufacturer. To detect CarF, 6 μg protein from each sample was subjected to Western analysis with anti-CarF antibodies (Cayuela et al., 2003).

\textbf{Sequence analysis and topological prediction.} For comparison of CarF with databases, the \textit{wu-BLASTP + BEAUTY} (http://searchlauncher.bcm.tmc.edu/seq-search/protein-search.html) and \textit{BLAST2-WU} (http://ebi.ac.uk/Tools/similarity.html) programs, provided by the BCM search launcher and EBI, respectively, were used. For the alignment of different peptides or proteins, the program \textit{CLUSTALW} 1.8 (http://searchlauncher.bcm.tmc.edu/multi-align/multi-align.html) from BCM was employed. For the prediction of transmembrane regions and CarF topology, the programs \textit{TMPPRED} and \textit{SOSUI} from the BCM search launcher (http://searchlauncher.bcm.tmc.edu/seq-search/struc-predict.html) and the programs \textit{TMHMMSS} (www.sbc.su.se/~melen/TMHMMSS/) and \textit{DAS} (www.sbc.su.se/~miklos/DAS/main.html) from the SBC server were used.

\textbf{Analysis of β-galactosidase.} Qualitative determination of β-galactosidase was performed on LB plates containing 40 μg X-Gal ml⁻¹. Quantitative analysis on liquid cultures was performed as described previously (Balsalobre et al., 1987). Enzyme-specific activities are given in nmol o-nitrophenol hydrolysed min⁻¹ (mg protein)⁻¹. For analysis of protein interactions with the BACTH system, overnight cultures, grown at 30 °C in LB medium with the appropriate antibiotics, were diluted and incubated up to OD_{550} ~ 0.5. At that point, IPTG (0.5 mM) was added and the cultures were incubated for 8 h at 30 °C.

\section*{RESULTS}

\textbf{CarF is located at the membrane in \textit{E. coli}.} Lack of the \textit{carF} gene product blocks the \textit{M. xanthus} carotenogenic response to blue light (\textit{Car}⁻ phenotype), whereas lack of the \textit{carR} gene product causes constitutive, light-independent carotenoid synthesis (\textit{Car}² phenotype). The fact that \textit{carF carR} double mutants show the same \textit{Car}² phenotype as the \textit{carR} single mutant strongly suggests that CarF participates in the light-mediated inactivation of CarR (Fontes et al., 2003; Fig. 1). CarR has been shown to be an integral membrane protein (Gorham et al., 1996; Browning et al., 2003), as may also be the case with CarF, since its amino acid sequence predicts at least three transmembrane domains (Fontes et al., 2003). To test this, we purified CarF from \textit{E. coli} cells carrying plasmid pET11b-\textit{carF}, and polyclonal anti-CarF antibodies were raised (see Methods). CarF was easily detected by Western analysis of the producing \textit{E. coli} strain, but could not be detected by the same analysis of the \textit{M. xanthus} wild-type strain DK1050, or the protease-deficient strain DK5077 (Petit & Guespin-Michel, 1992). To determine the subcellular localization of CarF in \textit{E. coli}, cell lysates were prepared from mid-exponential-phase cultures overexpressing CarF (see Methods). High-speed centrifugation was used to separate a soluble fraction (containing proteins from the cytoplasm and the periplasm) from an insoluble particulate fraction (containing cell membranes, wall components and ribosomes). The insoluble fraction was washed and incubated with 1 % Triton X-100. The solubilized membrane components were separated from the non-solubilized membrane fraction by high-speed centrifugation. As a control, the same procedure was performed with cultures of \textit{E. coli} carrying vector pET11b. Aliquots of each fraction (containing the same amount of total protein) were subjected to Western blotting and probed with polyclonal anti-CarF antibodies (Fig. 2). A band
of approximately 30 kDa (the expected size for CarF) was observed in the solubilized and non-solubilized membrane fractions. This indicates that CarF accumulates in the membrane and that it can be partially solubilized with 1% Triton X-100. The anti-CarF serum recognizes other E. coli proteins, probably due to impurities in the preparation of the CarF protein used to raise the antibodies (Fig. 2).

Membrane topology of CarF

Amino acid sequence analysis of CarF with different programs to predict transmembrane regions (TMHMM, SOSUI and DAS) suggests the presence of either three or four transmembrane domains in CarF. To assess the topology of CarF and the actual number of transmembrane domains, we used the pKY481 vector, which allows in-frame cloning of DNA fragments immediately upstream of the lacZ gene to make translational fusions (Cho & Zusman, 1999). Five different M. xanthus DNA fragments, each encoding a CarF protein containing one, two, three or four putative transmembrane domains, or the whole protein, were cloned into pKY481 (Fig. 3). β-Galactosidase is active only in the cytoplasm, as it is inhibited by the high pH of the periplasm (Gött & Boos, 1988). Thus, a positive result in the assay for β-galactosidase activity would indicate that the C terminus of a particular protein fusion faces the cytoplasm.

Previous experiments had revealed that the carF promoter is not active in E. coli. Thus, to express the translational CarF–LacZ fusions, we employed the promoter of the M. xanthus gene phoR1 that yields measurable activity in E. coli (Carrero-Lerida et al., 2005). A 781 bp fragment containing the phoR1 promoter was amplified by PCR from genomic M. xanthus DNA and cloned upstream of every one of the carF–lacZ fusions. The five constructions, and the vector pKY481 alone, were separately transferred into E. coli. The results of the β-galactosidase assays (Fig. 3) indicate that CarF might indeed contain four transmembrane domains. The first and third transmembrane domains of CarF should be oriented towards the periplasmic space, as LacZ fusions at amino acids 56 and 142 yielded only basal β-galactosidase activity. On the contrary, the second and fourth transmembrane motifs should be oriented to the cytoplasm, as fusions to amino acids 83 and 170 produced significantly higher β-galactosidase activities. It should be noticed that the β-galactosidase activity found when the entire CarF protein was fused to LacZ is similar to the activity obtained by the fusion to the fourth transmembrane motif.

Analysis of CarF–CarR interaction using yeast-based two-hybrid systems

As previously mentioned, genetic data points to CarF being involved in the light-driven inactivation of CarR (Fontes et al., 2003). The functional interaction between CarF and CarR might be mediated by a physical association between the two proteins. So, the yeast two-hybrid system described by Golemis & Serebriiskii (2001) was tried to assay a possible CarF–CarR physical interaction. Both CarF and CarR were used either as a ‘bait’ (fused to the DNA-binding domain of LexA) or as a ‘target’ (fused to the yeast activation domain B42) in independent experiments. In neither case did the bait constructions enter the nucleus of the yeast cell, probably due to the hydrophobic nature of both CarF and CarR. We then tested different, non-hydrophobic segments of CarF and CarR, but none of the bait fusions entered the yeast nucleus, maybe due to the instability of the peptides and/or their degradation by yeast proteases.

We also tested the putative interaction between CarF and CarR by employing the Cytotrap yeast two-hybrid system (Stratagene). In this system, one protein fusion serves to anchor a ‘target’ protein to the membrane. Upon interaction, the second protein fusion (‘bait’) is brought to the membrane to activate the Ras signalling pathway. In our case, both CarF and CarR, when used as ‘baits’, activated Ras signalling, even in the absence of the ‘target’ protein fusion.

Analysis of protein interactions using a BACTH system

Due to the restrictions of yeast two-hybrid systems to detect interaction between hydrophobic proteins, we employed a BACTH system (Karimova et al., 1998). This
system is based on the reconstitution of the cAMP signal transduction pathway in an E. coli adenylate-cyclase-deficient (cya−) strain. It takes advantage of the modular structure of the catalytic domain of the Bordetella pertussis adenylate cyclase, which consists of two complementary fragments (T25 and T18). When separately expressed in E. coli, they do not form an active enzyme, unless interacting polypeptides are genetically fused to both fragments. Restoration of enzymic activity results in cAMP synthesis to corroborate the CarR–CarR interaction. The LacZ+ Mal+ phenotype and the β-galactosidase specific activities generated by the appropriate constructions revealed that, as for CarR, different molecules of CarF can interact with each other (Fig. 4b and c).

The stoichiometry of CarQ and CarR is critical for the regulation of carQRS expression. This and other data suggest that, in the dark, CarQ and CarR might form a complex in the membrane (Martinez-Laborda & Murillo, 1989; Gorham et al., 1996; Browning et al., 2003). To test for CarQ–CarR interaction, Browning et al. (2003) used a yeast two-hybrid system. They could detect interaction only between some fragments of CarQ and CarR, but not between intact CarQ and CarR proteins. This could be due to the limitations of the yeast two-hybrid system for testing hydrophobic proteins. We have now used the BACTH system to test for interaction between intact CarQ and CarR. The carQ ORF was amplified by PCR and separately cloned into vectors pUT18C and pKT25. The resulting plasmids (pUT18CarQ and pKT25carQ) were separately introduced into E. coli, in combination with pUT18carR, pUT18CarR or pKT25carR. The LacZ+ Mal+ phenotypes and the β-galactosidase specific activities of the resulting transformants revealed that intact CarQ and CarR can interact with each other. The interaction was particularly strong when the CarQ C-terminal end remained free (Fig. 4d). The CarQ–CarR interaction in E. coli was also light-independent, as similar β-galactosidase specific activities were observed in cultures grown in the dark or in the light (data not shown).

Finally, the BACTH system was also used to test for a possible direct interaction between CarF and CarQ. No sign of interaction was detected in transformants carrying the pairs of plasmids pUT18CarQ+pKT25carF, pUT18carF+pKT25carQ or pUT18CarQ+pKT25carQ (data not shown).
CarF truncated versions

Comparative analysis of the predicted CarF protein sequence (281 aa) with other proteins in the databases shows that CarF shares strong similarity to members of a family of proteins, known as Kua, whose function is unknown (Fontes et al., 2003). These proteins were originally found only in eukaryotic organisms, but some prokaryotic members have been described recently (see supplementary Fig. S1 available with the online version of this paper). Most Kua proteins share several conserved domains, located between hydrophobic regions, whose importance was studied by generating truncated versions of carF and analysing their ability to phenotypically complement the deletion of carF (Fig. 5).

Three fragments of the carF gene were amplified by PCR and cloned into the EcoRI site of vector pMAR979, which carries a deletion of the entire carF ORF (Fontes et al., 2003; see Methods). The resulting constructions contained one N-terminal deletion (Met1–Ala25) and two C-terminal deletions (Val160–Pro281 and Gly241–Pro281). In addition, an internal deletion (from Ser86 to Ser139) was constructed by inverse PCR (Fig. 5; see Methods for details). The small C-terminal region deleted in pMAR2742 is not conserved in other Kua proteins, neither does it show similarity to other proteins or protein domains in the databases. The N-terminal region deleted in plasmid pMAR2728 shows a moderate conservation among Kua proteins. The large C-terminal region deleted in pMAR2731 and the internal region deleted in pMAR2727 contain sequences highly conserved in Kua proteins.

All indicated carF deletions were separately introduced into the M. xanthus carF mutant strain MR992, and more than 100 independent colonies from each transformation were tested for their Car phenotype. All colonies showed a Car− phenotype, except for those that had integrated the CarF version lacking the C-terminal 41 amino acids (Fig. 5). The Car+ phenotype of these latter colonies was indistinguishable from that of the M. xanthus wild-type strain DK1050.

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**Fig. 4.** Analysis of protein interactions using the BACTH system. The bait and target constructions employed for the interaction assays and their β-galactosidase activities [in nmol o-nitrophenol hydrolysed min⁻¹ (mg protein)⁻¹] are shown. The values of the self-activation controls are also shown. Data are means ± SD of at least three independent experiments.
Therefore, truncation of the fragments Met1–Ala25, Val160–Pro281 and Ser86–Ser139 affects CarF activity, conformation or stability, whereas the last 41 amino acids of CarF are not essential for the correct function of the protein. When the truncated versions of CarF were introduced into the wild-type strain DK1050, all the tested transformants accumulated carotenoids in the light like the parental strain. The dominance of the wild-type carF allele over the mutated ones indicates that none of the truncated CarF proteins can interfere with the normal action of wild-type CarF.

**Substitution of conserved His to Ala residues**

Comparison of the amino acid sequence of CarF with other Kua proteins shows a remarkable conservation of several His residues (supplementary Fig. S1). Eight of the 12 His residues of CarF are present, at the same relative position, in all Kua proteins. Three of the other His residues are present only in the animal Kua proteins, some of the prokaryotic ones, and that from *Leishmania major*. The remaining His residue is found exclusively in CarF. Previous experiments had shown that three of those conserved residues (His78, His103 and His113) are essential for the normal activity of CarF, while the non-conserved His218 is not (Fontes *et al.*, 2003).

We have extended the analysis of the functional relevance of CarF His residues by making His to Ala substitutions at positions 164 and 195, which are conserved in all Kua proteins (supplementary Fig. S1). Site-directed mutagenesis was performed by PCR overlap extension (see Methods) and the plasmids carrying the *carF* gene with the desired substitutions were transformed separately into the *M. xanthus* carF deletion mutant MR992. In each case, more than 100 transformants were tested for the colour phenotype. All the colonies carrying the H164A or H195A substitutions showed the same Car− phenotype as the parental strain MR992. Thus, the conserved His164 and His195 residues are essential for the normal activity of CarF, as shown with the previously tested conserved residues His78, His103 and His113. Substitutions H164A and H195A were also introduced separately into the *M. xanthus* wild-type strain. In both cases, the transformants were Car+, except for a few colonies (1%) that showed a Car− phenotype, probably the consequence of gene conversion (Stephens & Kaiser, 1987; Martinez-Laborda & Murillo, 1989).

**DISCUSSION**

Cells of the bacterium *M. xanthus* are known to accumulate carotenoid pigments only when stimulated by exposure to blue light or copper. Both responses depend on the activation at the transcriptional level of the structural genes for carotenoid synthesis. Our present model on how this is accomplished, based on many genetic and biochemical studies (see references in the Introduction), is summarized in Fig. 1. Both light- and copper-induced activation of the carotenogenic genes depend crucially on the inactivation of CarR, an antisigma factor that in the dark sequesters the ECF-sigma factor CarQ to the membrane. At present, nothing is known on how exposure to copper leads to inactivation of CarR. As for the light response, we have previously reported strong genetic evidence for CarR inactivation requiring the product of the *M. xanthus* gene *carF*, the only known element of the carotenogenic regulatory cascade that does not participate in the response to copper (Fontes *et al.*, 2003; Moraleda-Muñoz *et al.*, 2005).

To detect CarF, we raised anti-CarF antibodies. Unfortunately, we could not detect CarF by Western analysis of *M. xanthus* cell extracts. The low activity of the *carF* promoter may have contributed to our failure to detect CarF in *M. xanthus* cell extracts. The low activity of the *carF* promoter may have contributed to our failure to detect CarF in *M. xanthus* cell extracts, as we could easily detect the protein in the overproducing *E. coli* strain. A similar situation has been reported precisely for the *M. xanthus* antisigma factor CarR, which could not be detected by Western analysis neither in the wild-type nor in a strain in which expression of CarR was sixfold greater than in the wild-type (Browning *et al.*, 2003).
In *silico* sequence analysis predicts the presence in CarF of several transmembrane domains (Fontes *et al*., 2003; see also supplementary Fig. S1). Two lines of evidence presented in this paper confirm that CarF is a membrane protein. On the one hand, when expressed in *E. coli*, CarF accumulates almost exclusively in the membrane, from where it can be only partially solubilized with the detergent Triton X-100 (Fig. 2). On the other hand, a lacZ-based topological test for potential transmembrane domains detects four transmembrane domains in CarF, the first and third being oriented towards the periplasm, and the second and fourth towards the cytoplasm (Fig. 3). The fact that CarF alone, like CarR, led to Ras signalling when used as bait in the Cytotrap two-hybrid system (Stratagene) further confirms its membrane localization.

As mentioned before, CarR has been shown to be a membrane protein that sequesters the ECF-sigma factor CarQ in the dark. A previous study of CarR–CarQ interaction, using a yeast two-hybrid system that requires the fusion proteins to enter the nucleus, gave positive results only with some fragments of the proteins involved (Gorham *et al*., 1996; Browning *et al*., 2003). Our own results with the Cytotrap system mentioned above corroborate that CarR goes to the membrane when expressed in yeast cells. To test for protein interactions, we have used a BACTH system (Karimova *et al*., 1998, 2000) that, in principle, should work well with membrane proteins. As shown in Fig. 4(a, d), the bacterial test produced unequivocal evidence that CarF interacts physically with CarR, and corroborated that complete CarR and CarQ also interact physically with each other. Other results of the bacterial test indicate that CarF can form homomultimers (Fig. 4b), as does CarR (Fig. 4c). It should be noted that no interaction between CarF and CarQ was observed in the bacterial test.

Considering previous genetic data on the role of CarF in the light-activated regulatory cascade of *M. xanthus* (Fontes *et al*., 2003) and its physical interaction with CarR uncovered in this work, we propose that CarF is in fact an anti-antisigma factor for CarR. The interplay of the *M. xanthus* proteins CarF, CarR and CarQ reminds one of the *Bacillus subtilis* SpoIIAA, SpoIIAB and σ^F^ proteins, which have been subjected to extensive genetic and molecular analysis (for a review see Yudkin & Clarkson, 2005). In this case, a dimer of SpoIIAB binds a single molecule of σ^F^ and prevents it from acting as a transcription factor. During the process of sporulation, SpoIIAA binds the σ^F^-SpoIIAB^2^ complex and causes the release (activation) of σ^F^. As appears to happen with CarF, SpoIIAA does not interact physically with its cognate sigma factor, but docks on the σ^F^-SpoIIAB^2^ complex by binding a subunit of SpoIIA that is not in contact with σ^F^ (Ho *et al*., 2003).

Sequence comparison analysis gave no hint of what the molecular mechanism for CarF might be, and how it could be modulated by the light signal. When first described, CarF was reported to be very similar to a family of proteins of unknown function, named Kua, found only in eukaryotes (Thomson *et al*., 2000; Fontes *et al*., 2003). In a recent search, we have identified Kua-like proteins, also of unknown function, in prokaryotes and even in a virus (see supplementary Fig. S1). Transmembrane domains similar to those found in CarF are predicted for most Kua proteins. As seen in Fig. 6 and supplementary Fig. S1, the most conserved amino acid residues are grouped in two regions, one between the second and third transmembrane domains and another at the C terminus, following the fourth transmembrane domain. The importance of these conserved regions for the action, or stability, of CarF has been corroborated here by the inability of various truncated versions of CarF to complement a carF deletion. Singular to CarF among the Kua proteins is a C-terminal extension of about 40 aa. These appear to be dispensable, as deleting them does not affect CarF function in a significant way (Fig. 5).

Previous theoretical analysis revealed some similarities, but no overall sequence similarity, between Kua proteins and fatty acid desaturases and hydroxylases (Los & Murata, 1998; Thomson *et al*., 2000; Fontes *et al*., 2003). These enzymes contain two transmembrane domains that make the protein cross the membrane twice, and eight His residues, located at cytoplasmic domains, all of which play a critical role in the reaction mechanism (Mitchell & Martin, 1997; Shanklin *et al*., 1994). The topological assays carried out in this work show that CarF contains four transmembrane domains instead of two. The four transmembrane domains could be grouped in two pairs. In each one of them, one helix would face the periplasm and the other, only a few residues apart, would face the cytoplasm. Prominent among the conserved Kua residues are eight His residues that are found in all members of the family.
(supplementary Fig. S1), and they, as for the enzymes mentioned above, would be located at the cytoplasm (Fig. 6). All single His to Ala substitutions at five of the eight conserved His residues (His164 and His195, reported here, and His78, His103 and His113, reported by Fontes et al., 2003) resulted in the inactivation of CarF. For one of the hydroxylases mentioned above, it has been shown that the multiple His residues serve to form a catalytically essential di-iron cluster (Shanklin et al., 1997). The ability of CarF to bind transition metals is revealed by its ability to adsorb, in untagged form, Co^{2+} resins. However, as shown with other membrane proteins, purification of CarF in its native form has proven problematic. Certainly, more work will be needed to search for a possible prosthetic group in CarF and to unravel its possible role in triggering its anti-antisigma activity under blue light.

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