

Light-Induced Carotenogenesis in *Streptomyces coelicolor* A3(2): Identification of an Extracytoplasmic Function Sigma Factor That Directs Photodependent Transcription of the Carotenoid Biosynthesis Gene Cluster

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Carotenoids are produced by a variety of organisms, but the mechanisms that regulate gene expression leading to carotenoid biosynthesis have been characterized for only a few organisms. In this study, we found that *Streptomyces coelicolor* A3(2), a gram-positive filamentous bacterium, produces carotenoids under blue light induction. The carotenoid fraction isolated from the cell extract contained multiple compounds, including isorenieratene and β -carotene. The carotenoid biosynthesis gene cluster of *S. coelicolor* consists of two convergent operons, *crtEIBV* and *crtYTU*, as previously shown for *Streptomyces griseus*. The *crtEIBV* null mutant completely lost its ability to produce carotenoids. The *crt* gene cluster is flanked by a regulatory region that consists of two divergent operons, *litRQ* and *litSAB*. The *lit* (light-induced transcription) genes encode a MerR-type transcriptional regulator (LitR), a possible oxidoreductase (LitQ), an extracytoplasmic function sigma factor (σ^{LitS}), a putative lipoprotein (LitA), and a putative anti-sigma factor (LitB). S1 protection assay revealed that the promoters preceding *crtE* (*PcrtE*), *crtY* (*PcrtY*), *litR* (*PlitR*), and *litS* (*PlitS*) are activated upon illumination. A *litS* mutant lost both the ability to produce carotenoids and the activities of *PcrtE*, *PcrtY*, and *PlitS*, which suggested that σ^{LitS} directs light-induced transcription from these promoters. An RNA polymerase holo complex containing purified σ^{LitS} recombinant protein generated specific *PcrtE* and *PcrtY* transcripts in an in vitro runoff transcriptional assay. A *litR* mutant that had an insertion of the kanamycin resistance gene was defective both in the ability to produce carotenoids and in all of the light-dependent promoter activities. Overexpression of *litS* resulted in constitutive carotenoid production in both the wild type and the *litR* mutant. These results indicate that σ^{LitS} acts as a light-induced sigma factor that directs transcription of the *crt* biosynthesis gene cluster, whose activity is controlled by an unknown LitR function. This is the first report to describe light-inducible gene expression in *Streptomyces*.

Carotenoids are pigmented compounds that are widely produced by both eucaryotes and procaryotes (11). They are tetraterpenoids that consist of a polyene hydrocarbon chain derived from eight isoprene units. The C_{40} backbone is modified in several ways, such as cyclization and desaturation, to produce a variety of compounds with divergent chemical structures. Carotenoids have two major roles within the cell. First, in photoautotrophic organisms, they act as accessory pigments in the light-capturing complexes by absorbing light in the wavelength range of 400 to 500 nm and transferring the energy to chlorophyll. Second, in both phototrophic and nonphotoautotrophic organisms, they protect cells from photo-oxidative damage by scavenging harmful agents such as singlet and triplet molecular species produced upon illumination.

Carotenogenesis in prokaryotes is constitutive or photoinducible. Several prokaryotes, including *Erwinia herbicola* (1) and *Rhodobacter capsulatus* (3) produce carotenoids constitutively whereas organisms such as *Myxococcus xanthus* (9), *Flavobacterium dehydrogenans* (28), and *Sulfolobus* spp. (12) produce carotenoids in a photoinducible manner. The control mechanisms of carotenogenesis have been studied in phototro-

phic bacteria such as *Rhodobacter* spp., which has revealed the involvement of global signal transduction initiated by light capture in the bacteriochlorophyll (2). On the other hand, the molecular mechanism in nonphototrophic bacteria has not yet been fully studied except in *M. xanthus*, a gram-negative gliding bacterium characterized by a unique life cycle. In *M. xanthus*, extensive studies have revealed a major signaling pathway from light sensing to transcriptional control of biosynthetic genes, which involves a light-dependent extracytoplasmic function (ECF) sigma factor, CarQ, and its cognate membrane-associated anti-sigma factor, CarR (6, 8, 10, 23, 29).

We study carotenogenesis in *Streptomyces*, a gram-positive bacterial genus renowned for its ability to produce a variety of secondary metabolites. Although it is well known that the carotenoid-producing ability is widespread in this group of bacteria, the molecular mechanism involved in the regulation of carotenoid production is poorly characterized. We have found that carotenogenesis is light inducible in *Streptomyces coelicolor* A3(2), the model organism for genetic analysis. While the information from genomic sequence data (5) has already shown that this organism retains a carotenoid biosynthesis gene cluster, until now this has not been demonstrated to lead to carotenoid production. In normal strains of *S. coelicolor* A3(2), it is not clear whether the organisms produce carotenoids, since the marked production of actinorhodin and prodigiosin, the blue diffusible and red intracellular pigment an-

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tibiotics, respectively, prevents clear observation of the yellow pigment. Meanwhile, we noticed that an *S. coelicolor* strain harboring pKM284, a plasmid that has an activity which represses the production of the two pigmented antibiotics, accumulated the yellow pigment in the cell when it was illuminated with visible light. We also found that the photodependent carotenoid production occurs in the parental *S. coelicolor* strain. Here, we report the light-induced transcription of the carotenoid biosynthesis gene cluster of this organism along with the identification of a responsible sigma factor and related regulators.

MATERIALS AND METHODS

Bacterial strains, plasmids, and media. *S. coelicolor* A3(2) M145 (wild type), obtained from the John Innes Institute (Norwich, United Kingdom), was used as a parental strain and designated the wild type. The isolation of pKM284, the plasmid that repressed antibiotic production in wild-type *S. coelicolor*, was described previously (27). *Escherichia coli* JM109 (24) and BL21(DE3) (Novagen) were used as hosts for DNA manipulation and expression of recombinant proteins. pUC19 (30) was used for general DNA manipulation. pT7Blue (Novagen) was used for TA cloning of PCR products. pGEX4T-2 (Pharmacia) and pET26b(+) (Novagen) were used as vectors for effective expression of recombinant proteins. The conditions used for genetic manipulation in *E. coli* and *Streptomyces* have been described by Maniatis et al. (24) and Kieser et al. (18), respectively. All vector plasmids used were described by Kieser et al. (18). Cosmid StJ12, which contains the *crt* locus, was obtained from H. Kieser (John Innes Institute). *S. coelicolor* A3(2) strains were grown in Bennett's sugar medium (containing [in grams per liter] the following: yeast extract [Difco Laboratories, Detroit, Mich.], 1; meat extract [Kyokuto, Tokyo, Japan], 1; NZ amine [Wako Pure Chemical Industries, Ltd., Tokyo, Japan], 2; and an appropriate sugar [Kokusan, Tokyo, Japan], 10 [pH 7.2]) and YMP-sugar medium (containing [in grams per liter] the following: yeast extract [Difco], 2; meat extract [Kyokuto], 2; Bacto Peptone [Difco], 4; NaCl, 5; $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, 2; and an appropriate sugar [Kokusan], 10 [pH 7.2]). Solid media contained 1.5% agar (Kokusan). In order to select *E. coli* transformants, ampicillin (Wako) and kanamycin (Wako) were used at a final concentration of 50 $\mu\text{g}/\text{ml}$, and for selecting *S. coelicolor* A3(2) transformants, thiostrepton (Sigma Chemical Co., St. Louis, Mo.) and kanamycin were added to a final concentration of 20 $\mu\text{g}/\text{ml}$. Enzymes used for genetic manipulation were purchased from Takara-shuzo (Kyoto, Japan).

Production, extraction, and identification of carotenoids. In order to collect cells, *S. coelicolor* A3(2) strains were grown on cellophane on the surface of Bennett's glucose solid medium for 5 days at 28°C under light or dark conditions in a darkroom equipped with white light fluorescent lamps (Toshiba) (15 W). Under light conditions, white light was illuminated at approximately 2.4 $\mu\text{mol s}^{-1} \text{ m}^{-2}$ onto the solid culture. The same lamp, covered with a blue or red light filter, was used for illumination of blue (maximum transmittance of between 400 and 460 nm) or red (600 to 700 nm) light, respectively.

The carotenoid fraction was extracted by the method described by Schumann et al. (25). Cells collected from five agar plates were first lyophilized and then suspended in 10 ml of chloroform and incubated at 60°C for 15 min with occasional vortex mixing. The sample was then placed on ice for 5 min, filtered to remove cell debris, and evaporated. After drying, the sample was redissolved in 10 μl of chloroform. The concentrated preparation was subsequently purified on a silica gel column (Bond Elut; Varian) and eluted with hexane-toluene (85:15), according to the manufacturer's protocol, in order to obtain the final carotenoid fraction.

The resulting carotenoid fraction was subjected to photometric analysis with a UV spectrometer (UVmini-1240; Shimadzu). Qualitative analyses were performed by silica gel thin-layer chromatography with a solvent mixture of methanol-acetonitrile-2-propanol (85:10:5) for development and by silica gel column chromatography with high-performance liquid chromatography (HPLC) (LC-10; Shimadzu), as described by Schumann et al. (25). An authentic β -carotene sample was purchased from Funakoshi (Tokyo, Japan) and used as the standard control. Isorenieratene was isolated from cell extracts of *Mycobacterium phlei* JCM5865 as described previously (21).

S1 nuclease mapping. Transcriptional activities of the promoters preceding *crtE* (*PertE*), *crtY* (*PertY*), *litQ* (*PlitQ*), *litR* (*PlitR*), and *litS* (*PlitS*) were examined by S1 protection assays. Preparation of RNA from cells grown on cellophane on the surface of agar medium and S1 nuclease mapping were performed as de-

scribed by Kelemen et al. (17). Hybridization probes were prepared by PCR with various oligonucleotide primers (Table 1): PEF and PER (*PertE*, low resolution), PEF and PERH (*PertE*, high resolution), PYF and PYR (*PertY*, low resolution), PYF and PYRH (*PertY*, high resolution), PQF and PQR (*PlitQ*, low resolution), PQF and PQRH (*PlitQ*, high resolution), PRF and PRR (*PlitR*, low resolution), PRF and PRRH (*PlitR*, high resolution), PSF and PFR (*PlitS*, low resolution), PFF and PFRH (*PlitS*, high resolution), PAF and PAR (*litA*, readthrough), and PBF and PBR (*litB*, readthrough). Primers PRF and aph were used to generate the hybridization probe for investigating *PlitR* activity in the *litR* mutant that carries an *aphII* gene cassette downstream of the promoter region. In all probes, the downstream primers were labeled at the 5' ends with [γ - ^{32}P]ATP by using T4 polynucleotide kinase. S1-protected fragments were analyzed on 6% polyacrylamide gels. Maxam-Gilbert sequence ladders prepared from the labeled hybridization probe were used as standards for high-resolution analysis. Protected fragments were analyzed on a 6% polyacrylamide gel. The quality of RNA used for low-resolution analysis was checked by a control assay for *hrdB*, encoding a major sigma factor, using a probe described previously (19).

Gene disruption and overexpression. The kanamycin-resistant *crtEIBV*, *litR*, *litAB*, and *litB* mutants were generated by standard homologous recombination techniques that exchanged the wild-type allele with a mutated construct on a disruption plasmid. In order to generate the *crtEIBV* null mutant, the flanking DNA fragments were PCR amplified with the primers DcrtF-DcrtMR and DcrtMF-DcrtR (Table 1), and these fragments were digested with HindIII-BglII and BglII-EcoRI, respectively, and inserted between the EcoRI and HindIII sites of pUC19 by three-fragment ligation. The resulting plasmid was cleaved with BglII and ligated to a 0.9-kb *aphII* (kanamycin resistance) (4) cassette to generate the disruption plasmid. This construction replaced the entire *crtEIBV* region with *aphII*, which is transcribed in the opposite direction to *crtEIBV*. Similarly, each of the two DNA fragments prepared with primers DRF-DRMR and DRMF-DRR (for disruption of *litR*), DSABF-DSABMR and DBMF-DBR (for disruption of *litAB*), and DBF-DBMR and DBMF-DBR (for disruption of *litB*) was processed to generate kanamycin-resistant disruption plasmids. The disruption plasmids were introduced into *S. coelicolor* A3(2) wild-type cells by standard transformation; subsequently, we screened for kanamycin-resistant colonies. True recombination was confirmed by Southern hybridization. The resultant kanamycin-resistant disruptants carried an *aphII* cassette in each *lit* coding sequence in the opposite orientation (*litR*) and the same orientation (*litAB* and *litB*) as the disrupted *lit* gene. We employed REDIRECT technology for markerless disruption of *litS*; primers DSF and DSR were used to amplify the *litS* coding sequence from the cosmid StJ12, which was then processed as described by Gust et al. (13) to generate a markerless *litS* mutant. The resultant *litS* mutant lacked the entire *litS* coding sequence.

In order to overexpress LitS and LitR, the DNA fragments were amplified with primers HSF-HSR and HRF-HRR, respectively, and these fragments were digested with SphI and BglII and ligated between the SphI and BglII sites of pIJ702 (16). The resulting plasmids carried the promoterless coding sequences of *litR* and *litS* downstream of the *mel* promoter and in the same orientation, such that the promoter directs constitutive transcription of these genes.

Expression and purification of LitS by an *E. coli* host-vector system. The coding sequence of *litS* was amplified with primers rSF and rSR (Table 1) and cloned between the NdeI and HindIII sites of pET26b. This plasmid construct directed the expression of σ^{LitS} as a fusion protein with a C-terminal histidine tag in *E. coli* BL21(DE3). The *E. coli* cells harboring the expression plasmid were aerobically cultured at 28°C in 100 ml of Luria-Bertani liquid medium (24), and when the optical density at 600 nm reached 0.8, IPTG (isopropyl- β -D-thiogalactopyranoside) (1 mM) was added. Following IPTG addition, the cells were further grown for 4 h and then harvested by centrifugation. The resulting cell pellet was resuspended in an appropriate volume of phosphate-buffered saline buffer (24) and disrupted by sonication. The soluble recombinant proteins were purified from the cell extract by Ni affinity chromatography according to the manufacturer's instructions.

In vitro runoff transcription. An in vitro runoff transcription assay was performed by a previously described method (26). DNA fragments that contained the *PertE*, *PertY*, and *PlitS* regions were used as templates. These fragments were PCR amplified with the primers PEF and PER (*PertE*), PYF and PYR (*PertY*), and PSF and PSR (*PlitS*). These PCR products were mixed with the *E. coli* RNA polymerase core enzyme (AR Brown) and purified σ^{LitS} protein, which was prepared as described above. The transcripts that were generated were analyzed by polyacrylamide gel electrophoresis. A 100-bp ladder marker (Takara-shuzo), denatured by heat treatment, was used as a standard to estimate transcript sizes.

TABLE 1. Oligonucleotide primers used in this study

Primer	Sequence (5'→3') ^a	Restriction enzyme	Position (nucleotides) ^b
PEF	5'-AGCTGCTGGCCGGCAATCAGC		173554–173574
PER	5'-TCGACCGCGTTCGCCGCATCG		173886–173866
PERH	5'-ACCATCGGCGGCAGTCCAGG		173823–173804
PYF	5'-TACGTGTGCGGAGACCACCGG		182413–182393
PYR	5'-ACACTCTTCGGTTCCACGCTTC		182194–182215
PYRH	5'-TGCCACCGGGTTCACGCACAC		182178–182197
PQF	5'-TGACCCTGGGCCCTGGATGG		183896–183877
PQR	5'-TGATGCCAGGGGCAAAGG		183587–183605
PQRH	5'-TGCGAGTGCAGGGGAAACG		183675–183692
PRF	5'-TCGTGTACGCGTGTTCATGC		184975–184956
PRR	5'-ACCTCGCCGGTGGTCAATCC		184718–184737
PRRH	5'-TCGTGCGACCGGCTCGTCG		184757–184774
PSF	5'-ACTATACGACGCACAAACGACG		184804–184825
PSR	5'-TTCTCTCTGCCGTGCGTCGTACG		184952–184930
PSRH	5'-ACGTCGCCGGTCTACCTGTC		184901–184882
PAF	5'-TTCTACGAGGACCTGACGCAGG		185426–185447
PAR	5'-TTGGCGAAGACGTCGACCGTCG		185835–185814
PBF	5'-ACCAGACTCTGGCTCTGCCCG		186213–186233
PBR	5'-TCGCCACTGGCGTGACCGAGTGC		186541–186519
PhrdF	5'-CGGCCGCAAGGTACGAGTTGATGA		78385–78408
PhrdR	5'-GCCATGACAGAGACGGACTCGGCG		78726–78703
aph	5'-TGATATTGCTGAAGAGCTTGG		2228–2248
rSF	5'-GGAACGCATATGAACACCCGTACCCGCCCGAC	NdeI	184949–184980
rSR	5'-GGCCAAGCTTGACACACGTGTC	HindIII	185573–185552
DcrtF	5'-CTGGAAGCTTTACTGGTACT	HindIII	173081–173099
DcrtMR	5'-ATCAGATCTTCGTGGACCAGAGC	Bg/II	174132–174110
DcrtMF	5'-TTTAGATCTCGACGTCTCCTTCC	Bg/II	178146–178168
DcrtR	5'-CGCGGAATTCTGCTGATGTTCC	EcoRI	179422–179400
DRF	5'-CGTCGAATTCCTGCCGCCGAAG	EcoRI	186262–186241
DRMR	5'-TACGAGATCTATGCCCACTATAC	Bg/II	184787–184810
DRMF	5'-CCCGAGATCTGTTGTCGTCC	Bg/II	183799–183780
DRR	5'-GCACAAGCTTCAGGATGGCTCCG	HindIII	182651–182672
DSF	5'-CCCGTCCCCGTACGACGCACGGCAGGAGGAACGGGCATGATTCGGGGATCCGTCGACC		184922–184960
DSR	5'-GTGGATGCGCCCCGGGGCGTCTCTGCTCCGCCCGGTCATGTAGGCTGGAGCTGCTTC		185602–185564
DBF	5'-CCGGGAATTCACGAGGTACTC	EcoRI	185336–185356
DBMR	5'-GTGAAGATCTGCGCCCTCCTG	Bg/II	186491–186471
DBMF	5'-GAACAGATCTCCGAGGTCCG	Bg/II	186881–186900
DBR	5'-GAGGAAGCTTAGTACGTCCTCC	HindIII	188594–188573
DSABF	5'-CATAAAGCTTCTCACTCGACG	HindIII	183557–183577
DSABMR	5'-TACGAGATCTTCAGCCCGTTCC	Bg/II	184970–184952
HRF	5'-GTGGGCATGCACGCGTACACG	SphI	184805–184783
HRR	5'-GACAAGATCTCACGGGTACACC	Bg/II	183784–183805
HSF	5'-ACGGGCATGCACACGCGTAC	SphI	184952–184971
HSR	5'-TGGGAGATCTCGAAGGTGTTTCG	Bg/II	185640–185618

^a Restriction sites are underlined.^b All position numbers except those for primer aph correspond to the numbering in the *S. coelicolor* genomic sequence (http://www.sanger.ac.uk/Projects/S_coelicolor). For aph, the numbering corresponds to the description given by Beck et al.(4).

RESULTS

Characterization of photodependent carotenogenesis in *S. coelicolor* A3(2). During our study on the effect of pKM284, which was previously isolated by its repression of aerial growth of *Streptomyces griseus* (27), we noticed that the introduction of this plasmid into *S. coelicolor* A3(2) abolished the production of actinorhodin and prodigiosin, the pigmented antibiotics usually produced by this organism. The pigmented-antibiotic-deficient transformant in turn clearly exhibited the accumulation of yellow pigment only when it was illuminated (Fig. 1A). The yellow color and light induction suggested that the pigment was a carotenoid(s).

Subsequently, we found that the pigment production also occurs in wild-type *S. coelicolor*. Spectroscopic analysis of the carotenoid fraction extracted from the wild-type cells showed

that the strain accumulates the pigment upon cultivation under light conditions, while it produced no yellow pigment under dark conditions (Fig. 1B). Marked production of the pigment occurred after 2 days of growth under illuminated conditions. The production profile in the wild-type strain was the same as that in the antibiotic-deficient strain described above. This indicated that regulation of light-induced carotenoid production is independent of the effect of the introduction of pKM284. We further examined carotenoid production in the wild-type strain.

S. coelicolor produced carotenoids when illuminated with blue light ($\lambda = 400$ to 460 nm) but not when illuminated with red light ($\lambda = 600$ to 700 nm). Thin-layer chromatography analysis of the carotenoid fraction extracted from the wild-type cells illuminated with blue light revealed the presence of at

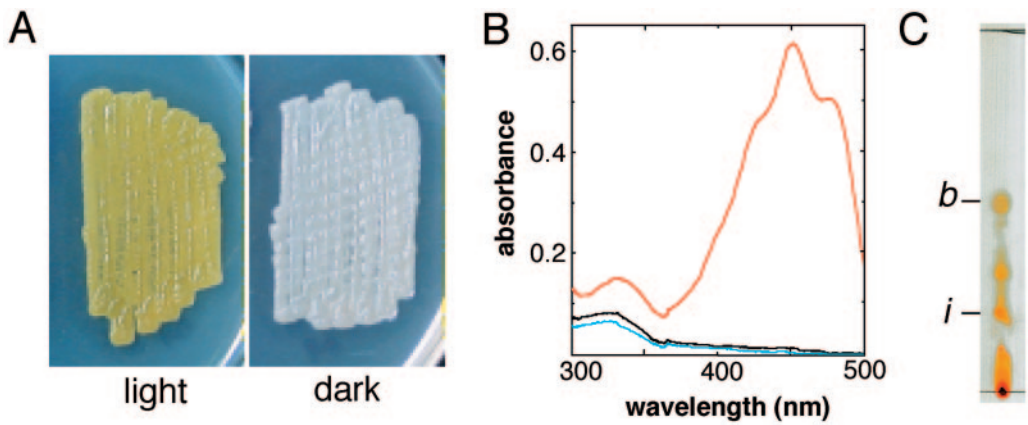


FIG. 1. Carotenoid production by *S. coelicolor* A3(2). (A) Colonies of wild-type *S. coelicolor* harboring pKM284. Colonies grown for 4 days on Bennett's glucose solid medium under light and dark conditions are shown. This strain is defective in the production of blue- and red-pigmented antibiotics due to the unknown effect of pKM284; consequently, it clearly exhibits the accumulation of yellow carotenoid pigments in the cells. (B) UV-visible spectrum of the purified carotenoid fraction extracted from *S. coelicolor* wild-type cells grown for 2 days under light (red line) and dark (black line) conditions. The same fraction extracted from the *crtEIBV* mutant cultured under light conditions was also analyzed (blue line). (C) Thin-layer chromatography analysis of the purified carotenoid fraction from *S. coelicolor* wild-type cells grown for 2 days under blue light illumination. The spots identified as β -carotene ($R_f = 0.54$) (b) and isorenieratene ($R_f = 0.24$) (i) by HPLC analysis are indicated.

least five yellow pigment spots that showed different R_f values (Fig. 1C). Among these spots, two were identified as β -carotene and isorenieratene by HPLC analysis with standard controls.

Carotenoid biosynthesis gene cluster of *S. coelicolor* A3(2). Figure 2A shows the carotenoid biosynthesis gene cluster and its flanking regions found in the genome sequence of *S. coelicolor* A3(2) (http://www.sanger.ac.uk/Projects/S_coelicolor) (5). The biosynthesis gene cluster consists of two convergent operon structures, *crtEIBV* and *crtYTU*. The gene organization is identical to that of the *crt* gene cluster in *S. griseus* (22). The *crt* gene cluster is also present in the genome of *Streptomyces avermitilis* (<http://avermitilis.ls.kitasato-u.ac.jp>) (14), which has the two operons in divergent orientations. The short distance

between the *crt* genes suggests that transcription of the *crt* biosynthesis gene cluster is directed by the promoters preceding *crtE* and *crtY*.

The *S. coelicolor* A3(2) *crt* gene cluster is flanked by a putative regulatory region that consists of two divergent operons, *litRQ* and *litSAB* (for light-induced transcription) (Fig. 2A). LitR (35 kDa) and LitS (22 kDa) are a MerR family transcriptional regulator and an RNA polymerase ECF sigma factor, respectively. LitA (23 kDa) is a possible lipoprotein. LitB (15 kDa) shows weak similarity to proteins known as anti-sigma factors, such as RsrR of *S. coelicolor*. LitQ (44 kDa) is a possible oxidoreductase. The *lit* cluster minus *litAB* is also present in the *S. avermitilis* genome but at a different locus from the *crt* cluster.

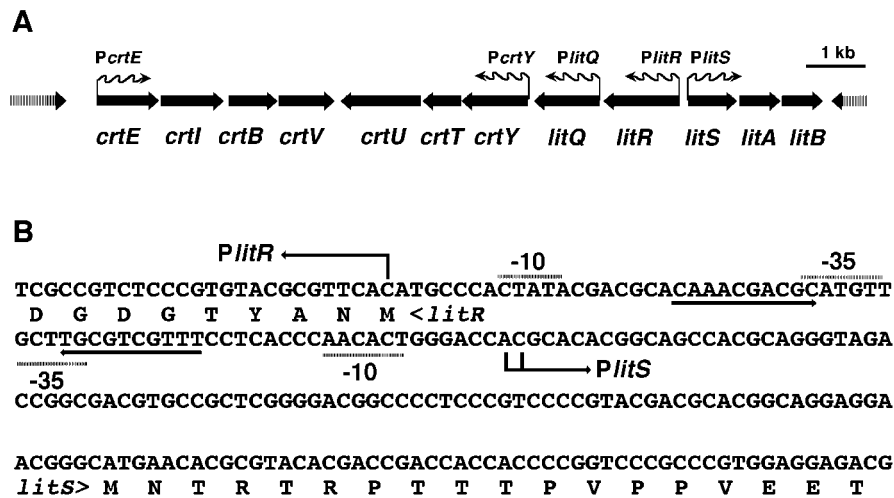


FIG. 2. Carotenoid biosynthesis gene cluster of *S. coelicolor* A3(2). (A) Schematic view of *crt* and the adjacent *lit* gene cluster. Wavy arrows indicate the orientations of the major transcripts. (B) Nucleotide sequence of the bidirectional promoter region between *litR* and *litS*. The N-terminal amino acid sequences of LitR and LitS are also shown. The transcriptional initiation sites determined by S1 mapping are indicated by bent arrows. An inverted repeat region is indicated by convergent arrows. Regions corresponding to the -35 and -10 sites are also indicated.

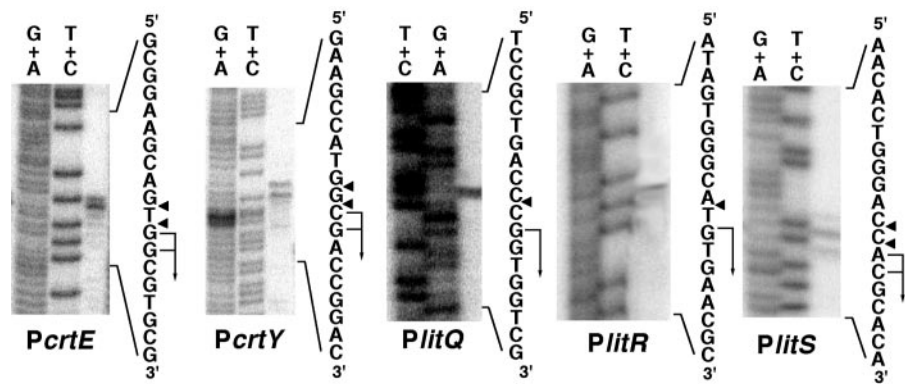


FIG. 3. Determination of transcriptional start sites by high-resolution S1 protection assay. Maxam-Gilbert sequencing ladders (G+A and T+C reactions) were generated from the same ³²P-labeled probe DNA fragments used in each S1 protection reaction. The positions of the S1-protected fragments are shown by the arrowheads, and the transcriptional initiation sites are assigned to the residues indicated by the bent arrows.

Inactivation of *crt* and *lit* genes and overexpression of *litS*. We generated *crt* and *lit* gene mutants in order to examine the roles of the genes in carotenogenesis in *S. coelicolor* A3(2). All mutants except the *litS* mutant have an insertion of a kanamycin resistance gene, which is oriented so as not to cause polar effects on expression of other *crt* and *lit* genes except *litR* (see Materials and Methods). The *crtEIBV* deletion mutant did not produce the yellow pigments, which confirmed that the pigments are carotenoids synthesized by the *crt* gene cluster (Fig. 1B). Inactivation of *litS* by markerless construction also abolished the ability to produce carotenoids. This strongly suggested that the sigma factor gene is essential for the transcription of the *crt* biosynthesis gene cluster. Meanwhile, *litAB* and *litB* inactivation did not affect carotenoid production. Carotenoid production in the *litS* mutant was restored by the introduction of a single copy of the intact *litS* gene on an integration plasmid, pIJ8660 (18).

We were unsuccessful in isolating markerless *litR* and *litQ* disruptants. In both genes, the single-crossover segregant that carries both an intact and a mutated coding sequence was successfully obtained; however, the double crossover, which removes one of the alleles, generated only revertants that exhibited the wild-type genotype. We also failed to obtain the mutants even when the cells were cultured in the dark. On the other hand, we could obtain a *litR* mutant by replacing the wild-type *litR* allele with a mutated construct carrying a kanamycin resistance gene in the middle of the *litR* coding sequence, in the direction opposite to that of *litR*. The mutant was defective in carotenoid production, while its growth, colony morphology, and antibiotic production were unaffected. Carotenoid production in this mutant was not restored when a single copy of an intact *litR* gene on a plasmid was introduced.

We also examined the carotenoid production phenotype resulting from *litS* and *litR* overexpression. The wild-type strain, which harbors a high-copy-number plasmid that carries *litS* downstream of the *mel* promoter (see Materials and Methods), produced carotenoids independent of illumination at the same level as the wild type cultured with illumination. The introduction of the *litS* expression plasmid also resulted in light-independent carotenoid production in the *litR* mutant. On the other hand, introduction of a *litR* expression plasmid that had a construction similar to that of *litS* did not affect the light-

dependent carotenoid production in the wild type or the carotenoid-deficient phenotype of any mutant strain.

S1 protection analysis. We examined the transcriptional activities of promoters preceding *crtE* (*PcrE*), *crtY* (*PcrY*), *litQ* (*PlitQ*), *litR* (*PlitR*), and *litS* (*PlitS*). High-resolution S1 protection analysis assigned a single transcriptional start site in each promoter (Fig. 3). It was noteworthy that the start site for *PlitR* was the first G residue of the translational initiation codon (GTG). The bidirectional promoter in the intergenic region between *litR* and *litS* had a symmetric structure centered on an inverted repeat that overlaps the -35 sequences of both promoters (Fig. 2B).

Quantification of transcripts by low-resolution S1 mapping showed that the majority of the promoters were light dependent (Fig. 4). In the wild-type strain, *PcrE*, *PcrY*, and *PlitS* activities were detected only under the light conditions. Meanwhile, *PlitQ* and *PlitR* showed activities under both dark and light conditions. *PlitQ* activity was relatively low and indepen-

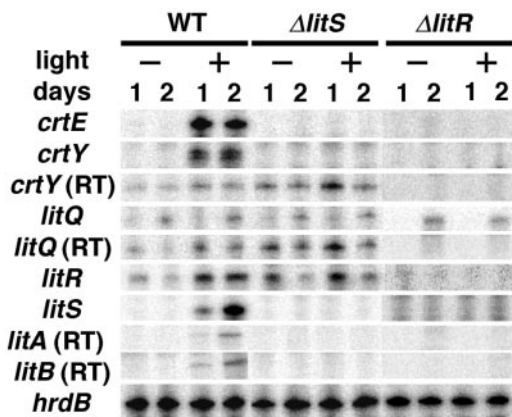


FIG. 4. Quantification of promoter activities by low-resolution S1 protection assay. The promoters preceding the coding sequences listed at the left were examined for their activities in the wild type (WT) and in the *S. coelicolor* A3(2) *litS* ($\Delta litS$) and *litR* ($\Delta litR$) mutants. RNA was isolated from cells cultured for 2 days under dark (-) or light (+) conditions. For *crtY*, *litQ*, *litA*, and *litB*, the signals derived from the readthrough transcripts (RT) are also shown. Transcription of *hrdB* was analyzed to confirm the quality and quantity of RNA.

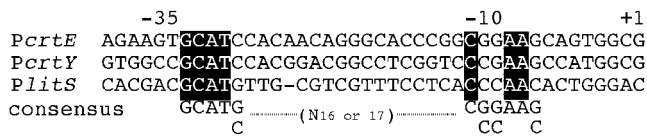


FIG. 5. Sequence alignment of the σ^{LitS} -dependent promoters. The solid boxes show a possible consensus sequence for recognition by $E\sigma^{\text{LitS}}$. Transcription initiation sites are underlined.

dent of illumination. *PlitR* showed lower activity under dark conditions than under light conditions. The probes for *PcrTE* and *PlitQ* also detected readthrough transcripts derived from *PlitR* (Fig. 2A). The probes for *PlitA* and *PlitB* detected only the readthrough transcript from *PlitS*, which indicates that *litAB* is cotranscribed with *litS*.

We also examined promoter activities in the *litS* and *litR* mutants (Fig. 4). In the markerless *litS* mutant, the activities of *PcrTE*, *PcrTY*, and *PlitS* were completely abolished. This suggested that *PcrTE*, *PcrTY*, and *PlitS* depend on σ^{LitS} . Comparison of the putative σ^{LitS} -dependent promoters revealed the presence of a possible consensus sequence, GCAT(G/C)-(16 or 17 bp)-C(G/C)(G/C)AA(G/C) at the -35 and -10 regions (Fig. 5). On the other hand, *litS* inactivation did not affect the activities of *PlitR* and *PlitQ*. The same analysis with the kanamycin-resistant *litR* mutant revealed that all promoter activities except that of *PlitQ* were completely abolished by the mutation. The S1 analysis with the probe for *PlitS* in the *litR* mutant detected a readthrough transcript derived from the promoter of the kanamycin resistance gene cassette, which had been inserted in the coding sequence of *litR* in the opposite orientation (data not shown). All activities of the *lit* and *crt* promoters in the *litAB* mutant were the same as those in the wild-type strain (data not shown).

In vitro runoff transcription analysis. We performed an in vitro runoff transcriptional assay in order to examine the dependence of *PcrTE*, *PcrTY*, and *PlitS* on σ^{LitS} . Recombinant σ^{LitS} protein was prepared by use of an *E. coli* expression-purification system (Fig. 6A). As shown in Fig. 6B, the RNA

polymerase holoenzyme containing the purified σ^{LitS} protein generated specific transcripts on the probe DNA fragments for both *PcrTE* and *PcrTY*. However, the holoenzyme did not generate a specific transcript on the probe DNA for the *PlitS* region, which led us to assume that the transcriptional initiation at *PlitS* requires an additional transcription factor, possibly LitR. Thus, we prepared purified LitR protein and assessed its effect on in vitro transcription by $E\sigma^{\text{LitS}}$ on a *PlitS* template; however, we failed to detect any specific transcript. In a gel mobility shift assay, the recombinant LitR protein did not bind DNA fragments containing *PlitS*.

DISCUSSION

In *Streptomyces*, carotenoid production is a widespread metabolic activity, which occurs in a constitutive, light-dependent, or cryptic manner (20). This implies the presence of a certain diversity in the molecular mechanisms of carotenoid production in this group of bacteria. To date, genetic studies on carotenoid production in *Streptomyces* have been described only for *Streptomyces setonii* (15) and *S. griseus* (22), both of which show cryptic carotenogenesis. Carotenoid production in these organisms is induced by an increased copy number of a gene that encodes a stress-response sigma factor, σ^{CrtS} . This suggests that this sigma factor is involved in the transcription of carotenoid biosynthesis genes in these species. On the other hand, *S. coelicolor* A3(2), examined in this study, does not retain the *crtS* homolog in its genome, and it was shown that an ECF sigma factor, σ^{LitS} , responsible for light-induced carotenoid production is present. The *litQRS* gene cluster is conserved in *S. avermitilis*, which also produces carotenoids in a light-dependent manner (H. Ikeda, personal communication). The presence of *lit* genes in the two phylogenetically divergent species raises the possibility that the regulatory mechanism is common to light-induced carotenoid production in this group of bacteria.

In terms of the molecular mechanism of light-dependent carotenogenesis, the best-studied nonphototrophic bacterium is *M. xanthus*, in which an ECF sigma factor is involved in the regulation of carotenoid production (8, 10, 23, 29). In *M. xanthus*, the light-inducible ECF sigma factor σ^{CarQ} directs transcription of the *carQRS* operon. Thus, expressed CarS acts as a positive regulator of the transcription of the carotenoid biosynthesis gene cluster. This transcription occurs through sequestration of a transcriptional repressor protein, CarA, which binds to the major promoter of the biosynthesis gene cluster to block transcription under dark conditions. The activity of σ^{CarQ} is negatively regulated by the adjacent anti-sigma factor CarR. CarR is a membrane protein that is unstable under illuminated conditions, and it is assumed to release active σ^{CarQ} upon illumination. Related studies have shown the presence of additional regulatory components involved in light-induced transcriptional control in *M. xanthus* (10).

The regulatory mechanism in *S. coelicolor* appears to be simple in comparison with the complex system in *M. xanthus*. *S. coelicolor* A3(2) LitS shows 21.3% amino acid sequence identity to *M. xanthus* σ^{CarQ} . However, the *S. coelicolor* genome does not contain homologous genes for CarR, CarS, or CarA of *M. xanthus*. While there is no in vitro evidence for transcription of *crt* genes by σ^{CarQ} in *M. xanthus* (8), *S. coelicolor* σ^{LitS}

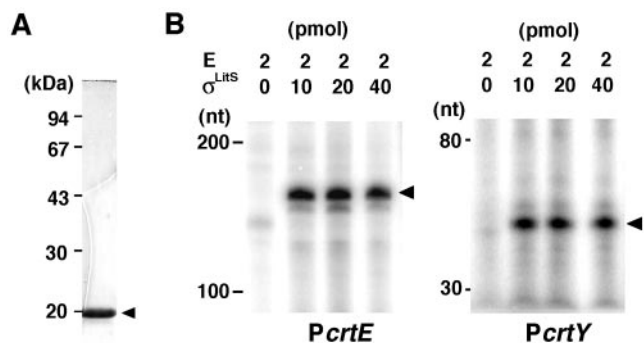


FIG. 6. In vitro runoff transcription by $E\sigma^{\text{LitS}}$. (A) Sodium dodecyl sulfate-polyacrylamide gel electrophoresis analysis of the σ^{LitS} protein expressed and purified by use of an *E. coli* host-vector system. (B) Polyacrylamide gel electrophoresis of σ^{LitS} -specific transcripts. The indicated amounts of an *E. coli* RNA polymerase core enzyme (E) and the recombinant σ^{LitS} protein were added to the reaction mixture with the promoter DNA fragments to generate transcripts specific for *crtE* (*PcrTE*) and *crtY* (*PcrTY*). Arrowheads indicate the specific transcripts. nt, nucleotides.

transcribes the *crt* operons. Such marked diversity in control mechanisms may reflect the ecological niches of those organisms.

The transcriptional analyses showed that light-induced transcriptional control underlies the photodependent carotenoid production in *S. coelicolor* A3(2). The S1 mapping result for the *litS* mutant strongly suggested that σ^{LitS} directs its own transcription. This indicates that *litS* comprises a positive feedback circuit, which may enable signal amplification that facilitates *crt* transcription upon receiving light. However, the σ^{LitS} -dependent transcription from the *litS* promoter was not reproduced in the in vitro transcriptional assay, while transcription from the *crt* promoters was successfully demonstrated. We speculate that transcription from the *litS* promoter requires an additional element(s), such as a transcriptional activator protein. The symmetric and compact structure of the bidirectional promoter region between *litS* and *litR* suggests that a photodependent regulator simultaneously controls the initiation of transcription in both directions. This may account for the photodependence of the *litR* promoter. The promoter region is highly conserved in the *S. avermitilis* counterpart, which suggests that an identical mechanism controls the light-dependent carotenoid production in this organism.

We were unsuccessful in isolating markerless mutants for *litR* and *litQ*, probably due to the lethal effect of the mutation. The *litR* mutant isolated in this study carries an *aphII* cassette that has a polar effect in the direction of *litS* and downstream genes. At present, we cannot clearly explain why the mutant construction did not affect viability. One possibility is that the constitutive readthrough transcription from the *aphII* promoter in the orientation toward *litS* compensates for the *litR* deficiency. Since polar transcription should confer low-level LitSAB expression, some functions of these proteins may be related to suppression of the lethality caused by *litR* inactivation. The suppression mechanism may also be related to the failure in the genetic complementation of the *litR* mutant, although we have no convincing working model for this mechanism.

Transcriptional analysis of the *litR* mutant demonstrated that all promoter activities, except *litQ*, are abolished in the mutant background. This result leads to a simple hypothesis for the light induction of *litS*: LitR acts as an essential transcriptional activator of the *litS* promoter under light conditions. This accounts for the effect of *litR* depletion on the promoter activities and carotenoid productivity. The lack of *litR* promoter activity in the *litR* mutant implies that LitR also acts as a positive regulator of its own promoter and thus comprises a positive feedback loop. Although this proposition appears to be justified, we believe it should be supported further by additional experimental results, particularly those using different genetic backgrounds. This is because we cannot overlook the possibility of the presence of an artifact caused by the polar effect in the *litR* mutant. Meanwhile, the effect of the high copy number of *litS*, which resulted in constitutive carotenoid production in the *litR* mutant, greatly supports the idea that *litS* is under *litR* control, such that *litS* overexpression can suppress the *litR* deficiency.

LitR shows an end-to-end similarity to the MerR-type transcriptional regulator, which usually acts as a transcriptional activator by binding to an inverted repeat structure localized in

the vicinity of the -35 region (7). The compact structure of the bidirectional promoter region between *litR* and *litS* appears to meet the condition for the action of this type of transcriptional regulator. Usually, MerR family regulators acquire DNA-binding ability by ligand binding (7). It is possible that the LitR recombinant protein prepared in this study requires the supply of a specific ligand in order to bind the intergenic promoter region between *litS* and *litR* in the in vitro DNA-binding assay. If this is the case, identification of the ligand molecule should serve as an important clue for uncovering the photodependent regulatory mechanism in *S. coelicolor*. We speculate that *litQ* may also be related to the control system, based both on the conserved features in two different *Streptomyces* species and on the lethality of its inactivation, as shown by *litR*. The amino acid sequence similarity search suggested that LitQ is an oxidoreductase, and its function may be related to the link between illumination and activation of the LitR protein. Studies on the function of Lit proteins will reveal a novel light-responding control mechanism in this group of bacteria.

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