Overexpression of the primary sigma factor gene sigA improved carotenoid production by *Corynebacterium glutamicum*: Application to production of β-carotene and the non-native linear C50 carotenoid bisanhydrobacterioruberin

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ABSTRACT

*Corynebacterium glutamicum* shows yellow pigmentation due to biosynthesis of the C50 carotenoid decaprenoxanthin and its glycosides. This bacterium has been engineered for production of various non-native cyclic C40 and C50 carotenoids such as β-carotene, astaxanthin or sardinaxanthin. In this study, the effect of overexpressing gene expression more broadly by overexpression of sigA factor genes on carotenoid production by *C. glutamicum* was characterized. Overexpression of the primary sigA factor gene sigA improved lycopene production by recombinant *C. glutamicum* wild type, overexpression of sigA led to 2-fold increased accumulation of the native carotenoid decaprenoxanthin in the stationary growth phase. Under these conditions, genes related to thiamine synthesis and aromatic compound degradation showed increased RNA levels and addition of thiamine and the aromatic iron chelator protocatechuic acid to the culture medium enhanced carotenoid production when sigA was overexpressed. Deletion of the gene for the alternative sigma factor SigB, which is expected to replace SigA in RNA polymerase holoenzymes during transition to the stationary growth phase, also increased carotenoid production. The strategy of sigA overexpression could be successfully transferred to production of the non-native carotenoids β-carotene and bisanhydrobacterioruberin (BABR). Production of the latter is the first demonstration that *C. glutamicum* may accumulate a non-native linear C50 carotenoid instead of the native cyclic C50 carotenoid decaprenoxanthin.

1. Introduction

Carotenoids are natural pigments which show various colors from yellow to red depending on their chemical structures (Britton et al., 2004). Because of versatile applications especially in food and feed industries, the demand for bio-based carotenoid production is increasing (Breithaupt, 2007; Scotter, 2011). In order to increase the production and efficiency, metabolic engineering has been applied in the natural carotenoid producers as well as in non-carotenogenic organisms (Ausch, 2009; Ye and Bhatia, 2012). The pathways and related enzymes for carotenogenesis are well understood. First, IPP (Isopentenyl pyrophosphate) and DMAPP (Dimethylallyl pyrophosphate) are synthesized either by the mevalonate (MEV) pathway or the mevalonate pathway (Chang et al., 2013). IPP and its isomer DMAPP are condensed to geranylgeranyl pyrophosphate (GGPP), of which lycopene is synthesized (Moise et al., 2014). Lycopene is a red C40 carotenoid that serves as precursor for the synthesis of other C40 and C50 carotenoids such as β-carotene, lutein or astaxanthin (Heider et al., 2014a; Misawa and Shimada, 1997).

*Corynebacterium glutamicum* is a non-pathogenic bacterium that is used for several decades for the million-ton-scale amino acid production, especially for L-glutamate and L-lysine (Eggeling and Bott, 2005). Furthermore, this bacterium has been engineered for production of various industrially relevant compounds (Wendisch, 2014). *C. glutamicum* naturally produces the C50 carotenoid decaprenoxanthin (Krabasik et al., 2001). Recently, *C. glutamicum* strains overproducing lycopene and other carotenoids including β-carotene and astaxanthin have been developed (Heider et al., 2014b, 2014c, 2014d, 2014e).
2012; Henke et al., 2016). C. glutamicum is a suitable production host for carotenoids; the volumetric productivity for astaxanthin of up to 0.4 mg L⁻¹ h⁻¹ is comparable to current algae-based production (Henke et al., 2016) and high cell density fermentation with up to 100 g cell dry weight (CDW) L⁻¹ is established (Knoll et al., 2007).

Production strain development may involve regulatory engineering, e.g. of sigma factor genes (Tripathi et al., 2014). Sigma factors are important for promoter recognition and transcription initiation as one of the subunits of RNA polymerase holoenzyme (Gruber and Gross, 2003). Bacteria often possess multiple sigma factor genes which encode primary and alternative sigma factors (Feklistov et al., 2014). Depending on the sigma factor targeted, sigma factor engineering may alter the transcription profile globally or may affect a subset of genes that share similar functions. For example, deletion of the general stress sigma factor gene rpoS improved putrescine production by recombinant E. coli (Qian et al., 2009). Overexpression of the nitrogen starvation sigma factor gene rpoN in E. coli improved heterologous expression of polyketide and non-ribosomal peptide biosynthesis gene clusters of Streptomyces rimosus, which may be due to the presence of RpoN promoter sequences in the majority of them (Stevens et al., 2013). E. coli strains with improved production characteristics such as ethanol tolerance could be selected from a population overexpressing randomly mutagenized RpoD (Alper and Stephanopoulos, 2007). In Synechocystis sp. PCC 6803, overexpression of alternative sigma factor gene sigE improved production of polyhydroxybutyrate (Osanai et al., 2013). In C. glutamicum, which possesses seven sigma factors SigA, SigB, SigC, SigD, SigE, SigH and SigM (Pátek and Nešerová, 2011; Toyota and Inui, 2015), overexpression of sigH resulted in overproduction of riboflavin and of flavin mononucleotide (FMN) when combined with overexpression of the endogenous gene encoding bifunctional riboflavinkinase/FMN adenyltransferase (Taniguchi and Wendisch, 2015). In this study, the effect of overexpression of sigma factor genes on carotenoid production was elucidated and overexpression of the general sigma factor gene sigA was shown to decrease overproduction of the C40 carotenoids lycopene and β-carotene as well as the C50 carotenoids decaprenoxanthin and bisanhydrobacterioruberin.

2. Material and methods

2.1. Bacterial strains, plasmids and oligonucleotides

The strains, plasmids and oligonucleotides (Metabion, Martinsried, Germany) used in this work are listed in Table 1. Plasmids were constructed based on pVWE1x or pEKEx3, which are both IPTG inducible expression vectors for E. coli and C. glutamicum (Peters-Wendisch et al., 2001; Stansen et al., 2005). Plasmid and strain construction was performed as described previously (Taniguchi and Wendisch, 2015). Briefly, the DNA fragment of the target gene was amplified with the respective oligonucleotide pairs in Table 1, and inserted into pVWE1x, alternatively pEKEx3, by Gibson assembly (Gibson et al., 2009). E. coli DH5α was used for cloning. The sequence of inserted DNA fragments was confirmed by sequencing (CeBiTec Sequencing Core Facility, Bielefeld, Germany). C. glutamicum competent cells were transformed by electroporation at 2.5 kV, 200 μΩ, and 25 μF (Eggeling and Bott, 2005; van der Rest et al., 1999).

2.2. Chromosomal deletion of sigB in C. glutamicum

For targeted deletion of sigB, the suicide vector pK19mobsacB was used (Schäfer et al., 1994). Genomic regions flanking cg2102 were amplified from the genomic DNA of C. glutamicum WT ATCC 13032 using oligonucleotide pairs sigB-A/B and sigB-C/D (Table 1), respectively. Afterwards, the purified PCR products were linked and cloned into BamHI digested pK19mobsacB via Gibson Assembly (Gibson et al., 2009). The resulting deletion vector pK19mobsacB-ΔsigB (Table 1) was confirmed via sequencing. Introduction of pK19mobsacB-ΔsigB into C. glutamicum was carried out via trans conjugation with E. coli S17-1 (Schäfer et al., 1994). Deletion of sigB via two-step homologous recombination as well as the selection for the first and second recombination events were carried out as described previously (Eggeling and Bott, 2005). Successful deletion of sigB was verified by PCR analysis of the constructed mutant using oligonucleotide pair sigB-E/F (Table 1) and sequencing of the PCR product.

2.3. Medium, growth condition and growth rate comparison

As far as not mentioned specifically, C. glutamicum was precultured in LB medium (Sambrook, 2001) with 56 mM of glucose overnight, washed once with CGXII medium (Eggeling and Bott, 2005) without carbon source and inoculated in CGXII with 222 mM of glucose at initial optical density (OD) (λ=600 nm) of 1. The OD was measured with UV-1202 spectrophotometer (Shimadzu, Duisburg, Germany) with suitable dilutions. When appropriate, 25 μg/mL of kanamycin and IPTG were added as indicated in the text. Growth experiment with Biolector® cultivation system (m2pLabs, Baesweiler, Germany) was performed in 1 mL of CGXII with 222 mM of glucose using FlowerPlate® (m2pLabs, Baesweiler, Germany) at 30 °C, 1100 rpm. Growth experiment with flask was performed in 50 mL of CGXII 222 mM of glucose using 500 mL of baffled flask at 30 °C, 120 rpm. For growth rate calculation, cell growth was monitored online every 10 min for 48 h with Biolector®. Maximum growth rate μ (h⁻¹) was calculated from 20 measuring points of arbitrary unit of the backscattering light (620 nm). Plate image was scanned with Perfection V750-M Pro scanner (Epson, Ludwigshafen am Rhein, Germany). Consumption of glucose was tested with Diabur 5000 glucose test strips (Roche Diagnostics, Mannheim, Germany) with a detection limit of 0.005% glucose.

2.4. Carotenoid extraction and quantification

For quantification of lycopene, decaprenoxanthin and β-carotene, 200–1000 μL of the cell culture was centrifuged and washed once with 500 μL of CGXII without carbon source. The cell pellet was frozen and stored at −20 °C until further use. The cell pellet was resuspended in 400–800 μL of acetone and carotenoids were extracted for 60 min at 50 °C and 1400 rpm. The supernatant was separated by centrifugation and absorption spectrum from 400 nm to 550 nm was measured by UV-1800 spectrophotometer (Shimadzu, Duisburg, Germany). The spectrum was normalized by the absorbance at wavelength of 550 nm and the maximum absorbance in the spectrum was used for quantification of each carotenoid (474 nm for lycopene, 440 nm for decaprenoxanthin, 454 nm for β-carotene). The quantification of bisanhydrobacterioruberin (BABR) was performed by high performance liquid chromatography (HPLC) with the Agilent 1200 series system (Agilent Technologies Sales & Services GmbH & Co. KG, Waldbronn). For HPLC analysis, pigments were extracted with 800 μL methanol: acetone (7:3) containing 0.05% BHT (Butylhydroxytoluol) at 60 °C for 15 min with careful vortexing every 5 min. The supernatant was separated by centrifugation and used for analysis. 50 μL of the sample was separated with a column system consisting of a precolumn (LiChrospher 100 RP18 EC-5, 40×4 mm, CS-Chromatographie, Langenwehe, Germany) and a main column (LiChrospher 100 RP18 EC-5, 125×4 mm, CS-Chromatographie) with methanol (A) and methanol/water (9:1) (B) as mobile phase. The following gradient was used at a flow rate of 1.5 mL/min; 0 min B: 0%, 10 min B: 100%, 32.5 min B: 100%. The UV/visible (Vis) spectrum was recorded with a diode array detector (DAD) and the amount of carotenoids was quantitated by the integration of the extracted wavelength chromatogram at 471 nm and by the analysis of the appropriate UV/Vis profiles. For quantification of lycopene and β-carotene, pure standards (Sigma-Aldrich, Germany) were used. Due to the lack of commercially available standards for
### Table 1
Bacterial strains, plasmids and oligonucleotides used in this study.

| Bacterial strain | Relevant characteristic | Reference or source |
|------------------|-------------------------|---------------------|
| **C. glutamicum** |                         |                     |
| WT               | Wild-type, ATCC 13032    |                     |
| WTΔsigB          | sigB deletion mutant of WT ATCC 13032 |                     |
| LVC5             | LVC5-P_{sig}-derivatives with insertion of crtEBI operon under the control of P_{sig} promoter integrated into the cgg2 cured region between cg1745 and cg1753 | ATCC this study (Henke et al., 2016) |
| LVC5ΔsigB        | sigB deletion mutant of LVC5 |                     |
| BETA3            | LVC5 derivatives with insertion of crvY from *P. ananatis* under the control of P_{sig} promoter integrated into the cgg2 cured region between cg1745 and cg1753 | this study (Henke et al., 2016) |
| BETA3ΔsigB       | sigB deletion mutant of BETA3 |                     |
| *E. coli*        |                         |                     |
| *E. coli* DH15a  | F-θi-1 endA ΔlacI7(r-– m-) supplE44 ΔlacU169 (Φ80lacZΔM15) recA1 gyrA96 relA1 | Bethesda Research Laboratories |

| **Plasmid**      | Relevant characteristic | References |
|------------------|-------------------------|------------|
| pVWEex1          | KanR: *E. coli*-C. glutamicum shuttle vector for regulated gene expression (P_{lac}, lacI, pCG1 oriVCg) | (Peters-Wendisch et al., 2001) |
| pVWEex1-sigA     | KanR, pVWEex1 with sigA from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigB     | KanR, pVWEex1 with sigB from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigC     | KanR, pVWEex1 with sigC from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigD     | KanR, pVWEex1 with sigD from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigE     | KanR, pVWEex1 with sigE from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigH     | KanR, pVWEex1 with sigH from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-sigM     | KanR, pVWEex1 with sigM from C. glutamicum WT | this study (Wendisch, 2015) |
| pVWEex1-lbtBC    | KanR, pVWEex1 with lbtBC from *Dietzia* sp. CQ4 | this study (Stansen et al., 2005) |
| pEEEx3           | SpecR: *E. coli*-C. glutamicum shuttle vector for regulated gene expression (P_{lac}, lacI, pCG1 oriVCg) | (Schäfer et al., 1994) |
| pEEEx3-sigA      | SpecR, pEEEx3-sigA from C. glutamicum WT |                     |
| pK19mobsacB      | KmR: *E. coli*-C. glutamicum shuttle vector for construction of insertion and deletion mutants in *C. glutamicum* (pK18 oriVEc sacB lacZ) | this study (Wendisch, 2015) |
| pK19mobsacB-ΔsigB| KmR: *E. coli*-C. glutamicum shuttle vector for construction of deletion mutant ΔsigB in *C. glutamicum* | this study (Wendisch, 2015) |

| **Oligonucleotide** | **Sequence (5'-3')** | References |
|---------------------|----------------------|------------|
| sigA-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigA-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigB-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigB-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigC-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigC-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigD-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigD-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigE-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigE-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigH-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigH-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigM-fwd            | GCCCTGCAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| sigM-rev            | CCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| lbtBC-fwd           | CATGCCCTGAGGAGACCTCTAGAGAAAGGAGGCCCTTCAGATGTTAGAAAAACACGTAGCAGAAAAAAGAGCGGTCG | (Taniguchi and Wendisch, 2015) |
| lbtBC-rev           | ATTCGACGACTTCGGTACCGGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigA                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigB                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigC                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigD                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigE                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigF                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigG                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigH                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigI                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigJ                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigK                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigL                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigM                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
| sigN                | GCGTACGCCGGGATTTTGTTGCTGAGTCGAGACCTGG | this study (Wendisch, 2015) |
decaprenoxanthin and BABR, the concentrations of these are given as β-carotene equivalents using standardization with pure β-carotene. The amount of each carotenoid production was normalized against cell weight/optical density.

2.5. Transcriptome analysis of the sigA overexpressing strain using DNA microarray

C. glutamicum strains WT(pVWEx1) and WT(pVWEx1-sigA) were cultured in LB medium with 56 mM of glucose and 25 µg/mL of kanamycin and inoculated into CGXII medium with 222 mM of glucose and 25 µg/mL of kanamycin for adaptation to glucose as carbon source. Cells were cultured overnight and inoculated into 50 mL of CGXII medium with 222 mM of glucose, 50 µM of IPTG and 25 µg/mL of kanamycin at the initial OD of 1. Cells were harvested 8 h after inoculation in the early exponential growth phase (OD between 6 and 8) and after 24 h in the stationary phase. RNA isolation was performed as described previously (Wendisch, 2003). The purified RNA was analyzed by spectrophotometer (NanoDrop) for quantity and gel electrophoresis for quality. The RNA sample was stored at −80 °C until further use. cDNA synthesis as well as DNA microarray hybridization were performed as described previously (Netzer et al., 2004; Polen et al., 2007). Normalization and evaluation of the data was done with the software package EMMA 2 (Dondrup et al., 2009). Genes which expression was upregulated or downregulated in WT(pVWEx1-sigA) were taken into account for further analysis (FDR < 0.05, M-value > 1 for upregulation, M-value < −1 for downregulation).

3. Results

3.1. Overexpression of sigA improved production of lycopene by C. glutamicum

Lycopene is a central intermediate of carotenogenesis in C. glutamicum (Fig. 1). In the lycopene producing strain L5C5 sigma factor genes were overexpressed and their influence on lycopene production was evaluated in Biolector microscale cultivations. The transformants were grown in minimal medium with glucose as carbon source and sigma factor gene expression from pVWEx1 was induced with 50 µM IPTG. Overexpression of sigma factor genes influenced the maximum growth rate differently (Fig. 2). Overexpression of sigA, sigD and sigE led to a slight decrease of the growth rate, and overexpression of sigB and sigM did not influence the growth rate. On the other hand, overexpression of sigC and sigH slowed cell growth considerably. Overexpression of sigma factor genes influenced the final biomass only to a small extent ( < ± 10%) except for sigH (−20%). Overexpression of sigH in L5C5 led to accumulation of riboflavin in the supernatant as reported previously for overexpression of sigH in C. glutamicum wild type (Taniguchi and Wendisch, 2015). Overexpression of sigA led to more reddish colored cells compared to the control strain, L5C5(pVWEx1) (Fig. 2). Quantification of lycopene showed that overexpression of sigC, sigD, sigH and sigM led to moderately increased lycopene production, while overexpression of sigB and sigE decreased production (Fig. 2). Furthermore, sigA overexpressing cells accumulated about 3.5-fold more lycopene than the control strain (Fig. 2).

To verify the effect of sigA overexpression on lycopene production, growth and lycopene accumulation was followed for 72 h during different growth phases using shake flasks. Cells reached the stationary growth phase (Fig. 3A) and glucose was consumed completely after 24 h. For the first 24 h of cultivation, lycopene accumulation was significantly but only slightly higher (about 1.5-fold) in the sigA overexpressing strain than in the control strain (Fig. 3B). However, the sigA overexpressing cells continued to produce lycopene in the stationary growth phase from 24 h to 72 h (Fig. 3B). After 72 h, the sigA overexpressing strain produced 0.82 ± 0.15 mg/g CDW lycopene, which was about 8-fold more than the control strain. Thus, overexpression of sigA encoding the primary sigma factor of C. glutamicum improved lycopene production particularly in the stationary growth phase.

3.2. C. glutamicum wild type overproduced decaprenoxanthin in the stationary growth phase as consequence of sigA overexpression

Overexpression of sigA affected lycopene accumulation most in the stationary growth phase (Fig. 3B). To determine if sigA overexpression affects biosynthesis of the native carotenoid decaprenoxanthin, C. glutamicum WT(pVWEx1-sigA) and WT(pVWEx1) were cultivated in glucose minimal medium in shake flasks for 72 h and decaprenoxanthin accumulation during the exponential and stationary growth phases were monitored. During exponential growth and until glucose was exhausted at about 24 h, decaprenoxanthin accumulation hardly differed between both strains (Fig. 3C). However, C. glutamicum WT(pVWEx1-sigA) continued to produce decaprenoxanthin in the stationary growth phase while WT(pVWEx1) did not (Fig. 3D). After 72 h, decaprenoxanthin production by WT(pVWEx1-sigA) was about twice as high as in the control strain (Fig. 3D). Thus, sigA overexpression increased decaprenoxanthin biosynthesis by wild type in the stationary growth phase.

3.3. Application of sigA overexpression to production of the non-native cyclic C40 carotenoid β-carotene

Lycopene is a precursor for several carotenoids and sigA overexpression improved its production. However, as lycopene production is not a reliable indicator of MEP pathway flux at least in E. coli (Bongers et al., 2015), we tested if sigA overexpression is beneficial for production of a lycopene-derived non-native carotenoid such as β-carotene. C. glutamicum WT does not synthesize β-carotene, but the recombinant strain C. glutamicum BETA3 produces β-carotene without accumulation of the precursor lycopene (Henke et al., 2016). This strain has been constructed based on C. glutamicum L5C5 by genomic integration of crtY from P. ananatis encoding lycopene cyclase under the control of a strong constitutive promoter (Fig. 1). A comparative growth experiment of C. glutamicum BETA3(pVWEx1) and BETA3(pVWEx1-sigA) revealed that both strains grew to similar biomass concentrations after glucose was completely consumed (Fig. 4A). In the stationary growth phase, BETA3(pVWEx1-sigA) produced 11.9 ± 1.5 mg/g CDW, which is about 3-fold more β-carotene than accumulated by the control strain BETA3(pVWEx1) (Fig. 4A).

3.4. Engineering of C. glutamicum for overproduction of a non-native linear C50 carotenoid

Carotenoids are believed to span the cytoplasmic membrane and the prevalence of C40 carotenoids may reflect the typical cytoplasmic membrane width (Abbes et al., 2013; Lazrak et al., 1987). Besides the native decaprenoxanthin (Heider et al., 2014c; Krubasik et al., 2001), we showed that recombinant C. glutamicum strains overproduced carotenoids of different lengths: linear and cyclic C40 carotenoids such as lycopene and β-carotene (Heider et al., 2014a; Henke et al., 2016) as well as cyclic C50 carotenoids such as sarcinaxanthin (Netzer et al., 2010). Here, we attempted to overproduce a non-native linear C50 carotenoid, namely bisanhydrobacterioruberin (BABR) (Norgård et al., 2010).
expressed from the IPTG inducible plasmid pVWEx1 in the lycopene overproducing strain C. glutamicum LYC5 (Fig. 1). To apply our finding that sigA overexpression improved carotenoid production, the resulting strain LYC5(pVWEx1-lbtBC) was transformed with pEKEEx3-sigA. After cultivation of this strain and an empty vector carrying control strain for 48 h in glucose minimal medium, carotenoids were extracted and quantified. Both C. glutamicum LYC5(pVWEx1-lbtBC) (pEKEEx3) and LYC5(pVWEx1-lbtBC)(pEKEEx3-sigA) accumulated the linear C50 carotenoid bisanhydrobacterioruberin to considerable concentrations. When BABR production was calculated based on a β-carotene standard due to the lack of a standard, C. glutamicum LYC5(pVWEx1-lbtBC)(pEKEEx3-sigA) produced 0.52 ± 0.12 mg/g CDW bisanhydrobacterioruberin. Thus, plasmid-driven sigA overexpression led to 1.8-fold higher BABR production (Fig. 4B).

3.5. Global gene expression changes due to sigA overexpression

Overexpression of the primary sigma factor gene sigA is expected to affect expression of many genes and this effect may differ in the exponential and stationary growth phases. In order to determine which genes are differentially expressed in the exponential and stationary growth phases due to sigA overexpression, C. glutamicum WT(pVWEx1) and WT(pVWEx1-sigA) were cultivated in glucose minimal medium and RNA was extracted in the mid-exponential growth phase at 8 h and at the beginning of the stationary growth phase when glucose was exhausted (at 24 h). Global gene expression analysis was determined by DNA microarray analysis (Table 2). The data were deposited at GEO (GSE86866). The analysis revealed significantly decreased RNA levels of 18 genes at 8 h and 32 genes at 24 h (false discovery rate < 0.05, M-value < –1) (Table S1, Table S2).

Only one gene (cg0612), which is annotated as putative aldo/keto reductase, showed decreased RNA levels both at 8 and 24 h (Table 2). In addition to the overexpressed sigA itself, RNA levels for 49 and 64 genes increased significantly at 8 h and 24 h, respectively (false discovery rate < 0.05, M-value > 1) (Table S1, Table S2). Notably, genes involved in carotenogenesis did not show increased RNA levels upon sigA overexpression. At both 8 h and 24 h, RNA levels of nine genes were increased (Table 2): seven genes encoding uncharacterized proteins, the cardiolipin synthase gene cls (Namposthiri et al., 2002) and the catechol 1,2-dioxygenase gene catA (Shen et al., 2004). Although catechol 1,2-dioxygenase is not involved in degradation of protocatechueic acid (PCA) (Shen et al., 2004), a medium component added as iron chelator, catA is among the genes induced during fast growth with glucose/PCA mixtures (Uththan et al., 2014). Thiamine is not added as medium component, however, thiamine biosynthesis genes were upregulated at both time points: thioS/G at 8 h and thiC and thiD1 at 24 h (Table S1, Table S2). Thiamine is necessary as a cofactor for 1-deoxy-D-xylulose-5-phosphate (DXP) synthase (Dxs), which catalyzes the first step of isopentenyl diphosphate (IPP) synthesis in the MEP pathway (Vranová et al., 2013) and in the thiamine biosynthesis pathway (Begley et al., 1999).

Based on the finding that catA and thiamine biosynthesis genes were induced when sigA was overexpressed, growth experiments in the presence of 10-fold increased PCA concentration (300 mg/L) or in the presence of 10 μg/L thiamine were performed. These supplements did not influence the final biomass of C. glutamicum WT(pVWEx1) and WT(pVWEx1-sigA) (Fig. 5A). Accumulation of decaprenoxanthin by WT(pVWEx1-sigA), but not by WT(pVWEx1), increased by about 10% when thiamine was added (Fig. 5B). In the presence of a 10-fold increased PCA concentration, decaprenoxanthin production by WT(pVWEx1-sigA), but not by WT(pVWEx1), increased by about 40% (Fig. 5B). Therefore, the finding that thiamine and PCA addition
supported decaprenoxanthin production by C. glutamicum WT(pVWEx1-sigA) is commensurate with the gene expression changes due to sigA overexpression.

3.6. Deletion of the sigma factor gene sigB increased carotenoid production

SigA of C. glutamicum is mostly associated to RNA polymerase in the exponential growth phase, while SigB is believed to replace SigA during the transition to the stationary growth phase (Kim et al., 2016; Larisch et al., 2007; Pátek and Nešvera, 2011). During the transition to the stationary growth phase, expression of sigA is reduced and expression of sigB increased (Larisch et al., 2007). Thus, we hypothesized that overexpression of sigA increases the proportion of SigA containing RNA polymerase holoenzymes in the stationary growth phase and that deletion of sigB might have a comparable effect. To test this hypothesis, sigB was deleted in carotenoid producer strains. The deletion and the control strains grew comparably in glucose minimal medium and consumed glucose within 24 h of cultivation. Extraction and quantification of carotenoids after 48 h of cultivation revealed that about 5-fold more lycopene was produced by LYCSΔsigB in comparison to C. glutamicum LYC5 (Fig. 6A), which is comparable to the about 8-fold increased lycopene production as consequence of sigA overexpression (Fig. 3). In a similar manner, deletion of sigB improved production of decaprenoxanthin, β-carotene and BABR by C. glutamicum WT, BETA3 and LYC5(pVWEx1-lbtBC), respectively (Fig. 6B–D). Thus, deletion of sigB is beneficial for carotenoid production by C. glutamicum, likely because a high proportion of SigA containing RNA polymerase holoenzymes can be maintained in the absence of sigma factor competition by SigB. Combination of deletion of sigB and overexpression of sigA did not further increase lycopene production but perturbed growth (data not shown).

4. Discussion

Production of endogenous and non-native carotenoids by C. glutamicum was improved by overexpression of the primary sigma factor gene sigA and by deletion of the alternative sigma factor gene sigB. Overexpression of sigma factor genes has also been used in metabolic engineering of E. coli for lycopene production (Alper and Stephanopoulos, 2007; Jin and Stephanopoulos, 2007; Kang et al., 2005). Overexpression of rpoS, the general stress sigma factor that regulates expression of hundreds of E. coli genes (Weber et al., 2005), increased the lycopene production by recombinant E. coli strains (Jin and Stephanopoulos, 2007; Kang et al., 2005). This effect is suggested to be caused by altering cellular oxidative status and preventing...
degradation of lycopene (Bongers et al., 2015). In an approach called global transcription machinery engineering (gTME), \textit{rpoD}, the gene for the principle sigma factor of \textit{E. coli}, was randomly mutated and overexpressed from a plasmid (Alper and Stephanopoulos, 2007). Libraries of \textit{E. coli} cells with mutated forms of \textit{rpoD} were subjected to selection/screening for ethanol tolerance and lycopene production.

Fig. 3. Effect of \textit{sigA} overexpression on accumulation of lycopene (A) and decaprenoxanthin (B) during growth. Growth (A) and lycopene production (B) of strains LYC5(pVWEx1) and LYC5(pVWEx1-sigA) are shown. Growth (C) and decaprenoxanthin production (D) of strains WT(pVWEx1) and WT(pVWEx1-sigA) are shown. Lycopene and decaprenoxanthin production were calculated based on the absorbance of carotenoid extracts at 474 nm for lycopene and 440 nm for decaprenoxanthin and were normalized by cell weight used for extraction. Cultivations were performed in baffled flasks. OD and carotenoid production are given as means of biological triplicates with standard deviations. \textsuperscript{†}Decaprenoxanthin production was calculated as \(\beta\)-carotene equivalent.

Fig. 4. Effect of \textit{sigA} overexpression on production of (A) \(\beta\)-carotene and (B) bisanhydrobacterioruberin (BABR) by recombinant \textit{C. glutamicum} strains. \(\beta\)-carotene production was calculated based on absorbance of carotenoid extracts at wavelength of 454 nm and normalized by cell weight used for extraction. Bisanhydrobacterioruberin production was calculated based on the HPLC analysis of the dominant peak at 471 nm for carotenoid extracts and normalized by cell weight. Cultivations were performed in baffled flasks. Carotenoid production is given as means of biological triplicates with standard deviations. \textsuperscript{†}BABR production was calculated as \(\beta\)-carotene equivalent.
and the rpoD sequences of the best performing mutants were analyzed (Alper and Stephanopoulos, 2007). Therefore, it is conceivable that the gTME approach may be applied in addition to overexpression of sigA for improved carotenoid producing C. glutamicum strains.

SigA is considered to be important for transcription of housekeeping genes, especially in the exponential growth phase (Pátek and Nešvera, 2011; Pfeifer-Sancar et al., 2013) and its transcript is more abundant in the exponential phase than in the stationary phase (Larisch et al., 2007). Some of the genes differentially expressed as consequence of sigA overexpression (Table 2, S1 and S2) were previously shown to be SigB-dependent (Larisch et al., 2007): cg0096, cg291, cg0753, cg0899, cg1109, cg1147, cg1930, cg2704, cg3022 and cg3330. These genes represent a subset of the 95 SigB-dependent genes (Larisch et al., 2007) were a subset of the 95 SigB-dependent genes (Larisch et al., 2007). In addition, two genes known to be differentially expressed in the stationary phase independent of SigB (Larisch et al., 2007) were affected by sigA overexpression: cg1227 and cg2378, two of 16 genes described to belong to this class of genes (Larisch et al., 2007). The finding that sigA overexpression affects at least a subset of known SigB-dependent genes is commensurate with the view that SigB replaces SigA in RNA polymerase holoenzymes during the transition from the exponential to the stationary growth phase. Accordingly, deletion of sigB increased carotenoid production (Fig. 5) and overexpression of sigB decreased lycopene production (Fig. 2). Mutagenesis of sigB has been used for metabolic engineering of C. glutamicum to improve secretion of green fluorescence protein (GFP) and α-amylase (Watanabe et al., 2012). Deletion of sigB, however, is known to impair growth of C. glutamicum at acidic pH (Jakob et al., 2007). Secretion of glutathione S-transferase was improved when using a mutated form of the SigB-dependent promoter of cg3141 (Kim et al., 2016). It is not known whether protein secretion or growth at low pH are also influenced by sigA overexpression.

Genes directly related to carotenogenesis were neither affected by sigA overexpression (Table 2, S1 and S2) nor are they known to be SigB dependent (Larisch et al., 2007). The transcriptional start sites of the carotenoid operons crtE-crg0722-crtB-cry2-cry1b (guanosine 114 nucleotides upstream of the first nucleotide of the ATG start codon) and carnitine uptake operon (guanosine thirteen nucleotides upstream of the first nucleotide of the ATG start codon) of C. glutamicum have been mapped (Heider et al., 2012). Typical SigA-dependent promoter sequence motifs (Pfeifer-Sancar et al., 2013) were found with the consensus −10 hexamer of C. glutamicum promoters (TANNNT) in the −15 to −10 region of crTE and a −35 motif of crTE sharing four identical nucleotides with the −35 consensus (TTGNCAG) hexamer (Heider et al., 2012).

**Table 2**

DNA microarray analysis of genes differentially expressed upon sigA overexpression after 8 h and after 24 h of cultivation.

| Gene IDa | Namea | Function of proteinb | At 8 h | M-valueb | FDRc | At 24 h | M-valueb | FDRc |
|----------|-------|----------------------|--------|----------|-------|--------|----------|-------|
| cg0612   | dkg   | Putative aldo/keto reductase | −1.2   | 5.6E−06  | 1.2   | 6.2E−04 |
| cg0998   | Trypsin-like serine protease | 1.0    | 1.3E−03  | 3.1    | 1.1E−03 |
| cg1096   | Hypothetical protein | 1.0    | 3.3E−04  | 1.3    | 3.6E−02 |
| cg1907   | Putative phosphophanthothienolyl synthetase/decarboxylase | 1.2    | 1.0E−08  | 2.8    | 4.5E−03 |
| cg2030   | Hypothetical protein | 1.2    | 2.0E−02  | 1.2    | 1.7E−02 |
| cg2092   | sigA  | RNA polymerase sigma factor SigA | 3.8    | 6.4E−11  | 5.4    | 3.4E−06 |
| cg2340   | ABC-type transporter, substrate-binding protein | 1.2    | 1.0E−03  | 1.2    | 1.7E−02 |
| cg2341   | Co/Zn/Cd cation efflux transporter | 1.0    | 4.1E−04  | 1.3    | 1.2E−03 |
| cg2636   | cateA | Catechol 1,2-dioxygenase | 1.2    | 1.0E−08  | 1.1    | 1.2E−02 |
| cg3037   | cls   | Cardiolipin synthase | 1.2    | 1.5E−06  | 4.6    | 1.1E−05 |

a Gene ID, gene name and function of proteins are given according to CoryneRegNet (http://coryneregnet.compbio.sdu.dk/v6/index.html) and from previous studies. b Relative RNA amount of sigA overexpressing strain against the control strain with the empty vector was shown as log 2 values (M-values). c FDR represents false discovery rate. 50 µM IPTG were added from the beginning of the cultivation.

**Fig. 5.** Effect of thiamine supplementation and increased addition of PCA on biomass formation (A) and decaprenoxanthin production (B). Cell weights after 48 h of cultivation were determined based on the optical density at 600 nm and normalized to the control cultivation of WT(pVWEx1) with regular CGXII minimal medium (Control). Either 10 µg/L of thiamine (Thiamine) was added or the PCA concentration was raised to 300 mg/L (10xPCA). Decaprenoxanthin production was calculated based on the absorption of carotenoid extracts at wavelength of 440 nm and normalized by cell weight used for extraction. Statistical significance was calculated with paired Student t-test (two-tailed). * and ** represent p-value less than 0.05 and 0.01, respectively. Microscale cultures were performed in Biolector Flower Plates. Carotenoid production is given as means of biological triplicates with standard deviations. †Decaprenoxanthin production was calculated as β-carotene equivalent.
overexpression of sigA was overexpressed at 8 h and 24 h (Table 2). Inactivation and overexpression of sigA have previously been shown to affect the lipid composition of the cellular membrane and temperature sensitivity (Nampoothiri et al., 2002). Thus, it is tempting to speculate that sigA overexpression may change the C. glutamicum cytoplasmic membrane, which in turn may be beneficial for carotenoid accumulation. However, it remains to be shown if modification of the lipid composition of the cytoplasmic membrane influences carotenoid accumulation.

Genes of carbon metabolism were differentially expressed when sigA was overexpressed. The genes for the glycolytic enzyme phosphofructokinase pfkA and for pyruvate quinone dehydrogenase pqo were shown to be upregulated during the transition and in the stationary phase (Ehira et al., 2008), which is downregulated at 24 h as consequence of sigA overexpression in this work (Table S2). This argues against increased availability of the glycolytic intermediates pyruvate and glyceraldehyde 3-phosphate for IPP synthesis as consequence of sigA overexpression.

The catechol 1,2-dioxygenase gene catA was upregulated both at 8 h and 24 h when sigA was overexpressed (Table 2). CatA and the dioxygenase required for PCA degradation are iron containing enzymes. At a 10-fold higher concentration of the iron chelator PCA, more carotenoid was produced (Fig. 4). Transcription of catA is directly repressed by iron homeostasis regulator RipA (Wennerhold et al., 2005). Besides catA, other members of the RipA regulon (Wennerhold and Bott, 2006), namely acn encoding the citric acid cycle iron sulfur cluster-containing enzyme aconitate, the heme biosynthesis gene hemH and ftn coding for an iron storage protein showed differential gene expression as consequence of sigA overexpression (increased RNA levels of acn and hemH, decreased RNA levels for ftn; Table S1 and S2). Iron availability is expected to influence carotenogenesis since two enzymes of the MEP pathway are iron containing proteins: IspG (Lee et al., 2010) and IspH (Gräwert et al., 2004). However, it has to be noted that genes of iron homeostasis tend to change expression levels in growth experiments with CGXII minimal medium due to the limited availability of iron and the iron chelator PCA (Krug et al., 2005; Liebl et al., 1989; Merkens et al., 2005; Unthan et al., 2014).

Carotenogenesis, pyridoxal (vitamin B6) biosynthesis (Hill et al., 1989) and thiamine biosynthesis (Begley et al., 1999) are interrelated as the condensation of glyceraldehyde 3-phosphate and pyruvate to 1-deoxyxylulose-5-phosphate (DXP) catalyzed by Dxs is the first step of the IPP, pyridoxal and thiamine biosynthetic pathways. Interestingly, expression of pyridoxal 5′-phosphate synthase genes (pdxST) was upregulated at 8 h as consequence of sigA overexpression (Table S1). Dxs requires thiamine pyrophosphate as cofactor (Sprenger et al., 1997) and is considered to be one of the rate-limiting steps of the MEP pathway (Estévez et al., 2001) as has also been shown for C. glutamicum (Heider et al., 2014c). Some genes of thiamine synthesis were shown to be upregulated at 8 h or 24 h upon sigA overexpression. Thiamine supplementation has been shown to be beneficial for carotenoid production were reported in Corynebacterium species, for example, Corynebacterium poinsettiae (Starr and Saperstein, 1953) and Corynebacterium michiganense (Saperstein et al., 1954). As shown here, thiamine was only limiting for carotenoid production when sigA was overexpressed (Fig. 4).

5. Conclusion

Taken together, overexpression of the gene for the primary sigma factor SigA in C. glutamicum led to pleiotropic effects, as expected. It has to be noted that it may be difficult to transfer this approach to other production strains and that it may be necessary to repeat screens of sigma factor gene overexpression for each production process. The beneficial effect of sigA overexpression on carotenoid production may be due to these pleiotropic effects involving direct metabolic links to thiamine and pyridoxal biosynthesis (DXP as common biosynthetic precursor) and indirect effects on cardiolipin biosynthesis (cytoplasmic membrane composition) as well as iron availability (iron sulfur cluster-containing enzymes IspG and IspH in the MEP pathway). These pleiotropic effects were shown to be beneficial for overproduction of lycopene, the endogenous decaprenoxanthin, and the non-native carotenoids bisanhydrobacterioruberin and β-carotene.
