

Review

Development and Perspective of Production of Terpenoids in Yeast

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ABSTRACT: Terpenoids are a large class of secondary metabolites known for their remarkable diverse biological activities, making them widely utilized in the pharmaceutical, food, cosmetic, biofuel and agricultural fields. However, the current production of terpenoids heavily relies on plant extraction and chemical synthesis, which brings about concerns regarding in-field, environmental and ecological issues. With the advancements in metabolic engineering and emerging synthetic biology tools, it is now possible to sustainably produce these high value-added terpenoids using microbial chassis. Among them, yeast has emerged as a promising candidate for the heterologous biosynthesis of terpenoids due to its inherent advantages, including robustness, safety, and the availability of sufficient precursor. This review focuses on the diverse strategies employed to enable terpenoids production in yeasts. These strategies encompass metabolic engineering approaches to optimize the mevalonate pathway, protein engineering techniques to improve terpenoid biosynthesis, the applications of organelles compartmentalization, high throughput screening and global approaches for the development of efficient cell factories. Furthermore, this review discusses the future prospects and challenges associated with yeast-based terpenoid production, while also emphasizing guidelines for future studies in this field.

Keywords: Terpenoids; Metabolic engineering; Protein engineering; Subcellular compartments; Global approaches; Yeast



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1. Introduction

Terpenoids represent a large and diverse group of isoprene-based natural products, with over 80,000 distinct structural types identified. Terpenoids are typically classified based on the number of isoprene units they contain, following the isoprene rule: monoterpenoids (C₁₀), sesquiterpenoids (C₁₅), diterpenoids (C₂₀), triterpenoids (C₃₀), tetraterpenoids (C₄₀) (Figure 1) [1]. Many terpenoids exhibit significant pharmacological functions and biological activities. For example, citronellol exhibits properties such as anti-inflammatory, anti-convulsant, anti-hyperalgesia and anti-cholesterol [2]. (+)-Nootkatone possesses activities such as anticancer, antiplatelet aggregation, antimicrobial, and anti-inflammation [3]. Forskolin is known for its anticancer, antiasthmatic and antihypertensive activities [4]. Protopanaxadiol exhibits a wide range of activities including anti-cancer, anti-inflammatory, anti-oxidant, hepatoprotectant, anti-lipogenic, wound-healing, and anti-obesity effects [5]; Violaxanthin has been found to display antiproliferative, anti-inflammatory and photoprotective effects [6]. Furthermore, the structural diversity of terpenoids also enables their extensive applications in biofuels [7], food [8,9], cosmetic [10,11] and agriculture industries [12]. This versatility offers immense potential and commercial value.

Currently, terpenoids are primarily obtained through plant extraction [13] or chemical synthesis [10,11]. However, plant extraction is hindered by long cultivation cycles and susceptibility to environmental factors. In addition, the concentration of synthetic terpenoids in plants is low, and structural analogs are commonly found, which makes it difficult to be separated and purified. On the other hand, chemical synthesis methods for terpenoids require a large amount of organic reagents, which is not environmentally friendly. Furthermore, due to the intricate stereoscopic structures and complexity of most terpenoids, their synthesis and separation using chemical methods are challenging [14]. In recent years, with the rapid development of metabolic engineering and synthetic biology, microorganisms are

gradually being developed as efficient platforms for the green industrial production of terpenoids [15]. Compared to traditional plant extraction and chemical synthesis methods, the utilization of fast-growing microorganisms offers several advantages. Notably, it significantly reduces the production cycle from months to just a few days, while also lowering production costs. Furthermore, the ease of separating the final products is enhanced as the target compounds are enriched within the microbial cell factory. Therefore, the microbial fermentation approach for terpenoid production not only enhances stability, controllability, and productivity but also promotes eco-friendliness [1,16].

Terpenoid biosynthesis involves two main pathways: the mevalonate (MVA) pathway and the methylerythritol phosphate (MEP) pathway [1]. Prokaryotes naturally possess the MEP pathway, while fungi predominantly use the MVA pathway. To date, various microorganisms have been used for producing terpenoids, with yeast being particularly advantageous and focused (Table 1). Yeast possesses several key attributes that make it versatile for terpenoid synthesis. Firstly, it exhibits tolerance to low pH and osmotic stress as well as resistance towards phage infections. More importantly, yeast can offer a wide range of precursors for most terpenoids through its endogenous MVA pathway, including GPP, FPP, GGPP and squalene (Figure 1). In addition, yeast possesses subcellular structures such as the endoplasmic reticulum, Golgi apparatus, and peroxisome, along with post-translational modification mechanisms, which are beneficial for the efficient expression of exogenous enzymes from animals and plants. Moreover, *Saccharomyces cerevisiae*, a commonly used yeast strain, is generally-recognized-as-safe (GRAS) host and has a well-characterized genetic background, making it widely used for food and alcohol production [17]. Considering all these factors, yeast emerges as an ideal host for terpenoids synthesis [1,16].

This review provides a comprehensive examination of the diverse strategies employed to enhance terpenoid production in yeasts. Specifically, we introduce several engineering strategies targeting the mevalonate pathway, as well as protein engineering strategies for terpene biosynthesis. Additionally, we thoroughly analyze the applications of various organelles, assessing both their advantages and limitations in terms of compartmentalization. Furthermore, we explore the development and prospects of global approaches and high-throughput methods for the creation of efficient cell factories. Finally, we extensively discuss the future perspectives and challenges associated with constructing yeast strains capable of producing terpenoids.

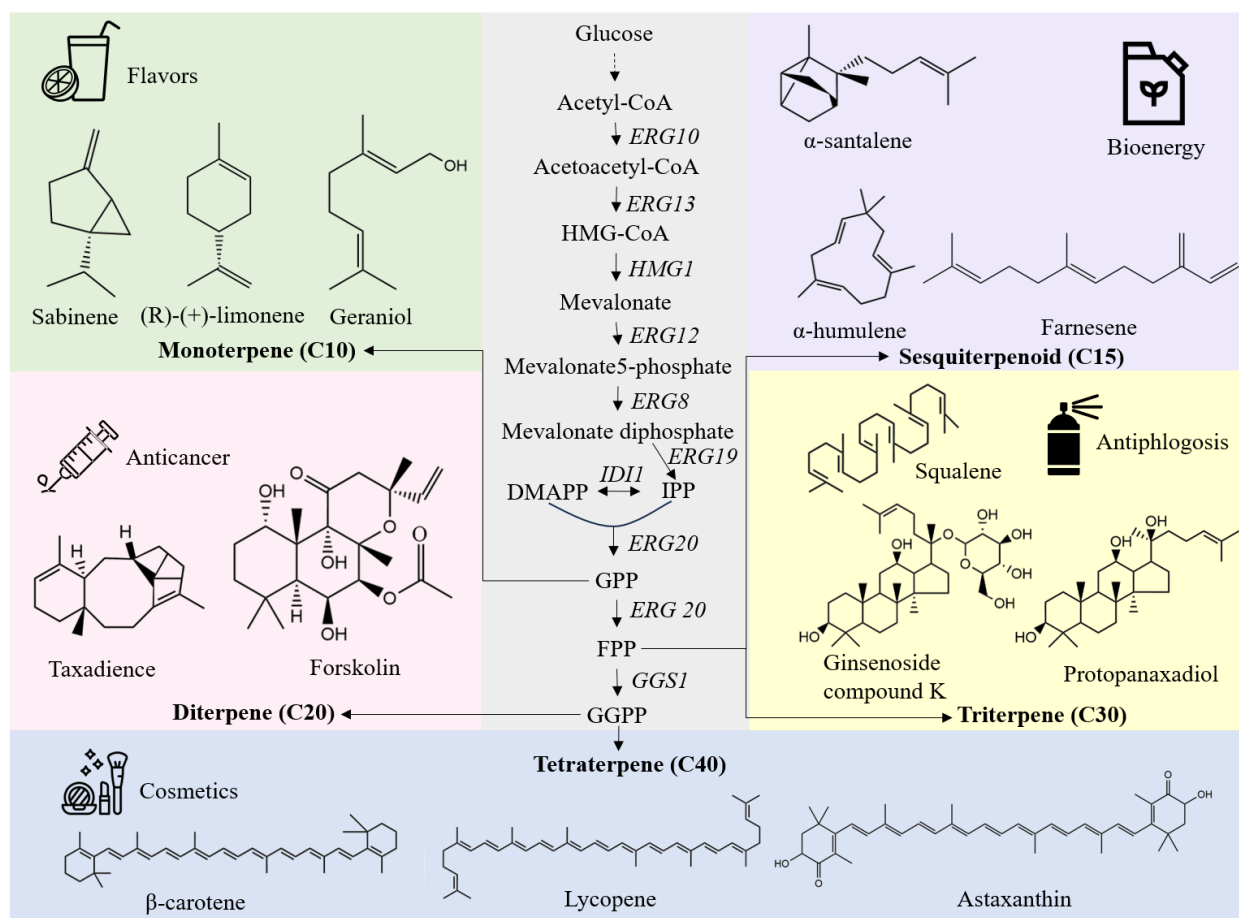


Figure 1. The biosynthesis of different classes of terpenoids in *S. cerevisiae* and *Y. lipolytica* using the mevalonate pathway.

Table 1. Examples of engineering yeasts for production of terpenoids.

Product	Host	Strategy Types *	Modifications and Features	Scale	Titer	Ref.
(R)-(+)-limonene	<i>Saccharomyces cerevisiae</i>	A, C	A combination of peroxisomal engineering and metabolic engineering strategies in the MVA pathway were employed to enhance the production of monoterpenes, resulting in an impressive 125-fold increase compared to cytosolic production.	Fed-batch fermentation	2.6 g/L	[7]
Limonene	<i>Saccharomyces cerevisiae</i>	A	The isopentenol utilization pathway (IUP) was introduced into <i>S. cerevisiae</i> to augment the native mevalonate pathway. The IUP significantly increased the IPP/DMAPP pool by 147-fold and geranylgeranyl diphosphate (GGPP) level by 374-fold compared with the native pathway.	Shake flasks	20.2 mg/L	[18]
Linalool	<i>Saccharomyces cerevisiae</i>	A, B	The efficiency of linalool biosynthesis was improved via overexpression of the complete mevalonate (MVA) pathway and utilizing a geranyl diphosphate (GPP) synthase variant (Erg20p ^{F96W/N127W}). Additionally, the process was accelerated by directed evolution of t67OMcLIS based on a lycopene-indicated high-throughput screening method.	Biphasic shake-flask culture	53.14 mg/L	[11]
Linalool	<i>Yarrowia lipolytica</i>	A, B	The linalool synthetic pathway was constructed by heterologously expressing a linalool synthase gene from <i>Actinidia arguta</i> in <i>Y. lipolytica</i> . Metabolic engineering was utilized to enhance linalool productivity by 76-fold, including overexpression of key genes involved in the mevalonate pathway and the mutant Erg20p ^{F88W-N119W} .	Shake flasks	6.96 mg/L	[19]
Sabinene	<i>Saccharomyces cerevisiae</i>	A, B	The yield of sabinene was improved by an impressive 340-fold through the protein engineering of Erg20p into a geranyl diphosphate synthase, fusion of the synthetic dominant negative Erg20p variant with the terpene synthase, and combined with yeast strain engineering.	Shake flasks	17.5 mg/L	[20]
Geraniol	<i>Saccharomyces cerevisiae</i>	C	Titers of geraniol were increased by 80% via compartmentalizing the biosynthetic enzymes for geraniol within the peroxisomes. Additionally, the deletion of <i>PEX30</i> and <i>PEX32</i> was employed to enhance peroxisome numbers and improve the yeast cells' tolerance to geraniol.	Shake flasks	2.75 mg/L	[10]
Geraniol	<i>Saccharomyces cerevisiae</i>	A, B	Efficient yeast strains for geraniol production were developed by regulating GPP flux and selecting highly efficient geraniol synthases. Through the global regulation of the MVA pathway and the fusion of a geraniol synthase (GES) with Erg20p(F96W-N127W) in <i>S. cerevisiae</i> , geraniol production reached an impressive level of 293 mg/L.	Fed-batch fermentation	293 mg/L	[21]
8-Hydroxygeraniol	<i>Saccharomyces cerevisiae</i>	C	The geraniol pathway was constructed in <i>S. cerevisiae</i> by compartmentalizing the entire MVA pathway, GPP synthase (GPPS), and geraniol synthase (GES) into the mitochondria. This approach resulted in a remarkable 6-fold increase in geraniol production compared to cytosolic producing strains. Furthermore, the integration of geraniol hydroxylase (G8H) into the geraniol-producing strain enabled the production of 8-hydroxygeraniol.	Fed-batch fermentation	227 mg/L	[22]
Citronellol	<i>Saccharomyces cerevisiae</i>	B	Titers of citronellol were significantly enhanced by 140-fold through several strategies compared to the starting strain. These included replacing the wild farnesyl diphosphate synthase (Erg20p) with the mutant Erg20g ^{F96W} to enhance geranyl diphosphate (GPP) production, improving the catalytic activity of reductases to increase geraniol production, employing synthetic protein scaffolds and protein fusion to colocalize four sequential enzymes for better substrate channeling, and deleting the intermediate degradation pathway gene <i>ATF1</i> .	Fed-batch fermentation	8.3 g/L	[2]
Cineole	<i>Saccharomyces cerevisiae</i>	A	To enhance plant monoterpene and sesquiterpene production, recyclable integration cassettes were utilized to target specific genes involved in the yeast sterol biosynthetic pathway, namely <i>HMG2</i> , <i>ERG20</i> and <i>ID11</i> , with several-fold improvements observed. Notably, the combination of chromosomal integrations and high copy plasmids expressing the two genes, <i>ID11</i> and <i>HMG2</i> (K6R) variant, led to an impressive 55-fold increase in cineole production compared to the wild-type parental cells.	Shake flasks	99 mg/g DW	[23]
α -Santalene	<i>Saccharomyces cerevisiae</i>	A, B	By compartmentalizing both the complete MVA pathway and the α -santalene synthesis pathway within the yeast mitochondria, a remarkable 3.7-fold improvement in α -santalene production was achieved.	Biphasic shake-flask culture	41 mg/L	[24]
(+)-Nootkatone	<i>Saccharomyces cerevisiae</i>	A, B	The biosynthesis pathway for (+)-nootkatone was successfully established in <i>S. cerevisiae</i> through the overexpression of several key enzymes for biosynthesis of (+)-nootkatone, and combined with various MVA pathway engineering strategies, such as the fusion expression of CnVS and Erg20p, overexpression of a truncated form of the rate-limiting enzyme 3-hydroxy-3-methylglutaryl-CoA (HMG-CoA) reductase (tHMG1), and downregulation of the squalene synthase enzyme (Erg9p).	Shake flasks	59.78 mg/L	[3]
(+)-Nootkatone	<i>Saccharomyces cerevisiae</i>	A, B, C, D	The candidate CYP71D55 (HPO) mutants were efficiently screened through a high-throughput screening (HTS) method, which relied on the color reaction between carbonyl compounds and 2,4-dinitrophenylhydrazine (DNPH). To further enhance the production of (+)-nootkatone, high-yield strains were developed by optimizing the pairing of HPO with cytochrome P450 reductase (CPR), and by employing a combination of metabolic engineering and ER engineering techniques.	Fed-batch fermentation	2.39 g/L	[25]
Farnesene	<i>Saccharomyces cerevisiae</i>	A	The central carbon metabolism of <i>S. cerevisiae</i> was effectively rewired through the incorporation of four non-native metabolic reactions. This rewiring enabled the biosynthesis of cytosolic acetyl-CoA with a reduced ATP requirement and minimized carbon loss through CO ₂ -emitting reactions, resulting in an improved pathway redox balance. Which exhibited a remarkable 25% increase in farnesene production, using the same quantity of sugar as control strains, while consuming 75% less oxygen.	Fed-batch fermentation in industrial bioreactors	130 g/L	[26]
α -Farnesene	<i>Yarrowia lipolytica</i>	A, E	Overexpression of all genes involved in the MVA pathway and α -farnesene synthase genes through non-homologous end-joining (NHEJ) mediated genome integration for enhancing α -farnesene production. Additionally, fermentation conditions were optimized, resulting in a significant increase in α -farnesene production, reaching an impressive 25.55 g/L.	Fed-batch fermentation	25.55 g/L	[27]

	α -Humulene	<i>Yarrowia lipolytica</i>	A, C	A series of multilevel metabolic engineering strategies were employed to produce α -humulene. These included compartmentalizing the α -humulene biosynthesis pathway within the peroxisome, enhancing ATP production by overexpressing the adenine nucleotide transporter gene, increasing acetyl-CoA supply through upregulation of β -oxidation and disruption of the competing glyoxylate cycle that consumes acetyl-CoA, and optimizing the gene copy numbers of rate-limiting enzymes.	Fed-batch fermentation	3.2 g/L	[28]
	Nerolidol	<i>Saccharomyces cerevisiae</i>	A, B	The production of nerolidol was significantly enhanced through the implementation of two key strategies. Firstly, the core mevalonate pathway was enhanced to optimize the synthesis of the precursor farnesyl pyrophosphate (FPP). Secondly, carbon flux was redirected from squalene to nerolidol by linking the degradation peptide Cln2p ^{PEST} to the C-terminus of Erg9p.	Shake flasks	~100mg/L	[29]
	β -Caryophyllene	<i>Saccharomyces cerevisiae</i>	A, B	An efficient yeast cell factory for β -caryophyllene production was developed through the implementation of combined engineering strategies. These strategies included enhancing the activity of β -caryophyllene synthase (CPS) through enzyme screening and directed evolution. Additionally, genes related to β -alanine metabolism and the mevalonate (MVA) pathway were overexpressed to enhance the synthesis of the precursor. Furthermore, an ATP-binding cassette transporter gene variant, <i>STE6^{T1025N}</i> , was introduced to improve the transmembrane transport of β -caryophyllene and mitigate potential cytotoxicity. The combined engineering of CPS and the yeast chassis resulted in a remarkable 2.93-fold increase in the yield of β -caryophyllene.	Fed-batch fermentation	594.05 mg/L	[30]
	Cubebol	<i>Saccharomyces cerevisiae</i>	E	Through the utilization of the metabolic model of <i>S. cerevisiae</i> , assessed using OptGene as the modeling framework and minimization of metabolic adjustments (MOMA) as the objective function, the deletion of NADPH-dependent glutamate dehydrogenase encoded by <i>GDH1</i> was identified as the most effective target gene for enhancing sesquiterpene biosynthesis. This genetic modification resulted in an impressive approximately 85% increase in cubebol titer, showcasing its significant impact on the production of this compound.	Batch fermentation	8.4 mg/L	[31]
	α -Bisabolene	<i>Yarrowia lipolytica</i>	A, E	The production of α -bisabolene was achieved by introducing the α -bisabolene synthase from <i>Abies grandis</i> through heterologous expression. To enhance the production, all endogenous mevalonate pathway genes were overexpressed and heterologous multidrug efflux transporters were introduced to improve the export of α -bisabolene from the cells.	Shake flasks	973.1 mg/L	[32]
	Geranylgeraniol	<i>Saccharomyces cerevisiae</i>	A	A combination of metabolic engineering strategies was adopted to effectively enhance the yield of geranylgeraniol. Firstly, exogenous GGPP synthase genes were overexpressed to enhance GGPP synthesis. Secondly, the metabolic flux of the MVA pathway was improved by deleting <i>ROX1</i> gene. Additionally, the supply of isopentenyl pyrophosphate (IPP) was enhanced by overexpressing the <i>AtIPK</i> gene and supplementing with isoprenol.	Fed-batch fermentation	5.07 g/L	[33]
	Miltiradiene	<i>Saccharomyces cerevisiae</i>	A, B	A push and pull strategy was implemented for improving miltiradiene production. This strategy involved two key approaches: increasing the supply of the precursor GGPP and enhancing the catalytic capacity of diterpene synthases through N-terminal truncation and fusion expression.	Fed-batch fermentation	3.5 g/L	[34]
Diterpene	Forskolin	<i>Saccharomyces cerevisiae</i>	A, B, C	Multiple metabolic engineering strategies were implemented to enhance the metabolic flow to forskolin, contains improving the catalytic efficiency of CfCYP76AhS by optimizing the adaptations between CfCYP76AhS, t66CfCPR and t30AaCYB5, enhancing the MVA pathway, regulating the supply of cofactors and expanding the ER area by overexpressing <i>INO2</i> .	Fed-batch fermentation	79.33 mg/L	[4]
	Sclareol	<i>Saccharomyces cerevisiae</i>	A, E	The modular approach was using to globally rewire the cellular metabolism for improving sclareol production to 11.4 g/L in <i>S. cerevisiae</i> . The first module focused on the central metabolism, which ensured an efficient supply of acetyl-CoA and cofactor NADPH. The second module encompassed the isoprenoid biosynthesis pathways, which were optimized to enable the efficient synthesis of sclareol. Lastly, the regulation factor module was designed to fine-tune and control the metabolic processes involved in sclareol production.	Fed-batch fermentation	11.4 g/L	[35]
	Taxadiene	<i>Saccharomyces cerevisiae</i>	B	The titers of taxadiene were improved by an impressive 22-fold through the multi-copy chromosomal integration of taxadiene synthase gene (<i>TASY</i>) and the combination with fusion expression of maltose binding protein (MBP) tagged <i>TASY-ERG20^{F96C}</i> .	Shake flasks	129 mg/L	[36]
	Squalene	<i>Saccharomyces cerevisiae</i>	A, C	A combinatorial strategy of cytoplasmic and mitochondrial engineering was established to alleviate the metabolic burden caused by the compartmentalized MVA pathway in mitochondria for improving cell growth and squalene production.	Fed-batch fermentation	21.1 g/L	[37]
	Squalene	<i>Saccharomyces cerevisiae</i>	A, C	The entire squalene synthesis pathway was compartmentalized into peroxisomes and achieved a 138-fold improvement in squalene titer.	Fed-batch fermentation	11.00 g/L	[38]
	Squalene	<i>Saccharomyces cerevisiae</i>	A, C	The production of squalene was increased by 71-fold through the engineering of the yeast <i>S. cerevisiae</i> . This was achieved by overexpressing <i>INO2</i> , a gene involved in the regulation of phospholipid biosynthesis, which led to the expansion of ER.	Shake flasks	175 mg/L	[39]
Triterpene	Protopanaxadiol	<i>Saccharomyces cerevisiae</i>	A, B	A modular engineering strategy was employed to optimize a multigene pathway for the production of protopanaxadiol (PPD) in <i>S. cerevisiae</i> , and the yield of PPD was significantly improved to 66.55 mg/L/OD600. Firstly, all genes in the MVA pathway were combinatorially overexpressed. Additionally, the metabolic balance in the triterpene biosynthesis module was optimized. Furthermore, the expression of the lanosterol synthase gene (<i>ERG7</i>) in the sterol biosynthesis module was fine-tuned using the TetR-TetO gene regulation system. Finally, to increase the supply of cytoplasmic acetyl-CoA, a mutant form of <i>Salmonella</i> acetyl-CoA synthetase (<i>ACS_{scl641F}</i>), was overexpressed.	Fed-batch fermentation	8.09 g/L	[40]
	Protopanaxadiol	<i>Saccharomyces cerevisiae</i>	A	A yeast cell factory for PPD production was engineered through the careful manipulation of enzyme expressions and the improvement of cofactor NADPH availability. This was achieved by utilizing different combinations of promoters (<i>P_{GPD}</i> , <i>P_{CCW12}</i> , and <i>P_{ADH2}</i>) and rerouting the redox metabolism, which resulted in more than an 11-fold increase in PPD titer over the initially constructed strain.	Flask fermentation	6.01 mg/L	[41]

Protopanaxadiol	<i>Saccharomyces cerevisiae</i>	A	In the engineered <i>S. cerevisiae</i> harboring the biosynthetic pathway for PPD, the suppression of lanosterol synthase using CRISPRi led to a remarkable increase in PPD production. This approach resulted in a 14.4-fold increase in PPD production during shake-flask fermentation and a 5.7-fold increase in a long-term batch-fed fermentation.	Batch-fed fermentation	294.5 mg/L	[5]	
Protopanaxadiol	<i>Saccharomyces cerevisiae</i>	C	The PPD pathway was successfully constructed and optimized in <i>S. cerevisiae</i> by genetically engineering the expression of three peroxisome biogenesis-related peroxins (Pex11p, Pex34p, and Atg36p). This peroxisome engineering approach resulted in a significant improvement in PPD yield, with a remarkable 78% increase compared to the wild-type strain.	Batch fermentation	4.1 mg/L	[42]	
Ginsenoside compound K	<i>Saccharomyces cerevisiae</i>	C	The biosynthetic pathway of ginsenoside compound K (CK) was successfully reconstituted in a PPD-producing chassis strain. This was achieved by targeting the cytochrome P450 enzyme protopanaxadiol synthase (PPDS), which is normally localized in the endoplasmic reticulum (ER), to lipid droplets (LDs) using the yeast LDs membrane protein PLN1. The volume of LDs was also increased by integrating rate-limiting enzyme genes involved in triacylglycerol (TAG) biosynthesis into the chromosomal GAL80 site. This strategic integration of genes led to a 4.4-fold increase in CK production compared to the native ER-expression strategy.	Fed-batch fermentation	4.88 g/L	[43]	
Ginsenoside	<i>Saccharomyces cerevisiae</i>	A, B	Cell factories for producing ginsenoside Rh2 were successfully established using a high PPD-producing chassis strain. This was achieved by overexpressing all genes involved in the MVA pathway and optimizing the expression levels of UGTPg45 in yeast. Additionally, the activity of UGTPg45 was improved through protein engineering using directed evolution techniques. Furthermore, efforts were made to identify novel UDP-glycosyltransferases (UGTs) with higher C3-OH glycosylation efficiencies from other plant species.	Fed-batch fermentation	2.25 g/L	[44]	
α -Amyrin	<i>Saccharomyces cerevisiae</i>	A, B	The production of α -amyrin was achieved through a combination of protein engineering and metabolic engineering approaches. The catalytic activity of α -amyrin synthase (MdOSC1) was optimized using a semi-rational protein engineering strategy that involved remodeling MdOSC1 to enhance its efficiency. Meanwhile, key genes involved in the MVA pathway (<i>tHMG1</i> , <i>ERG20</i> , <i>ERG9</i> and <i>ERG1</i>) were overexpressed. Furthermore, the overexpression of <i>DGA1</i> , which encodes diacylglycerol acyltransferase, expanded the intracellular storage capacity, facilitating the accumulation of α -amyrin. Finally, the yield of α -amyrin achieved 1.11 g/L by fed-batch fermentation, which was 106-fold higher than that of the original strain under the same conditions.	Fed-batch fermentation	1.11 g/L	[45]	
β -Amyrin	<i>Saccharomyces cerevisiae</i>	A	A combination of endogenous and heterologous acetyl-CoA synthesis pathways was employed to enhance β -amyrin production. The optimal strategy involved the synergistic coupling of the PK/PTA and A-ALD pathways, in conjunction with NADH-HMGR. This coupling ensured a balanced supply of redox cofactors, resulting in lower energy consumption and improved glucose utilization, which led to a remarkable 200% increase in β -amyrin production. Furthermore, the competing pathway of acetyl-CoA was disrupted led to a 330% increase in β -amyrin production as compared to the original strain.	Fed-batch fermentation	279 mg/L	[46]	
β -Amyrin	<i>Saccharomyces cerevisiae</i>	A, E	The N-degron-mediated protein degradation strategy was utilized to decrease the level of Erg7p by lowering its half-life with significantly increasing the supply of 2,3-oxidosqualene for triterpenoid biosynthesis. In addition, flux distribution between the ergosterol synthesis pathway and β -amyrin synthesis pathway was optimized by overexpressing all the genes in the pathway from acetic acid to squalene. To separate the growth stage from the production stage, the GAL-regulation system was modified. Through the implementation of these metabolic engineering strategies, the yield of β -amyrin was improved by a remarkable 47-fold.	Fed-fermentation	4.22 g/L	[47]	
Ganodermic acid	<i>Saccharomyces cerevisiae</i>	D	To obtain strains capable of producing ganoderic acid 3-hydroxy-lanosta-8,24-dien-26-oic acid (GA-HLDOA), a screening process utilizing fluorescence-activated cell sorting (FACS) was employed. Transformants integrated with the CYP5150L8 and iGLCPR expression cassettes at the rDNA loci of <i>S. cerevisiae</i> were screened using FACS, resulting in the identification of strain SC62. To further enhance the production of valuable compounds, the gene of CYP512W2 was expressed in strain SC62. This genetic modification led to the stable production of ganoderic acid Y (GA-Y) and ganodermic acid Jb (GA-Jb), both of which are highly desirable compounds.	Shake flasks	56.44mg/L	[48]	
Ursolic acid	<i>Saccharomyces cerevisiae</i>	A, C	The <i>de novo</i> syntheses of ursolic acid (UA) were achieved via the heterologous expression and optimization of CrAS, CrAO and AtCPR1. Metabolic flux was redirected by increasing the cytosolic acetyl-CoA level and tuning the copy numbers of <i>ERG1</i> and <i>CrAS</i> . Furthermore, the titer of UA was increased by ~50 times via the lipid droplet compartmentalization of CrAO and AtCPR1 alongside the strengthening of the NADPH regeneration system.	Fed-batch fermentation	1.13 g/L	[49]	
Medicagenic acid	<i>Saccharomyces cerevisiae</i>	C	The disruption of the phosphatidic acid phosphatase (encoding by <i>PAH1</i>) through CRISPR/Cas9 results in a remarkable proliferation of the endoplasmic reticulum (ER) and a significant accumulation of triterpenoids. Compared to the wild-type starter strain, the <i>pah1</i> strain exhibited a substantial increase in the accumulation of oleanane-type saponin β -amyrin and medicagenic acid, with levels elevated by 8-fold and 6-fold, respectively.	Shake flasks	27.1 mg/L	[50]	
Tetraterpene	Lycopene	<i>Saccharomyces cerevisiae</i>	A, C	By employing a systematic approach combining traditional engineering methods, the regulation of lipid oil-triacylglycerol (TAG) metabolism was optimized to enhance the production of lycopene. This involved several key strategies, including enhancing the MVA pathway, increasing the availability of acetyl-CoA precursors, facilitating the regeneration of the redox cofactor NADPH, fine-tuning the expression of lycopene-synthesis genes, disrupting functional bypass genes, overexpressing key genes involved in fatty acid synthesis and TAG production, modulating the fatty acyl composition of TAG, and regulating the size of lipid droplets.	Fed-batch fermentation	2.37 g/L	[8]

Lycopene	<i>Yarrowia lipolytica</i>	E	To identify suitable chemicals for lycopene production, a combination of flux balance analysis (FBA) and Plackett-Burman design was employed. This approach was applied to a metabolically engineered strain of <i>Y. lipolytica</i> that harbored <i>crtE</i> , <i>crtB</i> , and <i>crtI</i> genes. By utilizing FBA and Plackett-Burman design, the researchers were able to screen a range of chemicals and identify those that were most effective for lycopene production.	Fed-batch fermentation	242 mg/L	[51]
β -Carotene	<i>Saccharomyces cerevisiae</i>	C	Through the regulation of lipid metabolism pathways, a significant 2.4-fold improvement in β -carotene production was achieved. This improvement was achieved through the overexpression of sterol ester synthesis genes (<i>ARE1</i> and <i>ARE2</i>), as well as the deletion of phosphatidate phosphatase (PAP) genes (<i>PAH1</i> , <i>DPPI1</i> , and <i>LPP1</i>).	Shake flasks	8.98 mg/g DCW	[52]
β -Carotene	<i>Saccharomyces cerevisiae</i>	A	The initial carotenogenic strain was constructed via expressing <i>crtE</i> , <i>crtYB</i> , <i>crtI</i> , <i>HMG1</i> and another copy of <i>crtI</i> . Subsequently, <i>ZWF1</i> and <i>POS5</i> were individually overexpressed in the carotenoid-producing <i>S. cerevisiae</i> strain, and the final yield of β -carotene increased by 18.8% and 65.6%, respectively.	Shake flasks	0.53 mg/L	[53]
β -Carotene	<i>Yarrowia lipolytica</i>	D, E	High- β -carotene-producing strains were screened in a high-throughput manner using a fluorescence-activated cell-sorting approach from genome-scale mutagenesis libraries. In particular, the disruption of specific genes involved in fatty acid oxidation, lipid composition, and sterol transcriptional regulation was found to have a significant impact on increasing β -carotene production.	Fed-batch fermentation	9.4 g/L	[54]
Violaxanthin	<i>Saccharomyces cerevisiae</i>	A, B	Violaxanthin production was achieved by integration of the β -xanthophyll pathway in a β -carotenogenic yeast strain. Subsequent stepwise improvements, such as enzyme truncation, co-expression of redox partners, and adjustment of gene copy numbers, further enhanced violaxanthin yields, resulting in an impressive 18-fold increase.	Shake flasks	7.3 mg/g DCW	[6]
Lutein	<i>Saccharomyces cerevisiae</i>	B, C, E	Heterologous biosynthesis of lutein in <i>S. cerevisiae</i> was successfully achieved through combinatorial engineering, considering both temporal and spatial aspects. This was accomplished by utilizing a temperature-responsive Gal4M9- <i>P_{GAL1}</i> regulation system, which allowed for sequential expression of two lycopene cyclases, as well as the re-localization of the rate-limiting ϵ -cyclase to the plasma membrane. Furthermore, the metabolic flux was significantly enhanced through protein engineering of the ϵ -cyclase, employing directed evolution and saturation mutagenesis techniques.	Shake flasks	110.4 μ g/L	[9]
Astaxanthin	<i>Yarrowia lipolytica</i>	B, C	The fusion expression of β -carotene ketolase (CrtW) and β -carotene hydroxylase (CrtZ), along with the targeted localization of the astaxanthin pathway to subcellular organelles (lipid bodies, endoplasmic reticulum, and peroxisomes), led to a 141-fold increase in astaxanthin production compared to the initial strain.	Fed-batch fermentation	858 mg/L	[55]
Astaxanthin	<i>Yarrowia lipolytica</i>	A	Twelve different <i>Y. lipolytica</i> isolates were firstly screened for β -carotene production by introducing two genes, <i>crtYB</i> and <i>crtI</i> , from the red yeast <i>Xanthophyllomyces dendrorhous</i> . The β -carotene titer was further increased by optimizing the activities of 3-hydroxy-3-methylglutaryl-coenzyme A reductase and geranylgeranyl diphosphate synthase, and downregulating the competing squalene synthase. Subsequently, introducing β -carotene ketolase (CrtW) from <i>Paracoccus sp.</i> N81106 and hydroxylase (CrtZ) from <i>Pantoea ananatis</i> , and optimizing the copy numbers of <i>crtZ</i> and <i>crtW</i> , with achieving a yield of 3.5 mg/g DCW (54.6 mg/L) of astaxanthin in a microtiter plate cultivation.	Microtiter plate cultivation	54.6 mg/L	[56]
Astaxanthin	<i>P. rhodozyma</i>	E	A combination of metabolomics and transcriptomics approaches was employed to analyze the differential metabolites and expressed genes involved in astaxanthin synthesis induced by gibberellic acid (GA). The results showed that GA induction significantly increased fatty acid metabolism and ABC transporters, as evidenced by the upregulation of fatty acid desaturase and ABC transporter genes. This led to an elevation in the contents of unsaturated fatty acids and a subsequent 77.12% improvement in astaxanthin production.	Fed-batch fermentation	539.8 μ g/g	[57]

* Strategies and features can be mainly divided into five types: A represents metabolic engineering strategies for improving the mevalonate pathway; B represents protein engineering strategies for improving terpenoid biosynthesis; C represents the applications of organelles compartmentalization for improving terpenoid biosynthesis; D represents high throughput screening approaches for improving terpenoid biosynthesis; E represents the global approaches for improving terpenoid biosynthesis.

2. Metabolic Engineering Strategies for Improving the Mevalonate Pathway

The mevalonate (MVA) pathway in yeasts serves as the source of precursors for the synthesis of terpenoids. Terpenoids are synthesized mainly from two C5 precursors [23], namely isopentenyl diphosphate (IPP) and dimethylallyl diphosphate (DMAPP). As shown in Figure 2, IPP and DMAPP result from the multi-step conversion of acetyl-CoA through the MVA pathway. Notably, the conversion of HMG-CoA to mevalonate by HMG-CoA reductase represents the rate-limiting step of the MVA pathway [1,58]. These two five-carbon (C5) skeletons are then ligated in diverse proportions to produce different pyrophosphate precursors, such as geranyl pyrophosphate (GPP), farnesyl pyrophosphate (FPP) and geranylgeranyl pyrophosphate (GGPP). These precursors serve as substrates for a series of rearrangement, cyclization, methylation, and other reactions, ultimately leading to the diverse array of natural terpenoids (Figure 2) [15]. Considering the significance of the MVA pathway, the metabolic mechanism of the endogenous MVA pathway has been completely elucidated in yeast. Consequently, one of the primary approaches to significantly enhance the efficient synthesis of terpenoids is to focus on augmenting the flux of this pathway through the implementation of metabolic engineering techniques.

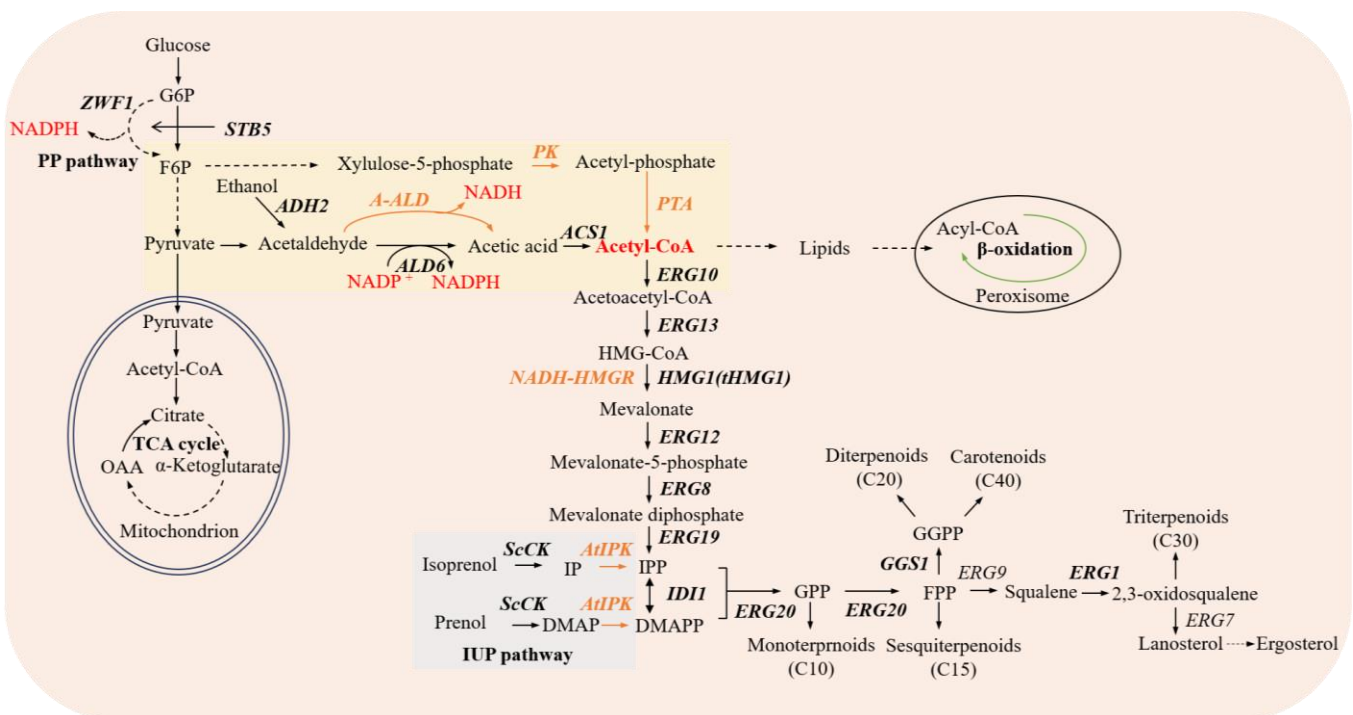


Figure 2. Manipulations of the mevalonate pathways for optimizing biosynthesis of terpenoids in yeast. Overexpressed endogenous genes are shown in black bold. Additionally, the introduction of heterologous acetyl-CoA synthesis pathways and the synthetic isopentenol utilization pathway (IUP) in yeast is represented in orange. The solid and dotted arrows indicate single and multiple steps, respectively. *PK*, phosphoketolase gene from *Leuconostoc mesenteroides*; *PTA*, phosphotransacetylase gene from *Clostridium kluyveri*; *A-ALD*, acetylating acetaldehyde dehydrogenase gene from *E. coli*; *NADH-HMGR*, NADH-dependent HMG-CoA reductase variant gene from *Silicibacter pomeroyi*; *AiIPK*, isopentenyl phosphate kinase gene from *Arabidopsis thaliana*; *ScCK*, choline kinase gene; *ZWF1*, glucose-6-phosphate dehydrogenase gene; *ADH2*, alcohol dehydrogenase gene; *ALD6*, acetaldehyde dehydrogenase gene; *ACS1*, acetyl-CoA synthase gene; *ERG10*, acetyl-CoA thiolase gene; *ERG13*, HMG-CoA synthase gene; *HMG1*, 3-hydroxy-3-methyl glutaryl coenzyme A reductase gene; *tHMG1*, truncated HMG1; *ERG12*, mevalonate kinase gene; *ERG8*, phosphomevalonate kinase gene; *ERG19*, mevalonate diphosphate decarboxylase gene; *IDII*, IPP isomerase gene; *ERG20*, geranyl/farnesyl diphosphate synthase gene; *GGS1*, geranylgeranyl diphosphate synthase gene; *ERG9*, squalene synthase gene; *ERG1*, squalene monooxygenase gene; *ERG7*, lanosterol synthase gene.

2.1. Improving the Supply of IPP and DMAPP Precursors

Enhancing precursor supply is one of the most common strategies to improve production of biochemicals. In the case of terpenoid production, overexpression of the key enzymes in the MVA pathway, such as HMG-CoA reductase (HMGR), is a widely used approach for improving IPP and DMAPP supply [1,58]. HMGR is a membrane-bound protein localized in the endoplasmic reticulum (ER), consisting of an N-terminal membrane anchor and a C-terminal catalytic domain [58]. Overexpression of either intact or N-terminal truncated HMGR has been shown to enhance the production of diverse terpenoids [1,15,56,59,60]. In previous studies, an N-terminal-truncated HMGR was usually suggested more effective, because the N-terminal truncation of HMGR can prevent self-degradation and make it more

stable in the cytoplasm [1,15,59]. However, Kildegaard et al. found that the native HMGR is superior for β -carotene production in *Y. lipolytica* [56]. IPP isomerase (IDI) is responsible for the interconversion between IPP and DMAPP and plays a crucial role in the flux distribution of GPP and FPP [59]. Overexpression of IDI is also a common strategy to enhance the biosynthesis of terpenoids. For example, over-expression of *IDI1* enhanced cineole by 24-fold in *S. cerevisiae* [23]. Besides HMGR and IDI, most of the pathway genes in the MVA pathway (*ERG10*, *ERG13*, *ERG12*, *ERG8* and *ERG19*) have also been over-expressed to increase production of terpenoids [32,44]. Such as, overexpression of all genes in MVA pathway improved the bisabolene production with 251-fold, 29-fold and 72-fold enhancement in α -bisabolene, β -bisabolene and γ -bisabolene production, respectively [32].

Furthermore, introducing heterologous synthesis pathways for improving IPP and DMAPP precursors is also a new strategy for yeasts to synthesize terpenoids. The isopentenol utilization pathway (IUP) was reported to provide universal C5 precursors for terpenoid synthesis, bypassing the lengthy and strictly regulated natural MVA pathway [15,61]. This pathway utilizes choline kinase (*ScCK*) from *S. cerevisiae* and isopentenyl phosphokinase (*AtIPK*) from *A. thaliana* to directly phosphorylate the secondary substrate isoprenoid alcohol into IPP and DMAPP at the expense of two molecules of ATP (Figure 2) [62]. Compared with the MVA pathway, the IUP pathway has the advantages of fewer steps and a higher metabolic flow [18,62]. For example, the introduction of a two-step isopentenol utilization pathway (IUP) into *S. cerevisiae* resulted in a 147-fold increase in the IPP/DMAPP pool compared to the native MVA pathway [18]. Cofeeding isoprenol and prenol further enhanced the geranyl diphosphate (GPP) content for monoterpene biosynthesis and elevated the GGPP level by 374-fold.

2.2. Improving the Supply of Acetyl-CoA and Cofactors

Increasing the supply of cytosolic acetyl-CoA is supposed to be an effective strategy to improve terpenoids yield [15]. In native *S. cerevisiae*, cytosolic acetyl-CoA is synthesized from the pyruvate dehydrogenase (PDH) bypass or during ethanol reoxidation, which involves pyruvate decarboxylase (PDC), alcohol dehydrogenase (ADH), aldehyde dehydrogenases (ALD) and acetyl-CoA synthetase (ACS) [1,46]. Common strategies for enhancing the cytosolic acetyl-CoA pool include manipulating the PDH-bypass, and removing the competitive glyoxylate shunt pathway by knocking out *MLS1* (encode malate synthase) and *CIT2* (encode peroxisomal citrate synthase) [1,46,49,63]. However, the endogenous PDH bypass for acetyl-CoA synthesis exhibits low efficiency owing to the high energy cost with the ATP-requiring reactions catalyzed by ACS (2 ATP for one acetyl-CoA formation) [1,46]. To address this limitation, Meadows et al. rewired the central carbon metabolism in *S. cerevisiae* using four non-native metabolic reactions by acetaldehyde dehydrogenase acylating (ADA, namely A-ALD), xylulose-5-phosphate (X5P)-specific phosphoketolase (xPK), phosphotransacetylase (PTA) and NADH-consuming HMG-CoA reductase (NADH-HMGR) (Figure 2) [26]. This rewiring generated cytosolic acetyl-CoA with lower ATP cost, reduced carbon loss to CO₂-emitting reactions and improved pathway redox balance, and resulted in a 25% increase in the production of β -farnesene while requiring 75% less oxygen. In another work, Liu et al. compared the effects of different acetyl-CoA synthesis pathways for β -amyrin synthesis [46]. Among these pathways, the combination of PK/PTA and A-ALD pathways together with NADH-HMGR has been found to result in a highly efficient glucose conversion to β -amyrin, achieving a high attainable yield of 0.296 g/g glucose. This combination allows for the satisfaction of ATP consumption in the PK/PTA pathway through the reoxidation of NADH generated in the A-ALD pathway. As a result, a more balanced redox cofactors and a much lower ATP consumption is achieved than any single pathway coupled β -amyrin synthesis.

The redox cofactor NADPH plays a critical role in the metabolism of *S. cerevisiae*. In addition to replacing the NADPH-dependent HMGR with the NADH-dependent NADH-HMGR to alter the cofactor dependence, replacing the NADH-generating enzyme with a NADPH-generating enzyme can also enhance the availability of NADPH (Figure 2). For example, the availability of NADPH was enhanced by 1.3-fold through replacing a NADH-generating enzyme (Ald2p) with a functionally equivalent enzyme (Ald6p) that generates NADPH instead, and protopanaxadiol (PPD) production was improved 4.5-fold in *S. cerevisiae* [41]. Furthermore, the supply of NADPH can be increased by promoting the pentose phosphate (PP) pathway, where the rate-limiting step is catalyzed by glucose-6-phosphate dehydrogenase (encoded by *ZWF1* gene in yeast) [53,64]. In a recent reported, it has been shown that overexpression of the *ZWF1* gene increased NADPH concentrations by 1.4-fold in *S. cerevisiae* [53]. This led to a 59.9% improvement in lycopene production and an 18.8% improvement in β -carotene production. In parallel, the *STB5* gene has been reported to be a basal regulator of the pentose phosphate (PP) pathway, and its overexpression increased NADPH concentrations by 1.2-fold in *S. cerevisiae* [41]. Moreover, NADH kinase is capable of directly phosphorylating NADH to NADPH without directly impacting the metabolic pathway. Overexpression of *POS5*, which encodes a mitochondrial NADH kinase that converts NADH to NADPH, significantly enhanced lycopene yield by 81.4% and β -carotene yield by 65.6% in *S. cerevisiae* [53]. When

controlled by P_{GAL7} , the overexpression of $POS5$ in *S. cerevisiae* improved the ursolic acid (UA) and oleanolic acid (OA) titers to 692.3 and 253.4 mg/L, respectively [49]. Consequently, this study offers a novel approach to construct cell factories for efficient terpenoid synthesis, particularly those that necessitate substantial quantities of NADPH.

2.3. Inhibition of Endogenous Competition Pathway

The precursor pool of GPP, FPP, and GGPP can also be promoted by down-regulating the downstream genes or genes in competitive pathways. For example, $ERG9$, which encodes squalene synthase responsible for squalene synthesis, is a key node in competition with FPP for the production of sesquiterpenoid [3], diterpenoid [34] and tetraterpenoid [56]. To increase the supply of FPP pool for the production of diterpenoid and tetraterpenoid, the native promoter of $ERG9$ was replaced it with a weak promoter P_{HXT1} or truncated, which has been shown to improve sclareol production to 310.5 mg/L [35] and β -carotene titer by 2.5 fold [56], respectively. In parallel, Peng et al. linked degradation peptide CLN2^{PEST} to the C-terminus of Erg9p for down regulating Erg9p at the translational level, which improved nerolidol titre by 86% to 100 mg L⁻¹ [29].

In a similar way, down-regulating $ERG7$ expression and reducing Erg7p activity are proven to be important strategies to increase triterpenoid titer by reducing 2,3-oxidosqualene flux into endogenous ergosterol pathway [65]. Various strategies have been employed to down-regulate $ERG7$ transcript expression in engineered yeasts, such as replacing the native $ERG7$ promoter with a methionine sensitive $MET3$ promoter [66] or copper-repressible $CTR3$ promoter [67], fine-tuning of $ERG7$ expression using CRISPR interference (CRISPRi) [5] and the tetracycline repressor-tetracycline operator (TetR-TetO)-based gene regulation system [40]. These transcript regulation strategies have proven to be effective in enhancing triterpenoid titer in yeast. Additionally, critical amino acid residues of Erg7p were engineered to decrease Erg7p activity, which resulted in 7.3- and 3-fold increases in the titers of dammarane-type and lupane-type triterpenoids, respectively [68]. Interestingly, a degron-mediated protein degradation strategy was employed to lower the half-life of Erg7p, resulting in 3 to 8-fold increase in 2,3-oxidosqualene-derived triterpenoid production and a 155-fold increase in 2,3:22,23-dioxidosqualene-derived triterpenoid production [47]. These studies demonstrated the effectiveness of deregulating the native competition within the ergosterol pathway to optimize flux in triterpenoid production and offered a novel perspective on the biosynthesis of triterpenoids.

3. Protein Engineering Strategies for Improving Terpenoid Biosynthesis in Yeasts

The construction of a *de novo* terpenoid synthesis pathway in yeasts typically involves the introduction of specific terpene synthases that can convert different precursors derived from the MVA pathway into target products. To improve synthesis efficiency and product yield, factors such as enzyme activity and selectivity need to be considered. Protein engineering strategies can be used to optimize and enhance enzyme properties, such as, rational design or semi-rational design, protein fusion expression and directed evolution of enzymes.

3.1. Rational Design or Semi-rational Design

For enzymes with known enzyme structure and catalytic mechanism, rational design or semi-rational design can effectively improve the specificity and activity of enzymes. In the terpenoid synthesis pathway, some enzymes are promiscuous and can catalyze the production of a wide range of natural and non-natural substrates. For example, the enzyme Erg20p functions as both a GPP synthase (GPPS) and FPP synthase (FPPS), posing challenges in separating the two synthases during monoterpenoids biosynthesis [11,59]. To increase the monoterpenoids titer, the mutant $ERG20^{F96W-N127W}$ has been employed to improve the GPP pool by rational protein engineering design of Erg20p in *S. cerevisiae*, achieved the 10-fold increase in sabinene production [20]. A similar strategy was also employed to modify Erg20p with introducing mutant $ERG20^{F88W-N119W}$ via rational protein engineering design in *Y. lipolytica*, and resulted in about 0.56 mg/L of linalool production [19]. For the majority of enzymes in the terpenoid biosynthetic pathway, their crystalline structure, active site residues and catalytic mechanism have not been resolved, posing challenges for rational design or semi-rational design approaches. Fortunately, molecular modeling tools can be employed to simulate the putative protein structure. Through homology modeling and docking, key residues were identified of CYP72A63 that is the P450 enzyme for licorice triterpenoid biosynthesis [69]. Then the structure model aided to rationally design the promiscuous CYP72A63 for improving the *in vivo* activity, regio- and chemoselectivity based on computation-guided mutations. The promiscuous CYP72A63 was successfully remodeled into an enzyme oxidizing a specific C-H bond into hydroxyl, aldehyde and carboxylic acid, respectively. In addition, the catalytic activity of MdOSC1 for α -amyrin biosynthesis was optimized via homology modeling and docking [45]. The triple mutant $MdOSC1^{N11T/P250H/P373A}$ was obtained by semi-rational protein engineering, and the yield of α -amyrin was increased to 11-fold higher than that of the control.

3.2. Protein Fusion Expression

Protein fusion is a convenient strategy for enhancing enzyme catalytic efficiency through increasing the concentration of local intermediates, and efficiently redirecting metabolic flux to the desired product [21,27,59]. For example, fusion of the double mutant Erg20p(F96W-N127W) with SpSabS1 resulted in a 3.5-fold increase in sabinene yield [20]. The design addresses a significant challenge in yeast monoterpene production by tackling the sequential nature of the yeast farnesyl diphosphate synthase reaction, without affecting the growth characteristics of the tested yeast strains. In another report, co-expression of Erg20p and α -farnesene synthase (FS) with a GGGS amino acid linker improves the catalytic efficiency of α -farnesene synthase on FPP, avoiding FPP branching metabolism pathways and improving α -farnesene titer by 4.8-fold [27]. The same group also indicated that fusion expression of Erg20p(F96W-N127W) and tVoGES after optimizing the linker led to a 23.5-fold increase in geraniol production compared to the strain with solely overexpressing the *tVoGES* gene [21]. The authors also observed a decrease in squalene accumulation compared to the strain overexpressing a single enzyme, indicating that the fusion of tVoGES-Erg20p(F96W-N127W) improved the GPP flux during geraniol production while reducing the GPP flux towards squalene synthesis pathway. Similarly, the fusion expression of GGPP synthase (Bts1p) and Erg20p^{F96C} with GGGS linker was adopted to increase the GGPP accumulation [4]. Meanwhile, the fusion expression of CfCYP76AH15, t66CfCPR and t30AaCYB5 not only decreased production of the intermediate 11-oxo-manoyl oxide, but also significantly increased the production of forskolin to 583.24 $\mu\text{g/L}$. Additionally, taxadiene production was enhanced up to 22-fold through fusion expression of maltose binding protein (MBP)-tagged taxadiene synthase (TASY) and Erg20p, along with effective copy number optimization [36]. This can be attributed to the enhancement of protein expression and solubility, along with the increased availability of FPP for the synthetic pathway. Furthermore, fusion of β -carotene ketolase (CrtW) from *Paracoccus* sp. with β -carotene hydroxylase (CrtZ) from *Haematococcus pluvialis* minimized the substrate-enzyme distance, increased reaction rates, and obtained 2.8-fold improvement in the production of astaxanthin [55]. The strain expressing the CrtW-Z fusion exhibited superior performance compared to the strain with the CrtZ-W fusion protein, indicating that the order of fusion proteins also plays a crucial role.

3.3. Directed Evolution of Enzymes

Directed evolution can also be used to improve the catalytic activity of enzymes through iterative mutations and screening of libraries. Mutant libraries can be constructed using traditional techniques such as error-prone PCR (ep-PCR) [30], DNA shuffling and the staggered extension process (StEP) to achieve random mutagenesis [61]. However, random mutagenesis-based directed evolution will generate a very large mutant library, encompassing numerous neutral and negative mutants alongside a few positive mutants. In order to accurately obtain positive mutants with desired traits from such a vast library, it is necessary to establish a fast, simple and reliable high-throughput screening method. Consequently, directed evolution finds its primary application in modifying reaction catalytic enzymes that can be linked to growth or pigment synthesis [11,30,61]. For instance, a high-throughput screening method was developed based on precursor DMAPP toxicity and used for isoprene synthase (ISPS) directed evolution towards enhanced catalytic activity for isoprene production [70]. Two positive mutants ISPS^{F340L/I478V} and ISPS^{A570T} were obtained from the mutant library generated by ep-PCR, and isoprene-forming activity were improved by 1.6-fold and 1.8-fold than the wild type, respectively. Another instance involves the engineering of a truncated linalool synthase (t67OMcLIS) from *M. citrata* through directed evolution, facilitated by the development of a competition-based color-indicated high-throughput screening method [11]. The variant t67OMcLIS^{E343D/E352H}, exhibiting increased enzyme activity, was successfully screened from a library generated by ep-PCR, resulting in a 52.7% increase of the linalool titer in *S. cerevisiae*.

4. Harnessing Yeast Subcellular Compartments for Improving Terpenoid Biosynthesis

Yeast harbors various subcellular compartments, which offer numerous advantages that cannot be overlooked. These compartments possess the ability to concentrate substrates and enzymes, and provide a unique cellular environment that can respond to biosynthesis pathways with different needs, such as separating target synthesis pathways from competitive pathways and serving as storage sites for intermediate products that mitigate the toxic effect of their accumulation on host cells [71]. Therefore, the use of subcellular compartments to synthesize terpenoids in yeast has attracted more and more attention.

4.1. Harnessing Mitochondria

Mitochondria, a double-membrane organelle, is rich in acetyl-CoA, ATP, and cofactors. Notably, the level of acetyl-CoA in mitochondria is 20–30 times higher than that in cytoplasm and nucleus, which is a promising site for terpenoid synthesis [72]. By targeting the geraniol biosynthetic pathway into mitochondria, the geraniol precursor GPP was protected from depletion by the cytoplasmic ergosterol pathway, resulting in a six-fold increase in geraniol production compared to strains confined to the cytoplasm [22]. In addition, SquM6 strain with squalene-specific titer of 79.78 mg/g DCW was obtained by locating the squalene synthesis pathway in mitochondria [37]. However, the strain's growth was significantly inhibited. To overcome this challenge, they successfully resolved the issue by overexpressing *tHMG1* in the cytoplasm. Finally, the squalene titer reached 21.1 g/L and the specific titer was 437.1 mg/g dcw after the two-stage fermentation process by combining cytoplasmic and mitochondrial engineering.

4.2. Harnessing Peroxisomes

Peroxisomes, single-layer membrane organelles, play a crucial role in generating a substantial pool of acetyl-CoA through β -oxidation. These organelles can also be genetically engineered to regulate their quantity and size [42,73], and their presence is not necessary for yeast to grow [73]. Consequently, peroxisomes have emerged as attractive synthetic sites for natural product synthesis [7]. By targeting 10 genes involved in the squalene synthesis pathway to the peroxisomes, the compartmentalization of squalene synthesis was achieved in *S. cerevisiae* [38]. Through further optimizing the supply of ATP, NADPH, acetyl-CoA in the peroxisome, and combining the dual cytoplasmic-peroxisomal engineering strategy, the squalene titer was significantly increased to 1698.02 mg/L and reached 11.00 g/L after a two-stage fed-batch fermentation. In another study, an increase in the number and copy number of peroxisomes was achieved by editing three peroxisome enzymes, Pex34p, Pex11p and Atg36p [42]. Then by reconstructing the protopanaxadiol biosynthesis pathway in the peroxisome engineered strain, the protopanaxadiol yield was about 78% higher than that of the parent strain.

4.3. Harnessing Endoplasmic Reticulum

Endoplasmic reticulum (ER) is the base of a series of important biological macromolecules, such as proteins, lipids, sugars, and nucleic acids. It also provides a natural environment for many pathway enzymes and P450 enzymes involved in terpenoid synthesis. The expansion of the ER space offers several benefits, including enhancing the synthesis and folding capacity of ER proteins, alleviating metabolic constraints caused by limited enzyme abundance, and increasing the capacity of ER-targeted P450 and CPR enzymes. Currently, ER can be expanded in *S. cerevisiae* by knocking out the phosphatidic acid phosphatase-encoding *PAH1* gene or by overexpressing *INO2*, a key regulator of ER size [39,50]. As an example, the deletion of *PAH1* gene stimulated the production of recombinant triterpene biosynthesis enzymes and resulted in a substantial increase in the accumulation of various triterpene skeletons [50]. In the ER-engineered *pah1* yeast, the production of β -amyrin, medicagenic acid, and medicagenic-28-*O*-glucoside was enhanced by eight-fold, six-fold, and 16-fold, respectively, compared to the wild-type strain. Furthermore, the impact of *PAH1* knockout was also evaluated for the production of other types of terpenoids. The results showed that *pah1* cells led to an approximately two-fold increase in sesquiterpene artemisinic acid production, but had an adverse effect on β -carotene production, which does not depend on ER-localized P450s. On the other hand, strains overexpressing *INO2* demonstrated a 71-fold increase in squalene production and an eight-fold increase in cytochrome P450-mediated protopanaxadiol production compared to strains without *INO2* overexpression [39]. Transcriptome analysis revealed that in the ER-expanded strain, there was a comprehensive reorganization of central carbon metabolism, along with significant alterations in the transcriptional levels of ER-resident chaperones. These changes were suggested to enhance glucose uptake, improve protein folding, and facilitate the assembly of functional metabolic pathways.

4.4. Harnessing Lipid Droplets

In yeast, lipid droplets (LDs), consisting of triacylglycerols (TAGs) and sterol esters (SEs), are promising storage organelles for lipophilic compounds. Due to their hydrophobic properties, lipid droplets were also acted as a reservoir of terpenoids, while anchoring specific enzymes onto their surface facilitates efficient production of terpenoid scaffolds. For example, lipid droplet is the storage organelle of the cytochrome P450 enzyme protopanaxadiol (PPD) synthase (PPDS) substrate dammarenediol-II (DD) [43]. By targeting PPDS to lipid droplets (LDs) using the yeast PLN1 protein, the efficiency of converting DD to PPD was significantly improved by 394%, with the conversion rate of DD increasing

from 17.4% to 86.0%. In another work, it was demonstrated that lipid droplets size can be regulated by overexpression of fatty acid desaturase (*OLE1*) and deletion of Seipin (*FLDI*), in addition to overexpressing key genes involved in fatty acid synthesis and TAG production. These modifications led to a 25% increase in lycopene production compared to the original high-yield strain. The yield of lycopene in fed-batch fermentation reached 2.37 g/L and 73.3 mg/g cdw, respectively. In addition, the regulation of lipid metabolic pathways can significantly impact the accumulation of lipophilic compounds in yeast. In the case of microbial production of β -carotene, overexpressing the sterol ester synthesis genes *ARE1* and *ARE2* resulted in a 1.5-fold increase, while deleting the phosphatidate phosphatase genes (*PAH1*, *DPP1*, and *LPP1*) led to a two-fold increase [52]. By combining these two strategies, β -carotene production was amplified by 2.4 times compared to the initial strain. This study also provided valuable insights into the accumulation of other lipophilic compounds in yeast.

5. The Potential of Incorporating High Throughput Approaches for Improving Terpenoid Biosynthesis

Developing high-throughput techniques is a critical part of strain engineering, which allows researchers to rapidly screen a large number of microbial strains or genetic modifications, saving time and resources compared to traditional methods [74]. One of the key advantages of high-throughput approaches is their ability to quickly and efficiently screen and identify optimal strains or genetic modifications that exhibit desired traits, such as higher productivity, yield, or stability. This enables researchers to quickly identify and select the most promising candidates for further development.

A commonly employed method for strain screening involves leveraging the inherent characteristics of the products, such as color, fluorescence, or growth status. For example, researchers established a fluorescence-activated cell sorting method that rapidly screened millions of mutants and identified novel gene targets involved in lipid metabolism, sterol metabolism, signal transduction, and stress response [54]. This was achieved by analyzing the β -carotene content and fluorescence intensity of cells using HPLC and FCM, respectively. Notably, disrupting genes that affect fatty acid oxidation, lipid composition, and sterol transcriptional regulation (*4CL-8*, *GCS*, and *YIsterTF*) increased β -carotene significantly. Finally, by engineering these targets in a high- β -carotene production, a strain that produced 9.4 g/L β -carotene was constructed. In the case of (+)-nootkatone, an important and expensive natural sesquiterpene compound, its microbial production is hindered by the low catalytic efficiency of its cytochrome P450-P450 reductase (HPO-CPR) system. To address this, a high-throughput screening (HTS) method was developed based on the color reaction between carbonyl compounds and 2,4-dinitrophenylhydrazine (DNPH) [25]. This method enabled the rapid screening of candidate HPO mutants for activity. By optimizing the pairing of HPO and CPR and employing semi-rational design, an optimal mutant HPO_M18 with a catalytic performance 2.54 times higher than the initial strain was obtained.

High-throughput approaches involve the use of automated and rapid techniques to screen a large number of microbial strains or genetic modifications for desired traits or characteristics. An automated high-throughput screening platform based on HPLC and UPLC-MS analytical techniques was established to functionally identify hundreds of *Ganoderma lucidum* cytochromes P450 in *S. cerevisiae* [48]. This platform enabled the successful achievement of a series of highly efficient heterologous biosynthesis of ganoderic acid in *S. cerevisiae*, along with the identification of key genes involved in the ganoderic acid biosynthetic pathway. Specifically, the catalytic reaction process of CYP512W2 and the formation of type II ganoderic acid were thoroughly investigated. Validation through shake flask fermentation of the high-yielding type II ganoderic acid strains, obtained after metabolic modification, showed that the yields of type II ganoderic acid reached more than 50 mg/L in all cases. This represents a significant improvement compared to the traditional artificial cultivation production method, with ganoderic acid yields increased by 1–4 orders of magnitude and the production efficiency enhanced by 2–5 orders of magnitude. Similarly, the screening progress of microbial colonies was expedited through the direct evaluation of microbial bioproduction colonies using supercritical fluid extraction-supercritical fluid chromatography-triple quadrupole mass spectrometry (SFE-SFC-MS/MS) [75]. The online SFE-SFC-MS/MS system offers great potential for high-throughput analyses, due to the automated extraction of metabolites without the need for pre-processing. The SFE-SFC-MS/MS system enables faster and easier screening of highly productive strains compared to conventional analyses, as demonstrated by the high-throughput screening of (–)-limonene bioproducers.

6. The Potential of Incorporating Global Approaches for Improving Terpenoid Biosynthesis

Global approaches aim to investigate biological systems in a holistic manner, involving the elucidation of mechanisms underlying the cellular behavior of specific cell factories or the simulation of the behavior of these systems. Incorporating global approaches has immense potential for improving the performance of microbial cell factories [76]. By adopting a holistic perspective and considering the entire biological system, these approaches can provide valuable insights and strategies for enhancing the performance of microbial cells in industrial settings. With the rapid

development of several advanced techniques, such as omics technologies and genome-scale models, several global approaches are employed for improving terpenoid biosynthesis.

Through extensive studies of transcriptomics, proteomics and metabolomics, we have gained a comprehensive understanding of the metabolic networks and regulatory mechanisms in both natural producers and engineered yeast strains. This approach enables the integration of genome-scale experimental data to uncover the underlying mechanisms, identify targets for optimization, and guide the design of genetic modifications or environmental conditions to enhance strain performance. For example, in yeast, out of the 1,470 presumed N⁶-Methyladenosine (m⁶A) peaks identified, it was observed that overexpression of IME4 (yeast m⁶A methyltransferase) led to significantly altered transcription levels in 94 pathway genes [77]. In particular, IME4 overexpression elevates the mRNA levels of the methylated genes in the glycolysis, acetyl-CoA synthesis and shikimate/aromatic amino acid synthesis modules, which provides essential precursors and acetyl-CoA for terpene synthesis. *Phaffia rhodozyma* is a potential strain for the biosynthesis of astaxanthin. However, the unclear metabolic characteristics of *P. rhodozyma* at different metabolic stages have hindered the efficient production of astaxanthin. To shed light on this, a quadrupole time-of-flight mass spectrometry metabolomics approach was employed to investigate the changes in metabolites [78]. The results showed that the down-regulation of purine, pyrimidine, amino acid synthesis and glycolysis pathways played a crucial role in astaxanthin biosynthesis. Similarly, a comprehensive analysis of differential metabolites and expression genes was conducted to explore the effects of gibberellin-induced astaxanthin synthesis, combining transcriptomics and metabolomics [57]. The results showed that the application of gibberellic acid (GA) significantly enhanced fatty acid metabolism and ABC transporters, and increased unsaturated fatty acid content. This favorable alteration facilitated the accumulation of astaxanthin, thereby promoting its synthesis.

Genome-scale metabolic models (GEMs) have emerged as a computational systems biology approach for effectively interpreting and integrating multi-omic data. By converting the organism's reactions related to metabolism, transcription, and translation into a mathematical formulation, GEMs can be modeled using optimization principles [79]. This modeling approach enables the optimization of microbial growth-associated traits by simulating biomass formation, predicting growth rates, and identifying the requirements for cell growth. Moreover, GEMs empower cells to enhance product formation by predicting gene knockout and overexpression targets using modeling tools. For instance, a GEMs was used to identify new target genes for enhancing biosynthesis of sesquiterpenes in the *S. cerevisiae* [31]. The impact of gene deletions on flux distributions in the metabolic model of *S. cerevisiae* was assessed using OptGene as the modeling framework and minimization of metabolic adjustments (MOMA) as objective function. The results revealed that the deletion of *GDH1* (gene encoding NADPH-dependent glutamate dehydrogenase) was the most promising target gene for improving sesquiterpene biosynthesis in yeast. This deletion increased the availability of NADPH in the cytosol for other NADPH requiring enzymes, including HMG-CoA reductase, resulting in an approximately 85% increase in the final cubebol titer. In another work, two prediction tools, namely FOCuS (a novel gene deletion tool) and OptKnock (a standard gene deletion prediction tool), were employed to identify knockout targets for the improvement of terpenoid pathway fluxes [80]. Notably, key knockout targets were predicted, including *LYS1*, *GAP1*, *AAT1*, *AAT2*, *THI7*, *KGD-m*, *MET14*, *PDC1* and *ACO1*, which are associated with fatty acid biosynthesis, amino acid synthesis pathways or nucleotide biosynthesis pathways. Similarly, overexpression targets such as *PFK1*, *FBA1*, *ZWF1*, *TDH1*, *PYC1*, *ALD6*, *TPII*, *PDX1* and *ENO1*, which are involved in glycolytic and pentose phosphate pathways, were identified. Each of these targets had plausible role in improving flux toward sterol pathway and appeared to be meaningful rather than artifacts. In this regard, an *in vitro* study was conducted to validate the predictions by overexpressing *ALD6* and *TPII*. The results demonstrated a 2.23- and 4.24-fold increase in squalene synthesis, respectively, compared to the control. In general, the rationale behind predicting these *in silico* targets was attributed to either increase the acetyl-CoA precursor pool or regenerate NADPH, both of which enhance the flux in the sterol pathway.

GEM can also be used to optimize fermentation media, offering a more efficient alternative to the laborious statistical design methods such as the Plackett-Burman design. By combining GEM (FBA) results with statistical methods such as the Plackett-Burman design, it becomes possible to develop a highly effective culture medium. Using this method, chemicals for lycopene production were screened in metabolically engineered strains [51]. In fed-batch cultivation mode, lycopene concentrations of 126 and 242 mg/L were achieved using the FBA-independent and FBA-assisted designed media, respectively. Transcriptional studies further revealed upregulations of heterologous genes in media design based on FBA, providing evidence for the predictive power of the model.

With the rapid increase in volume and complexity of data generated from omics studies, the prevalence of extensive data in the domain of multi-omics studies necessitates elaborate computational analysis. Machine learning (ML), a data-driven approach, serves as a computational tool that aims to analyze input data, identify patterns, and generate meaningful

insights in the output. ML has been applied to a wide range of biological problems [81], such as predicting pathway dynamics from omics data [82], determining transcription regulatory dynamics [83], and predicting genotype-phenotype associations [84]. Thus, the application of ML holds great promise in evaluating strain design strategies. For example, to investigate the individual contributions of five non-rate-limiting enzymes in the MVA pathway, a combinatorial library of 243 *S. cerevisiae* strains was created, each with an additional copy of the MVA pathway integrated into the genome and expressing the non-rate-limiting enzymes through a unique combination of promoters [85]. Through high-throughput screening combined with ML algorithm, it was revealed that Erg12p was the key enzyme in affecting the titer of the product. Optimal expression of *ERG12* was achieved using a medium-strength promoter, which represents the ideal ‘sweet spot’ for attaining high product yield. In addition, a platform strain was created by targeting the mevalonate pathway to both the cytosol and peroxisomes. By employing this dual localization approach, terpenoid production was synergistically enhanced, resulting in a remarkable 94-fold increase in monoterpene geraniol, a 60-fold increase in sesquiterpene α -humulene, and a 35-fold increase in triterpene squalene titers, respectively. This platform strain, with its enhanced capabilities, can serve as a versatile chassis for the production of various terpenoids and terpene derivatives.

7. Prospects and Conclusions

Terpenoids, which are natural secondary metabolites with various biological activities, possess immense potential and commercial value. The rapid advancements in synthetic biology, metabolic engineering, DNA sequencing and omics analysis have facilitated the sustainable and efficient production of a wide range of terpenoids through the heterologous expression of terpene synthases in various microbial chassis, such as *S. cerevisiae* and *Y. lipolytica*. However, there are still several challenges need to be addressed. Many high-value terpenoids and their derivatives with relatively complex structures and synthetic pathways have not been heterologous synthesized. Even for terpenoids that have been heterologously synthesized, there is still a significant gap between the production capacity of microbial cell factories and the requirements of industrial-scale production. The main obstacles to large-scale production of terpenoids mainly include the following aspects: (a) inadequate supply of precursor substrates, energy and cofactors; (b) low catalytic efficiency and poor substrate/product specificity of terpene synthases; (c) limited understanding of natural synthetic pathways and functions of key enzymes; (d) the incompatibility between heterologous pathways and chassis cells due to differences in intracellular microenvironment.

In response to these challenges, several strategies have demonstrated their effectiveness. Firstly, a combination of metabolic engineering approaches can be utilized. This includes enhancing precursor supply by introducing alternative heterologous acetyl-CoA pathways [26] and IUP pathway [18,33], optimizing the endogenous MVA pathway [23,32,44] and PP pathway [41,53], and improving and balancing the NADPH supply [26,41] to increase carbon fluxes toward terpenoid synthesis. Secondly, to optimize the expression of enzymes, successive enzyme colocalization strategies such as enzyme fusion [21,34,36], enzyme scaffolds [2], organelle localization [24,55] can be employed. These strategies aim to enhance substrate conversion efficiency and minimize the diversion of metabolic intermediates into competing pathways. Thirdly, a comprehensive understanding of the metabolic network of natural producers and the regulatory mechanism of engineering yeast strains can be achieved through the in-depth studies of omics techniques [37,78,79]. Fourthly, the adaptation mechanism between heterologous pathways and yeast cells can be analyzed to achieve a balance between cell metabolism and heterologous product synthesis through protein and metabolic engineering [9]. Additionally, the storage and transport mechanisms of synthesized terpene products in yeast can be investigated to maximize the utilization of intracellular resources [38,86]. Techniques such as subcellular localization [28] and compartmentalization regulation can be employed to create more space for storing hydrophobic terpenoids by expanding the membrane structure of yeast cells [4,8,42,43]. Exploring specific transporters for the efflux of toxic products can reduce metabolic stress on cells and improve production sustainability [86].

In the future, the development of more efficient multi-fragment and long-fragment DNA assembly, as well as genome editing techniques, can be used to realize heterologous synthesis of more complex terpenoids in yeast. Simultaneously, methods of high-throughput screening need to be developed for real-time detection of intermediate metabolites and intracellular products. Furthermore, the construction of an automated and intelligent synthetic biology platform can expedite the assembly and optimization process of complex synthetic pathways for terpenoid production in yeast cells. These platforms can shorten the design-build-test-learning cycle of cell factories, leading to the construction of efficient, stable, and controllable yeast cell factories for terpenoid biosynthesis.

Author Contributions

Y.X. and C.L. drafted and revised the manuscript; S.S. revised the manuscript; Y.X. and R.C. filled the table and drew figure; L.Q. drew figure.

Ethics Statement

Not applicable.

Informed Consent Statement

Not applicable.

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Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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