***De novo* production of the key branch point benzylisoquinoline alkaloid reticuline in yeast**

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Abstract

Microbial biosynthesis for plant-based natural products, such as the benzyloquinoline alkaloids (BIAs), has the potential to address limitations in plant-based supply of established drugs and make new molecules available for drug discovery. While yeast strains have been engineered to produce a variety of downstream BIAs including the opioids, these strains have relied on feeding an early BIA substrate. We describe the *de novo* synthesis of the major BIA branch point intermediate reticuline via norcoclaurine in *Saccharomyces cerevisiae*. Modifications were introduced into yeast central metabolism to increase supply of the BIA precursor tyrosine, allowing us to achieve a 60-fold increase in production of the early benzyloquinoline scaffold from fed dopamine with no supply of exogenous tyrosine. Yeast strains further engineered to express a mammalian tyrosine hydroxylase, four mammalian tetrahydrobiopterin biosynthesis and recycling enzymes, and a bacterial DOPA decarboxylase produced norcoclaurine *de novo*. We further increased production of early benzyloquinoline scaffolds by 160-fold through introducing mutant tyrosine hydroxylase enzymes, an optimized plant norcoclaurine synthase variant, and optimizing culture conditions. Finally, we incorporated five additional plant enzymes - three methyltransferases, a cytochrome P450, and its reductase partner - to achieve *de novo* production of the key branch point molecule reticuline with a titer of 19.2 µg/L. These strains and reconstructed pathways will serve as a platform for the biosynthesis of diverse natural and novel BIAs.

1.1 Introduction

The benzyloisoquinoline alkaloids (BIAs) are a large, diverse, and important family of plant secondary metabolites with valuable pharmacological properties, including analgesic (Hagel and Facchini, 2013), anti-tumor (Kemény-Beke et al., 2006), and anti-microbial (Orhana et al., 2007) activities. Some BIAs, such as the opiates morphine and codeine, are produced at large scale and are widely prescribed. However, other BIAs that hold promise as new drug molecules are only present in trace amounts within plant hosts and are unavailable at quantities necessary for drug development studies. Current BIA production methods are hampered by costly and inefficient extractions from plant material, and the structure of these molecules prevents their economical manufacture by chemical synthesis.

Microbial biosynthesis of BIAs has the potential to address limitations of plant-based production, stabilize the supply of established drugs, and make new molecules available for study as drug candidates. Efforts to reconstitute BIA biosynthesis pathways in microbes have yielded successes in bacterial (Minami et al., 2008; Nakagawa et al., 2014, 2011) and yeast hosts (Fossati et al., 2015, 2014; Hawkins and Smolke, 2008; Thodey et al., 2014; Trenchard and Smolke, 2015). While high titers of the critical BIA intermediate reticuline have been achieved in *Escherichia coli* (Nakagawa et al., 2011), there has been no follow up work to extend the BIA biosynthesis pathways to more complex end-products, such as the opiates, in this host. This is in part due to the limited ability of bacterial hosts to functionally express endomembrane-localized enzymes, such as plant cytochrome P450s, which are prevalent in BIA biosynthesis (Minami et al., 2008). In contrast, the eukaryotic model organism *Saccharomyces cerevisiae* is a good host for the functional expression of endomembrane-localized enzymes and offers the additional advantage of well established methods for the stable expression of many heterologous enzymes

as is needed for the extensive BIA biosynthesis pathways (Jensen et al., 2014; Shao et al., 2009; Siddiqui et al., 2014). Strains of *S. cerevisiae* have been engineered to produce a variety of complex BIA molecules including reticuline, sanguinarine, morphine, codeine, and various semisynthetic opioids from fed early BIA scaffolds (Fossati et al., 2015, 2014; Hawkins and Smolke, 2008; Thodey et al., 2014; Trenchard and Smolke, 2015).

The *de novo* production of BIAs from simpler precursors that are directly supplied by the yeast host remains a significant challenge and is crucial to the eventual goal of developing an economically viable microbial production process. BIA biosynthesis begins with two molecules of the amino acid tyrosine, which are converted to 4-hydroxyphenylacetaldehyde (4-HPAA) and dopamine, and subsequently condense to form norcoclaurine, the first BIA scaffold in plants. The pathway from central metabolism to 4-HPAA and dopamine has not been fully elucidated in plants. Thus, reconstruction of a *de novo* BIA biosynthesis pathway in microbial hosts requires identification of non-plant enzymes capable of connecting the host's central metabolism to 4-HPAA and dopamine. As yeast naturally synthesize 4-HPAA through the transamination and decarboxylation of tyrosine, the production of dopamine presents the major challenge for this host. One route to dopamine biosynthesis is through the selective hydroxylation of L-tyrosine at C3 to form L-DOPA, followed by the decarboxylation of L-DOPA to form dopamine. There are three types of enzymes known that can catalyze the C3 hydroxylation reaction: tyrosinases, mammalian tyrosine hydroxylases, and more recently a cytochrome P450 DOPA oxidase (CYP76AD1) (DeLoache et al., 2015). Tyrosinases are promiscuous oxygenases that can exhibit unwanted catechol oxidase activity which results in unwanted side products and reduced flux towards valuable end products (Nakagawa et al., 2014). CYP76AD1 is also a promiscuous oxygenase that leads to undesired side products, although mutations have been identified that

increase the specificity of this enzyme for the tyrosine hydroxylase activity (DeLoache et al., 2015). In contrast, mammalian tyrosine hydroxylases naturally exhibit high substrate and product specificity; however, this class of enzymes requires the electron carrier cosubstrate tetrahydrobiopterin (BH₄), which is non-native to microbial hosts.

In this work, we describe the *de novo* synthesis of the major BIA branch point intermediate reticuline via norcoclaurine in *S. cerevisiae*. We first engineered yeast strains with targeted modifications of central metabolism to improve flux to tyrosine biosynthesis to increase the supply of this critical BIA precursor. Our strain engineering efforts allowed us to achieve an approximate 60-fold increase in production of norcoclaurine from fed dopamine with no supply of exogenous tyrosine to the yeast host. We then engineered yeast strains for the *de novo* production of norcoclaurine through the heterologous expression of a mammalian tyrosine hydroxylase (TyrH), bacterial DOPA decarboxylase (DODC), and four enzymes associated with BH₄ biosynthesis and recycling. We further increased flux to norcoclaurine through introducing mutations into the tyrosine hydroxylase enzyme to alleviate product and substrate inhibition, selecting an optimized norcoclaurine synthase (NCS) variant, and optimizing culture conditions, resulting in an additional 160-fold improvement in norcoclaurine production levels. Finally, we incorporated five additional plant enzymes - three methyltransferases, a cytochrome P450, and its reductase partner - into our engineered yeast strain to achieve a microbial platform for *de novo* production of the key branch point molecule reticuline via norcoclaurine.

1.2 Materials and Methods

1.2.1 Yeast strain and plasmid construction

Oligonucleotides were synthesized by either the Stanford Protein and Nucleic Acid Facility (Stanford, CA) or Integrated DNA Technologies (Coralville, IA). Cloning was performed with chemically competent *Escherichia coli* (TOP10, LifeTech, F- mcrA Δ (mrr-hsdRMS-mcrBC) ϕ 80lacZ Δ M15 Δ lacX74 nupG recA1 araD139 Δ (ara-leu)7697 galE15 galK16 rpsL(Str^R) endA1 λ). *E. coli* were cultured in Luria-Bertani media (EMD Chemicals) with appropriate antibiotic: 100 μ g/mL ampicillin (EMD Chemicals) or 50 μ g/mL kanamycin (EMD Chemicals). Spin columns were used to purify plasmids from *E. coli* cultures according to the manufacturer's instructions (Epoch Life Science). Sequencing was performed by Elim Biopharmaceuticals (Hayward, CA). DNA polymerases used were either Hotstart PfuUltra (Agilent), PfuUltraII Fusion HS DNA Polymerase (Agilent), or Expand High Fidelity Polymerase (Roche). PCR products were cleaned up using QIAquick PCR purification kit (Qiagen).

Gene sequences for *RnPTPS* (AAH59140), *RnSepR* (AAA42130), *RnPCD* (NP_001007602), *RnQDHPR* (P11348), *RnDHFR* (AF318150), *TfNCS* (AAR22502), *CjNCS* (BAF45338), *PsNCS* (AAX56304), *PpDODC* (AAN68161) and *EcCYP80B1* (AF014801) were yeast codon-optimized and either synthesized by GeneArt (Life Technologies), IDT, or by *in vitro* PCR assembly as previously described (Hoover and Lubkowski, 2002). RnTyrH was obtained from the Fitzpatrick laboratory (Daubner et al., 2006). *Ps6OMT*, *PsCNMT*, *Ps4'OMT*, and *PsCPR* were obtained from previously engineered strains (Hawkins and Smolke, 2008; Trenchard and Smolke, 2015).

The engineered *Saccharomyces cerevisiae* strains described in this work are all derived from CEN.PK2-1D (MAT α ura3-52; trp1-289; leu2-3/112; his3 Δ 1; MAL2-8C; SUC2) (Entian and Kötter, 2007). Additional *S. cerevisiae* strains tested in Figure S1 are W303 α (MAT α leu2-3,112; trp1-1; can1-100; ura3-1; ade2-1; his3-11,15; [phi⁺]) (Ralser et al., 2012) and BY4742 (MAT α his3 Δ 1; leu2 Δ 0; lys2 Δ 0; ura3 Δ 0) (Brachmann et al., 1998). A standard lithium acetate protocol was used for transformation of plasmids (Gietz and Woods, 2006). Changes to central metabolism were performed with chromosomal integrations using integrating plasmids (Scherer and Davis, 1979; Siddiqui et al., 2014) or multi-fragment DNA assembly (Shao et al., 2009). BH₄ enzymes and core BIA enzymes were integrated using multi-fragment assembly (Shao et al., 2009) with the transformation of fragments into the yeast cells performed through electroporation (Chao et al., 2006). Yeast strains used in this study are listed in Table 1.

Table 1. Genotypes of yeast strains used in this study.

Strain	Genotype
CSY1030	CEN.PK2-1D; <i>fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1031	CEN.PK2-1D; <i>ARO9::P_{TEF1}-ARO9; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1032	CEN.PK2-1D; <i>ARO10::KanMX-P_{TEF1}-ARO10; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1033	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1034	CEN.PK2-1D; <i>his3Δ::KanMX-ARO7^{T226I}; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1035	CEN.PK2-1D; <i>zwf1Δ; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1036	CEN.PK2-1D; <i>TKL1::P_{GPD}-TKL1; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1037	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; TKL1::P_{GPD}-TKL1; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1038	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; zwf1Δ; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1002	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; zwf1Δ; TKL1::P_{GPD}-TKL1;</i>
CSY1039	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; zwf1Δ; TKL1::P_{GPD}-TKL1; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1042	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; zwf1Δ; TKL1::P_{GPD}-TKL1; P_{ARO10}::KanMX-P_{TEF1}; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1044	CEN.PK2-1D; <i>trp1Δ::KanMX-ARO4^{Q166K}; zwf1Δ; ARO10::KanMX-P_{TEF1}-ARO10; his3Δ::KanMX-ARO7^{T226I}; fcy1Δ::KIURA3-P_{GPD}-PpDODC-T_{ADHI}</i>
CSY1050	CSY1002; YBR197C Δ :: <i>P_{TPH1}-RnSepR-T_{STE2}; P_{TEF1}-RnPTPS-T_{CYC1}; KanMX, P_{GPD}-RnQDHPR-T_{ADHI}, P_{PGK1}-RnPCD-T_{PHO5}</i>
CSY1051	CSY1039; YBR197C Δ :: <i>P_{TPH1}-RnSepR-T_{STE2}; P_{TEF1}-RnPTPS-T_{CYC1}; KanMX, P_{GPD}-RnQDHPR-T_{ADHI}, P_{PGK1}-RnPCD-T_{PHO5}</i>
CSY1052	CSY1051; YDR514C Δ :: <i>P_{PYK1}-PsCNMT-T_{MFa1}; P_{PGK1}-Ps6OMT-T_{PHO5}; P_{GPD}-EcCYP80B1-T_{ADHI}; LEU2, P_{TEF1}-PsCPR-T_{CYC1}; P_{TPH1}-Ps4OMT-T_{STE2}</i>

Yeast plasmids described in this work were constructed with Gateway Cloning Technology (Life Technologies). Enzyme coding regions were PCR amplified and cloned into the pENTR vector through BP recombination reaction using BP clonase II and the pDONR221

vector (Life Technologies). All genes were subsequently recombined into selected pAG expression vectors from the Lindquist laboratory (available through Addgene) (Alberti et al., 2007) using LR Clonase II (Life Technologies) (Table S2). Site directed mutagenesis was performed using the QuikChange Site Directed Mutagenesis protocol and Hotstart PfuUltra DNA polymerase (Agilent).

Table 2. Plasmids used in this study.

Plasmid	Description	Source
pAG414GPD-ccdB	<i>Centromeric TRP, P_{GPD}-ccdB-T_{CYC1}</i> recombination cassette	(Alberti et al., 2007)
pAG413GPD-ccdB	<i>Centromeric HIS, P_{GPD}-ccdB-T_{CYC1}</i> recombination cassette	(Alberti et al., 2007)
pCS3231	pAG414- <i>P_{GPD}-RnTyrH-T_{CYC1}</i>	This work
pCS3232	pAG414- <i>P_{GPD}-RnTyrH^{W166Y}-T_{CYC1}</i>	This work
pCS3233	pAG414- <i>P_{GPD}-RnTyrH^{E332D}-T_{CYC1}</i>	This work
pCS3234	pAG414- <i>P_{GPD}-RnTyrH^{S40D}-T_{CYC1}</i>	This work
pCS3235	pAG414- <i>P_{GPD}-RnTyrH^{R37E,R38E}-T_{CYC1}</i>	This work
pCS3236	pAG414- <i>P_{GPD}-RnTyrH^{W166Y,E332D}-T_{CYC1}</i>	This work
pCS3237	pAG414- <i>P_{GPD}-RnTyrH^{S40D,W166Y}-T_{CYC1}</i>	This work
pCS3238	pAG414- <i>P_{GPD}-RnTyrH^{R37E,R38E,W166Y}-T_{CYC1}</i>	This work
pCS3239	pAG414- <i>P_{GPD}-RnTyrH^{R37E,R38E,W166Y,E332D}-T_{CYC1}</i>	This work
pCS3240	pAG413- <i>P_{GPD}-TfNCS-T_{CYC1}</i>	This work
pCS3241	pAG413- <i>P_{GPD}-CjNCS-T_{CYC1}</i>	This work
pCS3242	pAG413- <i>P_{GPD}-PsNCS-T_{CYC1}</i>	This work

1.2.2 Growth conditions for assays

Cultures were grown in deep-well 96-well plates covered with AeraSeal film (Excel Scientific) at 30 °C, 480 rpm, 80% humidity in a Kuhner Lab-Therm LX-T plate shaker. For dopamine and L-DOPA feeding assays, overnight yeast cultures were inoculated in the appropriate drop-out or synthetic complete YNB media with 2% dextrose (w/v) then back-diluted 75-100X into drop-out or synthetic complete liquid media with tyrosine excluded, unless otherwise indicated, in 96-well plates. Cultures were supplemented with either 100 mM dopamine (Alfa Aesar) or 2 mM L-DOPA (Sigma) and grown for 72 hours, unless otherwise indicated. For norcoclaurine and reticuline *de novo* production assays, yeast cultures were

inoculated in 96-well plates in appropriate selective YNB media with no tyrosine, 2% dextrose and 2 mM ascorbic acid (Sigma). Cultures were grown for 96 hours before analysis.

1.2.3 Analysis of metabolite production

Aliquots of yeast cultures were centrifuged at 13,000 rpm for 10 min and growth media samples were taken for analysis by LC-MS/MS with an Agilent 1260 Infinity Binary HPLC and an Agilent 6420 Triple Quadrupole LC-MS. Norcoclaurine and reticuline samples were separated on an Agilent EclipsePlus C18, 2.1x50 mm, 1.8 μ m column with 0.1% formic acid as solvent A and acetonitrile with 0.1% formic acid as solvent B at a constant flow rate of 0.4 mL/min and an injection volume of 5 μ L. For separation of norcoclaurine, the following method was used: 0-0.1 min, 7% B; 0.1-5 min, 10 to 20% B; 5-5.5 min, 20 to 90% B; 5.5-7 min, 90% B; 7-7.01 min 7% B; followed by a 3 minute equilibration at 7% solvent B. The LC eluent was directed to the MS for 1-2.5 min with ESI source gas temperature 350 $^{\circ}$ C, gas flow of 11 L/min, nebulizer pressure 40 PSI. For separation of reticuline, the following method was used: 0-0.1 min, 7% B; 0.1-5 min, 10 to 35% B; 5-5.5 min, 35 to 90% B; 5.5-7 min, 90% B; 7-7.01 min 7% B; followed by a 3 minute equilibration at 7% solvent B. The LC eluent was directed to the MS for 1-5 min with ESI source gas temperature 350 $^{\circ}$ C, gas flow of 11 L/min, nebulizer pressure 40 PSI.

L-DOPA, dopamine, BH₄, BH₂, and B samples were separated on an Agilent Pursuit 3 PFP, 150 x 2.0 mm, column with 0.1% formic acid as solvent A and acetonitrile with 0.1% formic acid as solvent B at a constant flow rate of 0.4 mL/min and an injection volume of 15 μ L. The following method was used: 0-0.5 min, 2% B; 0.5-1.5 min, 2 to 60% B; 1.5-4 min, 60% B; 4-4.01 min, 2% B; 4.01-7 min 2% B. The LC eluent was directed to the MS for 1-2.8 min with ESI source gas temperature 350 $^{\circ}$ C, gas flow of 11 L/min, nebulizer pressure 40 PSI.

For quantification, the MS was used in MRM mode to monitor the transitions in Table 3. Quantification of metabolites was based on integrated peak area MRM chromatogram using the Agilent MassHunter Workstation and reported as the mean $\pm$ s.d. of at least three independent experiments. We generated standard curves for L-DOPA (Sigma), norcoclaurine (Seqchem), and reticuline (Specs).

Table 3. MRM transitions used for quantification.

Compound	MRM	Fragmentor	CE	MRM Ref.
Norcoclaurine	272 $\rightarrow$ 107	135	25	(Schmidt and Raith, 2005)
Reticuline	330 $\rightarrow$ 137	135	35	(Schmidt and Raith, 2005)
L-DOPA	198 $\rightarrow$ 152	135	20	(Li et al., 2000)
Dopamine	154 $\rightarrow$ 91	135	20	(Li et al., 2000)
BH ₄	242 $\rightarrow$ 166	135	20	(Fismen et al., 2012)
BH ₂	240 $\rightarrow$ 165	135	20	(Fismen et al., 2012)
B	238 $\rightarrow$ 178	135	20	(Fismen et al., 2012)
Coclaurine	286 $\rightarrow$ 107	135	25	(Schmidt and Raith, 2005)
N-Methylcoclaurine	300 $\rightarrow$ 107	135	25	(Schmidt and Raith, 2005)
3'-Hydoxy-N-methylcoclaurine	316 $\rightarrow$ 192	135	25	(Schmidt and Raith, 2005)

1.3 Results

1.3.1 Production of norcoclaurine in yeast cultures fed dopamine

The biosynthetic route to the first BIA backbone in plants, norcoclaurine, has not been fully elucidated. Thus, we designed a route to the major BIA branch point intermediate, reticuline, via norcoclaurine from simple carbon and nitrogen sources in yeast through combining enzymes from plants, yeast, mammals, and bacteria (Fig. 1).

We first sought to confirm that *S. cerevisiae* provides sufficient levels of 4-HPAA to support synthesis of the first BIA backbone norcoclaurine. *S. cerevisiae* naturally synthesizes

tyrosine from sugar and ammonia, and can convert tyrosine to the BIA building block 4-HPAA via transamination (Aro8p, Aro9p) and decarboxylation (Aro10p) as part of the Ehrlich pathway for producing fusel alcohols from amino acids. Since the condensation reaction between 4-HPAA and dopamine to form norcoclaurine can proceed spontaneously, we assayed for norcoclaurine production from *S. cerevisiae* cultured in the presence of dopamine. Wild-type CEN.PK2, BY4742, and W303 strains were fed 100 mM dopamine and grown for 72 h at 30 °C in media without tyrosine. The growth media was analyzed for metabolite accumulation using high-performance liquid chromatography coupled to tandem mass spectrometry (LC-MS/MS), and production of norcoclaurine was confirmed by comparison of the MRM transition to a standard and to published spectra (Schmidt et al., 2005). We detected norcoclaurine in the growth media of all three strains, and CEN.PK2 produced the most norcoclaurine at 9.8 µg/L (Fig. S1). This result indicates that 4-HPAA produced in yeast can be directed to synthesis of the BIA scaffold norcoclaurine.

1.3.2 Engineering a tyrosine-overproducing yeast strain by deregulation of central metabolism to increase BIA production

To increase flux towards L-tyrosine biosynthesis, we next examined whether modifications to the regulation of yeast central metabolism would support increased titers of norcoclaurine from the engineered yeast strains. We examined modifications to yeast central metabolism that we hypothesized would increase the intracellular pools of the BIA precursors tyrosine and 4-HPAA and thus increase the production of norcoclaurine in engineered yeast cultures (Fig. 1A). We identified six genetic modifications: (i-ii) promoter replacement resulting in transcriptional upregulation of two genes associated with the Ehrlich pathway for transforming

aromatic amino acids to fusel alcohols ($P_{TEF1}ARO10$, $P_{TEF1}ARO9$); (iii) introduction of a feedback resistant mutant copy of the chorismate mutase gene designed to divert the aromatic amino acid biosynthesis pathway flux towards tyrosine and phenylalanine ($ARO7^{T226I}$) (Schmidheini et al., 1989); (iv-v) promoter replacement resulting in transcriptional upregulation of the *TKL1* gene and knockout of the *ZWF1* gene, previously shown to increase titers in a related engineered yeast biosynthesis pathway by modifying pentose phosphate pathway flux to favor the accumulation of erythrose-4-phosphate ($P_{GPD1}TKL1$, $\Delta zwf1$) (Curran et al., 2013) and (vi) introduction of a feedback resistant mutant copy of the *ARO4* gene to alleviate tyrosine feedback inhibition of the major control point in aromatic amino acid biosynthesis ($ARO4^{Q166K}$) (Fukuda et al., 1992a, 1992b, 1991).

The genetic modifications were made individually in a wild-type CEN.PK2 yeast strain via targeted chromosomal integration. Each yeast strain was grown in minimal SC media without tyrosine, fed 100 mM dopamine, and grown for 72 h at 30 °C. The growth media for each of the strains was analyzed for norcoclaurine accumulation using LC-MS/MS, and production of norcoclaurine was confirmed by comparison of the MRM transitions to a standard and to published spectra (Schmidt et al., 2005).

Of the individual genetic modifications tested, only $P_{TEF1}ARO10$ and $ARO4^{Q166K}$ resulted in substantial improvement in norcoclaurine production with 5- and 14-fold increases, respectively (Fig. 2). Next, we combined the $ARO4^{Q166K}$ modification with mutations directed to increasing flux from the pentose phosphate pathway to aromatic amino biosynthesis. While the *zwf1* deletion (CSY1035) and *TKL1* upregulation (CSY1036) modifications did not improve norcoclaurine production when introduced individually, the combination of the *zwf1* deletion and the $ARO4^{Q166K}$ modification (CSY1038) resulted in a 30-fold increase in norcoclaurine

production. An approximately 60-fold increase in norcoclaurine production levels was achieved upon combining the *ARO4*^{Q166K}, *zwf1* deletion, and *TKL1* upregulation modifications (CSY1039) when compared to a strain that does not harbor these changes to central metabolism (CSY1030), resulting in a norcoclaurine titer of 327 µg/L. The addition of the *ARO10* upregulation modification to this strain background (CSY1042) did not result in further increases in norcoclaurine levels. Further adding the *ARO7*^{T226I} modification (CSY1044) also did not result in additional increases to norcoclaurine production.

The results indicate that relieving feedback inhibition at the entry point of aromatic amino acid biosynthesis (*ARO4*^{Q166K}) improves the production of norcoclaurine through increased flux to the BIA precursors tyrosine and 4-HPAA, as expected. Additional modifications directed to increasing flux from the pentose phosphate pathway to aromatic amino biosynthesis (*P_{GPD1}TKL1*, *zwf1Δ*) are dependent on alleviation of the feedback inhibition of Aro4p. While the overexpression of Aro10p as an individual modification resulted in increased norcoclaurine levels relative to the base strain (CSY1030), the impact of this modification was not additive in the context of the highest norcoclaurine-producing yeast strain. Overall, the results indicate that modifications to central metabolism and aromatic amino acid biosynthesis to increase the production of tyrosine and 4-HPAA improve the production of norcoclaurine without the expression of heterologous enzymes.

Further analysis was performed on CSY1039 to verify norcoclaurine production in this strain background from fed L-DOPA. An expression cassette for *Pseudomonas putida* DODC (*PpDODC*) was chromosomally integrated into this yeast strain. The strain was grown in minimal SC media without tyrosine, fed 2 mM L-DOPA, and grown for 72 h at 30 °C. The growth media for the strain was analyzed for norcoclaurine accumulation using LC-MS/MS as

previously described. The engineered strain produced 49 $\mu\text{g/L}$ norcoclaurine from 2 mM L-DOPA, which indicated a higher conversion efficiency than from fed dopamine (Fig. S2). Based on these collective results, we selected CSY1039 as the base strain for building a yeast platform strain capable of synthesizing BIAs *de novo*.

1.3.3 Engineering a yeast strain for *de novo* production of the early BIA norcoclaurine through a tyrosine hydroxylase and BH_4 synthesis pathway

We next constructed a complete biosynthetic route to norcoclaurine from simple carbon and nitrogen sources, by incorporating the TyrH enzyme into our engineered yeast strain. Mammalian TyrHs catalyze the selective C3 hydroxylation of L-tyrosine to form L-DOPA in the presence of molecular oxygen and the co-substrate BH_4 , and are an attractive enzyme for the reconstructed BIA pathway due to their substrate and product specificity. However, yeast do not naturally synthesize BH_4 , and thus we further engineered our yeast strains to produce and recycle this required cosubstrate.

BH_4 biosynthesis begins with GTP, which is converted to dihydroneopterin triphosphate by GTP cyclohydrolase I (GTPCHI), an endogenous yeast enzyme involved in tetrahydrofolate synthesis (Fig. 3A). Two heterologous enzymes, 6-pyruvoyl-tetrahydropterin synthase (PTPS) and sepiapterin reductase (SepR), are required to convert dihydroneopterin triphosphate to BH_4 . BH_4 is hydroxylated during aromatic amino acid hydroxylation, and two additional heterologous enzymes, pterin-4 α -carbinolamine dehydratase (PCD) and quinonoid dihydropteridine reductase (QDHPR), are required to recycle it back to its active form (Fig 3A).

The BH_4 biosynthesis and recycling enzymes from *Rattus norvegicus* were yeast codon-optimized, synthesized, and directly assembled and integrated into a single locus of CSY1002 to

make strain CSY1050. BH₄ readily oxidizes and thus we expected to detect the oxidized products dihydrobiopterin (BH₂) and biopterin (B) in the growth media (Fismen et al., 2012; Valent and Tóth, 2004). The growth media was analyzed for BH₄ and the oxidized products using LC-MS/MS and compound production was confirmed by comparison of MRM transitions to a standard and to published spectra (Fismen et al., 2012). All three BH₄ products were detected in the standard and the samples (Fig. 3B).

We tested the ability of the strain with the integrated BH₄ biosynthesis and recycling (CSY1050) to support the activity of mammalian TyrHs, and thus *de novo* norcoclaurine production. The gene encoding TyrH from *R. norvegicus* (*RnTyrH*) was expressed from a low-copy yeast plasmid under the control of the *GPD* promoter (*P_{GPD}*). The base strain harboring the BH₄ synthesis and recycling genes and the *RnTyrH* expression construct was cultured in selective media without tyrosine and grown for 96 h at 30 °C. The growth media for the strain was analyzed for accumulation of L-DOPA using LC-MS/MS, and metabolite production was confirmed by comparison of the MRM transitions to a standard and to published spectra (Li et al., 2000). The strain expressing the wild-type *RnTyrH* produced L-DOPA at a titer of 63 µg/L (Fig. 3C).

We next examined whether increasing the availability of BH₄ would result in increased TyrH activity. Ascorbic acid has been shown to reduce the oxidation of BH₄ to BH₂ (Valent and Tóth, 2004), and thus the addition of ascorbic acid to the culture media may increase TyrH activity. CSY1050 was grown in selective media without tyrosine and supplemented with 2 mM ascorbic acid for 96 hours at 30 °C. L-DOPA and BH₄ intermediates were measured in the media by LC-MS/MS as described previously. The addition of ascorbic acid improves L-DOPA production by 1.5-fold over media lacking ascorbic acid (Fig. 3C, S3A). BH₄ concentration

increased by 1.7-fold with the addition of ascorbic acid (Fig. S3B). The increase in BH₄ concentration is likely a result of oxidation protection from the ascorbic acid resulting in the observed increase in TyrH activity.

Finally, we examined these engineered yeast strains for the *de novo* production of norcoclaurine in the absence of fed tyrosine and the impact of the NCS expression on norcoclaurine production levels. While 4-HPAA and dopamine can spontaneously condense to form norcoclaurine, we suspected that this condensation reaction would be limiting at the lower concentrations of dopamine present in these strains relative to previous dopamine feeding assays. We yeast codon-optimized and synthesized genes encoding NCS from *P. somniferum* (*PsNCS*), *T. flavum* (*TfNCS*) and *C. japonica* (*CjNCS*) and expressed each variant from a low-copy plasmid with *RnTyrH* from a separate low-copy plasmid in CSY1051 (CSY1050 with *PpDODC* integrated). The strains were grown in selective media without tyrosine and with 2 mM ascorbic acid for 96 hours at 30 °C, and norcoclaurine levels in the media were measured by LC-MS/MS as previously described. In the absence of NCS, the yeast strain produced 0.03 µg/L norcoclaurine. NCS expression improved *de novo* norcoclaurine production and the *CjNCS* variant resulted in the highest norcoclaurine production, increasing titers by over 15-fold to 0.56 µg/L (Fig. 4A, B).

1.3.4 Tyrosine hydroxylase mutants that relieve substrate and production inhibition improve BIA production

TyrH enzymes are reported to be both substrate (tyrosine) and product (catecholamine) inhibited. TyrH product inhibition has two modes, fast and slow, due to the competitive binding of catecholamines at the BH₄ binding site and the irreversible binding of catecholamines to the

iron center, respectively (Kumer and Vrana, 1996). We hypothesized that substrate and/or product inhibition may limit TyrH activity in the context of the engineered yeast strain. Thus, we examined whether variants of *RnTyrH* mutated to relieve these modes of inhibition would result in increased norcoclaurine production levels from the engineered yeast strains.

Mutations to *RnTyrH* have previously been reported that relieve different modes of inhibition. For example, substrate inhibition was shown to be relieved by a W166Y mutation (Quinsey et al., 2002), the fast product inhibition was shown to be relieved by the mutation S40D and the consecutive mutations R37ER38E (Nakashima et al., 2002), and the slow product inhibition was shown to be relieved by a E332D mutation (Briggs et al., 2011). Site directed mutagenesis was used to introduce each of these mutations into the wild-type *RnTyrH* coding sequence, three double mutants combining substrate and product inhibition mutations and one triple mutant combining the three modes of inhibition relief. Strain CSY1051 harboring a mutant enzyme variant expressed from a low-copy plasmid and *CjNCS* expressed from a low-copy plasmid was grown in selective media with 2 mM ascorbic acid for 96 hours at 30 °C. Norcoclaurine levels in the media were measured as previously described.

The data demonstrate that mutants that relieved fast product inhibition resulted in the greatest increases in norcoclaurine production (Fig. 4C). Specifically, the *RnTyrH* S40D mutant and the *RnTyrH* R37ER38E mutant resulted in a 1.7-fold and 3-fold increase in norcoclaurine levels, respectively. The slow inhibition mutant, E332D, reduced norcoclaurine production individually and in double and triple mutant combinations, which is likely explained by the reduced $V_{\max}$ of this particular *RnTyrH* mutant (Briggs et al., 2011). The substrate inhibition mutant, W166Y, exhibited no impact on norcoclaurine levels. The combination of the substrate inhibition mutant and a fast product inhibition mutant resulted in further increases to

norcoclaurine production levels. Specifically, norcoclaurine levels increased by 3-fold for W166Y, S40D (WS) and 9-fold for W166Y, R37ER38E (WR) to achieve a titer of to 5 µg/L.

1.3.5 Engineering a yeast strain for the de novo production of the key branch point BIA reticuline

Finally, we extended the reconstructed BIA biosynthetic pathway from the first BIA backbone molecule norcoclaurine to the key branch point BIA reticuline. Reticuline has previously been produced in yeast from the fed substrate norlaudanoline, which requires the expression of three methyltransferases (norcoclaurine 6-*O*-methyltransferase, 6OMT; coclaurine-*N*-methyltransferase, CNMT; 3'-hydroxy-*N*-methylcoclaurine 4'-*O*-methyltransferase, 4'OMT) (Hawkins and Smolke, 2008; Minami et al., 2008). Two additional enzymes, cytochrome P450 80B1 (CYP80B1) and a cytochrome P450 NADPH reductase (CPR), are required to make reticuline from norcoclaurine.

The five heterologous enzymes required for conversion of norcoclaurine to reticuline were synthesized, cloned into expression cassettes, and directly assembled and integrated into a single locus in CSY1051 resulting in CSY1052 (Fig. 5A). CYP80B1 from *E. californica* and CPR from *P. somniferum* were yeast codon-optimized, whereas the sequences for the methyltransferase genes were derived from *P. somniferum* (Hawkins and Smolke, 2008; Trenchard and Smolke, 2015). A panel of *RnTyrH* mutants and *CjNCS* were co-expressed from separate low-copy plasmids in CSY1052, and the strain was grown in selective media with 2 mM ascorbic acid for 96 hours at 30 °C. Reticuline levels in the media were measured by LC-MS/MS and confirmed by comparison of the MRM transitions to a standard and to published spectra (Schmidt et al., 2005). The data demonstrate that reticuline was produced *de novo* with no supply

of exogenous tyrosine from yeast engineered with the reconstructed pathways harboring each of the tested *RnTyrH* mutants (Fig. 5B, S4A). The best *RnTyrH* mutant improved reticuline titers by as much as 9-fold to 19.2 $\mu\text{g/L}$ under the assay conditions. The conversion efficiencies through the core of the pathway are high as reticuline is the most highly accumulated compound (Fig. 5C). There is an accumulation 3'-hydroxy-N-methylcoclaurine, which indicates that 4'OMT may be the least efficient enzyme in the core pathway. A substantial amount of dopamine accumulation is also observed (6 mg/L) indicating that the efficiency of the condensation reaction catalyzed by NCS is a target for further pathway improvement (Fig. S4B).

1.4 Discussion

We describe engineered yeast strains capable of *de novo* production (from sugar and ammonia) of the first BIA backbone molecule, norcoclaurine, and the key BIA branch point molecule, reticuline. We implemented a number of improvements into our engineered host strain and reconstructed pathway to increase flux towards BIA production, including introducing several modifications to central metabolism to increase supply of the early BIA precursors tyrosine and 4-HPAA. We also incorporated a mammalian TyrH and associated BH_4 biosynthesis and recycling enzymes to increase the specificity of the tyrosine hydroxylation step relative to earlier pathway reconstruction efforts. Flux to the early 1-benzylisoquinoline scaffolds was further increased by implementing mutant TyrHs to alleviate product and substrate inhibition of this enzyme, an optimized NCS variant, and optimized culture conditions, resulting in an approximate 160-fold improvement in *de novo* norcoclaurine production.

Our study reports the use of a mammalian TyrH in yeast to catalyze the conversion of L-tyrosine to L-DOPA, in contrast to earlier studies in *E. coli* that used a bacterial tyrosinase to

catalyze this reaction and the very recent work that was published while this manuscript was under review in which a plant cytochrome P450 DOPA oxidase was used to catalyze this reaction (DeLoache et al., 2015). Tyrosinases are substrate promiscuous oxygenases that exhibit unwanted catechol oxidase activity. In the earlier studies, the authors noted that the engineered bacterial strains were unable to stably produce pre-reticuline intermediates including norlaudanosoline, likely due to the inherent substrate promiscuity of the tyrosinase activity (Nakagawa et al., 2014). The recent work that used the plant cytochrome P450 similarly leveraged the promiscuity of this enzyme and applied a screening strategy to identify mutations that increased the enzyme's substrate specificity and activity; however, the enzyme still suffers from the production of undesired side products (DeLoache et al., 2015).

In contrast, TyrHs are characterized as having high substrate and product specificity, which are advantageous to directing greater flux toward the desired BIA precursor molecules and ultimately BIA products. However, TyrHs have the drawback of requiring the cosubstrate BH₄, which is non-native to microbial hosts. Therefore, we demonstrated the production of BH₄ in a yeast host to support TyrH activity through the expression of four additional heterologous enzymes for BH₄ biosynthesis and recycling. BH₄ is also a cosubstrate for other aromatic amino acid hydroxylases, such as the phenylalanine and tryptophan hydroxylases, and this BH₄ synthesis platform can be used more broadly to support the activity of these enzymes in a yeast host. We further improved TyrH activity and overall BIA production through the introduction of rationally chosen mutations to relieve substrate and production inhibition of this enzyme.

While the recent report describing a yeast strain engineered to produce reticuline incorporated one modification to central metabolism (Aro4^{FBR}) (DeLoache et al., 2015; Luttik et al., 2008), we explored a variety of modifications to central metabolism to increase tyrosine and

4-HPAA levels. These combined modifications resulted in significant improvements in early BIA production and enabled the yeast strains described in this work to produce reticuline in the absence of any fed tyrosine in the media (i.e., only from fed glucose). We also optimized NCS activity through testing several variants of NCS. We found that the *C. japonica* NCS was the most efficient variant in the context of the engineered pathway, whereas other work found an NCS variant from *P. somniferum* to be the most efficient of those tested (DeLoache et al., 2015).

The production of reticuline via norcoclaurine in an engineered yeast strain has several advantages over routes previously reported in *E. coli* that go through norlaudanosoline. First, 4-HPAA is a natural metabolite in yeast, and thus the expression of additional heterologous enzymes for 4-HPAA production is not required. Second, the native substrates of *Ps6OMT* and *PsCNMT* are norcoclaurine and coclaurine, respectively. Thus, by synthesizing reticuline via norcoclaurine these enzymes see their native substrates rather than chemical analogues, which may negatively affect turnover, flux, and product specificity within the engineered pathway. Finally, norcoclaurine biosynthesis requires one molecule of dopamine, whereas norlaudanosoline biosynthesis requires two molecules of dopamine, such that dopamine production may become limiting when producing reticuline via norlaudanosoline. The main drawback of synthesizing reticuline via norcoclaurine is the requirement to express two additional endomembrane-localized enzymes, a cytochrome P450 (CYP80B1) and its reductase partner. However, we found that we were able to achieve robust expression and activity of these two enzymes in our engineered yeast strains.

An analysis of intermediate accumulation in the *de novo* BIA biosynthesis pathway reported here, indicates several steps that are limiting to flux through the pathway. In particular, although NCS improves norcoclaurine production, a substantial amount of dopamine

accumulation is observed from these strains, suggesting further improvements can be made to increase the efficiency of this condensation reaction. One possibility is increasing 4-HPAA levels through further modifying the strain by knocking out yeast alcohol dehydrogenases (ADH) or aldehyde dehydrogenases (ALD) to prevent 4-HPAA degradation and increase its availability for condensation with dopamine. The production of L-DOPA is also likely a limiting reaction and further engineering efforts to improve TyrH activity and BH₄ availability as well as other approaches will be necessary to further increase flux through this step. Through the core of the pathway to reticuline, 4'OMT appears to be the least efficient enzyme, followed by the closely related enzyme 6OMT. Future work will focus on optimizing these steps through a number of strategies including targeted enzyme engineering and further strain optimization to increase the flux of BIA precursors into the pathway. In addition, fermentation optimization strategies can be applied to further improve the production process.

The yeast strains and associated enzymatic pathways described in this work will provide important platforms for the eventual development of economically viable microbial production processes for key BIA compounds and scaffolds, including the opioids. With further development such microbial production platforms have the exciting potential to address limitations of plant-based production, stabilize the supply of established drugs, and make new molecules available for study as drug candidates. For example, by providing a biosynthetic route through norcoclaurine, these yeast strains enable the synthesis of compounds comprising the large and important subfamily of bisBIAs, for which relatively little is currently known about their biosynthesis. A microbial platform for the biosynthesis of BIAs will also support combinations of enzymes that aren't present in natural biological organisms, enabling the efficient generation of novel and non-natural derivatives to fuel the discovery of new medicines.

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Author Contributions

IJT, MSS, and CDS conceived of the project, designed the experiments, analyzed the results, and wrote the manuscript. IJT, MSS, and KT performed the experiments.

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Figure Legends

Figure 1 Biosynthetic scheme for production of reticuline from L-tyrosine. (A) Engineered yeast aromatic amino acid synthesis for the production of BIA precursors L-tyrosine and 4-hydroxyphenylacetylaldehyde (4-HPAA). (B) Engineered biosynthetic pathway from L-tyrosine to reticuline. Enzymes labeled in black indicate unmodified yeast enzymes, red indicate modified yeast enzymes, blue indicate heterologous enzymes. Compounds labeled in green represent the connection between yeast central metabolism and BIA biosynthesis. PPP, pentose phosphate pathway; PEP, phosphoenolpyruvate; E4P, erythrose 4-phosphate; DAHP, 3-deoxy-D-arabino-heptulosonate-7-phosphate; EPSP, 5-enolpyruvateshikimate 3-phosphate; 4-HPP, 4-hydroxyphenylpyruvate; 4-HPAA, 4-hydroxyphenylacetylaldehyde; TyrH, tyrosine hydroxylase; DODC, DOPA decarboxylase; NCS, norcoclaurine synthase; 6OMT, 6-O-methyltransferase; CNMT, coclaurine-N-methyltransferase; CYP80B1, N-methylcoclaurine 3'-hydroxylase; CPR, cytochrome P450 NADPH-reductase; 4'OMT, 3'-hydroxy-N-methylcoclaurine 4'-O-methyltransferase.

Figure 2 Norcoclaurine production in engineered yeast strains as a function of modifications to central metabolism. Strains were cultured in synthetic complete YNB media with 100 mM dopamine for 72 hours, and norcoclaurine production was analyzed by LC-MS/MS. The number of central metabolism modifications introduced into the strain are indicated as: blue, one modification; red, two modifications; green, three modifications; orange, four modifications; purple, five modifications. Data is reported as the mean $\pm$ s.d. of at least 3 independent experiments.

Figure 3 Production of the TyrH cosubstrate BH₄ in engineered yeast. (A) Engineered biosynthetic pathway for production and recycling of BH₄. Enzymes labeled in black indicate endogenous yeast enzymes, blue indicate heterologous enzymes. Compounds labeled in green represent the connection between yeast central metabolism and BH₄ biosynthesis. GTPCHI, GTP cyclohydrolase I; PTPS, 6-pyruvoyl-tetrahydropterin synthase; SepR, sepiapterin reductase; PCD, pterin-4 α -carbinolamine dehydratase (PCD); QDHPR, quinonoid dihydropteridine reductase. (B) BH₄ production in yeast. Top, BH₄ standard analyzed with LC-MS/MS in MRM mode. Bottom, CSY1050 (CSY1039 with BH₄ biosynthesis genes integrated) media analyzed for BH₄, BH₂, and B with LC-MS/MS in MRM mode. Traces are color coded to match compound names in Fig. 3A. (C) L-DOPA production as a function of BH₄ and ascorbic acid. CSY1002 (no BH₄) and CSY1050 (BH₄ and BH₄ + ascorbic acid) each express the wild-type *RnTyrH* (pCS3231). Strains were grown in selective media lacking tyrosine for 96 hours before analysis. CSY1050 was grown in the presence and absence of 2 mM ascorbic acid. Data is reported as the mean $\pm$ s.d. of at least 3 independent experiments.

Figure 4 *De novo* production of norcoclaurine in engineered yeast. (A) LC-MS/MS MRM analysis of norcoclaurine production in CSY1051 with *RnTyrH* (pCS3231) and *CjNCS* (pCS3241). Norcoclaurine standard (2.5 nM) is shown in black. (B) Norcoclaurine production as a function of NCS species variant. CSY1051 strains harboring plasmids with *RnTyrH* (pCS3231) and either *TfNCS* (pCS3240), *PsNCS* (pCS3242), or *CjNCS* (pCS3241) were grown in selective media lacking tyrosine and supplemented with 2 mM ascorbic acid for 96 hours before analysis. (C) Norcoclaurine production as a function of TyrH mutant. CSY1051 strains harboring

plasmids with *RnTyrH* mutants (pCS3231-pCS3239) and *CjNCS* (pCS3241) were grown in selective media lacking tyrosine and supplemented with 2 mM ascorbic acid for 96 hours before analysis. Data is reported as the mean $\pm$ s.d. of at least 3 independent experiments.

Figure 5 *De novo* production of reticuline in engineered yeast. (A) Schematic depicting heterologous expression constructs in final strain design. Blue indicates heterologous enzymes, orange indicates selection marker. lcp, low-copy plasmid. (B) Reticuline production as a function of *RnTyrH* mutant. CSY1052 strains harboring plasmids with *RnTyrH* (pCS3231), *RnTyrH*^{R37ER38E} (pCS3235), or *RnTyrH*^{WR} (pCS3238) and *CjNCS* (pCS3241) were grown in selective media lacking tyrosine and supplemented with 2 mM ascorbic acid for 96 hours before analysis. Data is reported as the mean $\pm$ s.d. of at least 3 independent experiments. (C) LC-MS/MS MRM analysis of pathway intermediates from norcoclaaurine to reticuline in CSY1052 expressing *RnTyrH*^{WR} (pCS3228) and *CjNCS* (pCS3241). Cultures were grown in selective media lacking tyrosine and supplemented with 2 mM ascorbic acid for 96 hours before analysis. Reticuline standard (25 nM) shown in black.

Figure 1

Figure 2

Figure 3

Figure 4

Figure 5