Challenges and opportunities for engineering assembly-line polyketide biosynthesis in *Escherichia coli*

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**A B S T R A C T**

Assembly-line polyketide synthases generate natural products that have led to many live-saving drugs. The use of *E. coli* as a heterologous host for reconstituting these enormous and complex enzymatic machines has and will continue to be a critical strategy for understanding them. Here, we concisely summarize successful examples in exploiting *E. coli* for assembly-line polyketide biosynthesis as well as offer examples of new challenges in which this approach is primed to tackle.

Polyketide synthases (PKSs) manufacture truly remarkable natural products. Polyketides are structurally diverse and possess important medicinal and agricultural capabilities that include but are not limited to: antibacterial, anticancer, antiparasitic, immunosuppressant, herbicidal, insecticidal and fungicidal activities. Assembly-line PKSs are a subset of type I PKSs and are exceptionally complex molecular machines. For example, the extensively studied 6-deoxyerythronolide B synthase (DEBS) is a colossal homodimer (its molecular weight exceeds 2 MDa) that elaborately channels reactive intermediates through 22 distinct enzymatic domains to build 6-deoxyerythronolide B (6-DEB), a precursor of the antibiotic erythromycin A (Walsh, 2004; Khosla et al., 2007; Cortes et al., 1990; Donadio et al., 1991) (Fig. 1).

Unfortunately, assembly-line PKSs often are housed in poorly characterized and/or difficult to culture microbes. Further, many of the biosynthetic gene clusters (BGCs) involve with these PKSs are not expressed in laboratory conditions (Rutledge and Challis, 2015). Although game-changing tools such as the CRISPR/Cas system are being repurposed for *Streptomyces* – a treasure trove for assembly-line PKSs – (Cobb et al., 2015), bacteria like *Streptomyces* have slow growth rates, are labor- and time-intensive to genetically manipulate and have cluttered metabolic backgrounds. On the other hand, model strains of *Escherichia coli* are obvious “user-friendly” workhorse. They are easy to culture, grow fast and have a wealth of genetic tools. Importantly, their metabolic backgrounds are free from native complex natural product biosynthetic pathways, which lowers the potential for metabolic ‘cross-talk’ with heterologously introduced systems. Despite *E. coli*’s advantages, only a handful of assembly-line PKSs have been either fully or largely reconstituted in *E. coli*. Here in this brief perspective, we summarize the most successful case – DEBS – of using *E. coli* as a heterologous host for assembly-line polyketide biosynthesis. Finally, we propose examples of future challenges and opportunities that researchers exploiting *E. coli* as a tool may consider as avenues to further our understanding of these breathtaking enzymes.

Following the accomplishments of manipulating DEBS in both its native producer host *Saccharopolyspora erythraea* (Marsden et al., 1998) as well as *Streptomyces* hosts (Kao et al., 1994; Xue et al., 1999), the stage was set for its reconstituted biosynthesis in *E. coli*. Pfeifer et al. expressed the three large DEBS polyketides (their molecular weights each surpass 300 kDa, and their genes are each longer than 10 kb) in a highly engineered *E. coli* BL21(DE3) derivative named BAP1 (Pfeifer et al., 2001) and synthesized 6-DEB at titers of 20 mg/L in *E. coli*. BAP1 encodes *Bacillus subtilis* Sfp, a phosphopantetheinyl transferase that phosphopantetheinylates the seven acyl carrier protein domains in DEBS. To elevate intracellular concentrations of propionyl-CoA and methylmalonyl-CoA, Pfeifer et al. a) deleted the propionate catabolism genes *prpRBCD*, b) boosted expression of the propionyl-CoA synthetase PrpE and c) co-expressed Streptomyces coelicolor propionyl-CoA carboxylase, which generates methylmalonyl-CoA from propionyl-CoA. Lau et al. achieved titers that approached *Streptomyces* hosts (1.1 g/L) by optimizing the medium composition and fermentation conditions (Lau et al., 2004). Installing DEBS in *E. coli* enabled the engineers to rationally modify specific parts of DEBS to fabricate novel natural polyketides in vivo, a long-standing goal in natural products research. To manufacture a benzoate-containing analog of 6-DEB, Pfeifer et al. replaced the loading module of DEBS with the loading module from the rifamycin polyketide synthase, which accepts a benzoate starter unit. To completely produce
erythromycin A in *E. coli*, Zhang et al. impressively expressed 17 additional genes that are responsible for deoxysugar biosynthesis, macrolide tailoring, and resistance, which led to the biosynthesis of erythromycin A at titers of 0.6 mg/L (Zhang et al., 2010). P8/1-OG (an intermediate of rifamycin biosynthesis) (Watanabe et al., 2003) and epothilone C and D (Mutka et al., 2006) are two other working examples of engineered assembly-line polyketide biosynthesis in *E. coli*. While these two studies did not reconstitute the entire assembly-line, they are notable for utilizing chaperones and protein dissection to enhance the expression of massive PKS polypeptides: the 530 kDa RifA and 765 kDa EpoD proteins were each split into two polypeptides. Even assembly-line systems that did not reconstitute the entire assembly-line, they are notable for utilizing chaperones and protein dissection to enhance the expression of enormous PKS polypeptides: the 530 kDa RifA and 765 kDa EpoD proteins were each split into two polypeptides. Even assembly-line systems that are predominantly composed of non-ribosomal peptide synthase (NRPS) domains such as the yersiniabactin NRPS/PKS can be successfully reconstituted in *E. coli*; therefore, the lessons learned from the first fruitful attempts will be invaluable to researchers that desire to tap into the advantages that *E. coli* holds as a heterologous host.

Second, genomic sequencing has revealed large numbers (far surpassing known assembly-line PKS BGCs) of cryptic or “orphan” assembly-line PKS BGCs associated with no known polyketide products (O’Brien et al., 2014). This recent surge provides a vast resource for pharmaceutical and agrochemical natural products research. However, scientists must endeavor to decipher this embarrassment of riches in a timely fashion. Though platforms such as antiSMASH 4.0 (Blin et al., 2017), PRISM 3 (Skinnider et al., 2017) and TransATor (Helfrich et al., 2019) offer useful polyketide structural predictions, there is no substitute for interrogating PKSs and their repertoire of products either in vitro or in vivo. For example, the NOCardiosis-Associated Polyketide (NOCAP) synthase, an orphan assembly-line PKS found in clinical strains of the actinomycete *Nocardia*, exhibits iterative activity in one of its modules, a behavior which cannot be presently predicted in silico (Kuo et al., 2016). *E. coli* provides a way to quickly move from assembly-line PKS genes to natural products. *E. coli*’s aforementioned sizeable genetic toolkit, fast growth rate and uncluttered metabolic background should enable rapid and straightforward cloning of these clusters, polyketide biosynthesis and metabolic profiling. As in the case with trans-AT assembly-line PKSs, no assembly-line PKS identified in silico has been immediately “de-orphanized” by reconstitution in *E. coli*; this achievement would represent a new method of natural products discovery.

Third, although there have been major advances in the structural analysis of individual PKS domains, multidomain fragments and even intact modules (Robbins et al., 2016; Dutta et al., 2014; Edwards et al., 2014; Li et al., 2019), we know little about how these multienzyme machines behave as a whole in a cell. Transmission electron microscopy...

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**Fig. 1.** The 6-Deoxyerythronolide B Synthase (DEBS). Enzymatic domains: KS, ketosynthase; AT, acyltransferase; DH, dehydratase; KR, ketoreductase; ER, enoylreductase; ACP, acyl carrier protein; and TE, thioesterase.

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of *B. subtilis* cells revealed that the bacillaene hybrid PKS/NRPS fascinatingly assembles into a single organelle-like membrane-associated complex with approximate mass of 10–100 MDa (Straight et al., 2007). It would be most interesting to establish how many assembly-line PKSs organize into “megacomplexes”. Can microscopists study PKS complex dynamics with live-cell imaging, as oppose to electron microscopy which is limited to fixed cells? Do PKSs retain the ability to assemble as a megacomplex in a heterologous host like *E. coli*? If so, then pathways previously engineered to operate in *E. coli* could be easily retrofitted with a variety of fluorescent proteins or tags; therefore, permitting the monitoring of these enzymes with live-cell imaging tools that require fluorophores such as fluorescence or super-resolution microscopy. Once tagged, these proteins can be interrogated “coarsely” in the context of megacomplexes or “finely” in the context of domain-domain interactions with techniques such as FRET (Fluorescence Resonance Energy Transfer).

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