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Lessons from the Synthetic Chemist Nature

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This conceptual review examines the ideal multistep synthesis from the perspective of nature. We suggest that besides step- and redox economies, one other key to efficiency is steady state processing with intermediates that are immediately transformed to the next intermediate when formed. We discuss four of nature's strategies (multicatalysis, domino reactions, iteration and compartmentation) that commonly proceed via short-lived intermediates and show that these strategies are also part of the chemist's portfolio. We particularly focus on compartmentation which in nature is found microscopically within cells (organelles) and between cells and on a molecular level on multiprotein scaffolds (e.g. in polyketide synthases) and demonstrate how compartmentation is manifested in modern multistep flow synthesis.

1. Introduction

What is a perfect multistep synthesis? This is a key issue which has received much attention recently, despite the fact that as early as 1975 Hendrickson had already provided a definition for an “ideal synthesis”. According to him it is one which: *...creates a complex molecule...in a sequence of only construction reactions involving no intermediary refunctionalisations, and leading directly to the target, not only its skeleton but also its correctly placed functionality.*¹

Many years later this general statement was complemented with ideas such as atom,² step,³ and redox-economy⁴ that serve as standards by which the efficiency of chemical reactions and syntheses can be measured.⁵ However, some of these concepts are restricted to individual chemical transformations and the underlying mechanisms to achieve selectivity. The Baran group has discussed and provided criteria that allow for comparison and even quantification of multistep synthesis. They cited three specific criteria: a) overall yield, b) step count and c) percent ideality.⁵ The authors also acknowledge that the % ideality should be used with caution and is more of a simple tool than a true overarching measure of efficiency. Today, these straightforward concepts are generally accepted within the synthetic community as they reflect an important step toward striving for improved efficiency.

In several aspects, nature can be seen as an ideal synthetic chemist. Understanding the biosynthesis of a natural product is often a productive starting point for the design of highly efficient, biomimetic total syntheses.^[6] Still, chemists have yet to achieve the same exquisite levels of selectivity, particularly, chemoselectivity, in the design of multistep processes. However, analysis of biosynthetic pathways reveals that atom-, redox- and step economies are not always well fulfilled in nature although from the synthetic chemist’s point of view biosyntheses of complex natural products are regarded to be highly efficient. This may sound like a contradiction.

Herein, we discuss four fundamental strategies that are shared in both synthetic chemistry and nature, in so doing, we wish to shed light on this seeming contradiction. Based on these strategies, it is interesting to see the interplay between the chemist and nature, where the chemist can be seen to borrow from nature and in certain cases even improve on nature’s design. The hope of this highlight is to discuss these fundamental strategies in a broad context taking examples from biosynthesis and the state of the art in organic synthesis and presenting them side by side. To clarify the scope of this review is not about biomimetic synthesis and we have deliberately not chosen biomimetic chemical syntheses as this has been extensively reviewed elsewhere.⁶

In this unique setting we hope to inspire the synthetic chemist to think more on natural systems for inspiration. Too often does the chemist draw a line between themselves and biologists, we hope to encourage a blurring of the lines⁷

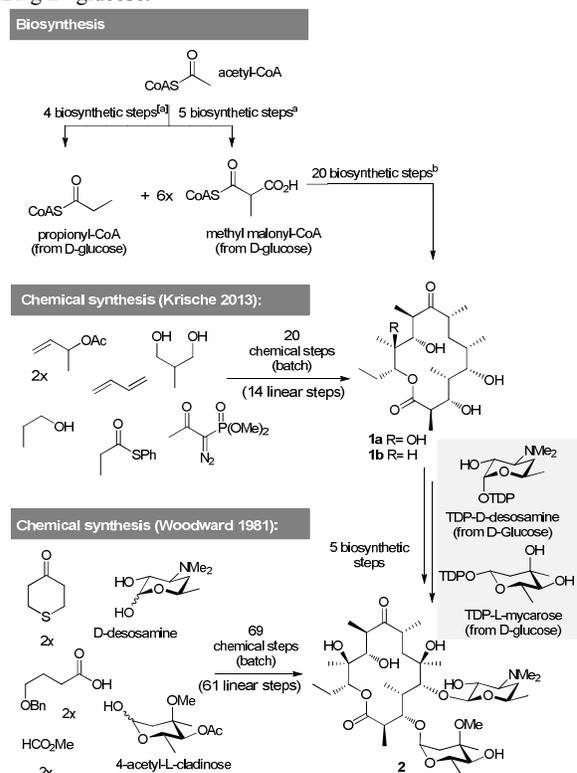
2. Step- and Redox-economies in Nature

In order to discuss the issue of step-economy in nature let us first take a look at erythronolide A (**1a**). It is composed of a macrolactone ring that is biosynthesised by a polyketide synthase type I (PKS). Starting from propionyl-CoA 14 transformations (additional four steps have to be added, if one starts counting from acetyl-CoA, additional six steps if transacylation steps from AT to ACP on the PKS are included, and additional seven steps if the activation and loading of the starter and elongation units are included) occur in a linear fashion to yield 6-deoxyerythronolide B (**1b**). After this, five additional biosynthetic tailoring steps fashion erythromycin A (**2**).⁸

So far, the first and only total synthesis of **2** was reported by Woodward and coworkers in 1981. This formidable synthesis was achieved in 69 (batch) steps (61 in the longest linear sequence) from commercially available starting materials.⁹ Several formal total syntheses of erythromycin precursors as well as of derivatives have

been published since then. The most recent synthesis by Krische *et al.* elegantly furnished 6-deoxyerythronolide B (**1**) in only 14 linear steps (20 overall steps) in the longest linear sequence, clearly in fewer steps than nature does it.¹⁰

However, the bioproduction of erythromycin A (**2**) is efficient because after about 60 hours *Saccharopolyspora erythraea* can provide the antibiotic in significant quantities. Examining the cyclic fed batch fermentation studies of Bushell and Lynch¹¹, a rough estimate on the productivity in *S. erythraea* is possible. About 0.18 – 0.53 % of the glucose fed ends up in erythromycin (**2**).¹² While this yield may not seem very impressive it is still significantly larger than the yield in the only total synthesis (less than 0.02 %). One gram of (dry) biomass produces up to 0.75 mg h⁻¹ of **2** starting from 0.21 g h⁻¹ glucose.

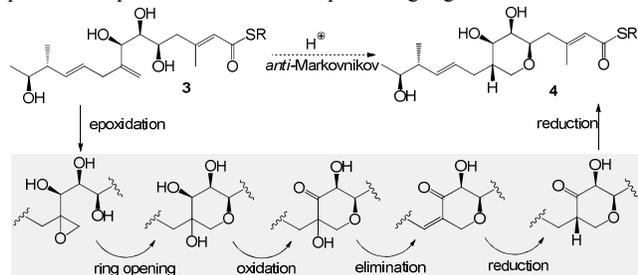


Scheme 1. Comparison of 6-deoxyerythronolide B (**1**) and erythromycin A (**2**) biosyntheses with Woodward’s total synthesis of **2** and Krische’s total synthesis of **1**; ^aStep counts may vary because different biosynthetic routes are known; ^b20 steps when transacylation from AT to ACP is included.

The biosynthesis¹³ of the pseudomonic acids, secondary metabolites originally isolated from *Pseudomonas fluorescens* NCIMB 10586, is an illustrative showcase for poor step- and redox-economy in nature (Scheme 2). A tetrahydropyran ring is a central structural element of the pseudomonic acids and starting from the advanced biosynthetic intermediate **3** it is formed by a six step sequence that includes two oxidations, two reductions an epoxide ring-opening and a dehydration to yield pyran **4**. A chemist would likely suggest a single step alcohol addition onto the olefinic double bond in an *anti*-Markovnikov fashion. Despite these facts pseudomonic acids are produced by fermentation and are used under the name mupirocin against *staphylococci* and *non-enteric streptococci*.

One may speculate why many biosyntheses are linear and rather long, occasionally with poor redox-economy. From a chemists perspective the repertoire of reactions and building blocks nature can rely on is rather small. Additionally, it has been hesitant to develop new biosynthetic pathways. Nature rather sticks to established multistep processes. Often iteration and the addition of “decorating” or “tailoring” steps lead to ever longer linear biosyntheses. From an evolutionary point of view long biosyntheses

originate from genes that were incorporated into biosynthetic gene clusters and became finally operative, when the new metabolite provided improved survival for the producing organism.¹⁴



Scheme 2. Biosynthesis of the pyran ring in pseudomonic acids.

Unlike most chemical syntheses, biosyntheses are not batch but steady state processes in which intermediates are directly processed as they are formed. We believe that this fact is important to approach the topic of a perfect multistep synthesis from nature's perspective. Therefore, we shall begin with a "Gedankenexperiment" of a conventional multistep batch synthesis in order to be able to mirror it with nature's multistep processes before we discuss the four synthetic strategies.

3. The flaw of Multistep Batch Synthesis is its Batch Character

Synthetic chemists commonly conduct multistep processes under batch conditions in which intermediates are formed in equimolar amounts and if necessary need to be isolated and/or further purified. Practically, batch mode synthesis is time consuming and uneconomical and here atom, step and redox- economies do matter.

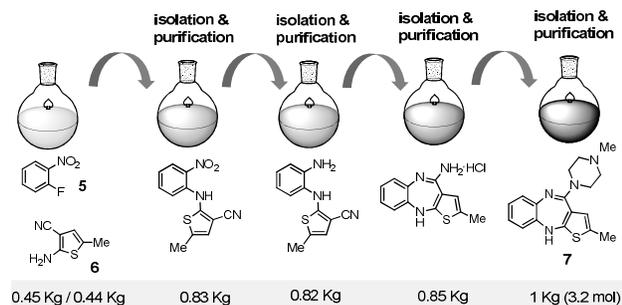


Figure 1. The concept of multistep batch synthesis exemplified in a "Gedankenexperiment" (amount of materials in case of quantitative transformations).¹⁵

In this context, the patented synthesis of Olanzapine (7) is illustrative.¹⁵ Olanzapine is commercialised under the name ZYPREXA[®] and was among the top 20 best selling drugs worldwide in 2011. ZYPREXA[®] is an atypical neuroleptic and is used for the treatment of bipolar disorders and schizophrenia.¹⁶ It is known to exert antagonistic activity towards the dopamine receptor type 4 (D₄-receptor) and the serotonin receptor type 2 (5-HT₂-receptor).¹⁷ The patent¹⁵ describes a four step batch protocol starting from 1-fluoro-2-nitrobenzene (5) and 2-amino-5-methylthiophene-3-carbonitrile (6). Assuming the preparation of 1 kg (3.2 mol) of 7 with quantitative yield for each step (Figure 1). This would require about 0.5 kg of starting materials 5 and 6, and more than 0.8 kg of three intermediates are formed that need to be isolated. As a consequence multistep reaction sequences should ideally avoid work-up and isolation of intermediates.

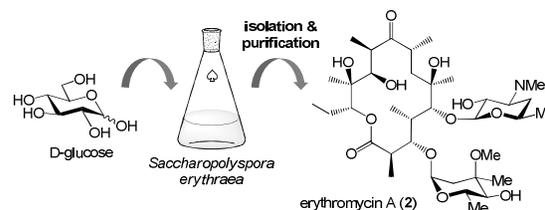


Figure 2. A simplistic view on the bioproduction of erythromycin A (2).

At first glance the bioproduction of complex natural products such as erythromycin A (2) may look like a true batch protocol (figure 2). D-Glucose is the starting material, *Saccharopolyspora erythraea* acts as the "catalytic" system¹⁸ and the fermenter resembles the flask or the reactor. Harvesting the bioreactor, yields the antibiotic 2, still being used in the clinic.

The gross simplification depicted in figure 2 treats *Saccharopolyspora erythraea* as a black box or a batch reactor. Using the tools of molecular biology and biochemistry we can pry into these multicatalytic biosystems. In fact, the biosynthesis of erythromycin A (2) is a highly orchestrated multistep and rather linear steady state process, catalyzed by enzymes, many being part of a catalytic megacomplex, the polyketide synthase (PKS, *vide infra*). In contrast to the multistep batch process depicted in figure 1, these catalytic architectures circumvent the stoichiometric formation of intermediates.

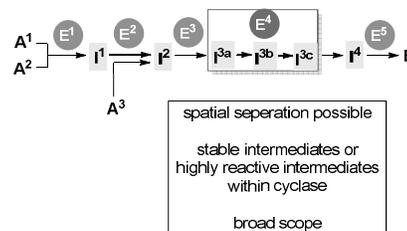


Figure 3. Enzyme-catalyzed multistep biosynthesis (A¹–A³= substrates, building blocks, B= product of biosynthesis/ natural product; I¹–I⁴= reactive intermediates, E¹–E⁵= enzymes, E⁴= enzyme that catalyzes a multistep process via highly reactive intermediates I^{3a}–I^{3c}).

Primary metabolism "feeds" the biosynthetic pathways of secondary metabolite production and is organised in a complex network of enzymatic transformations and routes. A biosynthetic intermediate is often generated from different sources and the intermediate can be the starting point for different biosynthetic routes as exemplified in figure 3. In most cases, building blocks A are activated carboxylic acids such as acetyl- or propionyl-CoA, aminoacyl phosphates or alkyl diphosphates.

Often tailoring transformations such as acylations, alkylations or glycosylations implement further "decoration" into the backbone (e.g. I⁴→B, figure 3). Intermediates I are difficult to detect or to track, so that first insights into biosynthetic pathways came from the isolation of biosynthetic end product B, while manifestation of intermediates I was necessary to provide a detailed description of the pathways.¹⁹ These biosyntheses either rely on a cocktail of individual enzymes (e.g. in terpene biosynthesis)²⁰ or are based on megaenzyme complexes such as the polyketide synthases that are e.g. part of erythromycin (2) biosynthesis, or non-ribosomal peptide synthases (NRPS). In both cases the templating property of PKS and NRPS guarantee spatial separation or compartmentation of each step (see also chapter 4.4).

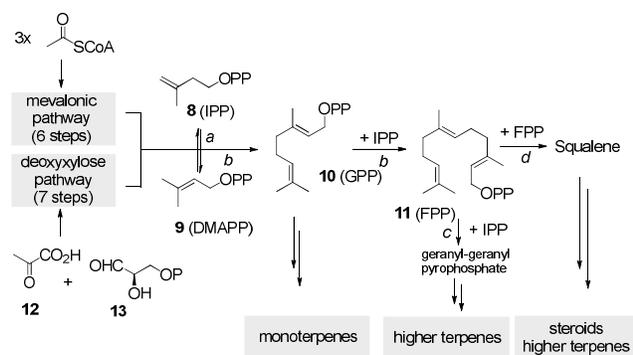
Now that we have provided an overview of secondary metabolite biosynthesis, with an emphasis on polyketides biosynthesis, we will go on to discuss four strategies that nature has developed in order to allow for efficient multistep synthesis with seeming disregard to the chemists notion of step count and atom economy. Typically, the strategies are based on steady state processes and are a)

multicatalysis, b) domino/cascade reactions, c) iteration and d) compartmentation. We discuss compartmentation in greater detail^{21,22} as this aspect can hardly be found in the literature yet.³ Overall, these four strategies add to those justified and often discussed arguments that are based on protecting group chemistry and the high chemo-, regio- and chemoselectivities of biocatalysts.²³

4. Nature's Synthetic Strategies are Based on Steady State Processes

4.1 Strategy 1: Continuous processing by multicatalysis

As previously discussed, nature relies on linear steady state processes, in which individual enzymes are responsible for single transformations. The early steps of terpene biosynthesis are a typical example for a multicatalytic biosynthetic pathway (Scheme 3). Two distinct biosynthetic pathways, namely the mevalonate and the deoxyxylulose route,²⁴ provide nature's isoprene analogues isopentenyl pyrophosphate [IPP (8)] and dimethylallyl pyrophosphate [DMAPP (9)]. The former pathway utilises six enzymes to reach IPP starting from SCoA activated acetic acid, while the latter route requires seven enzymes starting from pyruvate (12) and glyceraldehyde-3-phosphate (13). Then, an iterative process provides geranyl pyrophosphate [GPP (10)], farnesyl pyrophosphate [FPP (11)], geranyl geranyl pyrophosphate and squalene. Each of these advanced intermediate building blocks represents one of the terpene classes (e.g. mono-, di-, sesqui-, tri- and tetraterpenes).



Scheme 3. Overview on the biosynthesis of acyclic terpene pyrophosphates, a typical example of a biological multicatalytic system [IPP= isopentenyl pyrophosphate (8); DMAPP= dimethylallyl pyrophosphate (9); GPP= geranyl pyrophosphate (10); FPP= farnesyl pyrophosphate (11) (enzymes involved: a, Isopentyl-diphosphate delta-isomerase; b, farnesyl diphosphate synthase; c, geranylgeranyl diphosphate synthase; d, farnesyl-diphosphate farnesyltransferase).

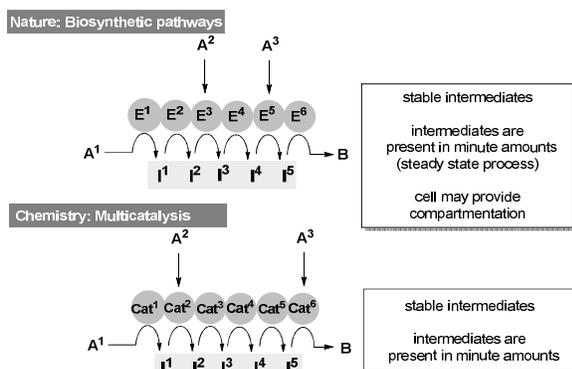
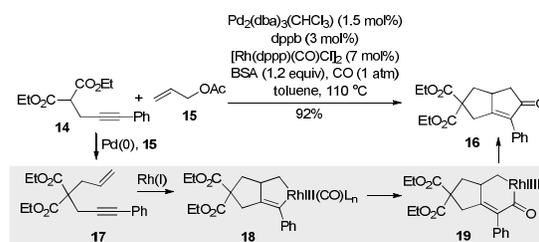


Figure 4. Multicatalytic systems in biosynthesis and chemical synthesis. Multicatalytic reactions and sequences have also emerged in organic synthesis. These systems could be purely based on biocatalysts and

synthetic catalysts or on biocatalyst / synthetic catalyst combinations.²⁵ Within the category of synthetic catalysts further subdivisions can be made based upon whether the catalyst is an organocatalyst or based on inorganic/organometallic catalysts (Figure 4).

The combination of multiple catalysts has several benefits such as reduced waste, reduced overall time to realise multistep syntheses, the ability to control unstable intermediates and the ability to generate molecular complexity from simple precursors. However, certain points must be taken into account, such as the compatibility of starting materials/intermediates and catalysts with each other and with the reaction conditions (temperature, solvent, pressure etc.). Additionally, the relative rates of the individual processes, and the selectivity of the individual catalysts must be considered, such that reactions occur in a definite sequence on precisely the intermediate which one intends. Despite many parameters needing to be controlled²⁶, researchers have begun to uncover reactions and strategies which already led to several impressive examples.

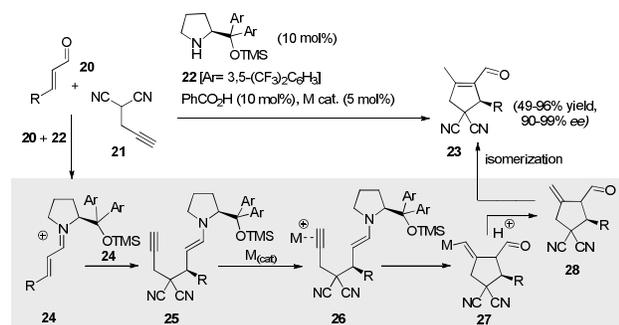
An exciting new field in the arena of multicatalysis is the use of several metal catalysts that operate orthogonally in one pot and provide highly complex molecular architectures from simple precursors. Jeong and co-workers²⁶ reported one of the first examples of a multicatalytic palladium/rhodium system with multiple C-C bonds being formed (Scheme 4). Initially, a palladium-catalyzed allylation between malonyl derivative 14 and allyl acetate 15 takes place to furnish intermediate 17. This intermediate enters a second catalytic cycle where rhodium triggers a Pauson-Khand type reaction (PKR). Via intermediates 18 and 19 the bicyclic ring system 16 is generated in excellent yield. The authors found that the chosen rhodium precatalyst is critical to the success of the multicatalytic cascade. This example demonstrates the multitude of opportunities chemists have compared to nature when developing multistep sequences, as there are no analogous catalytic reactions known in nature.



Scheme 4. Multicatalytic Pd-Rh-promoted allylic alkylation-PKR (for purposes of clarity, only key intermediates are depicted).

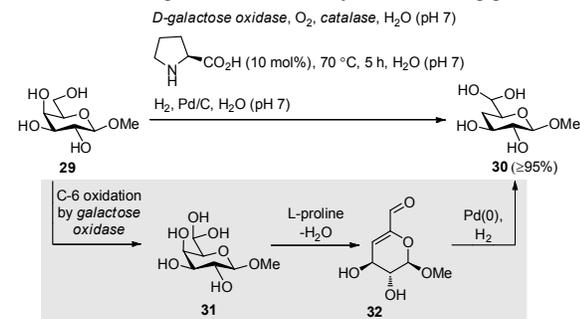
Several examples of cooperative utilisation of transition metal catalysts, organocatalysts and even enzymes in one-pot have appeared in the literature. Jørgensen *et al.* showed that proline-derivative 22 and a π -Lewis acid based on gold or copper, can affect the reaction of propargyl malononitrile 21 with α,β -unsaturated aldehydes 20, to furnish highly enantioenriched cyclopentene carbaldehydes 23 (Scheme 5).²⁷

Initially, the organocatalyst is responsible for iminium formation and directs highly selective 1,4-addition (intermediates 24 and 25). Then, the π -Lewis acid takes over (intermediate 26) disposing the alkyne toward intramolecular 5-*exo*-dig cyclisation to provide chiral cyclopentene carbaldehydes (intermediates 27 and 28). From control experiments it became evident that during transformation of 26 to 27 the two catalysts operate cooperatively.



Scheme 5. Dual metal-organocatalytic synthesis of cyclopentene carbaldehydes **23**; Mcat. = Cu(OTf)₂ (5 mol%)/PPh₃ (20 mol%), or CuOTf (5 mol%)/PPh₃ (20 mol%), or Au(NTf₂) (5 mol%) (for purposes of clarity, only key intermediates are depicted).

As early as 2002 a multicatalytic system was reported that utilises an enzyme, an organocatalyst and a metal catalyst in water as the solvent in one-pot (Scheme 6).²⁸ The conversion of methyl β-D-galactoside (**29**) to methyl 4-deoxy-6-aldehydo-β-D-galactoside (**30**) was initiated by the action of D-galactose oxidase that affects the oxidation at C-6. The resulting aldehyde (intermediate **31**), in equilibrium with its hydrate, was warmed up and L-proline promoted dehydration to form the corresponding enal (intermediate **32**). Finally, hydrogenation over palladium on charcoal gave the deoxygenated hexoside **30** in excellent yield. This process proceeded without purification or isolation of intermediates, only reagents are added and conditions are altered. Remarkably, this process does not require protection of hydroxyl groups and it demonstrates the power of multicatalysis in multistep processes.



Scheme 6. One-pot triple catalysis for the conversion of methyl β-D-galactoside (**29**) to methyl 4-deoxy-6-aldehydo-β-D-galactoside (**30**).

4.2 Strategy 2: Domino reactions

Domino or cascade reactions have emerged as an important tool for the organic chemist and over the past 20 years there has been a great deal of research in this area and a wealth of new processes have become available. As defined by Tietze, a domino reaction is a process whereby a starting material passes through two or more intermediates on its way to the final product, in a sequential manner, where functionality generated in the first step allows for further transformations to take place.²⁹ The whole process occurs under the same reaction conditions and commonly all components are present from the very beginning (figure 5).

Nature also relies on this concept and in this context terpene cyclases are excellent examples.³⁰ These enzymes have the ability to control the reactivity of intermediate carbocations of a domino sequence that results in the large number of known oligocyclic terpenes. During cyclisation the ring sizes and the configuration of stereogenic centers formed are dictated by the defined folding of the linear pyrophosphate precursor and governed by stereoelectronic effects in the enzyme active site, where “special” conformations can be stabilised. Commonly, the domino sequence is initiated by formation of a 3,3-disubstituted allyl cation which reacts intramolecularly with one of the trisubstituted alkene moieties of the

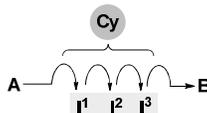
substrate. A new carbocation is formed which commonly has three options: a) internal or external trapping of a nucleophile, b) deprotonation and formation of an alkene or c) Wagner-Meerwein rearrangement and formation of a new carbocation (figure 5).³¹

Chemistry: domino/cascade reactions



short living, highly reactive intermediates
intermediates present in minute amount
no spacial compartmentation
interference into process difficult

Biosynthesis: e.g. terpene cyclases



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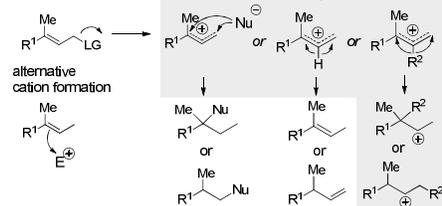
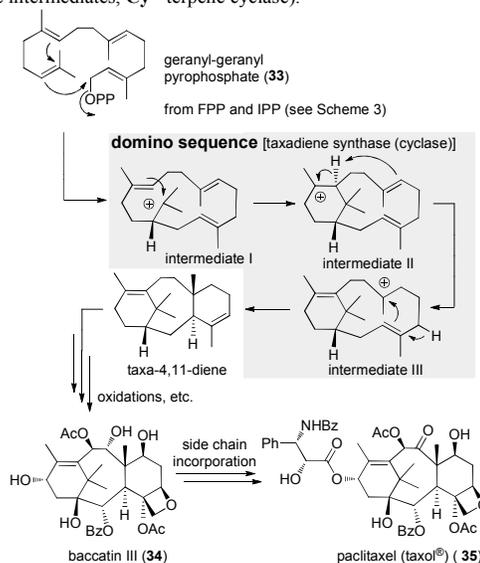


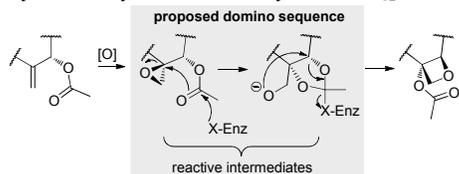
Figure 5. Principal concept of domino reactions and comparison to terpene cyclases; (A = substrate of terpene cyclases, B = cyclisation product; I¹-I³ = reactive intermediates, Cy = terpene cyclase).



Scheme 7. Biosynthesis of paclitaxel (**38**) (cationic intermediates are positioned on grey background).

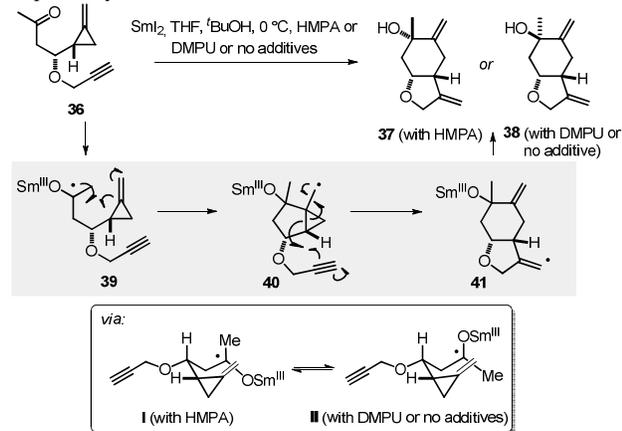
The taxadiene synthase that operates in the biosynthesis of the yew tree (*Taxus brevifolia*) metabolite paclitaxel (**35**, TAXOL[®]) is an illustrative example. Paclitaxel inhibits microtubule depolymerisation³² and commercially is one of the most successful anti-cancer compounds. This diterpene is based on a 6-8-6 tricyclic carbon backbone which is formed from geranyl-geranyl pyrophosphate (**33**) in an enzyme-catalyzed domino reaction (Scheme 7). The cascade proceeds through several cationic intermediates **I-III** formed either by π-solvolysis, elimination or H-shifts. A series of tailoring oxidations and esterifications provide paclitaxel via baccatin III (**34**). Within this biosynthesis, another remarkable domino sequence occurs during the tailoring phase of the taxane backbone to form the oxetane ring as depicted in Scheme 8.³³

Cationic intermediates also drive iminium biosyntheses of alkaloids. Anionic species govern cascade sequences in polyether and ladder polyether biosyntheses as studied in detail for monensin.³⁴ Radical cascade reactions are not very widespread in nature. Often these sequences are short and represent rearrangements commonly initiated by S-adenosyl methionin or by vitamin B₁₂.



Scheme 8. Proposed domino sequence in the formation of the oxetane ring in paclitaxel (**35**).

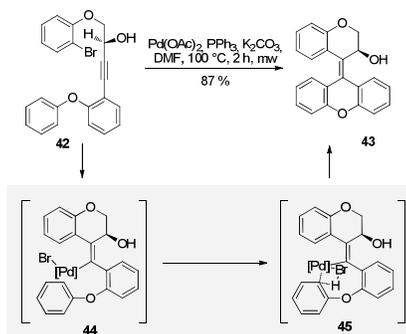
In a similar fashion chemists have utilised this concept of creating reactive intermediates based on carbocations, carbanions and C-centered radicals to build up several carbocyclic systems in one step. An impressive example was disclosed by the Kilburn group,³⁵ in which SmI₂ is used to reductively initiate a radical cascade of cyclisation-ring expansion-cyclisation via proposed intermediates **39-41** starting from cyclopropyl ketone **36**. The use of HMPA is supposed to influence the respective size of the coordinating samarium, thus stabilizing either intermediate **I** or **II** and hence controlling the relative stereochemistry of the product **37** and **38**, respectively.



Scheme 9. SmI₂ promoted cyclisation-ring expansion-cyclisation cascade by Kilburn *et al.*³⁵

Although this domino reaction differs from terpene cyclase-promoted biosyntheses in that radicals instead of carbocations are the highly reactive intermediates, the underlying principles of domino reactions are alike. In the case of terpene cyclases the resulting stereochemistry is controlled by the asymmetric environment provided by the biocatalyst which exerts a defined conformation of the substrate. In the Kilburn example the starting material preforms to a preferred conformer in solution, which can further be adjusted by choice of additional ligands on the samarium species.

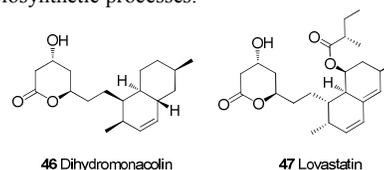
This concept of creating a reactive intermediate which initiates a second very different reaction is rather common in transition metal catalysis. Recently, Tietze *et al.* published an example of a palladium catalyzed carbopalladation-C-H activation sequence (Scheme 10).³⁶ The reaction starts with Pd insertion into the carbon halogen bond followed by a *6-exo-dig* cyclisation through carbometalation of the triple bond to furnish intermediate **44**. The close proximity of the aryl group to the metal center initiates a C-H insertion of the palladium, resulting in the formation of the product **43** after reductive elimination.



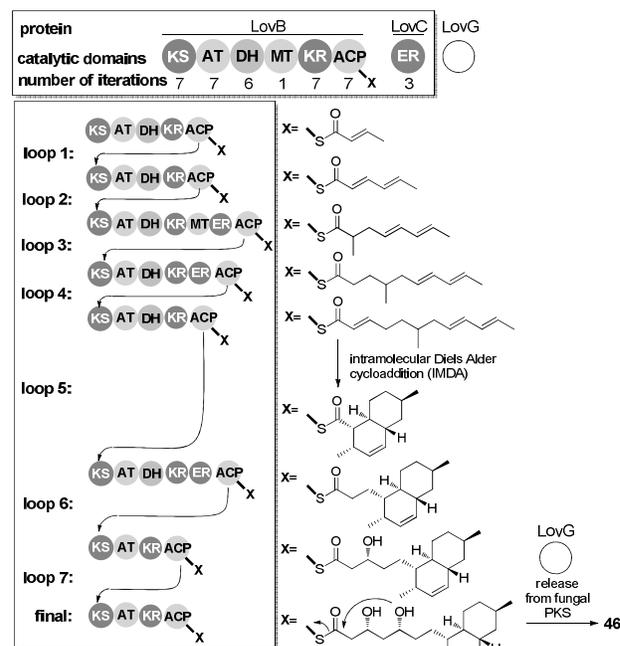
Scheme 10. A combined Heck/C-H activation.³⁶

4.3 Strategy 3: Iterative reactions

Iteration of steps as was discussed for the acyclic terpene pyrophosphates in Scheme 3, is a third concept how nature simplifies biosynthetic processes.

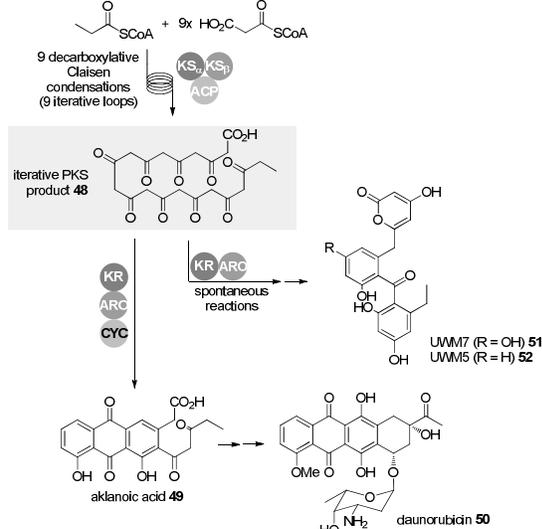


Indeed, modularity is widespread in nature and can also be found in polypeptides, oligonucleotides and oligosaccharides, indeed most natural small-molecules are biosynthesised via iterative coupling reactions. In contrast to bacterial type I polyketide synthases that are arranged in a complex modular fashion (see also erythromycin PKS in chapter 4.4), fungal polyketide synthases use only a simplified enzymatic machinery that is still able to create large and complex molecular architectures. The key for understanding these fungal PKS modules is their ability to operate iteratively.^[37] The exact mechanism of chain extension is largely unknown but their impressive iterative power is demonstrated in the reconstitution of the PKS of dihydromonacolin (**46**). Compound **46** is a precursor of the hypolipidemic drug lovastatin **47** that is produced by *Aspergillus terreus* (Scheme 11).^{38a}



Scheme 11. Dihydromonacolin (**46**), a precursor of the hypolipidemic drug lovastatin **47** and the iterative biosynthesis of **46** (KS= ketosynthase, AT= acyl transferase, DH= dehydratase, MT= methyl transferase, KR=

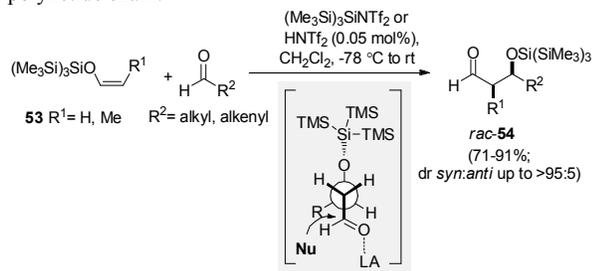
ketoreductase, ACP= acyl carrier protein, ER= enoyl reductase; TE= thioesterase, X= ACP-bound intermediates).



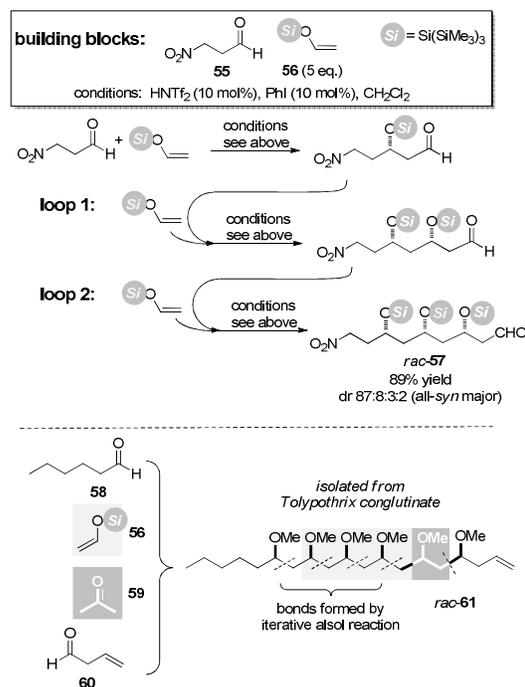
Scheme 12. Iterative biosynthesis of PKS type II decaketides to daunorubicin **50**. Lack of the cyclase DpsY results in formation of UWM5 (**52**) and UWM7 (**51**).

The catalytic domains are organised in the one multifunctional (LovB) and one monofunctional protein (LovC) that are responsible for the biosynthesis of **46**. The individual enzymatic units within the domains are iteratively used up to eight times before release from the PKS.^{38b} The final product **46** is formed after a bioprocess consisting of seven iterations that only differ in the number of enzymatic units operating and the length of the carbon backbone of the ACP-bound intermediates. As a result only seven different enzymes are required to perform 21 reactions. At first glance, the iterative process is not revealed in the structure of **46**. But the structural complexity is further increased because of an intervening intramolecular Diels-Alder cycloaddition, which most likely proceeds in a spontaneous manner (Scheme 11).

Polyketides of type II are formed by even more simplified PKS variants which are composed of only KS_{α} , KS_{β} and ACP domains.³⁹ The KS_{α} and the ACP domains work in an iterative sense, while KS_{β} , also known as chain initiation factor, programming the final length of the polyketide chain as KS_{β} converts malonyl ACP to acetyl CoA to form the starter unit. Optionally, other domains can be present such as ketoreductases (KR), cyclases (CYC) and aromatases (ARO). Interestingly, these enzymes are not an element of the PKS but are associated with it and serve to modify the linear polyketide chain.



Scheme 13. Iterative aldol reactions and proposed transition state for *syn*-product.⁴¹



Scheme 14. Iterative triple aldol reaction and in the total synthesis of the polymethoxy-1-alkene *rac*-**52**.⁴²

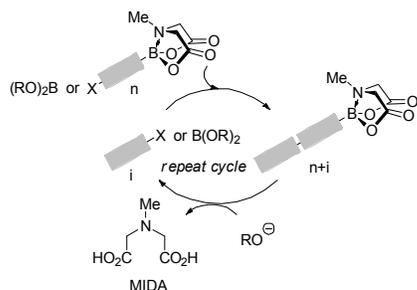
In Scheme 12 the iterative process for the biosynthesis of decaketides is summarised. Starting from one propionyl-CoA and nine malonyl-CoA extender units, nine iterative loops yield the intermediate polyketone which can be further processed to a variety of different carbon skeletons that are subsequently processed to anthracycline antibiotics such as daunorubicin (**50**). Although the factors and proteins governing and controlling the number of iterations are not fully understood yet, it is discussed that folding of the substrate in the enzyme pocket is crucial and once a certain chain length is reached the iterative process stops. Noteworthy, the biosynthesis nicely demonstrates the high reactivity of intermediate **48** – a lack of the cyclase results in spontaneous cyclisation to other products such as **51** and **52**.^{39,40}

Iterative sequences have also been developed by chemists. This strategy provides complexity in a modular fashion from simple precursors. In analogy to iteration by fungal PKS or type II PKS, the Yamamoto group found that supersilyl enol ethers **53** can be utilised for iterative aldol processes with aldehydes to yield aldol products **54** with high *syn*-selectivity (Scheme 13).⁴¹

Although a new aldehyde is formed, the second aldol reaction is retarded due to the large size of the supersilyl group. Additives such as PhI facilitated oligoaldol processes. For example one triple aldol reaction provided polyketide-type triol **57** from silyl enol ether **56** ($R = H$) and 3-nitro propanal **55** (Scheme 14). It was postulated that organoiodides react with $(Me_3Si)_3SiNTf_2$ to furnish a cationic, sterically less congested complex of the type $RI-Si(SiMe_3)_3^+ NTf_2^-$. The iterative process was successfully applied⁴³ in the synthesis of the polymethoxy-1-alkene **61** that was, among other metabolites, isolated from the polytoxin producing blue-green algae *Tolypothrix conglutinata*. Here, this aldol reaction proceeds three times before being terminated due to the steric congestion present in the product. 1-Iodo-2-phenylacetylene serves as a mediator.⁴⁴ The fact that structural parameters such as bulkiness lead to termination of the iterative cycle also likely play a role in type 2 PKS biosynthesis as discussed above.

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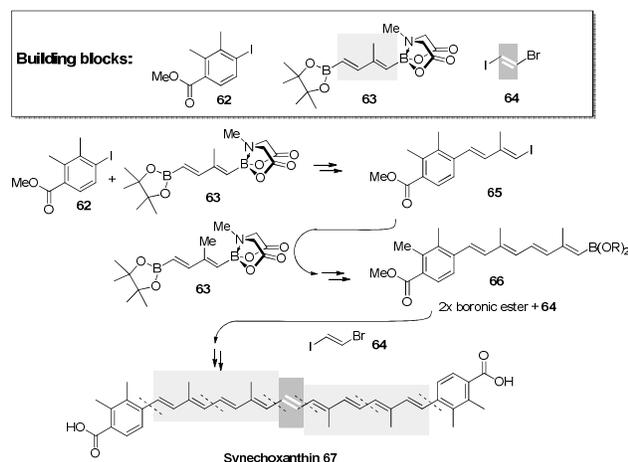
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Scheme 15. Concept of iterative cross-coupling with MIDA-boronates.

The group of Martin Burke developed a powerful methodology based upon iterative cross-coupling reactions of bifunctional MIDA-boronates (Scheme 15). In principal, the polarity of elements n or i can be flexibly chosen such that selective cross-coupling occurs in a stereoretentive sense to give the adduct $n+i$, deprotection and activation of the boronate then takes place after which time additional steps may invert the polarity of the substrate. Possible options are the conversion of the vinyl boronate into a vinyl halide or the formation of pinacol boronates. This cycle can be repeated several times to access fragments in stereochemically pure form bearing appropriate functionality for further reactions. The different principal options of this new methodology becomes evident when studying the total syntheses of the polyenes synechoxanthin (**67**) (Scheme 16).⁴⁵

The synthesis of synechoxanthin (**67**) started from aryl iodide **62** by Suzuki-Miyaura cross coupling with mixed boronate **63**. The resulting vinyl MIDA boronate moiety was transformed into the corresponding iodide **65** so that the same elongation process with building block **63** could be repeated again. The resulting polyenyl MIDA-boronate was activated as boronic ester **66** for the dimerisation step with dihalo alkene **64**. In a remarkable recent paper the Burke group demonstrated that 12 different MIDA building blocks are in principal sufficient to iteratively prepare about 75 % of all known polyene natural products.⁴⁶



Scheme 16. Synthesis of Synechoxanthin (**67**) using MIDA-boronate building blocks **62-64**.

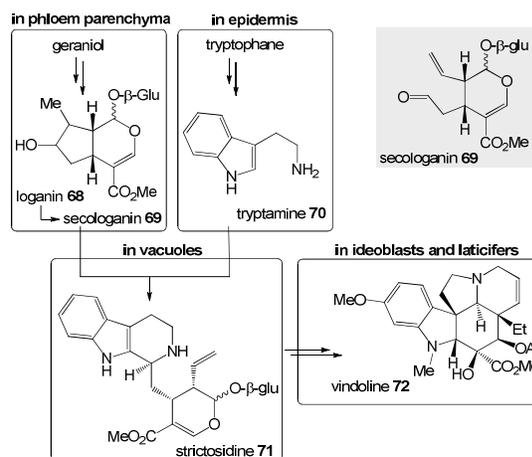
4.4 Strategy 4: Spatial compartmentation of individual steps – from polyketide biosynthesis to flow chemistry

The fourth strategy of nature's multistep biosynthesis is spatial compartmentation which can be discussed from two perspectives, a microscopic⁴⁷ and a molecular view. Microscopic compartmentation refers to cell organelles which have individual metabolic functions and which are characterised by different chemical conditions with respect to enzyme compositions and pH. Building blocks and metabolites are transferred between cell organelles or even between different cell types as found in plants by active transport systems or passively by e.g. osmosis. In that context organelles are chemical reactors which form a network with active mass transport among

them. Microscopic compartmentation has been studied in plants to some degree.²¹ For example, in *Catharanthus roseus* both intra- and intercellular compartmentation play an important role in the biosynthesis of different types of terpenoid indole alkaloid skeletons such as vindoline **72**, which is a precursor of the commercially important anticancer agents vinblastine and vincristine (Scheme 17).⁴⁸

The early terpenoid precursors such as loganin **68** from the mevalonate terpenoid pathway are made in internal phloem parenchyma, the other important precursor tryptamine **70** in the epidermis. The last step of the terpenoid precursor secologanin **69** occurs in epidermis cells. Strictosidine synthase is present in the vacuoles and catalyzes the condensation of tryptamine **70** and secologanin **69** to yield the intermediate strictosidine **71**. Finally, vindoline **72** is formed in other specialised cells, the ideoblasts and laticifers so that an intercellular transport of strictosidine must take place and the biosynthesis is compartmented. In essence, the flux through a pathway is controlled by transport from the site of production of a precursor to the site of the next enzyme.

The idea of compartmentation and passive transport is also found in synthetic chemistry at the interphase of two phases which can actively be promoted by phase transfer catalysts. E. g., this concept has been exploited for increasing the life time of free radicals using micelles as reactor compartments.²²



Scheme 17. Compartmentation in the biosynthesis of plant derived (*Catharanthus roseus*) indole alkaloid vindoline **72**.

Spatial separation can also be realised on a nanoscopic scale, where template structures based on e.g. multiple protein complexes are able to separate biosynthetic intermediates from each other. Such protein complexes are specifically folded bringing various catalytic domains in close proximity thereby allowing bound biosynthetic intermediates to be processed by transfer to the next catalytic domain (Figure 6).

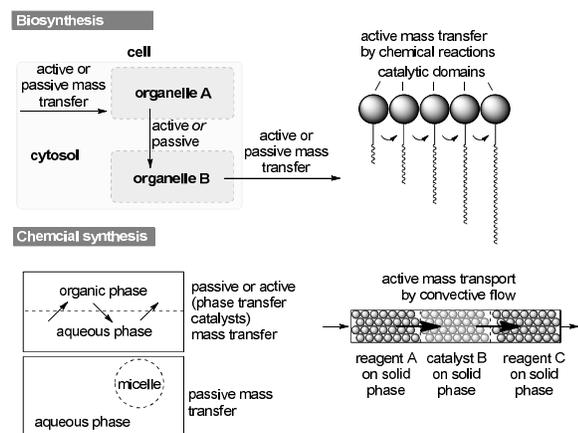


Figure 6. Examples of compartmentation in nature and in chemical synthesis.

As mentioned above, erythromycin A (**2**) is a classical example of a complex antibiotic produced by *Saccharopolyspora erythraea*. Its type 1 polyketide synthase (PKS) is able to carry out 20 steps required to form the complex carbon backbone of 6-deoxythronolide (**1**) the advanced precursor of erythromycin A (**2**) (Schemes 1 and 21). This type 1 polyketide synthases represents an examples of molecular compartmentation. This megaenzyme complex is based on three large proteins that are composed of several enzymes and which can span several so called PKS modules. Altogether, the erythromycin PKS is organised in seven modules (one starting and six extender modules) (Scheme 18).

In each module acyl transferases (AT) choose a starting building block (propionyl-CoA) or six extender units (methylmalonyl-CoA), respectively, and transfer them to the individual acyl carrier proteins (ACP) that play a central role for the following transformations. The ACP-bound intermediates are handed over from one catalytic site to the next and from one module to the next. A set of enzymatic domains (e.g. KS, KR, DH, ER), each module is responsible for C2-homologation and functional group manipulations (keto \rightarrow alcohol \rightarrow alkene \rightarrow alkane).

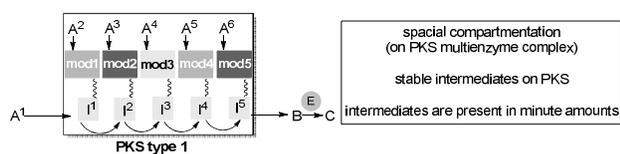
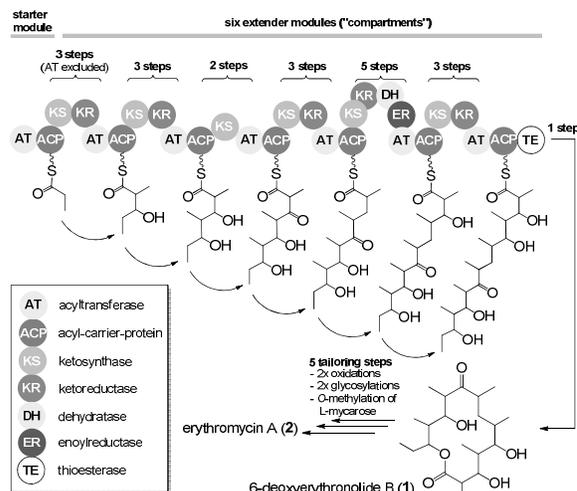


Figure 7. Polyketide biosynthesis – an example of spatial compartmentation on a molecular level.

This modular character and the permanent attachment of intermediates to the PKS lead to spatial separation of each chemical transformation. As the PKS operates continuously under steady state conditions all intermediates are only present in minute amounts (in theory one per protein).



Scheme 18. Biosynthesis of erythromycin A (**2**).

Besides the ribosomal machinery, the PKS 1 assembly line is probably the most sophisticated megaenzyme architecture nature has developed. It is highly modular and in order to smoothly operate the interaction of neighbouring catalytic domains are perfectly orchestrated. Determining the dynamic architecture of the PKS type 1 is still difficult due to the large size and high complexity. Staunton and Leadley developed a ‘double-helical’ model, that is based on the dimeric structure of subunits (Figure 8).⁴⁹ The model suggests that two identical proteins are wrapped against each other, which results in a line-up allowing all subunits to interact with one specific ACP domain. The underlying chemical concept of the PKS is compartmentation of single enzymatic steps and close spatial proximity that allow transfer of intermediates from one enzymatic domain to the next in a highly controlled manner (Figure 8). The twisting peptide chain model also explains why the ACP and KS domains, which require a close proximity in order to allow condensation, are usually found at opposite ends of the peptide chain (Figure 8, left-hand side). It also explains how a seemingly single module can be used more than once, since a dimer could be used twice in theory, if the folding is favorable. Conceptually, the PKS multistep process resembles a domino reaction with the difference that stable and not high energy intermediates are transformed from one module to the next.

While the chemical approaches towards erythromycin A (**2**) relied on multistep batch processes (see Scheme 1) the biosynthesis of erythromycin A (**2**) is a continuous steady state process in which stable intermediates are passed from one spatial environment to the next and in each compartment (module) a new composition of enzymatic activities provides a choice for individual elongation chemistry. Clearly, this free choice does not exist for domino-type multistep transformations as discussed for terpene cyclases and their reactive intermediates.

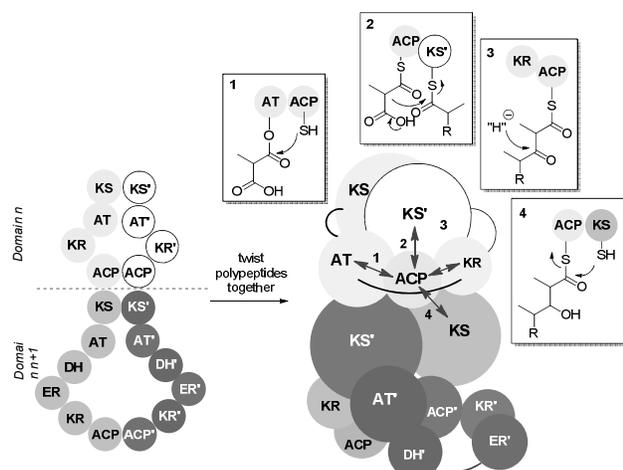


Figure 8. Double-helical model for modular polyketide synthases (PKs). The ACP is situated in the center to allow interaction with neighbouring domains AT, KS', KR and KS.

Conceptually, there is a counterpart in synthetic chemistry. Multistep flow synthesis using linearly assembled flow devices resembles such a continuous compartmented process.⁵⁰ Indeed, continuous flow processes are the best technical option to achieve spatial separation of individual transformations and the individual reactor may be regarded as an individual PKS module (Scheme 21 and Figure 9). In continuously operated multistep flow synthesis the burden of work-up is minimised and isolation protocols ideally take place after the last step. Furthermore, it is possible to set up flow systems in which two short linear synthetic flow sequences converge for allowing production of structurally complex products from two fragments. Technically, this concept is well established. E. g. one technical way to produce acrylic acid proceeds as a truly multistep continuous process. It starts with the steamcracking process of crude oil followed by a two step catalytic oxidation of propene via acrolein to acrylic acid. It is estimated that 5-10⁶ tons of acrylic acid are produced worldwide much of it in continuously.⁵¹

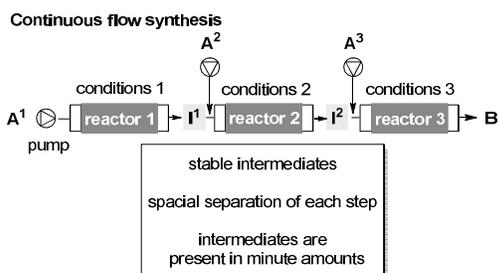
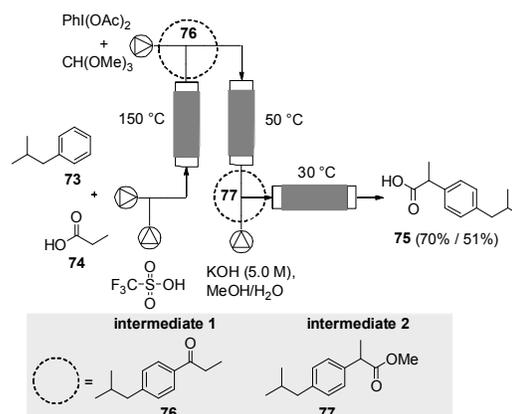


Figure 9. Principal concept of a multistep flow synthesis of product **B** starting from educt **A**¹ (**I**¹, **I**²= intermediates, **A**², **A**³= additional reactants).

Commonly, intermediates formed in continuous flow processes are chemically stable enough so that other components can be added to the stream of the reaction mixture and conditions required for the next reaction can be adapted in the next reactor (Figure 9). These options make multistep flow processes more flexible compared to multicomponent or cascade reactions.

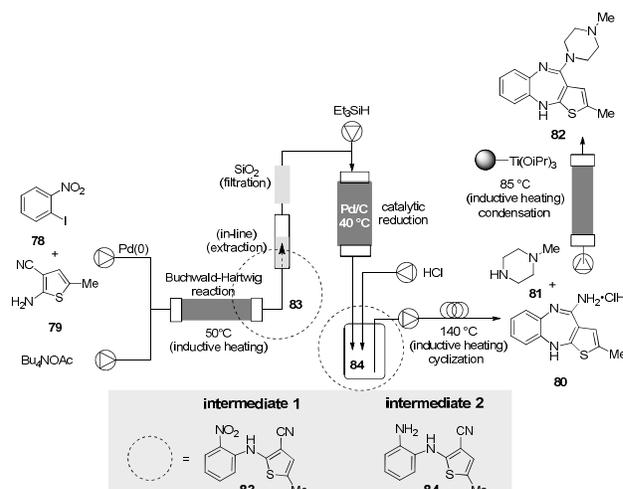
A state-of-the-art multistep flow protocol was reported by the McQuade group in 2009. Their three-step flow synthesis of ibuprofen (**75**), an anti-inflammatory and pain relieving drug, was performed continuously without purifying intermediates **76** or **77** (Scheme 19).⁵² The combined streams of neat isobutyl benzene (**73**), propionic acid (**74**) and triflic acid provided intermediate **76** (residence time: 5 min). Next, **76** was mixed with trimethyl orthoformate (TMOF) and iodobenzene diacetate by a third stream which initiated the desired 1,2 aryl migration and formation of intermediate **77**. Finally, saponification provided Ibuprofen (**75**)

which had to be purified by column chromatography (70%) or by simple recrystallisation of the crude solid (51%).



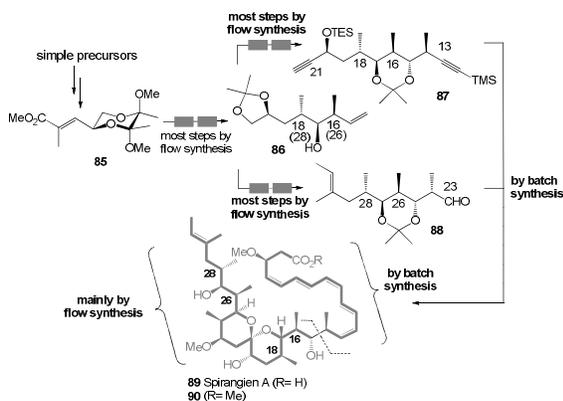
Scheme 19. McQuade's continuous three-step flow process of ibuprofen (**75**).

A second illustrative example, the continuous multistep flow synthesis of olanzapine (Zyprexa, **82**), was recently reported by our own laboratories (Scheme 20).⁵³ A continuous three-step synthesis of thieno[1,5]-benzodiazepine **80** started from aryl iodide **78** and thiophene **79** which were coupled via a Buchwald-Hartwig amination. The resulting intermediate **83** first had to be passed through an in-line extractor for removing base-derived ammonium salts then through a pad of silica that trapped palladium black before hydrogenation of the nitro group occurred in the second reactor. The aniline intermediate **84** that left this reactor was collected in a glass vessel where remaining hydrogen gas was liberated. At this point a stream of HCl in MeOH was added and the solution was injected into the tubular reactor that was heated inductively. As a result thieno[1,5]-benzodiazepine **80** was collected after 30 hours of continuous operation in 88% yield without the need of chromatographic purification. Finally, the synthesis of olanzapine **82** was achieved in 83% yield after coupling with piperazine **81** using an inductively heated fixed bed reactor containing a solid-phase bound titanium Lewis acid.



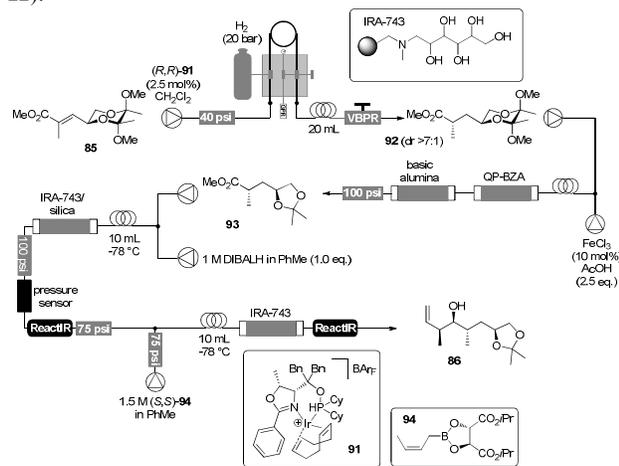
Scheme 20. The continuous multistep flow process of olanzapine (Zyprexa) (**82**).

A most remarkable work was published earlier this year by Ley and coworkers.⁵⁴ During the total synthesis of the methyl ester of spirangien A (**90**, Scheme 21) a polyketide-based natural product isolated from myxobacteria *Sorangium cellulosum* So ce90 several polyketide fragments and hence the western part were synthesised under continuous multistep flow conditions without isolation of intermediates.



Scheme 21. Ley's total synthesis of the methyl ester of Spirangien A (**90**)⁵³

Strategically, intermediate **85** was transformed into the key building block **86**. It contains a stereotriade that is found at C16-C18 and C26-C28, respectively, of spirangien A (**89**). Consequently, alkene **86** was transformed into the two advanced fragments **87** and **88**. These were merged convergently by nucleophilic addition of the alkynyl lithium derivative derived from **87** to the aldehyde present in **88**. At this point the authors switched from an overall flow process to the batch mode. The synthesis contains a lot of technical details and synthetic subtleties, so that only some key features of individual reactions and reaction sequences can be highlighted which demonstrate the state of the art of modern flow synthesis (Scheme 22).



Scheme 22. Multistep flow synthesis of polyketide fragment **86**.^[53] VBPR = variable back pressure regulator; QP = QuadraPure; BZA = benzyl amine BAR_F = tetrakis[3,5-bis(trifluoromethyl)phenyl]borate.

Utilizing the technically simple device of a tube-in-tube reactor suitable for continuous reactions with gases, the Ley group conducted an asymmetric hydrogenation using Pfaltz's catalyst (*R,R*)-**91** with alkene **85**, which yielded hydrogenation product **92** in excellent yield and with good selectivity (Scheme 22). The catalyst's special feature is associated with the fact that it can be applied for the hydrogenation of non functionalised alkenes without

the need of additional polar coordinating sites.⁵⁵ In the following, the protecting group was switched to an acetone and all reagents and byproducts were removed from product **92** by pumping the reaction mixture through a cartridge of polymer supported benzylamine and basic alumina oxide. Finally, the resulting ester **93** was selectively reduced to yield the corresponding aldehyde which was followed by the experimentally challenging Roush crotylation with *Z*-but-2-enylboronate **94** to furnish fragment **86**. It is noteworthy that aluminum containing byproducts were removed from the stream using a fixed bed material composed of a polyalcohol immobilised on the polymer IRA-743. It is now possible to perform the Roush crotylation directly by addition of the boronate to the flow system. Control of stoichiometry was achieved by measuring the concentration of the resulting aldehyde using a FlowIR cell. Data processing, allowed automatic adjustment of the flow rate of the pump that is responsible for the addition of boronate **94**.

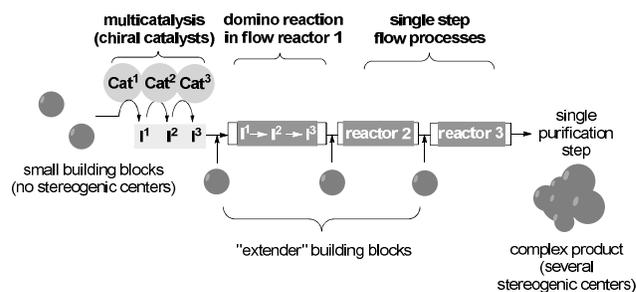
It has to be stressed, that in several aspects this total synthesis is not biomimetic, although continuous processes and compartmentation were exploited. Ley's synthesis does not follow the linear logic of the polyketide synthase but has the flexibility to dissect the spirangien A backbone into three major fragments which are merged in a convergent fashion, instead.⁵⁶

5. Outlook

This conceptual review has examined the ideal multistep synthesis by analyzing nature. This article is not about biomimetic natural product synthesis but we rather suggest that the key to efficiency is steady state processing with intermediates that are immediately transformed to the next intermediate when formed until the target secondary metabolite is formed. We discussed four strategies (multicatalysis, domino reactions, iteration and compartmentation) that commonly proceed via short-lived intermediates. Nature, unlike chemists, uses these four strategies flexibly, almost combinatorially like a juggler which is still a huge challenge for synthetic chemists. In essence, we believe that atom, step and redox-economy can only cover some major aspects of defining the ideal synthesis.

The flexible use of these and additional strategies²¹ should increase the efficiency of processes by minimizing work-up protocols of intermediates so that isolation of the final product is the only purification issue.

We believe that harnessing and enhancing these strategies should not only be a priority for academia but also for industry. We envision that a combination of these methods may one day change 'traditional' synthetic chemistry and open up rapid access to a world of natural product-like compounds with astonishing complexity with only one or two purification steps (Scheme 23). We are aware that it is still a long way to go to develop such processes, but we believe we should not shy from such an opportunity to access a larger part of the chemical space in a more economical fashion.



Scheme 23. Multistep processing towards complex molecular targets by combination of different steady state techniques.

Synthetic chemists can learn from nature by closely studying – not only individual transformations– but also how *in vivo* multistep

processes are conducted. With the multitude of reactions discovered since Wöhler's first preparation of urea in 1828, synthetic chemists have the potential to develop more powerful multistep processes than nature, if only we can manage to put them into use effectively.

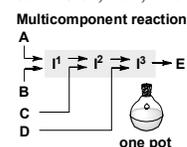
5. Notes and References

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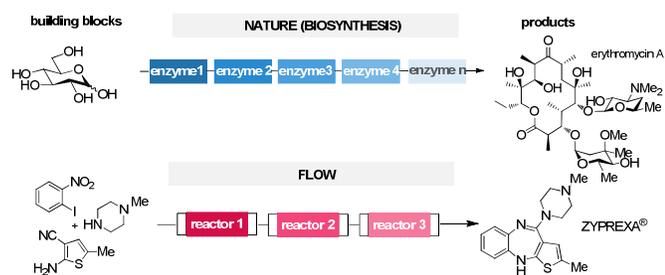
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Graphical abstract



Nature's strategy of performing ideal multistep (bio)synthesis are based on multistep, domino reactions, iteration and compartmentation. These are discussed and compared with chemical synthesis in this conceptual review.