Recent development and applications of semipinacol rearrangement reactions

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As has been well-recognized, semipinacol rearrangement functions as an exceptionally useful methodology in the synthesis of β-functionalized ketones, creation of quaternary carbon centers, and construction of challenging carbocycles. Due to their versatile utilities in organic synthesis, development of novel rearrangement reactions has been a vibrant topic that continues to shape the research field. Recent breakthroughs in novel electrophiles, tandem processes, and enantioselective catalytic transformations further enrich the toolbox of this chemistry and spur the strategic applications of this methodology in natural product synthesis. These achievements will be discussed in this minireview.

Introduction

The highly efficient formation of carbon–carbon bonds and construction of corresponding molecular skeletons are two representative key scientific topics in organic synthesis. Therefore, rearrangement reaction related synthetic strategy development, which can simultaneously achieve the above two synthetic goals in a very efficient manner, has been an evergreen topic in the synthetic chemistry community. Among the rearrangement reactions developed so far, semipinacol rearrangement, as a typical 1,2-migration process, has also attracted the broad attention of synthetic chemists. Since the inception of the term “semipinacol” to describe a specific subclass of pinacol rearrangement reactions by Tiffeneau in 1923,1 the definition of this research area has continuously extended with advancement of novel types of reactions.2–4 Generally, despite the diverse reaction patterns semipinacol rearrangement may involve, a typical process often entails the generation of a carbonyl group through a 1,2-C–C or C–H bond migration from an oxygen-containing carbon to a vicinal carbon center, which is archetypally an electrophilic carbon1 though a C-radical center has also been suggested in recent emerging radical rearrangements (Scheme 1).4 A noteworthy aspect of this class of 1,2-migration reactions is that it provides a reliable and efficient approach to access synthetically useful β-functionalized ketones. In addition, the semipinacol rearrangement also serves as a potent and versatile method for the construction of quaternary carbon centers5 that are challenging to create but pervasive in natural molecules and pharmaceuticals.6–15 These remarkable features render this rearrangement reaction a powerful tool in natural product synthesis,5,14,15 and the utilization of this rearrangement to assemble intricate carbocycles from easily accessible ones, probably the most exciting application of such reactions, has become a subject of particular interest.

As the development of semipinacol rearrangement in the last two decades has gradually become mature, summarization of a certain aspect of the venerable reaction has been documented in several previous reviews.2–5,14–17 Nonetheless, recent advances of semipinacol rearrangement in terms of novel electrophiles, radical processes,4 tandem reactions, and catalytic
enantioselective transformations\(^1\text{6,}1\text{7}\) have further enriched the repertoire of this chemistry. These breakthroughs are accompanied by the prevailing application of the rearrangement reactions as a key step/strategy in the assembly of complex natural products. In some cases, a semipinacolase is even elucidated to be responsible for catalyzing the key rearrangement in the biosynthetic pathway of certain natural molecules.\(^1\text{8}\) Therefore, a summary of these contributions, as presented in this review, would provide not only an up-to-date overview of this research field but also guidance for its future development.

The new patterns and applications of semipinacol rearrangement covered in this review will be classified on the basis of four substrate types—epoxides, allylic alcohols, \(\beta\)-hetero-substituted alcohols, and \(\alpha\)-hydroxy ketones and imines. Although some of the basic transformations of each type of substrate have been well-established, recent progress has focused on tandem and asymmetric rearrangement modes. These representative examples along with the extension of radial initiated rearrangement of allylic alcohols will be discussed herein, with an emphasis of the recent synthetic applications in each section.

### Epoxides

Epoxides are a classical type of substrate in semipinacol rearrangement. In the presence of an acid, ring-opening of oxirane would result in the formation of an \(\alpha\)-hydroxy carbocation that could induce an ensuing 1,2-migration to produce a carbonyl product. Meanwhile, the resulting aldehydes or ketones could also participate in various tandem processes, such as the Tishchenko reaction, alkylation, propargylation, and the Schmidt reaction, to produce structurally complex 1,3-diols or amides.\(^5\) In these cases, pre-preparation of epoxides from the corresponding alkenes or allylic alcohols is generally required. Thus, the development of a one-pot or tandem epoxidation/semipinacol reaction would be more step-economic since an alkene substrate could be directly used in such a cascade. In this regard, Jiao and coworkers recently developed an oxygenation/semipinacol rearrangement of cyclobutyl-derived allylic carbinol 1 (Scheme 2a).\(^1\text{9}\) Notably, oxygen was used as the epoxidation reagent and \(\text{Cu(OTf)}_2\) as the catalyst in this reaction, and a catalytic amount of Lewis acid was sufficient to induce the desired rearrangement. Apart from allylic alcohols, a cascade process of ortho-hydroxy styrenes 4 was devised by Sun’s group for the synthesis of hydrodibenzofurans 6 (Scheme 2b).\(^2\text{0}\) The formation of an ortho-quinone methide intermediate 5 was suggested after the epoxidation, which would trigger the ring-expansion of the resulting cyclopentanol to generate 6 with two consecutive quaternary stereocenters.

As the 1,2-migration of 2,3-epoxy alcohols often proceeds via a stereospecific process with the migrating group antiperiplanar to the C–O bond of the oxirane, access to enantiopure \(\beta\)-hydroxy ketone products with these reactions predominantly relies on the initial preparation of enantiopure epoxides. However, for racemic epoxide substrates with only one stereogenic center, regioselective ring-opening of the oxirane would form a prochiral \(\alpha\)-hydroxy carbocation intermediate, and thus an ensuing enantioselective migration could potentially take place to produce valuable enantiopure ketones or aldehydes. Such a catalytic stereoconvergent process has independently been realized by Zhu’s and Sun’s groups at the same time.

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Under the catalysis of either chiral N-trifyl phosphoramidate (S)-cat. 1 or phosphorous acids (R)-cat. 2 or (S)-cat. 3, racemic epoxides 7, 9, or 11 could be converted to enantioenriched α-quaternary ketones 8, 10 or 12, respectively, with high yields and enantioselectivities. Mechanistically, formation of a chiral counteranion of the deprotonated catalyst and the α-hydroxy carbocation was deemed to be crucial for inducing an enantioselective rearrangement.

The semipinacol rearrangement of epoxides has found diverse applications in natural product synthesis during the last two decades. The utility of this reaction for quaternary carbon center formation with stereochemical control has been well-recognized and continues to play an essential role in the synthesis of key intermediates of natural molecules. For a recent example, the 1,2-migration of 2,3-epoxy tertiary alcohol 13 would provide α-quaternary-β-hydroxy cyclohexanone 14 that serves as a key building block in the total synthesis of (−)-isocelorbic by Ogura and coworkers (Scheme 4). The correct installation of the C10 quaternary carbon center is attributed to a stereoselective rearrangement of the vinyl group anti to the epoxide.

In 2020, Chen and Yang’s group proposed an elegant total synthesis of (−)-spirochensilide A. During their early-stage synthesis, a two-step epoxidation/semipinacol ring-contraction consequence has been developed to convert a trans-octahydroporphalene system 15 to an octahydroindene carbocycle 17 with vicinal quaternary carbon centers (Scheme 5). Of note, only a single diastereoisomer was obtained in the rearrangement reaction of 16, indicating a highly regio- and diastereocchemical control of the ring-contraction process, and the aldehyde functionality introduced through the rearrangement could be further elaborated as a functional handle.

The skeletal 1,2-rearrangement could also be applied as a crucial strategy to guide synthetic planning. For the assembly of highly strained or complex carbocycles, alternation of the cyclic ring system via a retro-1,2-migration might provide an easily accessible precursor from the retrosynthetic viewpoint. As a representative example, such a strategy has been ingeniously applied in the total synthesis of (−)-illisimonin A (Scheme 6). As the trans 5–5 system of the molecule presents a well-recognized synthetic challenge, Rychnovsky and coworkers devised a key skeletal rearrangement to access the trans ring-system from its cis counterpart since it is calculated to be less strained (7.03 kcal mol$^{-1}$ more favorable). Furthermore, this cis-
carbocycle 21 could be readily prepared from 2-methylcyclopentane-1,3-dione 18 in three steps via an intramolecular Diels−Alder reaction of α,β-unsaturated cyclopentenone 19. The late-stage semipinacol rearrangement of 2,3-epoxy alcohol 22 derived from ester 21 would provide the tricyclic ketone 23 with the establishment of the strained trans 5-5 ring. Finally, the application of White’s acid-directed C−H oxidation of 24 enabled the construction of the bridging lactone and completed the total synthesis.26,27

**All allylic alcohols**

Allylic alcohols are a privileged type of substrate in the development of novel semipinacol rearrangements because of the availability of these compounds from simple chemical feedstocks. In classical reactions, activation of the alkene group by an electrophile would generate an α-hydroxy carbocation intermediate, thus triggering the subsequent 1,2-migration to produce β-functionalized ketones (Table 1). Alternatively, addition of a radical species to the double bond is also feasible. In this case, an intermediary α-hydroxy radical is initially generated, which in most cases is further oxidized to the α-hydroxy carbeneium prior to the rearrangement (Table 1, path a). However, when the migrating group is an arene or heteroarene, a neophyl-type 1,2-radical migration followed by oxidation is usually suggested (Table 1, path b). Although a wide range of electrophiles capable of initiating the rearrangement have been well-documented, interest in the discovery of novel types of electrophiles and radical initiators continues. Among these breakthroughs, novel catalytic techniques, such as photocatalysis and electrocatalysis, have also enabled reactions under environmentally benign conditions, thus preventing the use of excess oxidants or toxic reagents.

Recent advancements of electrophiles and radicals in the semipinacol rearrangement reactions of allylic alcohols are summarized in Table 1. For a nitrogen group element, nitrosonium (NO+) was proven to be able to promote the rearrangement reaction, producing oximes and oxime ethers with a spirocyclic skeleton. This methodology is a rare example of oxime synthesis from non-ketone or hydroxylamine substrates. Besides N-centered electrophiles, an N-radical induced rearrangement has been reported wherein N-fluorobenzenesulfonylimide (NFSI) was used as the radical source under the catalysis of Cu(i). Additionally, with the assistance of photoredox-catalysis, the phosphorylation/semipinacol rearrangement could be realized either with a diaryl(alkyl)-phosphine oxide or oxime phosphonate substrate.

Chalcogen, specifically sulfur and selenium, related electrophiles and radicals are also viable species in promoting the rearrangement reactions of allylic alcohols. For example, the SCF₃ radical could be generated under oxidative conditions, while the SCF₃, SCN, or SeR electrophile was commonly released from certain electrophilic reagents such as PhNHSCF₃, N-trifluoromethylthiosaccharin, (PhSO₂)₂NSCN, PhSeBr, or RSeR in the presence of an acidic promoter. In 2019, a sulfenylation/semipinacol rearrangement has been established independently by several groups. Among these reactions, various types of reagents, such as ArN₂BF₄ and (DABCO)−(SO₂)₃, ArSO₂H, and ArSO₂Cl were used to generate arylsulfonyl radicals by means of photoredox-catalysis. Of note, electrochemical conditions have also been applied for the generation of arylsulfonyl radicals from either ArSO₂NH₂-NH₂ or PhSO₄Na, which provide mild and convenient protocols for the synthesis of β-sulfonated ketones.

The halogenation/semipinacol rearrangement of allylic alcohols has found wide application in the synthesis of β-halogenated ketones. Although electrophilic halogenation reagents have been pervasively used in such transformations, these reagents are generally expensive and toxic. In comparison, the use of inorganic halide salts as halogen sources would be more appealing since they are low-cost and eco-friendly. These approaches have recently been realized taking advantage of either oxidative conditions (PIDA) or electrocatalysis. In each case, sodium or magnesium halide was used respectively to generate the halogenium in situ to induce the rearrangement.

Apart from heteroatom-centered electrophiles and radicals, carbon-centered radicals are equally competent in inducing migration reactions. The alkyl radicals generated from RBr, RB(OH)₂, or N-aclyloxypythalimides under photoredox-catalytic conditions could be utilized for the development of allylative semipinacol rearrangement reactions. These methodologies provide efficient routes to β-functionalized ketones under mild and environmentally benign conditions. Similarly, trifluoromethylation/rearrangement reactions could also be conducted under photolysis or electrocatalytic conditions to produce a wide range of β-trifluoromethyl ketones with high efficiency.

Notably, a catalytic radical-polar cross over reaction has been developed in 2018 in which the generation of a hydrogen radical was responsible for promoting the semipinacol rearrangement of allylic alcohols. Compared with the proton induced reactions, this approach features a much broader substrate scope that could be a useful complement to the existing methods.

Advancement of electrophile induced enantioselective semipinacol rearrangement of allylic alcohols has also received considerable attention from the synthetic community with continued development of novel types of catalytic modes. In 2019, Gong and coworkers developed a notable catalytic enantioselective tandem allylic alkylation/oxidative rearrangement using a chiral iodine catalyst (Scheme 7). Oxidation of cat. 6 by selectfluor and TsoH would in situ generate a hypervalent iodine(III) species, which serves as the real catalyst for the enantioselective oxidative rearrangement. Under the optimal conditions, various β-alkoxylated ketones 27 were obtained with good yields and excellent enantioselectivities.

Taking advantage of Lewis base catalysis, Tu’s group accomplished an asymmetric sulfonylation/semipinacol rearrangement that provides a straightforward methodology for the synthesis of a wide range of α-quaternary carbon-containing β-arylhthio ketones 30 from allylic alcohols 28 (Scheme 8). The best enantioselection was achieved when selenide (R)-cat. 7 was used as the Lewis base catalyst and chiral phosphoric acid (S)-CPA as a cocatalyst. The indispensable role of (S)-CPA for a highly enantioselective transformation was studied by...
computational methods which indicate a steric repulsion of \((R)\)-cat. 7 and \((S)\)-CPA to be responsible for the facile selection during the sulfonyl group transfer event.

In the past two decades, significant studies have been devoted to the development of catalytic enantioselective halogenation/rearrangement of allylic alcohols, and precedent
reactions focusing on the establishment of stereogenic sp<sup>3</sup> carbon centers with point chirality were well-established. Recently, Yeung and co-workers further advanced this methodology with an application of the rearrangement for the introduction of axial chirality (Scheme 9). As the methyl substituents in the 4,5-dimethylfluorene system (31) are spatially remote, rotation of the diaryl C–C bond is facile, resulting in the racemization of the atropisomers. However, when 4,5-dimethylphenanthrene product 32 was generated via ring expansion of 31, proximity of the methyl substituents in 32 would render a high rotation barrier of the diaryl C–C bond. In this case, two conformationally stable atropisomers could be obtained. With the application of semipinacol rearrangement, the authors developed a dynamic-kinetic resolution strategy to simultaneously assemble the quaternary stereogenic carbon center as well as the axial chirality of 32 from the halogenation/rearrangement of racemic alcohol 31 using (DHQD)<sub>2</sub>PHAL as the catalyst (cat. 8). The diastereoselectivity of the rearrangement can be rationalized by initial formation of a preorganized intermediate A or B from each atropisomer of 31. The less steric interaction between the catalyst and the substrate in A would favour the formation of 32.

Compared with heteroatom-electrophiles, asymmetric semipinacol rearrangement reactions initiated by carbon-electrophiles are rather limited, although they are synthetically more useful in generating complex ketone scaffolds. In 2017, Gaunt and coworkers developed an elegant enantioselective arylation/semipinacol rearrangement of tertiary allylic alcohols (Scheme 10a). When Cu(OtF)<sub>2</sub> was used as the catalyst in combination with a Box ligand L2, the reactions of 33 or 36 with diaryliodonium salts 34 could proceed smoothly in the presence of 2,6-di-tert-butylpyridine (DTBP) to produce β-aryl spirocyclic ketones 35 or 37, respectively, in high yields and enantiomeric ratios. Notably, the method features a broad substrate scope in terms of diaryliodonium salts and indene or dihydropyran derived allylic alcohols. Ring expansions of cyclobutyl, cyclopentyl, oxetane, and azetidine-containing substrates were all feasible. In the same year, Zhu’s group also described their independent development of the tandem reaction (Scheme 10b). The CuCl catalyst with another Box ligand L3 was used in this case. Similarly, high yields and enantioselectivities were obtained with mainly indene-derived allylic alcohols.

In addition to cascade reactions initiated by electrophiles and radicals, the semipinacol rearrangement of allylic alcohols could be incorporated into other more complex tandem or one-pot processes. These often involve the combination of other reaction types with the rearrangement for the purpose of generating molecular complexity in an expeditious manner. For example, a tandem hydroacylation/semipinacol rearrangement of alkynyl cyclobutanols with salicylaldehydes has been established by Zhang and co-workers, offering an efficient protocol for the preparation of 2-(2-oxo-2-phenylethyl)cyclopentanones with multiple substitutions (Scheme 11).
The transformation was carried out with a catalytic amount of rhodium(i) catalyst and a PPh₃ ligand. While the reactions of terminal alkynes 41 with chelating aldehydes 42 could directly afford 1,4-diketones 43, a step-wise procedure was needed for the transformation of disubstituted alkynes 44.

A recent report by Zhang and Tu describes a one-pot semipinacol rearrangement/Michael addition/Henry reaction that enables rapid assembly of polyfunctionalized carbocycles from vinylogous α-ketols 46 and nitroalkenes 47 (Scheme 12).³⁶ Initial addition of TMSOTf would trigger the ring-expansion of 46 to form a rearranged enolate intermediate that could then add to the nitroolefin partner upon activation by TiCl₄. Once the adduct was formed, an intramolecular Henry cyclization would take place to deliver tricyclic ketones 48 and 48¹ as a pair of stereoisomers.

In 2020, Yu and Wang established a Au(i)-catalyzed tandem cyclization/seminapicol rearrangement of allene-substituted alcohols 49 (Scheme 13a),³⁹ which provides a direct synthetic approach for the preparation of quaternary carbon-containing cyclohexenes 51. On the basis of this work, Wang and Tu further developed a two-step Castro–Stephens coupling/tandem acyloxy shift/cyclization/seminapicol rearrangement sequence for the synthesis of such cyclohexenes in a more modular fashion (Scheme 13b).⁴⁰ The coupling of terminal alkynes 52 with allylic bromides 53 was followed by a Au(i)-catalyzed cascade mechanistically similar to the allene-cyclization/rearrangement process. Specifically, activation of the alkyne group of 54 by Au(i) would trigger an acyloxy shift to generate an allylic carbenium 56 that could induce the cyclization/seminapicol rearrangement, as in the reaction of allene 50, to finally give rise to the spirocyclic ketones 57. Besides the ring-expansion rearrangement, migration of aryl groups was also feasible in this cascade transformation. In these cases, a wide range of cyclohexenes 59 with an aryl substituted quaternary carbon center were obtained. A synthetic application of this highly efficient procedure was also demonstrated in the assembly of the core carbon skeleton of waihoensene, a diterpene compound with a unique tetracyclic ring-system and consecutive quaternary-center arrangement. Cyclobutanol 61 prepared from 60 in 6 steps could be transformed into tricyclic ketone 62 facilely using this methodology. Late-stage functional group manipulations followed by an aldol cyclization finally gave ketone 63 with consistent relative configuration with the natural product.

An exciting direction surrounding tandem reactions is their application in natural product synthesis. Accordingly, development of such rearrangement-involving cascades for total synthesis purpose has been an ongoing topic in this research field. Recently, Zhang and Tu developed an enantioselective aldehyde α-alkylation/seminapicol rearrangement taking advantage of organo-SOMO catalysis.⁴¹ As depicted in Scheme 14, oxidation of the enamine generated from aldehyde 65 and secondary amine catalyst cat. 9 (ref. 62–64) would form a radical cation that could add to the alkene of allylic alcohol 64, thus promoting the ring-enlargement rearrangement. A broad range of enantioenriched α-quaternary cyclopentanones can be accessed using this method. Notably, a synthetic utility of this approach has been demonstrated in the catalytic asymmetric synthesis of (+)-cerapicol. After preparation of aldehyde 68 from bromide 67 in 5 steps, it was transformed to bicyclic 1,5-dicarbonyl compound 69 through an intramolecular version of the tandem process. Late-stage aldol cyclization and functional group manipulations finally completed the total synthesis.

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**Scheme 11** Tandem hydroacylation/seminapicol rearrangement.

**Scheme 12** Semipinacol rearrangement/Michael addition/Henry reaction.

**Scheme 13** Tandem cyclization/seminapicol and Castro–Stephens coupling/tandem acyloxy shift/cyclization/seminapicol rearrangement.
The simultaneous creation of vicinal all-carbon quaternary carbon centers presents a nontrivial task in synthetic chemistry due to their congested chemical environment, although such structural moieties broadly exist in diverse natural molecules, especially indole derived alkaloids. Aiming at assembling a bis(spirocyclic) indole framework with continuous quaternary carbon centers, Tu’s group established a Bischler–Napieralski/semipinacol rearrangement that enables the transformation of tryptamine-derived allylic alcohols to tetracyclic indoles in a highly diastereoselective fashion (Scheme 15). Upon treatment of bis(spirocyclic) indole under acidic conditions, the following intramolecular Mannich reaction would produce bridge[2.2.1]cyclic ketone. This advanced pentacyclic intermediate could be applied in the collective total synthesis of four aspidofractinine alkaloids, further showcasing the synthetic utility of this methodology.

In terpene synthesis, due to the intrinsic complexity of certain carbon skeletons, direct assembly of ring-systems using cyclization methods would sometimes be arduous. In this case, alternation of the carbocycle to an easily accessible one by means of 1,2-carbon migration thereby provides a synthetic solution. Such a strategy has been elegantly performed by Luo and coworkers in the total synthesis of (-)-oridonin (Scheme 16). In order to construct the central tetracyclic carbocycles in a rapid fashion, a Nazarov/Hosomi–Sakurai cascade of (prepared from the coupling of segments 77 and 79) followed by a singlet oxygen ene reaction was developed for the synthesis of ketone (81). This highly efficient tandem reaction enables the establishment of a central cyclopentanone and the bicyclo[3.2.1]octene motif as well as the creation of four consecutive stereogenic centers. Although both carbocycles assembled are not consistent with the ones embedded in (-)-oridonin, the authors suggested that later skeletal rearrangement via 1,2-carbon bond migration would adjust the ring-system. Therefore, ketone (81) was transformed into allylic alcohol (82) in a 3-step sequence, which was subjected to a bromination/semipinacol rearrangement to produce β-bromide ketone (83) with alternation of the central cyclopentanone to a cyclohexane scaffold. Subsequently, after protecting and functional group manipulations, the resulting epoxy (84) could undergo another semipinacol rearrangement in the presence of Lewis acid EtAlCl2 to give (85) with settlement of the functional bridge ring-system. From this advanced intermediate, the total synthesis can be accomplished in another 5-step transformations.

As another strategic application, the ring-expansion semipinacol rearrangement of a dihydrofuran derived cyclobutanol has been used in the total synthesis of natural alkaloid lycoponcumin A. In Zhang and Tu’s studies (Scheme 17), a highly functionalized allylic alcohol (87) was prepared from aldehyde (86) in five steps and could be converted to cyclopentanone (88) via a highly regio- and diastereoselective ring-enlargement rearrangement. This key transformation established the typical cis-fused[6.5] ring-system of a fawcettinimine-type Lycopodium alkaloid. After functional group interconversions, a late-stage transannular 1,3-dipolar cyclization of (89) was developed to produce pentacyclic compound (90) with the construction of the tetrahydroisoxazole ring. Finally, another four transformations culminate in the total synthesis. It is worth noting that from advanced intermediate (88) total synthesis of sieboldine A could also be achieved.
The 1,2-migration of β-heterosubstituted alcohols with loss of a leaving group has found wide application in total synthesis, though the development of novel types of such reactions is relatively limited. Strategic use of this type of semipinacol rearrangement to transform an easily accessible carbon skeleton into a synthetically complicated one would provide an overall convenient route in natural product synthesis, as represented in the elegant total synthesis of (-)-gardmultimine A by She’s group (Scheme 18). From known indole ester 91, they developed a synthetic sequence including a C–H borylation to prepare the tetracyclic ketone 92. Next, an oxidative rearrangement of 92 was designed and realized to transform the fused tetrahydro-pyrido indole ring system into a spirooxindole compound 96. From precedent studies, this cascade reaction may involve the initial formation of β-chlorinated hemiaminal 94 from indole chlorination product 93. Then, an intermediate benzylic carbocation 95 would presumably be formed from 94 and trigger the following ring-contraction semipinacol rearrangement to afford 96 in 75% yield. The total synthesis of (-)-gardmultimine A could then be accomplished in another 11 steps.

In 2020, Ma and coworkers reported the first total synthesis of kopsinitarine E wherein the semipinacol rearrangement of a β-iodinated alcohol was also applied as a crucial strategic step (Scheme 19). Their synthesis began with the preparation of tetracyclic bromide 98 from known Boc-protected carbazolone 97 in 9 steps. Then, a highly efficient SmI2-mediated Dieckmann condensation/intramolecular Prins-type cyclization of 98 was developed to assemble the complex 8-membered bridge ring of the natural product and generate a β-hydroxy ketone compound 99. At this stage, switch of the synthetically accessible central N-hetero[3.3.1]bridge cycle to a desired N-hetero[4.2.1]bridge ring system should be achieved. Though challenging, this skeletal reorganization could be successfully accomplished using the semipinacol rearrangement of β-iodinated alcohol 100 prepared from iodination of 99. Late-stage formation of the hemiaminal and Mannich cyclization finally enabled the total synthesis.

Skeletal rearranged steroids pinnigorgiols B and E are appealing targets for chemical synthesis due to their intriguing tricyclic decane framework and potential cytotoxic bioactivities. In 2018, Li’s group had developed an effective strategy to construct such core structures in their elegant total synthesis of aplysiasecosterol A, though the biosynthetic pathway of the scaffold from a steroid precursor is still yet to be elucidated. Inspired by a proposed biosynthetic α-ketol rearrangement cascade, Gui and coworkers ingeniously accomplished a concise synthesis of pinnigorgiols B and E from an inexpensive steroid. As depicted in Scheme 20, dehydroergosterol 102 was converted to mesylate 103 in 5 steps. Subsequent semipinacol rearrangement of 103 under basic conditions would
afford ketone 104 with the conversion of the trans-bicyclo[4.4.0]decane to the cis-bicyclo[3.3.0]decane ring system. After cleavage of the alkene and cycloheptanone functionality via a 7-step consequence to give thiolester 105, a highly efficient acyl radical cyclization cascade of 105 was developed that enabled the formation of the key tricyclic decane framework and completed the synthesis of pinnigorgiol E, which was then converted to pinnigorgiol B via deacetylation.

**α-Hydroxy ketones and imines**

The semipinacol rearrangement of α-hydroxy carbonyls and imines, also known as acylin rearrangement, is a rare type of reversible 1,2-migration reaction, which favors the formation of a thermodynamically more stable product from an α-ketol precursor. Because of the reversibility of the rearrangement process, development of such a catalytic enantioselective reaction presents a challenging task in asymmetric catalysis. Breakthroughs from Maruoka’s and co-workers in 2017. When α-hydroxy acetals 112 might be controlled by the formation of an ion pair between the in situ formed oxocarbenium ion and the conjugate anion of the catalyst.

Compared with the wide substrate patterns of acylin rearrangements of α-hydroxy carbonyls, studies on the reactions of α-hydroxy imines mainly focus on the rearrangement of indol-3-ols, which represents a powerful methodology to access 2,2-disubstituted indolin-3-ones. Although stepwise preparation of indol-3-ols and semipinacol rearrangement has been well-documented, direct reaction of 2,3-disubstituted indolin-3-ones in a tandem oxidation/migration process would be more efficient. Such an oxidation/semipinacol rearrangement has been independently investigated by Brasholz’s and Lu and Xiao’s groups (Scheme 23a and b). With the aid of photocatalysis, oxygen could be used as an oxidant for the tandem transformation, producing a wide range of 2,2-disubstituted indolin-3-ones (116 or 121) in both catalytic systems. Recently, a catalytic enantioselective variant of the tandem process has been realized by Zhao and Jiang’s group (Scheme 23c). The SPINOL-derived CPA cat. 11 was found to be an optimal catalyst in this reaction when combined with a dicyanopyrazine-derived chromophore photoredox catalyst. Generally, moderate to high product

**Scheme 22** Enantioselective acylin rearrangement of α-hydroxy acetals.

**Scheme 23** Aerobic oxidation/semipinacol rearrangement of 2,3-disubstituted indoles.
Conclusion and perspectives

With recent significant advancement in the tandem process and enantioselective catalysis, myriad transformations regarding semipinacol rearrangement have been continuously developed that inarguably demonstrate the vitality of this reaction in chemical synthesis. Given the widely recognized feats of β-functionalized ketone synthesis, quaternary carbon center formation, and skeletal conversion of carbocycles, such a rearrangement reaction has been employed as a guiding strategy in natural product synthesis, inspiring novel disconnection logics in synthesis design. While these impressive contributions have largely boosted the research field, there are still exciting new frontiers to further exploit the reaction potential. For instance, development of carbon-electrophile-initiated enantioselective rearrangements, though in its infancy, is highly appealing because of the molecular complexity that the reaction can generate from simple precursors. Furthermore, most studies focus on the simple use of a 1,2-C–C bond migration reaction, and the application of novel tandem rearrangement reactions in total synthesis is still sparse, in particular, the construction of polycyclic rings or systems with multiple stereogenic centers. We hope that our summarization of the existing achievements in this review with an updated outlook of the research field would inspire future innovations on semipinacol rearrangement.

Author contributions

All authors conceptualised the review. X.-M. Z., B.-S. L. and S.-H. W. wrote and revised the manuscript.

Conflicts of interest

There are no conflicts to declare.

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Notes and references