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FeCl₂-Mediated Rearrangement of Allylic Alcohols
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Supporting Information

ABSTRACT: A mild, one-pot procedure to produce 3-substituted allylic alcohols from α,β-unsaturated ketones is described. The addition of an organolithium nucleophile produces a tertiary allylic alcohol as an intermediate, which undergoes a 1,3-OH-migration assisted by FeCl₂. The proposed mechanism indicates that a syn-facial migration occurs for the major product. Yields as high as 98% for the one-pot reaction are reported.

INTRODUCTION

Allylic alcohols are indispensable functional groups in the synthesis of structurally complex organic molecules.1−8 3,3'-Disubstituted allylic alcohols, in particular, are often more difficult to synthesize than their monosubstituted counterparts. Though methods such as the conjugate addition of nucleophiles to ynoles and the addition of organometallic reagents to propargylic alcohols have been described,9−13 one method that has received little attention is the 1,3-migration of the hydroxy group in tertiary allylic alcohols. This transformation has been primarily catalyzed by oxo catalysts,14−16 though oxidative palladium catalysis has also been employed to perform the migration and oxidize the allylic alcohol to a β-disubstituted enone.17 This rearrangement can be performed by enzymes to form enones.2 A rhenium-assisted rearrangement has also been used in the synthesis of semisquarates.18 Trifluoroacetic acid has also been used to isomerize allylic alcohols, and this method has been applied to the synthesis of valerenic acid, which binds to both the GABA_A and 5-HT_3A receptors, and is used as a treatment of insomnia.19 Acid-assisted allylic alcohol rearrangement was also used in the synthesis of two quinolone natural products isolated from Pseudonocardia sp.20 Additionally, one can envision that this rearrangement could be used to create artemisinin-like antimalarial drugs via Singh’s synthetic scheme (Scheme 1).21

Iron has recently been studied as a catalyst for a number of coupling reactions.12−28 The most widely used of these processes are alternatives to traditional palladium-catalyzed transformations, such as the Kumada coupling and C−H arylation and alkylation reactions.22,29,30 Even a few examples of substitution reactions via π-allyl iron intermediates have been reported.31−35 Iron is preferable to late transition metal catalysts due to its low expense and toxicity. Additionally, iron catalysts are often capable of performing synthetic steps that conventional late transition metal catalysts cannot perform, such as breaking and functionalizing strong C−O bonds. Herein, we describe a one-pot addition of an organolithium reagent to an α,β-un saturated ketone, followed by an iron-mediated 1,3-rearrangement reaction. We propose that the novel reaction proceeds via the formal cleavage of a C−O bond and the intermediacy of an allylic cation.

RESULTS AND DISCUSSION

During the course of our studies on iron-mediated reactions involving organolithium and organomagnesium nucleophiles,36 we discovered that the addition of an organolithium reagent to an α,β-un saturated ketone in the presence of an iron salt does not result in the expected 1,4-addition products, but rather an isomeric allylic alcohol is formed (2).

Cyclohex-2-enone, (1) was chosen as a reliable and simple substrate for reaction optimization (Table 1). A number of
both iron(II) and iron(III) salts were selected, out of which FeCl₂ was determined as the most efficient reagent. In general, iron(III) salts were inferior to iron(II) salts. Diethyl ether (with or without butylated hydroxytoluene (BHT) stabilization) was the most desirable solvent for the rearrangement, and the other common ether solvents, THF and Me-THF, (with or without butylated hydroxytoluene (BHT) stabilization) was the most desirable solvent for the rearrangement, and that commercial, stabilized ether was a suitable solvent for the addition/migration. The reduction of the reaction, and that commercial, stabilized ether was a suitable solvent for the rearrangement, and the other common ether solvents, THF and Me-THF, (with or without butylated hydroxytoluene (BHT) stabilization) was the most desirable solvent for the rearrangement, and the other common ether solvents, THF and Me-THF, produced low or no yields of the desired rearranged product, 2. Although the reaction proceeded in both inhibited and purified Et₂O, the addition of 1 equiv of BHT further hindered the reaction (entry 11), indicating that the BHT preservative was not involved in the reaction, and that commercial, stabilized ether was a suitable solvent for the addition/migration. The reduction of the loading of FeCl₂ from 1 equiv to 10 mol % of FeCl₂ resulted in an 8% yield (entry 13), indicating that the reaction required a stoichiometric amount of the iron reagent.

The scope and limitations of the addition-rearrangement with respect to both linear and cyclic α,β-unsaturated ketone substrates were also investigated (Table 2). Cycloalkenones were found to be the best substrates, with cyclohex-2-enone (1) giving the highest yield. Linear α,β-unsaturated ketones (entries 3–5) produced only the 1,2-addition product.

We then investigated the scope and limitation of organolithium and Grignard nucleophiles (Table 3). Alkyl lithium reagents did not yield any results, presumably due to their basicity. The naphthyllithium reagent produced only the 1,2-addition product 15, indicating that the 1,3 migration may be inhibited due to steric hindrance. Only trace amounts of the product were obtained from the Grignard reagent (entry 6). We investigated the feasibility of this method using heterocyclic aryllithiums, and we obtained only a trace amount of products 20 and 22.

Table 1. Evaluation of Catalytic Conditions* 

| entry | iron species | solvent | temp (°C) | yield (%) | 1 yield (%) | 2 yield (%) |
|-------|--------------|---------|-----------|-----------|-------------|-------------|
| 1a    | FeCl₂        | Et₂O    | 25        | 55        | 4           |
| 2     | FeCl₂        | Et₂O    | 25        | 87        | 17          |
| 3     | Fe(OAc)₂     | Et₂O    | 25        | 35        | 27          |
| 4     | Fe₃         | Et₂O    | 25        | 25        | 40          |
| 5     | Fe(acac)₂    | Et₂O    | 25        | 41        | 44          |
| 6     | Fe(acac)₂    | Et₂O    | 25        | 30        | 29          |
| 7     | FeCl₂        | THF     | 25        | 5         |
| 8     | FeCl₂        | 2-methyl THF | 25 | 4 | 91 |
| 9     | FeCl₂        | MBTE    | 25        | 23        | 32          |
| 10    | FeCl₂        | Et₂O    | −78       | 73        |
| 11b   | FeCl₂        | Et₂O    | 25        | 36        |
| 12b   | FeCl₂        | Et₂O    | 25        | 8         | 83          |
| 13b   | FeCl₂        | Et₂O    | 25        | 71        |

*Standard conditions: 0.2 M in Et₂O (stabilized with 5 ppm BHT), −78 °C to room temperature (rt), 1 equiv of FeCl₂ and 3 equiv of nucleophile. a1 equiv of the aryllithium was used. b1 equiv of BHT was added. d10 mol % of FeCl₂. eSolvent was diethyl ether without BHT.

Table 2. Substrate Scope* 

| En | Substrate | 1,3-Migration | 1,2-Addition |
|----|-----------|---------------|--------------|
| 1a |           |               | 2 97%        |
| 2  |           |               | 3 19%        |
| 3  |           |               | 4 60%        |
| 4  |           |               | 5 52%        |
| 5  |           |               | 6 50%        |

*Conditions: 0.2 M in Et₂O (stabilized with 5 ppm BHT), −78 °C to rt, 1 equiv of FeCl₂ and 3 equiv of nucleophile.

To probe the mechanism of the reaction, the biphenyl lithium reagent was added to 1 and stirred for 3 h. Purification by flash chromatography gave 3. FeCl₂ (1 equiv) was then added to a solution of 3 (1 equiv), and the reaction was stirred at room temperature for 3 h, providing 2 after column chromatography (Scheme 2). This confirmed that the tertiary allylic alcohol (3) was an intermediate for the rearrangement.

Furthermore, the use of a chiral α,β-unsaturated ketone indicated that the OH-migration was diastereoselective. 6-Methylcyclohexenone, 23, was reacted with phenyllithium to produce a 1:1 mixture of diastereomers 24 and 25 (as judged by 1H NMR of the crude reaction mixture). Of the two possible diastereomers, only 24 could be isolated by flash chromatography. When this stereoisomer was reacted with FeCl₂, a 2:1 mixture of the two diastereomeric migration products 26 and 27) was obtained. NOESY NMR spectroscopy revealed that the major diastereomer retained the anti-relationship between the methyl group and the alcohol. This indicates that the 1,3-migration proceed primarily via a syn-facial pathway due to less energy needed for the iron-oxo species to approach from the same face of the allylic cation than to approach from the opposite face as the methyl group to give the syn product (Scheme 3).

Based on these data and the previous work by McCubbin, we propose the mechanism shown in Scheme 4. The organolithium reagent reacts with the α,β-unsaturated ketone to give the tertiary alkoxide (1,2-addition). The FeCl₂ coordinates to the alkoxide (28), and LiCl is formed. The iron-oxo species (29) cleaves the C–O bond, forming an allylic carbocation (30), and the iron-oxo species then attacks the 3-position of the allylic cation, forming a new C–O bond. The major product of this process arises from the migration of the iron-oxo species. We hypothesize that the intimate
pair shown in Scheme 4 (30) could explain the formation of both diastereomers (26 and 27) and the preference for the trans isomer (26). Density functional theory calculations indicate that both the trans (26) and cis (27) rearranged products have similar ground-state energies, so the observed 2:1 selectivity likely arises from kinetic control. When aryllithium nucleophiles are used, the final allylic alcohol is conjugated which we believe is the overall driving force for the reaction.

Finally, the extent of the OH-migration was investigated. Phenyllithium was added to a solution of the conjugated dieneone, 32, then after an hour FeCl2 was added. The mixture was then stirred at room temperature for 24 h, producing 33, a 1,2-addition product, and 34, the product of a 1,5-OH-migration. The 1,3-migration product (35) was not observed, likely because it was less conjugated than 33 or 34. This result corroborates the proposed OH-migration mechanism and confirms that the overall driving force for the reaction is the creation of an extended conjugated system (Scheme 5).

**CONCLUSIONS**

In summary, we have developed a novel iron-mediated process that isomerizes allylic alcohols. The system can be used to effect the transformation of cyclic α,β-unsaturated enones to 3,3′-disubstituted allylic alcohols. Future work in this field could involve the enhancement of the diastereoselectivity of the process and its application to the synthesis of medicinal compounds.

**EXPERIMENTAL SECTION**

All reactions were carried out in an oven-dried glassware under a nitrogen atmosphere, unless stated otherwise. Yields refer to chromatographically and spectroscopically pure compounds unless stated otherwise. 1H NMR and 13C NMR spectra were recorded on Bruker Avance 400 MHz and Bruker Avance 300.
MHZ spectrometers. NMR spectra were measured in dimethyl sulfoxide (DMSO) and CDCl₃ solutions. The chemical shifts δ are reported relative to the residual solvent peaks (δ H, δ DMSO = 2.50 ppm; 13C, δ DMSO = 39.52 ppm; 1H, δ CDCl₃ = 7.26 ppm; 13C, δ CDCl₃ = 77.16 ppm). All 1H and 13C shifts are given in ppm (s = singlet; d = doublet; t = triplet; q = quadruplet; m = multiplet; br = broad signal). High-resolution mass spectrometry was performed using a Thermo Scientific LTQ Orbitrap XL instrument.

**General Procedure (Method A).** n-Butyllithium (0.32 mL, 0.6 mmol, 3 equiv) was slowly added to a solution of aryllaldehyde (1.0 equiv) in anhydrous diethyl ether (0.20 M) precooled to −78 °C. The resulting slightly turbid solution was stirred for 30 min allowing it to warm to ambient temperature. Then, the aryllithium solution was added to the substrate (1 equiv) dissolved in anhydrous diethyl ether in a glovebox droptwise and the reaction mixture was stirred for 3 h, and then iron species (1 equiv) were added to the reaction mixture. The reaction was allowed to run overnight at room temperature. Silica was then added to the reaction. Purification by flash column chromatography using hexanes and ethyl acetate provided the desired 1,3-rearranged allylic alcohol.

**General Procedure (Method B).** α-β-Unsaturated ketone (1.0 equiv) was dissolved in an anhydrous diethyl ether (0.20 M) in a glovebox droptwise. Then, aryllithium solution (3.0 equiv) was added to the substrate solution in a dropwise manner, and the reaction mixture was stirred for 3 h and FeCl₂ (1.0 equiv) was added. The reaction was allowed to run at room temperature overnight. Silica was then added to the reaction. Purification by flash column chromatography using hexane and ethyl acetate provided the corresponding desired 1,3-rearranged allylic alcohols.

**3-Biphenylcyclohex-2-en-1-ol (2).** According to method A, 1.44 mL of n-butyllithium (1.6 M, 0.9 mmol) was slowly added to a solution of 4-bromobiphenyl (210 mg, 0.9 mmol) dissolved in anhydrous diethyl ether (0.20 M) and precooled to −78 °C. After 30 min, the reaction solution was added to 2-cyclohexen-1-one (97%, 29.1 μmol, 0.3 mmol) dissolved in an anhydrous diethyl ether (0.20 M) and precooled to −78 °C. After stirring for 3 h, FeCl₂ (anhydrous, 37.9 mg, 0.3 mmol) was added to the reaction mixture. After 12 h, purification (column chromatography: 17% ethyl acetate in hexane) of reaction mixture gave 216 mg of 3-biphenylcyclohex-2-en-1-ol (2) as a white powder (87% yield).

**Scheme 5. Extent of Migration**
mmol) was slowly added to a solution of 4-bromobiphenyl (1250 mg, 6 mmol) dissolved in an anhydrous diethyl ether (0.20 M) and precooled to −78 °C. 4-Methyl-2-penten-1-one (98%, 294 μL, 2 mmol) and FeCl₃ (anhydrous, 255 mg, 2 mmol) were added. After 12 h, purification (column chromatography: 14% ethyl acetate in hexane) of reaction mixture gave 460 mg (1.5 mmol) of (E)-2-[(1′,1′-biphenyl)-4-yl)-4-phenylbut-3-en-2-ol (7) as a white powder. (77% yield). ³H NMR (400 MHz, DMSO-d₆) δ 7.31 (s, 4H), 6.13 (dt, J = 3.6, 1.8, 1.8 Hz, 1H), 4.29 (bs, 1H), 2.35 (m, 1H), 2.27 (m, 1H), 1.93 (s, 1H), 1.81 (s, 1H), 1.61 (m, 2H). ¹³C{¹H} NMR (101 MHz, chloroform-d) δ 141.40, 140.03, 128.33, 127.43, 126.67, 125.41, 66.34, 31.69, 27.52, 19.53. HRMS (ESI-TOF) m/z: [M + Na]⁺ calcd for C₂₁H₁₇O₃Na+: 323.1412; found: 323.1407.

3-Phenylcyclohex-2-ene-1-ol (9). According to method B, 2-cyclohexen-1-one (97%, 29.1 μL, 0.3 mmol) was dissolved in 2 mL of ether (stabilized with BHT) reacted with phenyllithium (2.1 mL, 1.9 M, 0.2 mmol) and precooled to −78 °C. 4-Bromotoluene (0.7 mL, 6 mmol) was slowly added to a solution of 1-bromonaphthalene (311 mg, 1.5 mmol) dissolved in an anhydrous diethyl ether (0.20 M) and precooled to −78 °C. After 30 min, this solution was added to 2-cyclohexen-1-one (97%, 49.0 μL, 0.5 mmol) and stirred for 3 h, and then FeCl₃ (anhydrous, 63 mg, 0.5 mmol) was added. After 12 h, purification (column chromatography: 14% ethyl acetate in hexane) of the reaction mixture gave 97 mg (0.43 mmol) of (naphthalen-1-yl)cyclohex-2-ene-1-ol (15) as a white powder. (87% yield). ³H NMR (400 MHz, chloroform-d) δ 8.67 (dd, J = 7.4, 2.8 Hz, 1H), 7.89 (m, 1H), 7.80 (m, 2H), 7.47 (m, 3H), 6.11 (ddd, J = 10.0, 3.7, 3.7 Hz, 1H), 5.99 (dt, J = 10.0, 2.3, 2.3 Hz, 1H), 2.49 (ddd, J = 13.7, 10.6, 3.2 Hz, 1H), 2.24 (m, 3H), 2.11 (m, 1H), 1.91 (m, 1H), 1.64 (m, 1H). ¹³C{¹H} NMR (101 MHz, chloroform-d) δ 142.15, 134.80, 133.87, 130.45, 129.97, 128.49, 126.54, 125.20, 125.13, 124.89, 124.73, 73.51, 37.21, 37.20, 31.72, 29.61. HRMS (ESI-TOF) m/z: [M − OH]⁺ calcd for C₁₄H₁₄O: 207.1174, found 207.1179.

6-Methylcyclohex-2-ene-1-ol (23). Dissopropylamine (1.5 equiv) was dissolved in 3.2 mL of THF and placed in a sealed nitrogen-filled vial with a stir bar. The vial was cooled to 0 °C, and n-butyllithium (1.6 M, 10.5 mmol) was added dropwise. After stirring for 20 min, the reaction was then cooled to −78 °C; then, the cyclohexenone (1 equiv) was added dropwise, and the reaction was stirred for 30 min. At the same temperature, 1.3 mL of methyl iodide was added, and the reaction was then stirred for 2 h. The reaction was then warmed to 0 °C before 8 mL of ether was added, and the organic layer was washed with saturated ammonium chloride (5 mL × 3) and saturated sodium chloride (5 mL × 3) and then dried over sodium sulfate. The product 6-methylcyclohex-2-ene-1-one (23, 542 mg) was obtained after purification by flash chromatography (10:1 Hex/EtOAc) as a clear solid. (47% yield). ³H NMR (400 MHz, chloroform-d) δ 6.91 (dd, J = 10.0, 1.9, 1.8 Hz, 1H), 1.37 (m, 3H), 2.04 (ddd, J = 16.7, 13.2, 6.2, 4.9 Hz, 1H), 1.71 (ddd, J = 13.4, 12.0, 8.3, 6.8 Hz, 1H), 1.12 (dd, J = 6.8, 1.1 Hz, 3H). ¹³C{¹H} NMR (101 MHz, chloroform-d) δ 202.35, 149.71, 129.34, 41.60, 30.80, 25.50, 15.04. HRMS (ESI-TOF) m/z: [M + Na]⁺ calcd for C₁₆H₁₄O⁺: 111.0811; [M + H]⁺ found: 111.0804.

2-Methyl-3,4-dihydro-[1′,1′-biphenyl]-1-(2H)-ol (24). According to method B, 0.2 mmol (22 μL) of 23 was dissolved in 1 mL of anhydrous diethyl ether (stabilized with BHT). Phenyllithium (316 μL, 1.9 M, 0.6 mmol) was added dropwise to the reaction mixture and stirred for 3 h at room temperature. Purification (column chromatography: 17% ethyl acetate in hexane) of reaction mixture gave 15 mg of 2-methyl-3,4-dihydro-[1′,1′-biphenyl]-1-(2H)-ol as a clear liquid (37% yield). ³H NMR (400 MHz, chloroform-d) δ 7.45 (m, 2H), 7.35 (m, 2H), 7.26 (m, 1H), 6.00 (m, 1H), 5.76 (dt, J = 9.9, 2.1, 2.1 Hz, 1H), 2.22 (m, 1H), 1.88 (m, 1H), 1.64 (dt, J = 3.9, 2.4, 2.4 Hz, 2H), 0.85 (dd, J = 6.7 Hz, 3H). LRMS (ESI, m/z): [M + H]⁺ calcd for C₁₅H₁₄O: 189.13; [M + H]⁺ found: 189.14.
6-Methyl-3,4,5,6-tetrahydro-[1,1'-biphenyl]-3-ol (26) and (27). In 1 mL of anhydrous diethyl ether, 0.2 mmol of 24 was dissolved, and FeCl$_2$ (25 mg, 0.2 mmol) was then added in the glovebox. The reaction was stirred overnight at room temperature. Purification (column chromatography; 17% ethyl acetate in hexane) of the reaction mixture gave 20 mg of (3R,6R)-6-methyl-3,4,5,6-tetrahydro-[1,1'-biphenyl]-3-ol (26) and (3S,6R)-6-methyl-3,4,5,6-tetrahydro-[1,1'-biphenyl]-3-ol (27) as a clear solid with 52% yield. This diastereomeric mixture was impossible to separate by flash chromatography, but 1H NMR analysis of the crude mixture indicated a 2:1 ratio of the two isomers, 26 and 27 (determined by comparing the integrals of the two peaks corresponding to the methyl groups in each isomer). 1H NMR (400 MHz, chloroform-$d$) $\delta$ 7.24 (m, 5H) (both), 5.82 (dd, $J = 3.7, 1.5$ Hz, 2H) (26), 5.79 (dd, $J = 3.7, 1.5$ Hz 1H) (27), 4.26 (m, 1H) (both), 2.75 (m, 1H) (both), 1.97 (m, 1H) (both), 1.85 (m, 1H) (both), 1.60 (m, 1H) (both), 1.41 (m, 1H) (both), 0.90 (dd, $J = 7.1$ Hz, 3H) (27), 0.83 (dd, $J = 7.1$ Hz, 6H) (26). $^{13}$C{1H} NMR (101 MHz, chloroform-$d$) $\delta$ 146.38, 141.52, 128.24, 127.24, 127.20, 126.53, 66.26, 30.93, 29.03, 19.42. LRMS (ESI-TOF) $m/z$: [M + Na]$^+$: 335.1412; found: 335.1406.

(E)-1,5,5-Triphenylpenta-2,4-dien-1-ol (33) and (2E,4E)-1,1,5-Triphenylpenta-2,4-dien-1-ol (34). Anhydrous diethyl ether (2 mL) was added to cinnamylideneacetophenone (234 mg, 1 mmol), and the solution was cooled to 0°C. Phenyllithium (1.6 mL, 3 mmol, 1.9 M) was then added dropwise, and the reaction mixture was stirred for 1 h before FeCl$_2$ (126 mg, 1 mmol) was added in the glovebox. The reaction was stirred overnight at room temperature. Purification (column chromatography, 17% ethyl acetate in hexane) of reaction mixture isolated two isomers. 1H NMR analysis of the crude mixture indicated a 2:1 ratio of (2) and (3).

Notes

The authors declare no competing financial interest.

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