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# Total syntheses of spiroviolene and spirograterpene A: a structural reassignment with biosynthetic implications†

Hyung Min Chi,  ‡ Charles J. F. Cole,  Pengfei Hu,  Cooper A. Taylor  and Scott A. Snyder  \*

The recent natural product isolates spiroviolene and spirograterpene A are two relatively non-functionalized linear triquinane terpenes with a large number of structural homologies. Nevertheless, three significant areas of structural disparity exist based on their original assignments, one of which implies a key stereochemical divergence early in their respective biosyntheses. Herein, using two known bicyclic ketone intermediates, a core Pd-catalyzed Heck cyclization sequence, and several chemoselective transformations, we describe concise total syntheses of both natural product targets and propose that the structure of spiroviolene should be reassigned. As a result, these natural products possess greater homology than previously anticipated.

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## Introduction

In recent years, chemists have devoted much attention to addressing the synthetic challenges posed by terpenes, such as 1–3 (Scheme 1), which possess a modicum of traditionally reactive functional groups.<sup>1–4</sup> Indeed, efforts with such molecules have suggested that since functional groups are typically required to forge C–C bonds, they should be chosen strategically and used for multiple purposes to limit step counts since they often are target superfluous.<sup>5</sup> In addition, when multiple quaternary centers are present,<sup>6</sup> they should be used as an essential part of retrosynthetic planning, focused on the question of how the presence of one might assist in the formation of others.<sup>2b</sup>

It was against this backdrop that we became interested in the structures of spiroviolene (4)<sup>7</sup> and spirograterpene A (5),<sup>8</sup> two recently disclosed linear triquinanes characterized largely on the basis of NMR data. The first of these was isolated by Dickschat and co-workers from the treatment of a terpene cyclase analog (termed spiroviolene synthase, SvS) with geranylgeranyl diphosphate (GGPP).<sup>7</sup> The other was obtained by the Yang group from *Penicillium granatum* MCCC 3A00475 and was shown to possess modest anti-allergic activity.<sup>8</sup> From a structural perspective, these molecules possess a high degree of structural homology, with both containing the same core

carbon skeleton comprised of 4 rings and 3 all-carbon quaternary centers. However, they also have three main differences: their overall oxidation level, with 5 possessing an additional hydroxyl and carboxylic acid group as opposed to the saturated hydrocarbon units of 4, the specific placement of their central alkene, and the stereochemical disposition of a methyl group at the highlighted position. It was the latter of these differences that especially caught our attention since its accuracy would require a divergence in how those chiral centers were originally set as part of their respective biosyntheses. Indeed, in the Dickschat work<sup>7</sup> it was postulated that this key stereocenter might arise from intermediate 7 via a facially selective 1,3-hydride shift to produce 8; deuterium-labelling studies supported that stereochemical assessment, assuming that all stereocenters were set as drawn in the initial cyclization of linear GGPP leading to this polycycle.<sup>9</sup> Subsequent proton elimination would then afford spiroviolene (4).<sup>10</sup> By contrast, as postulated by Yang,<sup>8</sup> if diastereomer 9 could also be obtained from a cyclization process involving GGPP, then loss of an alternate hydride could afford the distinctive alkene of 10, ultimately leading to spirograterpene A (5) through further oxidations. Herein, we delineate a unified approach capable of achieving the expeditious total synthesis of both natural products. As a result, we also provide a body of evidence implicating that the structure of spiroviolene is in error and has a stereochemical disposition of its highlighted chiral center which matches that of spirograterpene A.

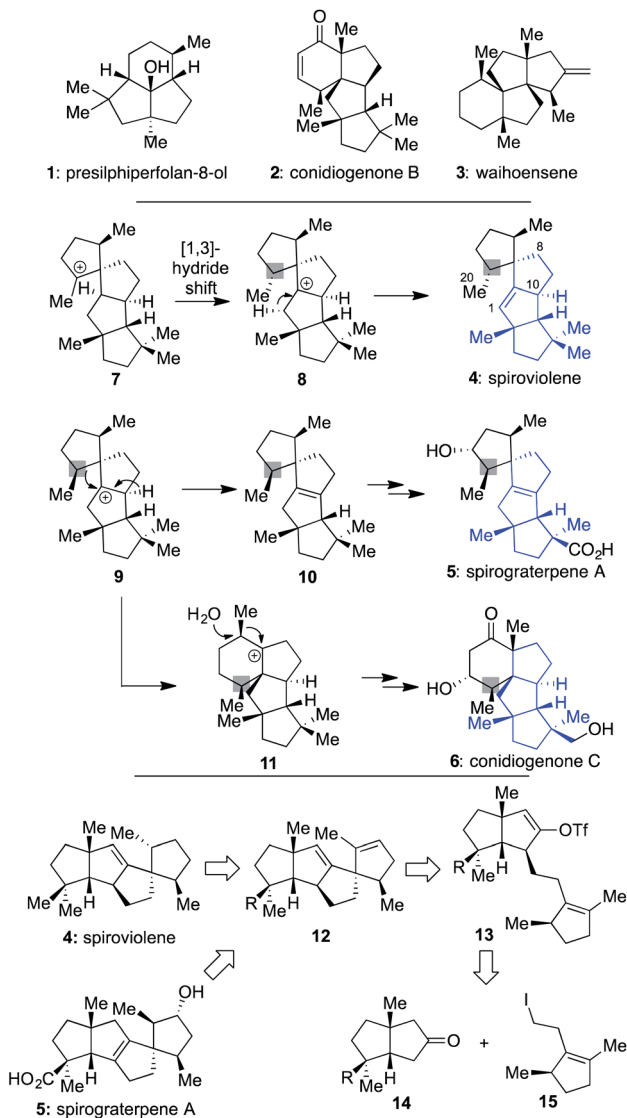
## Results and discussion

Given that we had already synthesized several members of the conidiogenones as single enantiomers and verified their

Department of Chemistry, University of Chicago, 5735 S. Ellis Avenue, Chicago, IL, 60637, USA. E-mail: [sasnyder@uchicago.edu](mailto:sasnyder@uchicago.edu)

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‡ Present address: Department of Chemistry, Pohang University of Science and Technology, South Korea.



**Scheme 1** Unique terpene natural products, the conserved linear triquinane core of spiroviolene (4), spirograterpene A (5), and the conidiogenones (2 and 6), and a proposed synthesis of both 4 and 5 via a common pathway and set of intermediates to clarify the stereochemical identity of the chiral center denoted with a gray box.

absolute configuration in the process, including 2 and 6,<sup>2b</sup> we surmised that the stereochemical assignments for spirograterpene A (5) were likely accurate given that it was co-isolated with 6 and other members of the class. Biosynthetically, what was required to obtain these conidiogenone architectures from 9 was a 1,2-alkyl shift, instead of a  $\beta$ -hydride elimination, to afford reactive intermediate 11. That material, following a further 1,2-alkyl shift and attack by water, could afford most of the structural elements of 6.<sup>8</sup> As such, to determine whether or not 4 and 5 were accurately assigned, we developed the retrosynthetic approach shown in the bottom of Scheme 1; the structures of 4 and 5 have been rotated here from their initial presentation. Our hope was that intermediates of type 12, differing solely in terms of their R group, could afford the

means to access both targets and address any stereochemical ambiguities. In that regard, we assumed that the central alkene within 12 could be isomerized to that of its tetrasubstituted variant in spirograterpene A under thermodynamic control, and that the distal alkene could be chemoselectively functionalized through hydrogenation or hydroboration, respectively, to afford the final patterning of both natural products. We hoped that the latter of these functionalizations could be effected without complete facial control to provide access to both diastereomers to aid in structural assignments. In turn, the key quaternary spirocyclic stereocenter within 12 was anticipated to arise *via* a Heck cyclization<sup>11</sup> from vinyl triflate 13, itself prepared from the regiospecific merger of bicyclic ketone 14 and alkyl iodide 15. These final steps would mirror operations used in our syntheses of both 1 and 2,<sup>1,2b</sup> commencing from an intermediate set (14) already in hand and used as part of our conidiogenone syntheses;<sup>2b</sup> as such, these bicyclic ketones could be envisioned as common intermediates leading to several different classes of natural products if successfully advanced to 4 and 5.

As shown in Scheme 2, our efforts commenced with the preparation of enantioenriched 15. Using a procedure developed by Piva and co-workers,<sup>12</sup> we were able to access  $\gamma$ -hydroxy ester 16 in 3 steps and in  $\sim 90\%$  ee. From here, subsequent regioselective elimination, as promoted by  $\text{SOCl}_2$ , led to  $\beta,\gamma$ -unsaturated ester 17 in near quantitative yield. The desired alkyl iodide was then completed by reduction of the ester to a primary alcohol using  $\text{LiAlH}_4$  followed by an Appel reaction; these operations afforded 15 in 79% overall yield.



**Scheme 2** Total synthesis of spiroviolene (4) starting from known intermediate 18: (a) pyridine (1.2 equiv.),  $\text{CH}_2\text{Cl}_2$ , then  $\text{SOCl}_2$  (1.1 equiv.),  $0^\circ\text{C}$  to  $40^\circ\text{C}$ , 12 h, 98%; (b)  $\text{LiAlH}_4$  (2.0 equiv.),  $\text{Et}_2\text{O}$ ,  $0^\circ\text{C}$ , then 17,  $\text{Et}_2\text{O}$ ,  $0^\circ\text{C}$  to  $23^\circ\text{C}$ , 30 min; (c)  $\text{Ph}_3\text{P}$  (1.2 equiv.), imidazole (1.5 equiv.),  $\text{CH}_2\text{Cl}_2$ ,  $23^\circ\text{C}$ , 5 min, then  $\text{I}_2$  (1.2 equiv.),  $0^\circ\text{C}$  to  $23^\circ\text{C}$ , 1 h, 79% over 2 steps; (d)  $\text{H}_2\text{NNMe}_2$  (5.0 equiv.),  $23^\circ\text{C}$ , 12 h, 96%; (e) LDA (1.3 equiv.), THF,  $-78^\circ\text{C}$ , 15 min, then 18, THF,  $0^\circ\text{C}$ , 2 h, then HMPA (1.3 equiv.),  $-78^\circ\text{C}$ , 10 min, then 15 (1.2 equiv.), THF,  $-78^\circ\text{C}$  to  $23^\circ\text{C}$ , 15 h, then 1 M  $\text{HCl}/\text{THF}$  (1 : 1),  $23^\circ\text{C}$ , 12 h, 65%; (f) KHMDS (1.3 equiv.), THF,  $-78^\circ\text{C}$  to  $0^\circ\text{C}$ , 2 h, then Comins' reagent (1.1 equiv.), THF,  $-78^\circ\text{C}$ , 1 h, 87%; (g)  $\text{Pd}(\text{OAc})_2$  (10 mol%),  $\text{Ph}_3\text{P}$  (20 mol%), toluene,  $23^\circ\text{C}$ , then  $\text{Et}_3\text{N}$  (2.0 equiv.),  $90^\circ\text{C}$ , 20 h, 88%; (h)  $\text{H}_2$  (balloon pressure),  $\text{Pd}/\text{C}$  (10 wt%),  $\text{EtOH}$ ,  $23^\circ\text{C}$ , 30 min,  $>99\%$ .



Next, following the conversion of known bicyclic ketone **18** to its corresponding hydrazone derivative (**19**), a subsequent regiospecific alkylation with **15** was achieved through the action of LDA and HMPA, using an acidic treatment (1 M HCl/THF) to cleave the hydrazone and afford ketone **20** in 65% isolated yield. In this event, the quaternary center uniting the two five-membered rings is presumed to afford sufficient steric bulk to drive its regioselectivity.<sup>2b</sup> From here, treatment with KHMDS and Comins' reagent led to the regiospecific formation of vinyl triflate **21**, setting the stage for a stereospecific Heck cyclization leading to the requisite 1,3-*trans* stereochemical arrangement of substituents in tetracycle **22**.<sup>11</sup> These operations proceeded in 87% overall yield. At this point, all that remained was controlled reduction of one of its two alkenes to reach the putative structure of spiroviolene (*i.e.* **4**). Given that exploration of molecular models suggested standard hydrogenation might afford the undesired diastereomer preferentially, initial attempts focused on hydrogen atom transfer (HAT)-processes in a radical-type manifold.<sup>13</sup> However, all efforts under such a regime failed. As such, an attempt was made under more standard conditions using a H<sub>2</sub> atmosphere with Pd/C in EtOH. To our surprise, this event led to the smooth formation of a single product in near quantitative yield (98%). Its spectral properties (<sup>1</sup>H and <sup>13</sup>C NMR) and optical rotation fully matched those reported by Dickschat and co-workers.<sup>7</sup> Unclear, however, was the identity of the new chiral center within this synthetic material as the face of addition could not be confirmed.



**Scheme 3** Total synthesis of spirograterpene A (**5**) starting from known intermediate **23**: (a) H<sub>2</sub>NNMe<sub>2</sub> (3.0 equiv.), 23 °C, 12 h, 96%; (b) LDA (1.5 equiv.), THF, −78 °C, 15 min, then **23**, THF, 0 °C, 2 h, then HMPA (1.5 equiv.), −78 °C, 30 min, then **15** (1.2 equiv.), THF, −78 °C to 23 °C, 15 h, then 1 M HCl, 23 °C, 12 h, 63%; (c) KHMDS (1.3 equiv.), THF, −78 °C to 0 °C, 2 h, then Comins' reagent (1.1 equiv.), THF, −78 °C, 1 h, 65%; (d) Pd(OAc)<sub>2</sub> (20 mol%), Ph<sub>3</sub>P (40 mol%), Et<sub>3</sub>N (3.0 equiv.), toluene, 23 °C, then **25**, toluene, 23 °C to 90 °C, 16 h, 55%; (e) BH<sub>3</sub>·THF (1.0 equiv.), THF, 0 °C, 5 h, then NaOH/H<sub>2</sub>O<sub>2</sub> (1 : 1), 23 °C, 1 h, 66%; (f) BzCl (3.0 equiv.), Et<sub>3</sub>N (4.0 equiv.), 4-DMAP (1.0 equiv.), CH<sub>2</sub>Cl<sub>2</sub>, 23 °C, 2 h, 96%; (g) 6 M HCl/THF (1 : 2), 50 °C, 4 h, 92%; (h) Dess–Martin periodinane (1.5 equiv.), NaHCO<sub>3</sub> (5.0 equiv.), CH<sub>2</sub>Cl<sub>2</sub>, 23 °C, 1 h, 85%; (i) 2-methyl-2-butene (10 equiv.), *t*-BuOH, then NaClO<sub>2</sub> (35 equiv.), NaH<sub>2</sub>PO<sub>4</sub> (50 equiv.), H<sub>2</sub>O, 23 °C, 1 h, 96%; (j) HCl (4 M in 1,4-dioxane), 80 °C, 48 h, 58%; (k) K<sub>2</sub>CO<sub>3</sub> (3.0 equiv.), MeOH, 55 °C, 15 h, 52%.

As such, we endeavored next to synthesize spirograterpene A (**5**, *cf.* Scheme 1). As indicated in Scheme 3, the initial elements of our route were effected following the same general sequence as for spiroviolene, starting with the MOM-protected **23**.<sup>2b</sup> Alternate modes of protection provided distinct challenges in appending alkyl iodide **15**, but with this group in place, that event, as well as subsequent vinyl triflate formation and Heck cyclization, afforded smooth access to tetracycle **26**. At this juncture, the remaining major operations consisted of olefin isomerization and a series of oxidative manipulations, events that, in principle, could be conducted in either order. In practice, however, we were unable to effect olefin migration to its arguably more stable tetrasubstituted isomer at this stage as a series of undesired rearrangements were observed under a variety of conditions. As such, we exposed **26** to a standard hydroboration/oxidation protocol. This event afforded a single product in 66% yield, drawn here as **27** to match the stereochemistry proposed for spirograterpene A, though again of unknown configuration at this stage (*vide infra*).

With this material in hand, efforts were once again made to effect olefin isomerization, but either no transformation or decomposition was observed. As a result, we protected the secondary alcohol as a benzoate ester and adjusted the oxidation state of the other alcohol motif to that of a carboxylic acid over three additional steps to deliver **29**.<sup>14,15</sup> As shown in Table 1, the alkene within this material proved equally challenging to migrate, with a number of transition metals and photochemical conditions failing to afford any desired product.<sup>16</sup> Under acidic conditions, including treatment with *p*-TsOH or TFOH, we observed the recovery of starting material, even at elevated temperatures. However, when 4 M HCl in 1,4-dioxane was used at 55 °C over the course of 12 h, 15% conversion to the desired product was detected by <sup>1</sup>H NMR analysis based on the presence of a diagnostic allylic proton peak. By increasing the temperature (80 °C) and prolonging the reaction time (72 h), the desired alkene could be obtained in 58% isolated yield.<sup>17</sup> Finally, methanolysis of the benzoate ester with K<sub>2</sub>CO<sub>3</sub> at 55 °C afforded material which matched both the spectral (<sup>1</sup>H and <sup>13</sup>C NMR) and optical rotation data obtained by the Yang group for spirograterpene A (**5**).<sup>8</sup>

With both targets in hand, and both events proceeding with a fully facially selective alkene functionalization (*i.e.* hydrogenation and hydroboration, respectively) on intermediates differing solely by the presence of a remote MOM-protected alcohol in the case of **26** (*cf.* Scheme 3), our conclusion was that the one of the two natural products was misassigned and that the configuration about their highlighted centers as denoted in Scheme 1 is conserved. Unfortunately, despite multiple efforts encompassing a range of synthetic derivatives, we were unable to obtain suitable material for X-ray diffraction to aid in such a reassignment.<sup>18</sup> As such, computational, chemical, and spectroscopic means were explored instead.

First, DFT transition state analysis for the hydroboration of **26** (see ESI†) revealed that in the potential energy surface for the transformation, there is a difference of 8.0 kcal mol<sup>−1</sup> favoring hydroboration from the  $\alpha$ -face, thereby supporting the stereochemical outcome originally proposed for spirograterpene A. In addition, NOE experiments with **27** (Scheme 4) revealed an

Table 1 Conditions screened to effect the critical olefin isomerization

|  |   |                             |
|--|---|-----------------------------|
| Entry  | Conditions  | Result                      |
| 1  | [Pd(CH <sub>3</sub> CN) <sub>4</sub> ](BF <sub>4</sub> ) <sub>2</sub> , CH <sub>3</sub> CN, 23 °C, 12 h | No reaction                 |
| 2  | [Pd(CH <sub>3</sub> CN) <sub>4</sub> ](BF <sub>4</sub> ) <sub>2</sub> , CH <sub>3</sub> CN, 80 °C, 12 h | No reaction                 |
| 3  | UV (<365 nm), Et <sub>2</sub> O, 23 °C, 1 h   | No reaction                 |
| 4  | UV (<365 nm), Et <sub>2</sub> O, 23 °C, 12 h  | Decomposition               |
| 5  | <i>p</i> -TsOH, toluene, 100 °C, 12 h   | No reaction                 |
| 6  | <i>p</i> -TsOH, 1,4-dioxane, 100 °C, 12 h   | No reaction                 |
| 7  | TfOH, 1,4-dioxane, 100 °C, 12 h   | No reaction                 |
| 8  | HCl (4 M in 1,4-dioxane), 23 °C, 12 h   | No reaction                 |
| 9  | HCl (4 M in 1,4-dioxane), 55 °C, 12 h   | <15% conversion             |
| 10   | HCl (4 M in 1,4-dioxane), 80 °C, 72 h   | Full conversion (58% yield) |



**Scheme 4** Synthetic and NMR support for the structural reassignment of spiroviolene (**4**): (a) BH<sub>3</sub>·THF (1.0 equiv.), THF, 0 °C, 5 h, then NaOH/H<sub>2</sub>O<sub>2</sub> (1 : 1), 23 °C, 1 h, 58%; (b) TCDI (3.0 equiv.), 4-DMAP (0.5 equiv.), CH<sub>2</sub>Cl<sub>2</sub>/pyridine (1 : 1), 23 °C, 12 h; (c) *n*-Bu<sub>3</sub>SnH (2.0 equiv.), AIBN (0.15 equiv.), toluene, 110 °C, 10 min, 47%; (d) 6 M HCl/THF (1 : 2), 50 °C, 4 h, >99%; (e) TCDI (4.0 equiv.), 4-DMAP (0.5 equiv.), CH<sub>2</sub>Cl<sub>2</sub>/pyridine (1 : 1), 23 °C, 12 h; (f) (TMS)<sub>3</sub>SiH (4.0 equiv.), AIBN (0.2 equiv.), toluene, 110 °C, 30 min.

interaction between the vinylic proton and the proton geminal to the hydroxyl group, one that would be possible only had the hydroboration taken place from the indicated, and computationally expected, face. These results are in agreement with the Mosher's ester analysis performed by the isolation team, which indicated that the absolute stereochemistry of the secondary alcohol is as drawn.<sup>8</sup>

Further, as shown in Scheme 4, to assess whether the MOM-protecting group of **26** played any role in leading to a differential facial preference for functionalization, alkene **22** was subjected

to the same hydroboration/oxidation sequence to afford **31**. Subsequent treatment with 1,1'-thiocarbonyldiimidazole (TCDI) and subjection of the resultant crude thiocarbamate intermediate to typical Barton–McCombie deoxygenation conditions<sup>19</sup> afforded a sample with spectral properties that matched that of spiroviolene. Finally, intermediate **27**, following conversion to diol **32** and a similar double Barton–McCombie deoxygenation, also afforded material with spectral properties that matched spiroviolene.<sup>20</sup>

Finally, we also considered the collated body of spectral data for natural spiroviolene collected by the Dickschat group. In particular, we examined key NOE correlations that served as a core element for the initial structural assignment. Specifically, the observed interaction between H-10 and C-20 (see Scheme 1 for positional numbering) as well as the interaction between H-8 $\alpha$  and C-20, would appear to support both the original and revised structures. However, there is an additional interaction, albeit relatively weak, between the H-1 vinyl proton and C-20, one that could only be possible with the relative stereochemistry displayed in the revised structure.

As such, based on this collated evidence, we believe that the structure of spiroviolene should be reassigned to that of structure **4** as drawn within Scheme 4, with stereochemistry at the key center matching that of the originally assigned spirograterpene A. This result would thus suggest that **9** (*cf.* Scheme 1), not **8**, is a likely biosynthetic precursor for **4**, with implications for the stereochemical outcome of cyclizations and/or other events starting from GGPP to account for its formation. At what point in that biosynthesis this alteration would occur is unclear, and it may be distinct for the route used to reach **9** for spirograterpene A given that these natural products are produced by bacteria and fungi, respectively.

## Conclusions

In conclusion, we have achieved concise and enantioselective syntheses of both spiroviolene and spirograterpene A, in **10** and





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- 14 Attempts were made to selectively oxidize the primary alcohol in the presence of the secondary alcohol using some of the methods cited here, but ultimately proved unsuccessful. See: (a) P. L. Anelli, C. B. F. Montanari and S. Quici, *J. Org. Chem.*, 1987, **52**, 2559; (b) J. B. Epp and T. S. Widlanski, *J. Org. Chem.*, 1999, **64**, 293; (c) M. Zhao, J. Li, E. Mano, Z. J. Song, D. M. Tschaen, E. J. J. Grabowski and P. J. Reider, *J. Org. Chem.*, 1999, **64**, 2564; (d) M. Zhao, J. Li, E. Mano, Z. J. Song and D. M. Tschaen, *Org. Synth.*, 2004, **81**, 195; (e) M. Shibuya, T. Sato, M. Tmoizawa and Y. Iwabuchi, *Chem. Commun.*, 2009, 1739.
- 15 The most challenging of these operations was the initial MOM cleavage, as significant decomposition when observed when this operation was conducted on scale in the presence of trace water (such as from *p*-TsOH·H<sub>2</sub>O); only 6 M HCl in THF afforded consistent results.
- 16 For examples of metal-mediated olefin isomerization, see: (a) P. A. Greico, M. Nishizawa, N. Marinovic and W. J. Ehmann, *J. Am. Chem. Soc.*, 1976, **98**, 7102; (b) J. Andrieux, D. H. R. Barton and P. Henri, *J. Chem. Soc., Perkin Trans. 1*, 1977, **4**, 359; (c) L. A. Paquette, W. E. Fristad, D. S. Dime and T. R. Bailey, *J. Org. Chem.*, 1980, **45**, 3017; (d) A. Sen and T.-W. Lai, *Inorg. Chem.*, 1984, **23**, 3257; (e) A. Sen, T.-W. Lai and R. T. Thomas, *J. Organomet. Chem.*, 1988, **358**, 567.
- 17 This step required strictly anhydrous conditions to avoid hydrolysis of the benzoate, as not only was the resultant free alcohol prone to decomposition, but the resulting benzoic acid by-product proved extremely difficult to remove, often requiring several rounds of preparative TLC.
- 18 To date, several unsuccessful attempts have been made to obtain a crystal of suitable quality under a variety of solvent combinations and crystallization techniques. The compounds used toward this end include **5**, **S6** (and the tosyl hydrazone derivative thereof), the bis(thionoimidazole) derivative of **32**, and **S5** (as well as the *p*-bromobenzoate and ferrocenecarboxylate analogues).
- 19 For the original report of the Barton-McCombie deoxygenation, see: (a) D. H. R. Barton and S. W. McCombie, *J. Chem. Soc., Perkin Trans. 1*, 1975, **16**, 1574. For an example of a similar application, see: (b) W.-C. Liu and C.-C. Liao, *Chem. Commun.*, 1999, 117.
- 20 In these latter experiments, stereochemical scrambling at the key reassigned center could only have occurred if there was a 1,2-hydrogen shift to form a tertiary radical followed by H<sup>•</sup> capture from the more hindered face (*i.e.* the opposite face of addition observed from hydrogenation and hydroboration of an alkene at that position). We do not believe such a sequence to be likely.

