

# A Diastereoselective Formal Synthesis of Berkelic Acid

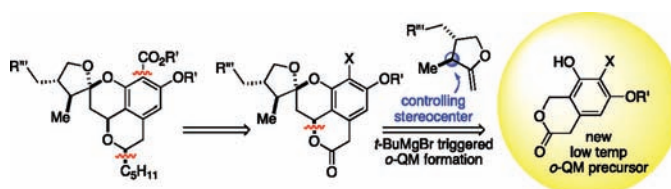
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## ABSTRACT



A formal synthesis of berkelic acid is reported. The strategy employs the combination of a chiral exocyclic enol ether and an achiral isochromanone to afford the chroman spiroketal core via a base-triggered generation and cycloaddition of an *o*-quinone methide intermediate. Other key steps include equilibration of the spiroketal, intramolecular benzylic oxidation, and lactone addition/hemiketal reduction; all occur with good diastereoselectivity.

The natural product berkelic acid, isolated by Stierle et al. in 2006, is a secondary metabolite produced by a species of *Penicillium* that thrives in Berkely Pit Lake outside of Butte, Montana.<sup>1</sup> The toxic lake, which is an abandoned copper mine that has since filled with over 40 billion gallons of contaminated groundwater, was designated in 1994 as an EPA superfund site.<sup>2</sup> Given the proximity of other extreme environments, such as hot springs of Yellowstone National Park some hundred miles to the south, it is doubtful that the extremophile evolved in the waste pit. However, researchers have speculated that some natural products produced in extreme environments serve to protect their progenitor. Berkelic acid, in particular, emerged from an assay-driven study aimed at identifying metal dependent enzyme inhibitors. The compound was reported to inhibit the cysteine protease caspase-1 (98  $\mu\text{M}$ ) and matrix metalloproteinase-3 (MMP-3, 1.87  $\mu\text{M}$ ). In further studies with the NCI-60 cancer cell panel, potent and selective cytotoxicity was observed toward a particularly aggressive ovarian cancer cell line (OVCAR-3, 91 nM).

Originally assigned as compound **1**, berkelic acid belongs to a very select group of chroman spiroketal natural products of considerable synthetic interest and biological importance (Scheme 1).<sup>3</sup> The unknown relative stereochemistry at the quaternary carbon, however, gave us pause. We first envisioned a diastereoselective plan that might address this question toward the end of the synthesis. Late stage introduction of the aryl acid utilizing an aryl-X substituent of compound **2**, in principle, made this functionality more distinguishable from the methyl ester. Furthermore, researchers had theorized that MMP inhibitors less dependent upon metal chelation might prove more selective toward validated isoforms of the enzyme; this would allow for analog development along the synthetic pathway.<sup>4</sup> We further imagined construction of the isobenzopyran ring by utilization of a stereocontrolled benzylic oxidation of the benzopyran with concomitant reduction of the ketone functionality within the R'' residue of compound **3**. To construct the chroman spiroketal motif, we sought to employ chiral exocyclic enol ethers, such as **5**, in a diastereoselective

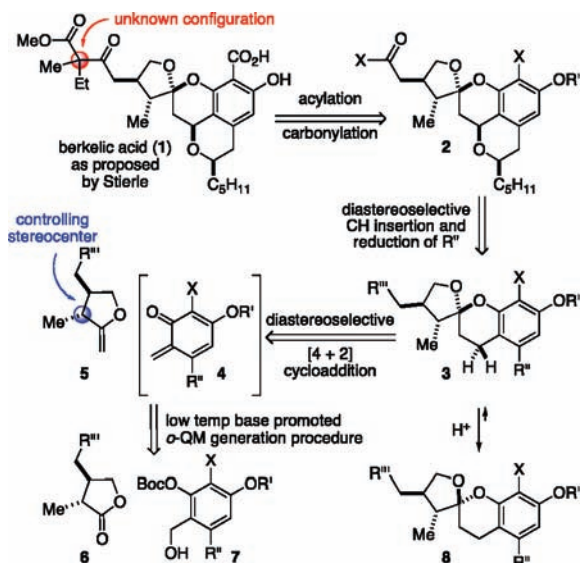
(1) Stierle, A. A.; Stierle, D. B.; Kelly, K. J. *Org. Chem.* **2006**, *71*, 5357.

(2) EPA ID: MTD980502777.

(3) Sperry, J.; Wilson, Z. E.; Rathwell, D. C. K.; Brimble, M. A. *Nat. Prod. Rep.* **2010**, *27*, 1117.

(4) (a) Itoh, Y.; Nagase, H. *Essays Biochem.* **2002**, *38*, 21. (b) Zucker, S.; Cao, J.; Chen, W. T. *Oncogene* **2000**, *19*, 6642.

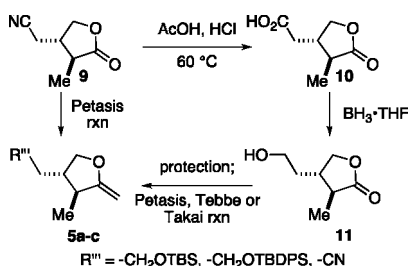
### Scheme 1. An Early Retrosynthetic Plan



inverse demand Diels–Alder reaction with an *o*-quinone methide (*o*-QM) **4** generated at low temperature using our magnesium base-triggering methods in conjunction with various *o*-OBoc compounds **7**.<sup>5</sup>

Hence, we began by devising a short synthesis of exocyclic enol ethers (**5a–c**) derived from  $\gamma$ -lactone **9**, which could be produced later in the desired enantiomeric form (Scheme 2).<sup>6</sup> These enol ethers were then evaluated in conjunction

### Scheme 2. Construction of Enol Ether Coupling Partners



with assorted *o*-QM precursors **7** (R'' = -H; X = -H, -Br; R' = -Me, -Bn) using our standard magnesium base-triggering methodology.

Our early experiments showed that the low temperature cycloaddition afforded what we believed to be the desired relative stereochemistry, whereby the methyl residue in the enol ether had controlled the spiroketal stereocenter (>4:1) in compound **3**.<sup>7</sup> We further observed and reported that this

kinetic spiroketal epimer could be equilibrated to the corresponding thermodynamic epimer spiroketal **8** (>1:2) with a protic acid.<sup>8</sup> A short time thereafter, Fürstner reported the synthesis of berkeleyic acid methyl ester **12** by coupling three separate chiral components together. Their observations led to the revision of three stereocenters and an assignment of the relative stereochemistry including the quaternary stereocenter (Figure 1).<sup>9</sup> This landmark effort paved the way

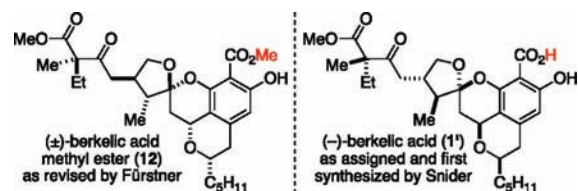


Figure 1. Subsequent revisions by Fürstner and Snider.

for future synthetic successes, and soon afterward, Snider reported the first total synthesis establishing the absolute structure of berkeleyic acid as its enantiomer (–)-**1**.<sup>10</sup> The Snider synthesis, which involved an acid-catalyzed coupling of two chiral components followed by late-stage chromatographic resolution of the quaternary center, is masterful in its overall efficiency. Sometime later, De Brabander reported a biomimetic total synthesis combining spiciferin and pulvilloric acid surrogates.<sup>11</sup>

These successes and structural revisions forced us to reevaluate our strategy with the prime intention of preserving our initial diastereoselective approach. While we remained confident that enol ethers **5a–c** would react to form chroman spiroketals,<sup>12</sup> we needed to redesign our *o*-QM precursor so that a new substituent could be elaborated into the desired tetracycle. We postulated that compound **17** might prove to be a novel *o*-QM precursor participating in our base-triggering procedure and thereby lead to the *o*-QM carboxylate **16** and the spiroketal **15** (Scheme 3). Unfortunately, the cycloaddition was expected to afford the kinetic diastereomer. Although our prior equilibration studies were encouraging, the final (2:1) ratio would be a challenge to improve. However, we hoped that the added carboxylic acid residue and aryl iodide might bolster the ratio afforded by a thermodynamic equilibration. While the specifics of the sequence needed for isobenzopyran formation remained

(7) Marsini, M. A.; Huang, Y.; Lindsey, C. C.; Wu, K.-L.; Pettus, T. R. R. *Org. Lett.* **2008**, *10*, 1477.

(8) Huang, Y.; Pettus, T. R. R. *Synlett* **2008**, 1353.

(9) (a) Buchgraber, P.; Snaddon, T. N.; Wirtz, C.; Mynott, R.; Goddard, R.; Fürstner, A. *Angew. Chem., Int. Ed.* **2008**, *47*, 8450. (b) Snaddon, T. N.; Buchgraber, P.; Schulthoff, S.; Wirtz, C.; Mynott, R.; Fürstner, A. *Chem.–Eur. J.* **2010**, *16*, 12133.

(10) (a) Wu, X.; Zhou, J.; Snider, B. B. *Angew. Chem., Int. Ed.* **2009**, *48*, 1283. (b) Wu, X.; Zhou, J.; Snider, B. B. *J. Org. Chem.* **2009**, *74*, 6245.

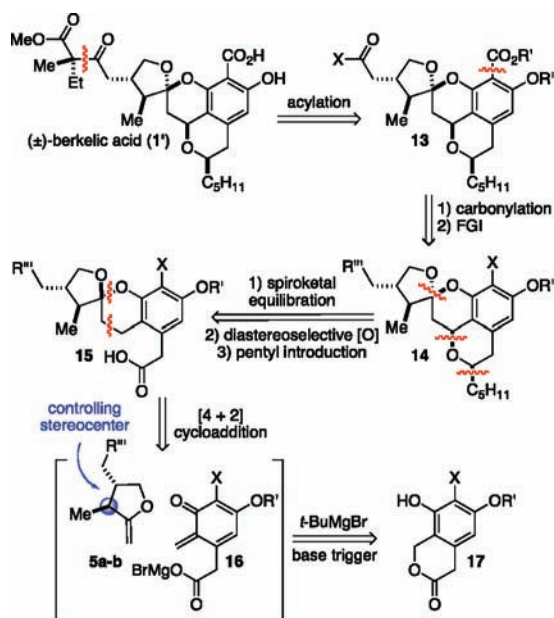
(11) Bender, C. F.; Yoshimoto, F. K.; Paradise, C. L.; De Brabander, J. K. *J. Am. Chem. Soc.* **2009**, *131*, 11350.

(12) Compound **5b** was made by Snider and coworkers in enantiopure form by 1,4-addition of metallated chiral phosphoramides: see ref 10a and Hanessian, S.; Gomtsyan, A.; Malek, N. *J. Org. Chem.* **2000**, *65*, 5623.

(5) (a) Selenski, C.; Pettus, T. R. R. *J. Org. Chem.* **2004**, *69*, 9196. (b) Jones, R. M.; Van de Water, R. W.; Lindsey, C. C.; Pettus, T. R. R. *J. Org. Chem.* **2001**, *66*, 3435.

(6) See Supporting Information and Pirrung, M. C.; Dunlap, S. E.; Trinks, U. P. *Helv. Chim. Acta* **1989**, *72*, 1301.

Scheme 3. Revised Diastereoselective Strategy

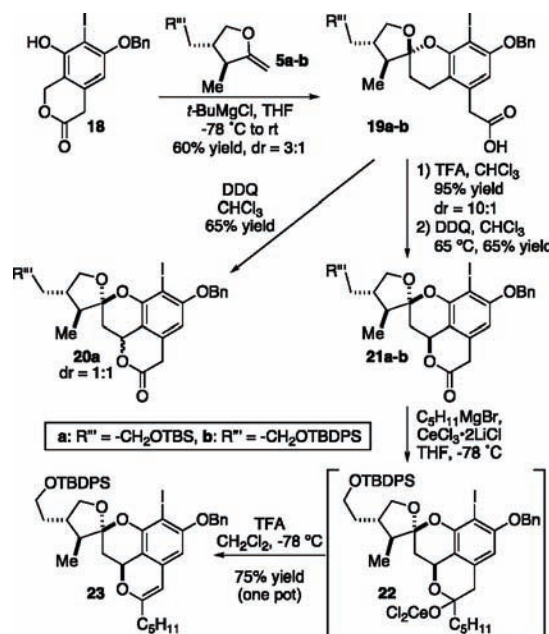


undetermined, it mandated stereoselective benzylic oxidation, etherification, and stereoselective incorporation of the pentyl residue to produce the fully elaborated tetracycle **13**. Acylation could then be used to introduce the quaternary center and methyl ester. The exact timing for carboxylation, however, depended on the concluding events and surrounding functionality.

Proceeding in the forward direction, *tert*-butyl magnesium chloride (1.05 equiv) is added to the lactone **18** (0.1 M in THF,  $-78\text{ }^{\circ}\text{C}$ ), which we had previously prepared in seven steps from commercial materials,<sup>13</sup> followed by the addition of the enol ether **5a** or **5b** (neat, 2.1 equiv). Subsequent warming of the mixture to room temperature affords the respective spiroketals **19a–b** in 60–65% yield with a 3:1 ratio of spiroketal epimers favoring the endo isomer (Scheme 4). Treatment of the mixture **19a** (0.01 M in  $\text{CHCl}_3$ ,  $65\text{ }^{\circ}\text{C}$ ) with DDQ afforded the spiroketal **20a** of opposite configuration with a 1:1 ratio at the oxidized benzylic carbon atom. This result indicated to us that, under the reaction conditions, equilibration of the spiroketal was occurring more slowly than benzylic oxidation. To circumvent this issue, equilibration was carried out before exposure to the oxidant. Relative energy calculations for all of the possible diastereomers of spiroketal **19a** indicated that the desired spiroketal epimer **19b** is more stable by greater than 4.5 kcal/mol.<sup>14</sup> Gratifyingly, treatment of the 3:1 mixture of **19a** or **19b** (0.06 M in  $\text{CHCl}_3$ , rt) with TFA (0.3% in  $\text{CHCl}_3$ ) confirms these calculations and results in a nearly quantitative yield of the corresponding thermodynamic spiroketals in a 10:1 diastereomeric ratio. Separation and intramolecular benzylic oxidation using the prior conditions produced the respective lactones **21a–b** as single diastereomers.

(13) See the Supporting Information for experimental details.

(14) Calculations were performed using MacroModel MM3 Monte Carlo conformational analysis.

Scheme 4. Diastereoselective Construction of Enol Ether **23**

Our next hurdle involved selective *mono* addition of the desired pentyl alkyl chain to the lactone scaffold **21**. The methylene adjoining the lactone carbonyl in compound **21** was problematic—addition of most organometallic species resulted in deprotonation and returned starting material upon workup. Weinreb amides derived from **21** were also investigated in the hope that an acyclic system would prove less acidic and undergo mono addition to afford the corresponding ketone. However, we eventually uncovered observations by Cohen et al.<sup>15</sup> that describe the selective mono addition of organocerium species, prepared by the normal Imamoto process, to lactones.<sup>16</sup> After confirming the integrity of the desired pentylcerium reagent (generated by the method of Knochel)<sup>17</sup> by testing its reaction with 1,3-diphenylacetone, we were pleased to observe its selective mono addition to lactone **21** produces the corresponding hemiketal **22** in greater than 80% yield. The hemiketal **22**, however, exists in equilibrium with its ketone counterpart, which thwarted its complete spectroscopic identification. We therefore sought to reduce it in situ. Conventional reduction tactics, such as the combined action of triethylsilane and protic or Lewis acids, afforded a mixture of isobenzopyran diastereomers favoring the undesired stereochemistry. In the absence of a reductant, however, treatment of the cerium alkoxy hemiketal **22** (0.02 M in  $\text{CH}_2\text{Cl}_2$ ,  $-78\text{ }^{\circ}\text{C}$ ) in situ with TFA resulted in smooth formation of the corresponding enol ether **23** in 75% yield over the one-pot operation from lactone **21**.

We had planned to conclude our synthesis with carbonylation, and compound **23** enabled us to investigate a variety

(15) (a) Ahn, Y.; Cohen, T. *J. Org. Chem.* **1994**, *59*, 3142. (b) Mudryk, B.; Shook, C. A.; Cohen, T. *J. Am. Chem. Soc.* **1990**, *112*, 6389.

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of plausible, mild conditions that might prove compatible with the  $\beta$ -keto ester functionality. Unfortunately, all palladium- and zinc-mediated coupling processes failed in our hands. In addition, we found that all homogeneous and heterogeneous hydrogenations of the enol ether in **23** caused hydrogenolysis of the aryl iodo functionality. These problems forced us to proceed with carbonylation before reduction of the enol ether, much earlier than originally planned. Lithium-halogen exchange proceeds with iodide **23** (0.01 M in THF,  $-110\text{ }^\circ\text{C}$ ) upon addition of *t*-BuLi (2 equiv), whereupon cannulation onto carbon dioxide affords the desired acid upon acidic workup (Scheme 5). Subsequent hydrogenation of the

Snider's, thereby confirming all of the stereochemical assignments and resulting in a formal synthesis of berkelic acid with the synthesis of the acid **24**.

Nevertheless, we sought to shorten the potential endgame for our route by one additional step. Thus, the iodo compound **23** was subjected to lithium-halogen exchange and addition of carbon dioxide as before. In this instance, however, the carboxylate anion was intercepted with benzyl bromide to produce the corresponding benzyl ester in 84% yield. We further discovered that this material (0.025 M in toluene) could be selectively reduced with rhodium on carbon at 65 psi of hydrogen pressure with the benzyl ether and ester remaining intact, thereby yielding compound **26** in 87% yield. In view of the newly reported synthesis of (–)-**1'** by Fürstner in which a benzyl ester is used for orthogonal deprotection in the presence of a methyl ester, the benzylated material **26** should prove as viable in the Snider endgame as ester **25**.

In conclusion, we have successfully accomplished a diastereoselective formal synthesis of berkelic acid by interception of Snider's previously uncharacterized salicylic acid **24**. We believe that our strategy has merit since our tactical construction of the isobenzopyran ring is orthogonal to previous syntheses; all the prior strategies utilize the combination of various chiral components together with various chiral isobenzopyran derivatives to arrive at the tetracycle. In our case, the relative stereochemistry of the tetracyclic core is efficiently relayed from the stereocenters derived from the original enol ether *o*-QM coupling partner. Furthermore, the flexibility of this route allows for the introduction of various scaffold modifications of the tetracyclic core (acid removal, spiroketal epimerization, alkyl derivatization) that may shed light on the mechanism of action of this interesting natural product.<sup>19</sup>

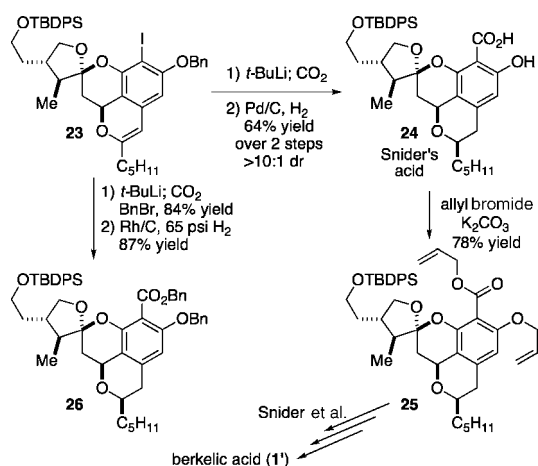
**Acknowledgment.** This work was funded with support from the National Institute of General Medical Sciences (GM064831).

**Supporting Information Available:** Experimental details for the preparation of new compounds and complete spectral data for new compounds accessible and stable as single entities (**5a–c**, **18**, **19a–b**, **21a–b**, **22–26**). This material is available free of charge via the Internet at <http://pubs.acs.org>.

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(19) Fischbach, M. A.; Walsh, C. T. *Science* **2009**, 325, 1089.

**Scheme 5.** Formal Synthesis of (±)-Berkelic Acid (**1'**)



crude acid (0.01 M in EtOAc) over palladium on carbon (10%, 1 atm H<sub>2</sub>) occurs from the less hindered face to afford salicylic acid **24** in a 64% yield over two steps with >10:1 dr.<sup>18</sup>

The salicylic acid **24** had been a key intermediate in the Snider synthesis, and it is positioned six steps from the natural product. However, this acid had not been fully characterized. We therefore constructed the published allyl derivative **25** from acid **24**. Spectroscopic comparison of the <sup>1</sup>H NMR showed our synthetic material to be the same as

(18) Reduction of hemiacetal analogs is reported to stereoselectively provide the corresponding 2,6-*cis*-tetrahydropyran. See: Nicolaou, K. C.; Cole, K. P.; Frederick, M. O.; Aversa, R. J.; Denton, R. M. *Angew. Chem., Int. Ed.* **2007**, 119, 9031.