



Published in final edited form as:

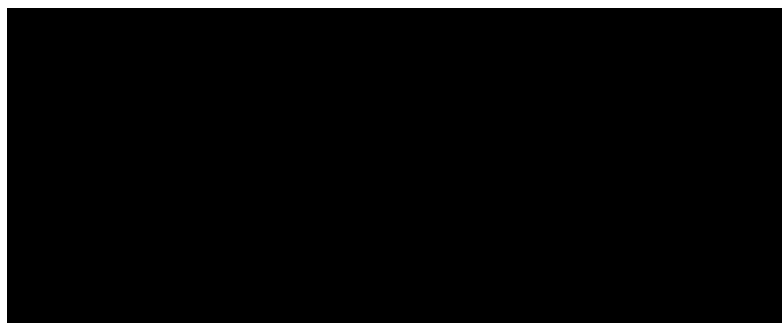
*Org Lett.* 2002 September 5; 4(18): 3131–3134.

## Directed Reduction of $\beta$ -Amino Ketones to Syn or Anti 1,3-Amino Alcohol Derivatives

Gary E. Keck and Anh P. Truong

Department of Chemistry, University of Utah, 315 South 1400 East RM 2020, Salt Lake City, Utah 84112-0850 keck@chemistry.utah.edu

### Abstract



The reduction of  $\beta$ -amino ketones with samarium(II) iodide has been investigated. Either the syn or anti 1,3-amino alcohols can be obtained as the major product due to a divergence in selectivity with different N-protecting groups.

The development of methodology for the efficient generation of relative stereochemistry in acyclic systems continues to be a subject of major importance in synthetic organic chemistry. Recently, a protocol for the reduction of  $\beta$ -hydroxy and certain  $\beta$ -alkoxy ketones with  $\text{SmI}_2$  was developed in our laboratories. Anti 1,3-diols and certain anti 1,3-alkoxy alcohols can be produced with high selectivity in this electron-transfer reduction.<sup>1</sup> The product-determining event was suggested to be protonation of an intermediate samarium carbanion in a favored conformation of a chelate formed by complexation of a second samarium between the  $\beta$ -oxygen and the carbonyl oxygen. The  $\beta$ -oxygen was also shown to be responsible for a significant rate enhancement in these reductions. We record herein the results of an investigation of  $\beta$ -nitrogen substituents in such reductions. In this context, it should be noted that 1,3-amino alcohol fragments are of interest due to their presence in several antibiotics and other biologically active natural products.<sup>2</sup> Thus, several reduction methods for accessing such structures have previously been developed.<sup>3</sup> For example, Narasaka achieved good syn diastereoselectivity in the  $\text{LiAlH}_4$ - $\text{NaOMe}$  reduction of *O*-benzyl oximes derived from  $\beta$ -hydroxy ketones.<sup>4</sup> Pilli and co-workers have also developed a stereoselective route to anti and syn *N*-aryl- $\beta$ -amino alcohols via reduction with  $\text{Et}_3\text{BHLi}$  and  $\text{Zn}(\text{BH}_4)_2$ , respectively.<sup>5</sup>

The first set of  $\beta$ -amino ketones used in this study was prepared by the sequence shown in Scheme 1. Allylation of cyclohexylcarboxaldehyde with methallyl-tri-*n*-butylstannane gave the corresponding homoallylic alcohol.<sup>6</sup> Conversion to the mesylate and displacement with sodium azide gave the desired homoallylic azide **1** in 57% overall yield. Reduction of azide

**1** was accomplished with LiAlH<sub>4</sub> in THF at 0 °C. The resulting amine was then protected as either the trifluoroacetamide, acetamide, *tert*-butyl carbamate, or benzyl carbamate using standard protocols. Subsequent ozonolysis provided the corresponding *N*-protected amino ketones in good yields (Scheme 1).

Each of these *N*-acyl- $\beta$ -amino ketones (**2a–d**) was first reduced with NaBH<sub>4</sub> to afford a mixture containing roughly equal amounts of the 1,3-*syn* and 1,3-*anti* diastereomers. These mixtures were then used to develop an analytical HPLC separation of the reduction products expected from each substrate. Reductions with SmI<sub>2</sub> were then investigated. In a typical reaction, the ketone and 20 equiv of MeOH were dissolved in THF and added to a freshly prepared solution (0.1–0.5 M) of SmI<sub>2</sub> in THF at 0 °C.

The results of these experiments (Table 1) showed that high yields and reasonable levels of stereoselectivity are obtained for the reduction of these protected  $\beta$ -amino ketones. However, at this point the stereochemistry of the products was not known.

The structure of the major diastereomer formed in these reactions was unambiguously established by an X-ray crystal structure determination of the major product obtained from **2a**. This was followed by chemical correlation of that material with the major products obtained from **2b–d**. Thus, removal of the trifluoroacetate<sup>7</sup> (K<sub>2</sub>CO<sub>3</sub>/MeOH/H<sub>2</sub>O) from both the major (*syn*) and minor (*anti*) diastereomers of **3a** provided samples of the free amino alcohols.

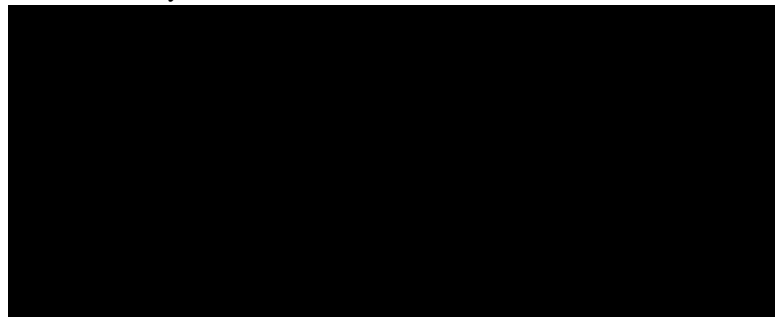
These were then compared with the amino alcohols obtained by removal of the protecting groups present in the major diastereomers of **3b–d**. In all cases, the major diastereomer was 1,3-*syn*. The observed stereochemical outcome is initially quite surprising given the *anti* selectivity previously observed in reductions of both  $\beta$ -hydroxy and  $\beta$ -alkoxy ketones under these conditions. A different class of protected  $\beta$ -amino ketones was also synthesized using the methodology of Kobayashi.<sup>8</sup> This reaction was applied successfully to the synthesis of a number of *N*-aryl- $\beta$ -amino ketones (Scheme 2). Again, each of these substrates (**4a–f**) was reduced with NaBH<sub>4</sub> to secure mixtures containing comparable amounts of the expected 1,3-*syn* and 1,3-*anti* reduction products; these were then used to develop analytical HPLC separations for each substrate as before.

Under our standard conditions, these *N*-aryl- $\beta$ -amino ketones were also reduced to the corresponding 1,3-amino alcohols in excellent yield (90–99%). Stereochemical assignments for the products of these reactions were again made unambiguously through use of the X-ray crystal structure determination of **3a**. This involved the removal of the *o*-MeOPh (catalytic amount of silver nitrate and ammonium persulfate in aqueous CH<sub>3</sub>CN)<sup>9</sup> or *p*-MeOPh (CAN in aqueous CH<sub>3</sub>CN)<sup>10</sup> groups from each of the reduction products **5a–e**.<sup>11</sup> The resulting free  $\beta$ -amino alcohols were then directly compared (<sup>13</sup>C NMR) to the free  $\beta$ -amino alcohols that were obtained from the major (*syn*) and minor (*anti*) diastereomers of **3a**. *It was found that a complete reversal of stereoselectivity for the reduction occurs between the two classes of substrates: whereas the N-acyl derivatives all reduced to afford the 1,3-syn diastereomer as the major product, the N-aryl derivatives all afforded the 1,3-anti diastereomer as the major product.*

Thus, these *N*-aryl  $\beta$ -amino ketones are reduced in the same sense (1,3-*anti* diastereoselectivity) as the previously studied  $\beta$ -hydroxy and  $\beta$ -alkoxy ketones. We suggest that the overall mechanism is as previously proposed for those cases. However, the present situation seems more complex than with the oxygen series. Reductions of two *N*-alkyl- $\beta$ -amino ketones were examined in order to determine the extent to which these reductions paralleled those of the  $\beta$ -alkoxy ketones.

In contrast to the corresponding *O*-benzyl derivative, the *N*-benzyl derivative was reduced but with a low level of stereoselectivity (3.5:1). The *N*-methyl derivative also reduced, but, in contrast to previous observations in the oxygen series, afforded a 1:1 mixture of diastereomers. Reductions of such  $\beta$ -methoxy ketones in our previous study showed very high levels of stereoselectivity; note eq 2. Although the substituents differ in the cases shown (cyclohexyl vs phenyl) in eqs 1 and 2, inspection of Table 2 shows that the level of asymmetric induction realized is relatively independent of this substituent. Cases in which this substituent was a phenyl group were not examined in the present study due to the low stability of these materials toward  $\beta$ -elimination.

Thus *N*-substitution by an electron-rich aryl ring seems to be required to realize high levels of stereoselectivity in



these reductions. In this context, reduction of substrate **4f**, which lacks any additional electron-releasing substituents on the aryl ring other than nitrogen, leads to a somewhat lower level of stereoselectivity than is observed with the methoxy-substituted compounds. Moreover, since both the ortho and para methoxy compounds are reduced with similarly high levels of diastereoselectivity, a chelation motif involving the nitrogen and the methoxy group itself cannot be involved as a control element in determining the stereochemical outcome.<sup>8,12</sup> However, it does seem plausible based upon the available data that complexation of the electron-rich aryl ring by samarium may influence the stereochemical outcome of the reduction in these cases.

At first glance, the *syn* stereochemistry observed in reduction of the *N*-acyl derivatives would seem to preclude a mechanism involving chelation as a control element; however, these results are easily accommodated by such a mechanism. Since samarium(II) is believed to be a hard oxophilic Lewis acid,<sup>13</sup> it is reasonable to expect that the oxygen of the amidate, rather than the nitrogen, is involved in initial coordination to samarium. The reduction could then occur via the formation of an intermediate eight-membered ring chelated structure as shown in Scheme 3. Initial coordination of samarium to the oxygen of the amidate is followed by a one-electron transfer to the carbonyl to give **III**. A second equivalent of Sm(II) donates another electron to the resulting radical leading to the samarium carbanion **IV**. Finally, protonation of this anion then leads to the *syn* amino alcohol **V** in the stereochemistry-determining step.

The *syn* stereochemical outcome predicted by this scheme relies upon extrapolation of the known conformational preferences in eight-membered carbocyclic rings<sup>14</sup> to this case involving samarium. The *A* value associated with the substituent "R" in structure **IV** is known to be very large in eight-membered ring carbocycles.

In summary, the reduction of  $\beta$ -amino ketones using samarium(II) iodide provides a versatile synthetic method for the synthesis of 1,3-amino alcohols. High levels of stereoselectivity and excellent yields are possible in the reduction of *N*-aryl  $\beta$ -amino ketones, while a variety of *N*-acyl derivatives are reduced in good yield and with moderate levels of 1,3-*syn* stereoselectivity.

One especially attractive feature of the process is that by employing different protecting groups, one can easily access both syn and anti diastereomers of these  $\beta$ -amino alcohols.

## Supplementary Material

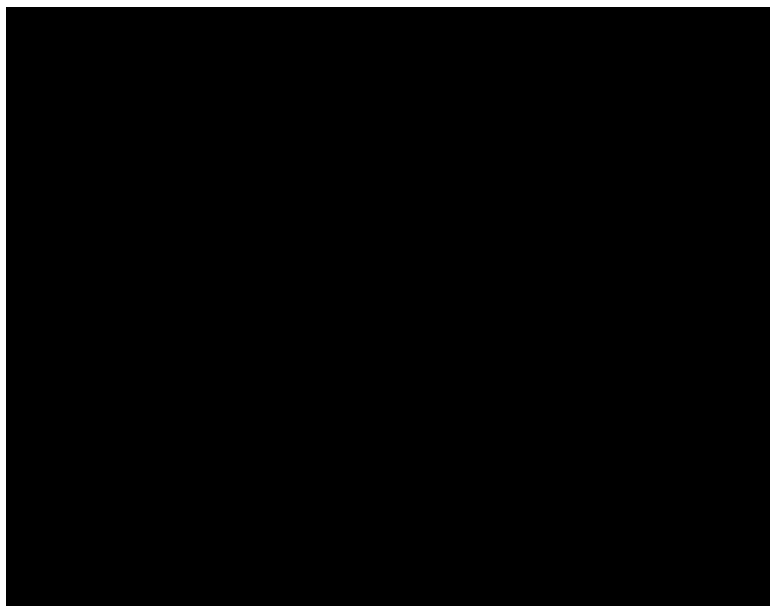
Refer to Web version on PubMed Central for supplementary material.

### Acknowledgment

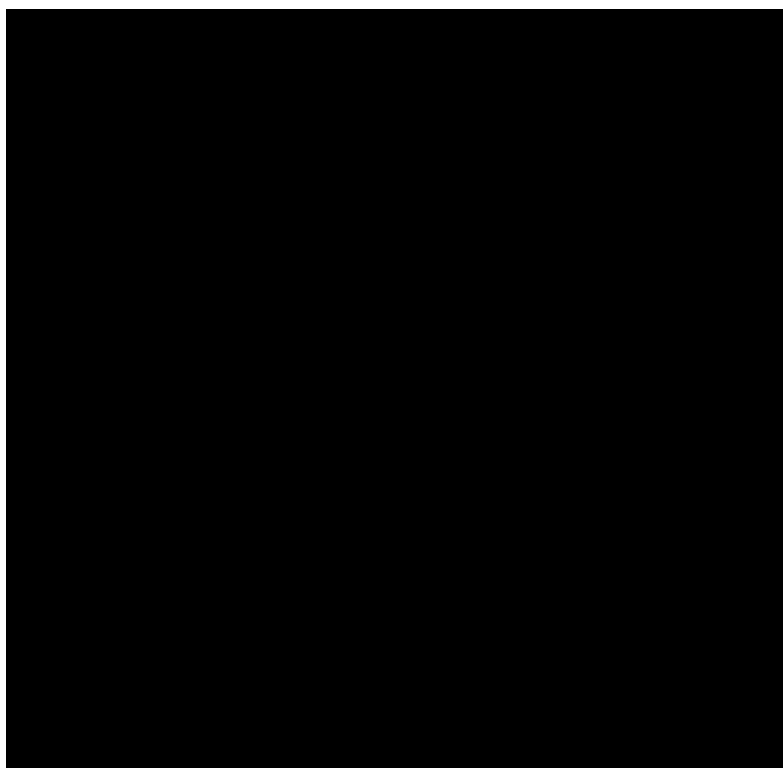
Financial assistance provided by the National Institutes of Health (Grant GM-28961) is gratefully acknowledged.

## References

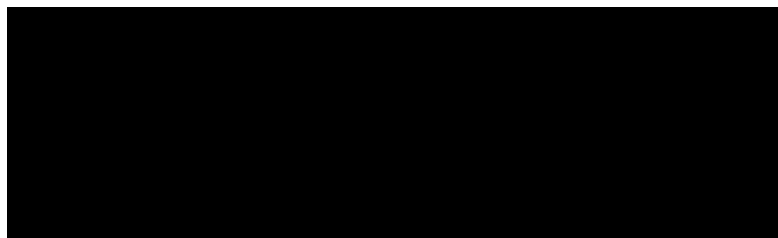
1. (a) Keck GE, Wager CA, Sell T, Wager TT. *J. Org. Chem.* 1999;64:2172. (b) Keck GE, Wager CA. *Org. Lett.* 2000;2:2307. [PubMed: 10930270]
2. Nogradi, M. VCH: Weinheim; Germany: 1987. *Stereoselective Synthesis*. Chapters 3, 5, 6
3. Barluenga J, Aguilar E, Fustero S, Olano B, Viado A. *J. Org. Chem.* 1992;57:1219. and references therein
4. Narasaka K, Ukaji Y, Yamazaki S. *Bull. Chem. Soc. Jpn.* 1986;59:525.
5. (a) Pilli RA, Russowsky D, Dias LC. *Chem. Commun.* 1987:1053. (b) Pilli RA, Russowsky D, Dias LC. *J. Chem. Soc., Perkin Trans. 1* 1990:1213.
6. Keck GE, Krishnamurthy D, Grier MC. *J. Org. Chem.* 1993;58:6543. For an asymmetric version of this reaction, see:
7. Boger DL, Yohannes D. *J. Org. Chem.* 1989;54:2498.
8. Kobayashi S, Ueno M, Suzuki R, Ishitani H, Kim H-S, Wataya Y. *J. Org. Chem.* 1999;64:6833. [PubMed: 11674693]
9. Saito SS, Hatanaka K, Yamamoto H. *Org. Lett.* 2000;2:1891. [PubMed: 10891184]
10. Kronenthal DR, Han CY, Taylor MK. *J. Org. Chem.* 1982;47:2765.
11. The stereochemistry of product **5f** has not been unambiguously established but is assumed on the basis of the results with the other *N*-aryl compounds and the similarity of the  $^1\text{H}$  and  $^{13}\text{C}$  NMR data
12. Ishitani H, Ueno M, Kobayashi S. *J. Am. Chem. Soc.* 2000;122:8180. For previous proposals of such a complexation mode in reactions involving *o*-hydroxy anilines and derivatives, see: and references therein
13. Molander GA. *Org. React.* 1994;46:211.
14. Still WC, Galynker I. *Tetrahedron* 1981;37:3981.



**Scheme 1.**  
Preparation of *N*-Acyl  $\beta$ -Amino Ketones



**Figure 1.**  
Structure of the major diastereomer of *N*-trifluoroacetamide **3a** by X-ray analysis.




**Scheme 2.**  
Preparation of *N*-Aryl  $\beta$ -Amino Ketones



**Scheme 3.**




**Table 1**  
Reduction of *N*-Acyl  $\beta$ -Amino Ketones Using SmI<sub>2</sub>-MeOH



| entry | P   | time (h) | yield of 3 (%) | ratio (syn:anti) |
|-------|-----|----------|----------------|------------------|
| 1     | TFA | 4.75     | 86             | 90:10            |
| 2     | Ac  | 6        | 96             | 89:11            |
| 3     | Boc | 4        | quant          | 81:19            |
| 4     | Cbz | 7        | 84             | 84:16            |

Table 2

Reduction of *N*-Aryl  $\beta$ -Amino Ketones

| entry | R                                       | P               | yield of 5 (%) | ratio (anti:syn) |
|-------|---|-----------------|----------------|------------------|
| 1     | <i>n</i> -C <sub>3</sub> H <sub>7</sub> | <i>o</i> -MeOPh | 95             | 97:3             |
| 2     | C <sub>6</sub> H <sub>11</sub>          | <i>o</i> -MeOPh | 100            | 97:3             |
| 3     | PhCH[unk]CH                             | <i>o</i> -MeOPh | 90             | 97:3             |
| 4     | <i>n</i> -C <sub>3</sub> H <sub>7</sub> | <i>p</i> -MeOPh | 95             | 94:6             |
| 5     | C <sub>6</sub> H <sub>11</sub>          | <i>p</i> -MeOPh | 98             | >99:1            |
| 7     | <i>n</i> -C <sub>3</sub> H <sub>7</sub> | Ph              | 96             | 85:15            |