

PREPARATION AND A NOVEL REARRANGEMENT REACTION OF 1,2,3,4-TETRAHYDRO-9-HYDROXY- β -CARBOLINE, AND THEIR APPLICATIONS FOR THE TOTAL SYNTHESIS OF ($\pm$)-COERULESCINE¹

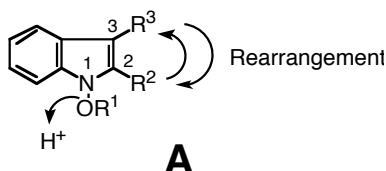
Masanori Somei,* Koichi Noguchi, Ryutaro Yamagami, Yumiko Kawada, Koji Yamada, and Fumio Yamada

Faculty of Pharmaceutical Sciences, Kanazawa University,
13-1 Takara-machi, Kanazawa 920-0934, Japan

Abstract — Novel 9-hydroxy- β -carboline derivatives were produced for the first time. A novel rearrangement reaction of 1,2,3,4-tetrahydro-9-hydroxy- β -carboline was discovered to give 3,3-disubstituted oxindoles, which was successfully applied to the total synthesis of ($\pm$)-coerulescine.

1-Hydroxyindoles undergo various types of reactions depending on the structures and reaction conditions.²

Figure 1



We have designed 1,2,3,4-tetrahydro-9-hydroxy- β -carboline (**1a-d**, Scheme 1) as key substrates to elucidate whether the rearrangement of the 2-substituent to the 3-position^{3a} or *vice versa*, predicted by our 1-hydroxyindole hypotheses,³ occurs or no as illustrated in a general formula **A** of 1-hydroxyindoles (Figure 1).

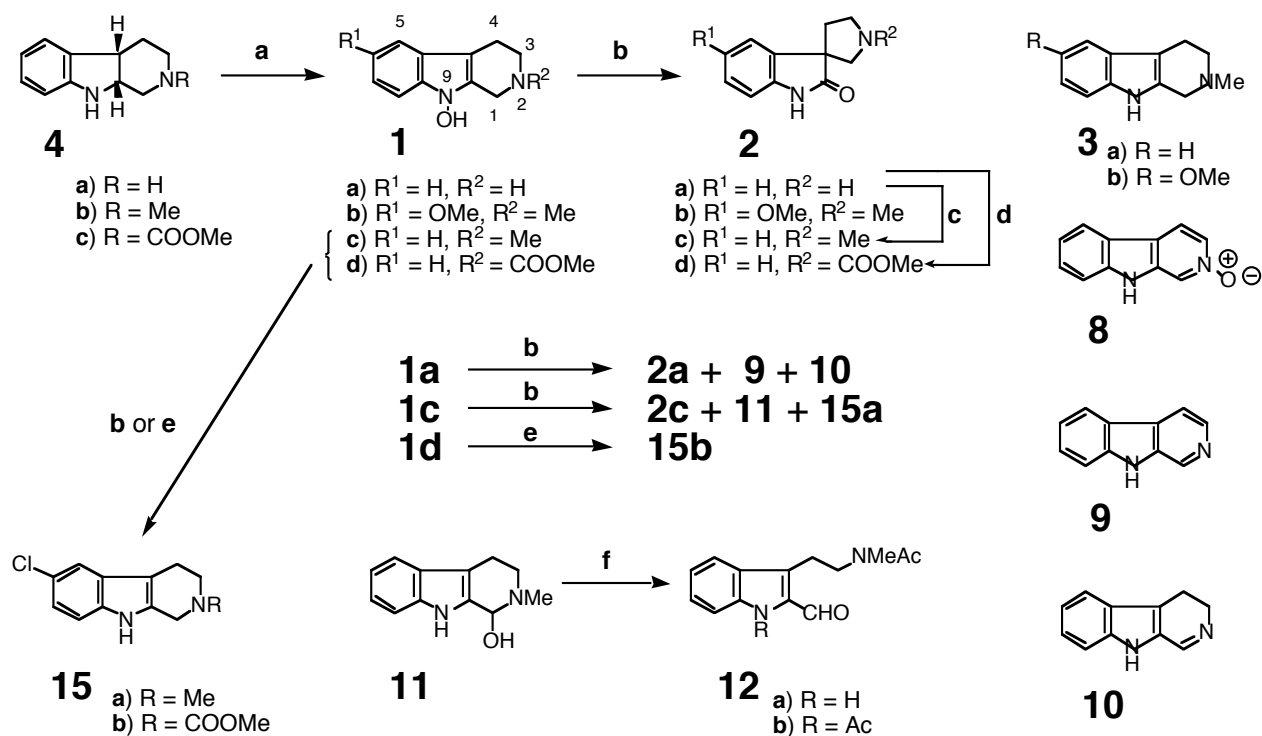
Quite recently, Colegate and co-workers⁴ isolated and determined (–)-coerulescine (**2c**) from *Phalaris coerulescens*. We have noticed that *Phalaris* species generally contain 1,2,3,4-tetrahydro- β -car-

bolines (**3a,b**), because our hypotheses assumed^{3a} biosyntheses of the related alkaloid, (–)-horsfiline⁵ (**2b**) and **2c** from **3b** and **3a**, respectively, through the corresponding 9-hydroxy- β -carboline (**1b** and **1c**).

Now, we wish to report successful preparations of **1a** and **1c**, and their novel rearrangement achieving a total synthesis of 3,3-disubstituted oxindole alkaloid, ($\pm$)-coerulescine (**2c**).

We have already developed the preparation of 1,2,3,4-tetrahydro-9-hydroxy-2-methoxycarbonyl- β -carboline⁶ (**1d**) from the corresponding 1,2,3,4,4a,9a-hexahydro compound (**4c**) by Na₂WO₄·2H₂O–30% H₂O₂ method.⁷ According to the procedure, oxidation of **4a** was examined and typical results are summarized in Table 1. As can be seen from the Table, reaction time was found to be an important factor for preparing **1a**. In cases (Entries 1 and 2) where reaction times were 25 min, even a trace amount of **1a** was not detected in the reaction mixture, instead 9-hydroxy- β -carboline (**5**), 3,4-dihydro-9-hydroxy- β -carboline *N*-oxide (**6**), and 9-hydroxy- β -carboline *N*-oxide (**7**) were produced. When the reaction time was longer than 25 min, total yield of the three 9-hydroxy- β -carboline (**5–7**) decreased, while the production of **1a** was observed in 11% yield when it was shortened to 10 min (Entry 3). Better yield of **1a** (55%) was attained by reacting **4a** for only 5 min as shown in Entry 4. As for oxidizing reagent, urea hydrogen peroxide was a reagent of choice to give satisfactory result (65%, Entry 5) as discovered in our previous report.⁸ Similarly,

Scheme 1



a) Na₂WO₄·2H₂O, 30% H₂O₂; b) MeOH-c-HCl (1:3, v/v), reflux;
c) HCHO/AcOH/NaBH₃CN; d) ClCOOMe, Et₃N; e) MeOH-c-HCl,
under various reaction conditions; f) Ac₂O, pyridine.

Table 1

Entry	Oxidizing Reagent (mol eq)	Reaction Solvent System	Conditions Temp. (°C)	Time (min)	Yield (%) of 1a	Yield (%) of 5	Yield (%) of 6	Yield (%) of 7
1	30% H ₂ O ₂ (10)	MeOH-H ₂ O (10:1, v/v)	19	25	0	14	21	7
2	"	<i>t</i> -BuOH-H ₂ O (4:1, v/v)	22	25	0	5	25	21
3	"	"	21	10	11	trace	22	15
4	"	MeOH-H ₂ O (9:1, v/v)	27	5	55	trace	5	5
5	98% Urea·H ₂ O ₂ (10)	"	25.5	5	65	trace	5	2

oxidation of 1,2,3,4,4a,9a-hexahydro-2-methyl- β -carboline (**4b**) afforded 1,2,3,4-tetrahydro-9-hydroxy-2-methyl- β -carboline (**1c**) in 69% yield.

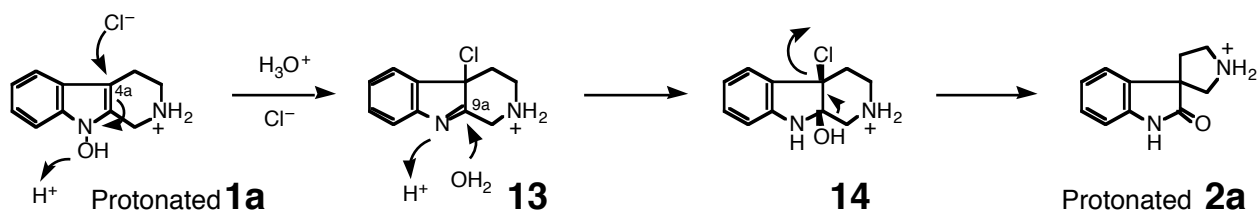
On the other hand, oxidation of **5** with MCPBA afforded **7** in 29% yield in addition to β -carboline *N*-oxide⁹ (**8**) and β -carboline (**9**) in 31 and 11% yields, respectively. Dehydrogenation of **6** with DDQ provided 75% yield of **7**. Interestingly, treatment of **6** with chloroacetyl chloride produced **8** and **9** in 60 and 33% yields, respectively. Thus, four novel 9-hydroxy- β -carboline compounds (**1a** and **5–7**) and **8** are now readily available. Since these compounds are useful building blocks for β -carboline alkaloids syntheses, their applications are currently in progress.

With **1a,c** and **1d** in hand, we treated them with MeOH-*c*-HCl (1:3,v/v). Although no reaction took place at room temperature, the desired reaction of **1a** occurred under refluxing for 1 h resulting in the formation of **2a** in 47% yield together with **9** and **10** in 2 and 36% yields, respectively. Subsequent methylation of **2a** with HCHO-AcOH-NaBH₃CN afforded ($\pm$)-**2c** in 91% yield, while the reaction of **2a** with methyl chloroformate gave **2d** in 99% yield. Similarly, treatment of **1c** with refluxing MeOH-*c*-HCl (1:3,v/v) for 3 h produced ($\pm$)-**2c**, **11**, and 6-chloro-1,2,3,4-tetrahydro-2-methyl- β -carboline (**15a**) in 42, 46, and 9% yields, respectively.

The structure of ($\pm$)-**2c** was confirmed by the direct comparison with the authentic sample prepared alternatively according to the reported procedure.^{4c} Spectral data of our samples, authentic sample, and the reported ones of (–)-**2c** were identical in every respects. The structure of relatively unstable **11** was deduced based on its spectral data and the fact that it yielded formyl compounds (**12a** and **12b**) by the reaction with Ac₂O-pyridine.

The mechanism of the rearrangement would be explained as shown in Scheme 2. Initial protonation of 9-hydroxy group, followed by its elimination and chloride attack at the 4a carbon generates chloroindolenine (**13**). After addition of water to the 9a imine carbon atom of **13** giving **14**, concerted elimination of chloride and rearrangement of the alkyl side chain attached to the 9a carbon atom result in 3,3-disubstituted oxindole structure(**2a**).

Scheme 2



It should be noted that **2d** was not formed in the reaction of **1d** with methanolic HCl under various examined reaction conditions. In these cases, nucleophilic substitution reaction took place affording 6-chloro-1,2,3,4-tetrahydro-2-methoxycarbonyl- β -carboline (**15b**, 8–16%) together with 1,2,3,4-tetrahydro-2-methoxycarbonyl- β -carboline (21–30%).¹⁰

In conclusion, we could prepare novel 9-hydroxy- β -carbolines which would be utilized as useful building blocks for the syntheses of β -carboline compounds. We have also discovered a novel rearrangement

reaction on 1-hydroxyindole skeletons providing valuable means for the preparation of 3,3-disubstituted oxindole alkaloids. The present findings suggest chemists to be very careful to use acids for isolation of indole alkaloids from natural resources, otherwise oxindole alkaloids might be obtained as artifacts of genuine 1-hydroxy- or 1-methoxyindole alkaloids, if they happen to exist.^{1b,2b,3a}

ACKNOWLEDGMENT

This work is supported in part by a Grant-in-Aid for Scientific Research from the Ministry of Education, Science, Sports, and Culture, Japan, which is gratefully acknowledged.

REFERENCES AND NOTES

1. a) This is Part 95 of a series entitled "The Chemistry of Indoles". b) Part 94: M. Hasegawa, K. Yamada, Y. Nagahama, and M. Somei, *Heterocycles*, 2000, submitted. All new compounds gave satisfactory spectral and elemental analysis or high-resolution MS data for crystals or gums, respectively. **1a**, mp 171–175°C (decomp); **1c**, mp 181–183°C (decomp); **1d**, gum; **2a**, gum; **2c**, mp 111–112°C; **5**, mp 205.0–206.5°C; **6**, mp 193.5–195.0°C (decomp); **7**, mp 254.0–265.5 (decomp); **11**, mp 179.5–181.0°C; **15a**, mp 207–209°C (decomp); **15b**, mp 194–195°C.
2. a) M. Somei, N. Oshikiri, M. Hasegawa, and F. Yamada, *Heterocycles*, 1999, **51**, 1237; b) Review; M. Somei, *ibid.*, 1999, **50**, 1157 and references cited therein.
3. a) Review; M. Somei, *J. Synth. Org. Chem. Jpn.*, 1991, **49**, 205; b) M. Somei and M. Natsume, *Tetrahedron Lett.*, 1973, 2451; c) M. Somei and Y. Fukui, *Heterocycles*, 1993, **36**, 1859. See also reference 2b.
4. a) N. Anderton, P. A. Cockrum, S. M. Colegate, J. A. Edgar, K. Flower, I. Vit, and R. I. Willing, *Phytochemistry*, 1998, **48**, 437. b) S-I. Bascop, J. Sapi, J-Y. Laronze, and J. Levy, *Heterocycles*, 1994, **38**, 725. c) M. E. Kuehne, D. M. Roland, and R. Hafter, *J. Org. Chem.*, 1978, **43**, 3705.
5. C. Pellegrini, C. Strassler, M. Weber, and H-J. Borschberg, *Tetrahedron: Asymmetry*, 1994, **5**, 1979; G. Lakshmajah, T. Kawabata, M. Shang, and K. Fuji, *J. Org. Chem.*, 1999, **64**, 1699. See also reference 4b.
6. M. Somei, K. Kobayashi, K. Tanii, T. Mochizuki, Y. Kawada, and Y. Fukui, *Heterocycles*, 1995, **40**, 119.
7. M. Somei and T. Kawasaki, *Heterocycles*, 1989, **29**, 1251; M. Somei, T. Kawasaki, K. Shimizu, Y. Fukui, and T. Ohta, *Chem. Pharm. Bull.*, 1991, **39**, 1905. See also references 2b and 8. Our synthetic method for 1-hydroxy- and 1-alkoxytryptophans has been utilized by chemists without citing us fairly. One recent example is the following. D. L. Boger, H. Keim, B. Oberhauser, E. P. Schreiner, and C. A. Foster, *J. Am. Chem. Soc.*, 1999, **121**, 6197.
8. M. Somei and T. Kobayashi, *Heterocycles*, 1992, **34**, 1295.
9. Synthesis of **8** from **9** in 78% yield: M. Natsume and R. Tanabe, *Ann. Rept. ITSUU Lab.*, 1968, **15**, 21 and references cited therein.
10. Remarkable differences observed in the reactivities of **1a,c** and **1d** might be explained by the protonation of β -nitrogen. The detailed elucidation of the results is an interesting future subject.