

Total syntheses of indole alkaloids, annonidine A and 7-(3-methyle-2-buten-1-yl)indole

著者	Somei Masanori, Funamoto Tetsuo, Ohta Toshiharu
journal or publication title	Heterocycles
volume	26
number	7
page range	1783-1784
year	1987-01-01
URL	http://hdl.handle.net/2297/4311

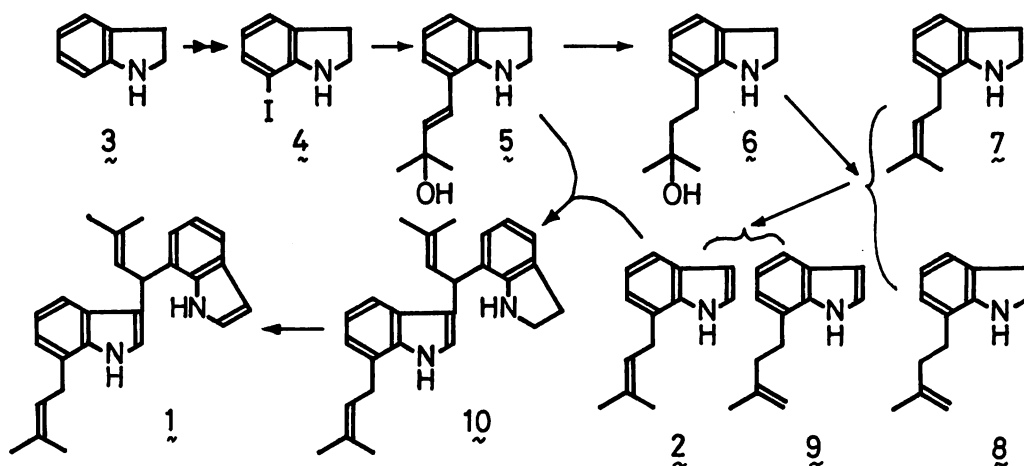
TOTAL SYNTHESSES OF INDOLE ALKALOIDS, ANNONIDINE A
AND 7-(3-METHYL-2-BUTEN-1-YL)INDOLE¹

Masanori Somei,* Tetsuo Funamoto, and Toshiharu Ohta
Faculty of Pharmaceutical Sciences, Kanazawa University,
13-1 Takara-machi, Kanazawa 920, Japan

Abstract ——— Practical total syntheses of annonidine A and
7-(3-methyl-2-buten-1-yl)indole are reported.

Achenbach and co-workers² isolated annonidine A (1) and 7-(3-methyl-2-buten-1-yl)-indole^{3a} (2) from the stem bark of the west african medicinal plant, Annonidium Mannii Engl. & Diels (Annonaceae) in 1985. We are much interested in annonidine A because of its hitherto unknown prenylated bisindole structure. In this report, we describe the total syntheses of annonidine A and 7-(3-methyl-2-buten-1-yl)indole.^{3b} We have already reported a convenient and practical synthetic method for 2,3-dihydro-7-iodoindole (4) from 2,3-dihydroindole (3) in three steps in 62% overall yield.⁴ Heck reaction of 4 with 2-methyl-3-buten-2-ol in the presence of a catalytic amount of palladium acetate in N,N-dimethylformamide afforded 4-(2,3-dihydroindol-7-yl)-2-methyl-3-buten-2-ol (5, mp 85.0-86.5°C) in 74% yield.^{4b} Catalytic hydrogenation of 5 over 10% palladium/carbon at an atmospheric pressure produced 4-(2,3-dihydroindol-7-yl)-2-methyl-3-butanol (6, mp 88-90°C) in 87% yield. Subsequent treatment of 6 with p-toluenesulfonic acid in refluxing benzene afforded 95% yield of an inseparable mixture of 2,3-dihydro-7-(3-methyl-2-buten-1-yl)indole (7) and its double bond isomer (8) in the ratio of 13:1. Oxidation of the mixture with dioxygen in the presence of a catalytic amount of salcomine⁵ in methanol at room temperature for 2 h afforded 7-(3-methyl-2-buten-1-yl)indole [2, mp 43.5-44.0°C (lit.² mp 43-44°C)] and its double bond isomer (9, mp 18-19°C) in 75% and 3% yields, respectively. Condensation of 2 and 5 in tetrahydrofuran by the action of 2N-hydrochloric acid produced regiospecifically 3-[1-(2,3-dihydroindol-7-yl)-3-methyl-2-buten-1-yl]-7-(3-methyl-2-buten-1-yl)indole (10, viscose oil) in 78% yield. Subsequent salcomine⁵ catalyzed oxidation with dioxygen afforded 3-[1-(indol-7-yl)-3-methyl-2-buten-1-yl]-

7-(3-methyl-2-buten-1-yl)indole [1, mp 105-108°C (lit.² mp 106-108°C)] in 65% yield. Spectral data of 1 and 2 were identical with those of the natural alkaloids. Since the present method is simple and practical, synthesis of various derivatives and evaluations of their pharmacological activities are currently in progress.



ACKNOWLEDGEMENT

The authors express their cordial gratitude to Prof. Achenbach for kindly providing us with spectral data of natural alkaloids, annonidine A and 7-(3-methyl-2-buten-1-yl)indole.

REFERENCES AND NOTES

1. This report is part XLI of a series entitled "The Chemistry of Indoles." Part XL: M. Somei, F. Yamada, and K. Naka, Chem. Pharm. Bull., **35**, 1322 (1987).
2. H. Achenbach and C. Renner, Heterocycles, **23**, 2075 (1985); H. Achenbach and D. Franke, Arch. Pharm. (Weinheim), **320**, 91 (1987).
3. a) V. Benesova, Z. Samek, V. Herout, and F. Sorm, Collect. Czech. Chem. Commun., **34**, 1807 (1969). b) The first synthesis of 7-(3-methyl-2-buten-1-yl)indole (2) was reported by Natsume and co-workers: H. Muratake and M. Natsume, Heterocycles, **24**, 261 (1986). Other related 7-substituted indole syntheses: S. Nakatsuka, T. Masuda, and T. Goto, Tetrahedron Lett., **27**, 6245 (1986); M. Akagi and K. Ozaki, Heterocycles, **26**, 61 (1985); T. Martin and C.J. Moody, J. Chem. Soc., Chem. Commun., **1985**, 1391; A.P. Kozikowski and K. Isobe, ibid., **1978**, 1076; S. Inoue, N. Takamatsu, and Y. Kishi, Yakugaku Zasshi, **97**, 558 (1977) and see also references cited in the above literatures.
4. a) M. Somei and Y. Saida, Heterocycles, **23**, 3113 (1985). b) M. Somei, Y. Saida, T. Funamoto, and T. Ohta, Chem. Pharm. Bull., **35**, No. 8 (1987), in press.
5. A. Inada, Y. Nakamura, and Y. Morita, Chem. Lett., **1980**, 1287.

Received, 3rd April, 1987