

HETEROCYCLES, Vol. 77, No. 2, 2009, pp. 1163 – 1170. © The Japan Institute of Heterocyclic Chemistry
Received, 4th August, 2008, Accepted, 18th September, 2008, Published online, 22nd September, 2008
DOI: 10.3987/COM-08-S(F)95

DEHALOGENATION AND BARBIER-TYPE HYDROXYALKYLATION OF π -DEFICIENT HALOHETEROCYCLES USING INDIUM

Eri Fukuda, Yukiko Takahashi, Natsumi Hirasawa, Osamu Sugimoto,* and
Ken-ichi Tanji*

Laboratory of Organic Chemistry, School of Food and Nutritional Sciences,
University of Shizuoka, 52-1 Yada, Suruga-ku, Shizuoka 422-8526, Japan
osamu@smail.u-shizuoka-ken.ac.jp; tanji@smail.u-shizuoka-ken.ac.jp

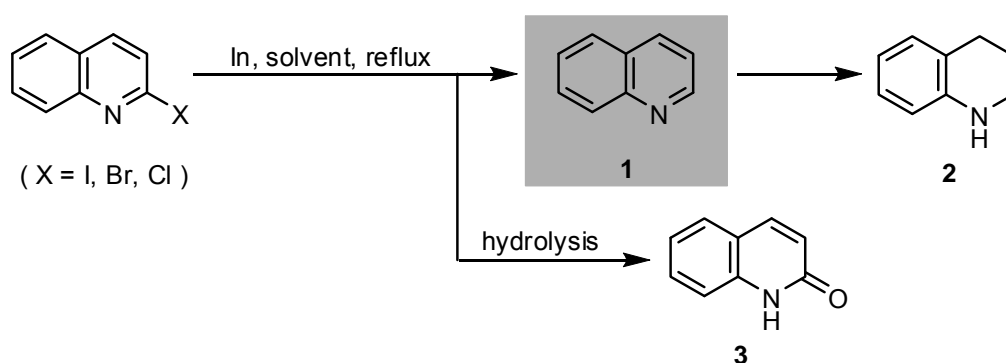
Abstract – The reaction of π -deficient haloheterocycles with indium metal in water gave corresponding dehalogenated heterocycles. The use of diluted hydrochloric acid instead of water accelerated the reductive reactivity of indium metal. Furthermore, Barbier-type additions proceeded by reactions of α -iodoheterocycles with indium in the presence of pivalaldehyde.

Indium (In, atomic number 49) is one of various rare metals discovered by Reich and Richter in 1863. Recently, indium has been widely used in organic syntheses. For example, organic halides react with acetylenes,¹ or carbonyl compounds,² in the presence of indium to create corresponding carbon-carbon bonded products. Iodoalkanes or iodobenzenes are converted into bialkyls or biphenyls by reactions with indium in dry DMF.^{3,4} It has been reported that nitrogen-containing heterocycles, *e. g.* quinolines, isoquinolines, quinoxalines, and indoles are reduced by indium in aqueous ethanolic ammonium chloride in order to create dihydro- or tetrahydroheterocycles.⁵ To our knowledge, however, the study of haloheterocycles as a substrate which uses indium has not been mentioned except in a sole report of the conversion of 2-chloroquinoline into 1,2,3,4-tetrahydroquinoline using indium.⁵ Here we report the deiodination of nitrogen-containing π -deficient haloheterocycles.⁶

Pitts *et al.* have reported that the reaction of 2-chloroquinoline with excess indium in aqueous ethanolic ammonium chloride under refluxing for several days creates 1,2,3,4-tetrahydroquinoline with a 79% yield.⁵ In this reaction, it has been speculated that the substrate, 2-chloroquinoline is dechlorinated to afford quinoline, which is further reduced to give 1,2,3,4-tetrahydroquinoline because quinoline is converted into 1,2,3,4-tetrahydroquinoline under the same condition. So we planned a selective dehalogenation of haloheterocycles using indium metal.

This paper is dedicated to Professor Emeritus Keiichiro Fukumoto on the occasion of his 75th birthday.

Table 1. The reaction of 2-haloquinolines with indium under several conditions



Entry	Solvent	X	Indium (eq)	Time (h)	Yields (%)			Recovery
					1	2	3	
1	aq. NH ₄ Cl	I	10	48	8	73		
2	aq. NH ₄ Cl	I	1	24	84			
3	aq. NH ₄ Cl	I	1	2.5	64			
4	H ₂ O	I	10	2	70			Trace
5	H ₂ O	I	10	48	84	Trace		
6	H ₂ O	I	1	1	57			33
7	H ₂ O	I	1	2	91			7
8	H ₂ O	I	1	3	83			6
9	H ₂ O	I	1	5	91			2
10	H ₂ O	I	1	11.5	92			4
11	H ₂ O	I	2	5	92			4
12	H ₂ O	I	2	10	83			3
13	H ₂ O	I	3	10	81			6
14	H ₂ O	Br	1	2	68			15
15	H ₂ O	Br	1	5	88			3
16	H ₂ O	Cl	1	2	51			39
17	H ₂ O	Cl	1	5	60			15
18	0.5 N HCl	Cl	1	5	69		21	
19	0.25 N HCl	Cl	2	2	66	3	3	12
20	0.5 N HCl	Cl	2	5	75	4		
21	0.5 N HCl	Cl	2	2	75	1	10	
22	1 N HCl	Cl	2	2	67		17	Trace
23	0.5 N HCl	Cl	3	2.5	66	10	5	

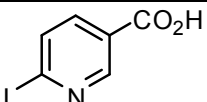
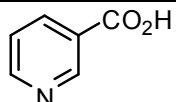
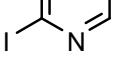

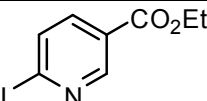
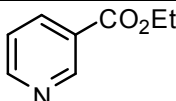
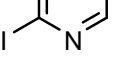

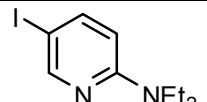
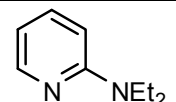
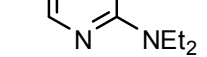
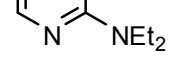
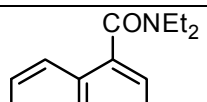
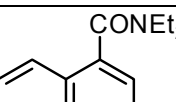
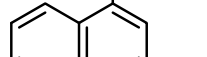
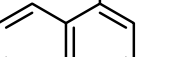
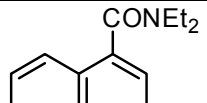
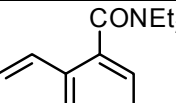
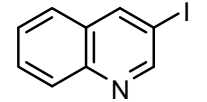
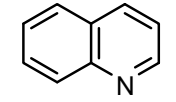
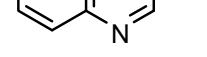
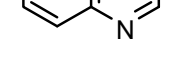
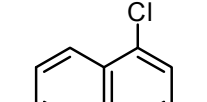
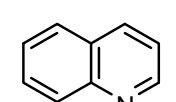
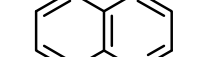
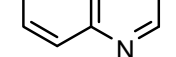
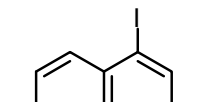
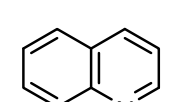
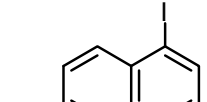
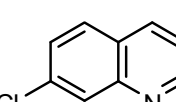
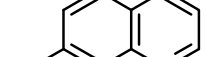
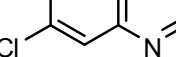
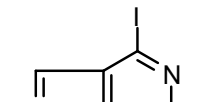
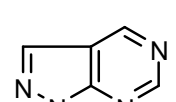
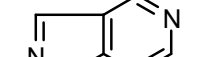
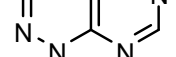
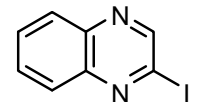
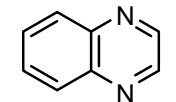
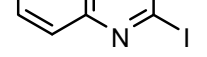
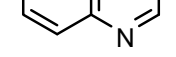
Initially, reactions of 2-haloquinolines with indium under several conditions were performed as shown in Table 1. A mixture of 2-iodoquinoline and indium powder (10 eq) in aqueous ammonium chloride was heated to reflux for 48 h and produced 1,2,3,4-tetrahydroquinoline (**2**) with a 73% yield and quinoline (**1**) with an 8% yield (entry 1). These results are identical to the results reported by Pitts *et al.* Conversely and

interestingly, a selective deiodination proceeded to create **1** under mild conditions such as the use of a stoichiometric amount of indium (entries 2,3), or the use of water as a solvent instead of aqueous ammonium chloride (entries 4,5). Next, the deiodination of 2-iodoquinoline in water was performed with various amounts of indium and the reaction times. It was discovered that 2-iodoquinoline was successfully deiodinated to create **1** with the use of a stoichiometric amount of indium in water for 2 h (entry 7). Dehalogenation of 2-bromoquinoline and 2-chloroquinoline was performed (entries 14,15) in order to discover the different reactivities with 2-iodoquinoline. Although both 2-bromoquinoline and 2-chloroquinoline were dehalogenated to create **1**, it was made clear that 2-bromoquinoline and 2-chloroquinoline are less reactive compared to 2-iodoquinoline, and more drastic conditions would be required to dehalogenate successfully. Subsequently, the dechlorination of 2-chloroquinoline using indium in diluted hydrochloric acid was performed (entries 16-21) because it is known that the reactivity of indium is accelerated in acidic solvent. As shown in entries 18 and 19, 2-chloroquinoline was smoothly dechlorinated with 2 eq of indium in 0.5 *N* hydriodic acid and refluxing for 2-5 h to produce a 75% yield of quinoline.

Table 2 shows the dehalogenation of some haloheterocycles using indium. Water or 0.5 *N* hydrochloric acid was used as solvents to compare the effects on dehalogenation reactivity. In many cases, the use of 0.5 *N* hydrochloric acid as a solvent accelerated the dehalogenation of halopyridines and haloquinolines. Especially, chloroquinolines such as *N,N*-diethyl 2-chloroquinoline-4-carboxamide (entries 7,8) and 4-chloroquinoline (entries 12,13), were dechlorinated smoothly, using 0.5 *N* hydrochloric acid to produce corresponding dechlorinated products. It should be noted that a reaction of 7-chloro-4-iodoquinoline with indium preceded deiodination at the 4-position of the quinoline ring selectively and dechlorination at the 7-position did not proceed (entries 15,16). This result indicates that ring nitrogen of the quinoline ring affects the reactivity of dehalogenation using indium. Deiodination of iododiazines, 4-iodo-1-phenyl-1*H*-pyrazolo[3,4-*d*]pyrimidine (entries 17,18) and 2-iodoquinoxaline (entries 19,20), in 0.5 *N* hydrochloric acid did not produce any corresponding deiodinated compounds whereas deiodination in water created products in 30% and 44%, respectively. It was found that aqueous hydrochloric acid is not suitable for indium mediated deiodination of iododiazines because diazines tend to dimerize under acidic conditions.

Plausible mechanisms for the dehalogenation of haloheterocycles using indium were shown in Scheme 1. Pitts *et al.* have reported that quinoline derivatives are reduced to give 1,2,3,4-tetrahydroquinolines under drastic conditions, using excess indium in aqueous ammonium chloride for several days.

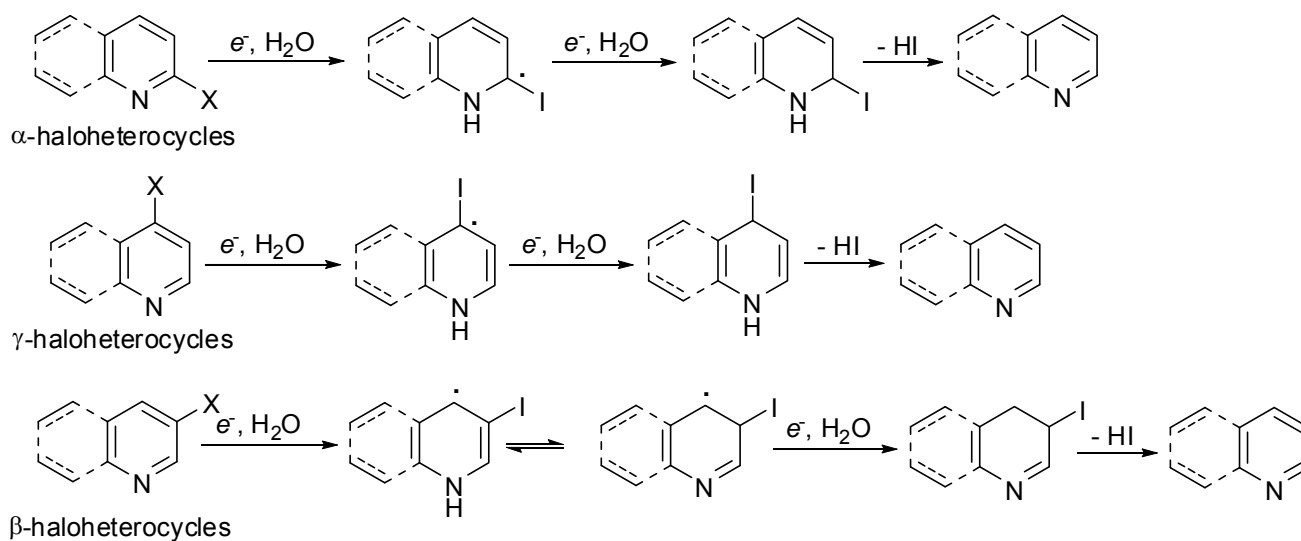
Table 2. Dehalogenation of various haloheterocycles using indium metal

Entry	Substrate (Het-X)	Indium (eq)	Solvent	Time (h)	Product (Het-H)	Yields (%)	
						Het-H	Recovery
1		1	H ₂ O	5		60	
2		2	0.5 N HCl	2		80	
3		1	H ₂ O	10		64	
4		2	0.5 N HCl	2		8 ¹	
5		1	H ₂ O	10		49	20
6		2	0.5 N HCl	10		53	24
7		1	H ₂ O	2		24	55
8		2	0.5 N HCl	2		55	
9		1	H ₂ O	2		91	
10		1	H ₂ O	8		52	36
11		2	0.5 N HCl	5.5		80	
12		1	H ₂ O	5		60	19
13		2	0.5 N HCl	2		86	
14		1	H ₂ O	5		69	
15		1	H ₂ O	15		59	16
16		2	0.5 N HCl	2		55	
17		1	H ₂ O	2		30	9
18		2	0.5 N HCl	2		0	
19		1	H ₂ O	2		56	14
20		2	0.5 N HCl	2		0	

1. Nicotinic acid was obtained

There was also speculation that a single electron transfer would proceed from indium to the carbon-nitrogen bond of the quinoline ring. Thus we have speculated that the dehalogenation would proceed through a similar mechanism, that is, a single electron transfer from indium to the carbon-nitrogen bond.

Scheme 1. Plausible mechanisms for dehalogenation of haloheterocycles



Since π -deficient heterocycles are less reactive with electrophiles, lithiation⁷⁻⁹ are frequently used to introduce electrophiles into π -deficient heterocycles. However, metalation of π -deficient heterocycles usually requires low temperature to avoid some side reactions. Through the examination of this study, we planned the novel introduction of electrophiles into π -deficient heterocycles *via* a reaction of iodoheterocycles with carbonyl compounds in the presence of indium without protonic solvents such as water. Table 3 shows the reaction of iodopyridines and iodoquinolines with indium in the presence of pivalaldehyde. The reaction of 2-iodoquinoline with indium in the presence of pivalaldehyde under refluxing created the corresponding adduct, 2,2-dimethyl-1-(2-quinolinyl)-1-propanol in moderate yields as well as the deiodinated product, quinoline (entries 1-5). 2-Iodopyridine was also converted to 2,2-dimethyl-1-(2-pyridinyl)-1-propanol using a similar method (entries 6-12). Interestingly, a Barbier-type hydroxyalkylation was facilitated using α -iodoheterocycles (entries 13,14). It is evident that more studies using various kinds of iodoheterocycles are necessary in order to clarify the potentiality and limitation of this reaction.

In conclusion, we have found a novel method of the deiodination of π -deficient haloheterocycles using indium in water to create corresponding dehalogenated heterocycles. The use of diluted hydrochloric acid instead of water accelerated the reductive reactivity of indium. Barbier-type coupling reactions were preceded by the reaction of α -iodoheterocycles with indium in the presence of pivalaldehyde.

Table 3. The reaction of iodopyridines and iodoquinolines with indium in the presence of pivalaldehyde

Substrate $\xrightarrow[75\text{ }^\circ\text{C (solvent-free)}]{\text{In, } t\text{BuCHO (excess)}}$ Product $\left[+ \text{Quinoline} \right]$

Entry	Substrate	In (eq)	Time (h)	Product	Yields (%)		
					Product	Quinoline	Recovery
1		1	18		47	42	
2		4	3		33	23	30
3		4	6		59	37	
4		4	24		51	36	
5		4	30		46	43	
6		2	4.5		7		45
7		2	8		23		14
8		3	8		38		18
9		3	24		41		Trace
10		3	66.5		27		Trace
11		4	8		29		33
12		4	24		56		
13		1	18		2		9
14		1	18		0	76	

EXPERIMENTAL

All melting points were not corrected. $^1\text{H-NMR}$ spectra was measured with Hitachi R-90H spectrometer (90 MHz) using tetramethylsilane as an internal standard.

The dehalogenation of haloheterocycles using In in aqueous media (General procedure 1) : A mixture of In, haloheterocycles, and solvent was heated to reflux. The reaction mixture was alkalinized with 1 *N* NaOH and extracted with EtOAc. The organic layer was dried over Na_2SO_4 and treated with silica gel column chromatography to give the corresponding product.

1,2,3,4-Tetrahydroquinoline : Eluted with hexane - EtOAc (2:1). Yellow liquids. $^1\text{H-NMR}$ (CDCl_3), δ : 1.72-2.13 (2H, *m*, $\text{C}^3\text{-H}$), 2.60-2.91 (2H, *m*, $\text{C}^4\text{-H}$), 3.15-3.47 (2H, *m*, $\text{C}^2\text{-H}$), 3.70 (1H, *brs*, NH),

6.32-6.72 (2H, *m*, C⁶ and C⁸-H), 6.78-7.11 (2H, *m*, C⁵ and C⁷-H).

Quinoline : Eluted with hexane - EtOAc (2:1). Yellow oil. ¹H-NMR (CDCl₃), δ : 7.38 (1H, *dd*, J = 8.3 Hz, 4.2 Hz, C³-H), 7.45-7.90 (3H, *m*, C⁵, C⁶, and C⁷-H), 8.13 (2H, *d*, J = 8.3 Hz, C⁴ and C⁸-H), 8.90 (1H, *dd*, J = 4.2 Hz, 1.4 Hz, C²-H).

2(1*H*)-Quinolinone : Eluted with EtOAc-EtOH (20:1). White solids. Mp 199.5-200.5 °C (lit.,¹⁰ 196-197 °C). ¹H-NMR (CDCl₃), δ : 6.72 (1H, *d*, J = 9.5 Hz, C³-H), 7.02-7.32 (1H, *m*, C⁶-H), 7.32-7.68 (3H, *m*, C⁵, C⁷, and C⁸-H), 7.81 (1H, *d*, J = 9.5 Hz, C⁴-H), 12.50 (1H, *brs*, NH).

Nicotinic acid : Eluted with EtOAc-EtOH (4:1). Yellow solids. Mp 237.7-239.1 °C (lit.,¹¹ 236-239 °C). ¹H-NMR (DMSO-*d*₆), δ : 7.50 (1H, *dd*, J = 7.6 Hz, 4.7 Hz, C⁵-H), 8.25 (1H, *d*, J = 7.6 Hz, C⁴-H), 8.74 (1H, *d*, J = 4.7 Hz, C⁶-H), 9.08 (1H, *s*, C²-H).

Ethyl nicotinate : Eluted with hexane - EtOAc (1:1). Pale yellow liquids. ¹H-NMR (CDCl₃), δ : 1.41 (3H, *t*, J = 7.1 Hz, CH₃), 4.42 (2H, *q*, J = 7.1 Hz, CH₂), 7.38 (1H, *dd*, J = 7.9 Hz, 4.8 Hz, C⁵-H), 8.29 (1H, *dt*, J = 7.9 Hz, 1.7 Hz, C⁴-H), 8.76 (1H, *dd*, J = 4.8 Hz, 1.7 Hz, C⁶-H), 9.22 (1H, *d*, J = 1.7 Hz, C²-H).

2-Diethylaminopyridine : Eluted with hexane - EtOAc (15:1). Slight violet liquids. ¹H-NMR (CDCl₃), δ : 1.18 (6H, *t*, J=7.0 Hz, CH₃ x 2), 3.51 (4H, *q*, J=7.0 Hz, CH₂ x 2), 6.32-6.57 (2H, *m*, C³ and C⁵-H), 7.38 (1H, *td*, J=7.8 Hz, 1.9 Hz, C⁴-H), 8.13 (1H, *dd*, J=5.5 Hz, 1.9 Hz, C⁶-H).

N,N-Diethyl-4-quinolinecarboxamide : Eluted with EtOAc. White oil. ¹H-NMR (CDCl₃), δ : 1.02 (3H, *t*, J = 7.1 Hz, CH₃), 1.38 (3H, *t*, J = 7.1 Hz, CH₃), 3.09 (2H, *q*, J = 7.1 Hz, CH₂), 3.43-3.95 (2H, *m*, CH₂), 7.29 (1H, *d*, J = 4.4 Hz, C³-H), 7.41-7.90 (3H, *m*, C⁵, C⁶, and C⁷-H), 8.14 (1H, *dd*, J = 8.4 Hz, 1.5 Hz, C⁸-H), 8.94 (1H, *d*, J = 4.4 Hz, C²-H).

7-Chloroquinoline : Eluted with hexane - EtOAc (2:1). Yellow liquids. ¹H-NMR (CDCl₃), δ : 7.25-7.60 (2H, *m*, C³ and C⁶-H), 7.74 (1H, *d*, J = 8.8 Hz, C⁵-H), 7.95-8.24 (2H, *m*, C⁴ and C⁸-H), 8.90 (1H, *dd*, J = 4.2 Hz, 1.6 Hz, C²-H).

1-Phenyl-1*H*-pyrazolo[3,4-*d*]pyrimidine : Eluted with hexane - EtOAc (2:1). Pale yellow solids. Mp 75-79 °C (lit.,¹² 80 °C). ¹H-NMR (CDCl₃), δ : 7.25-7.70 (3H, *m*, phenyl-H), 8.08-8.38 (2H, *m*, phenyl-H), 8.32 (1H, *s*, C³-H), 9.12 (1H, *s*, C⁶-H), 9.26 (1H, *s*, C⁴-H).

Quinoxaline : Eluted with hexane - EtOAc (2:1). Brown needles. Mp 28.5 °C (lit.,¹³ 29-30 °C). ¹H-NMR (CDCl₃), δ : 7.66-7.91 (2H, *m*, C⁵ and C⁸-H), 7.91-8.22 (2H, *m*, C⁶ and C⁷-H), 8.85 (2H, *s*, C² and C³-H).

The reaction of iodoheterocycles with In in the presence of pivalaldehyde (General procedure 2) : A mixture of iodoheterocycles, In, and excess pivalaldehyde was heated to reflux. The reaction mixture was diluted with EtOAc, and insoluble solids were filtered off. After the filtrate was diluted with water, the mixture was neutralized with 1 *N* NaOH and extracted with EtOAc. The organic layer was dried over Na₂SO₄ and treated with silica gel column chromatography to give the corresponding product.

2,2-Dimethyl-1-(2-quinoliny)-1-propanol : Eluted with hexane - EtOAc (5:1). Pale yellow solids. Mp 70-72 °C (lit.,¹² 73-75 °C). ¹H-NMR (CDCl₃), δ : 0.98 (9H, s, ^tBu), 4.50 (1H, brs, CH), 4.52-4.85 (1H, br, OH), 7.32 (1H, d, J = 8.6 Hz, C³-H), 7.38-7.66 (1H, m, C⁶-H), 7.66-7.90 (2H, m, C⁵ and C⁷-H), 8.08 (2H, d, J = 8.6 Hz, C⁴ and C⁸-H).

2,2-Dimethyl-1-(2-pyridinyl)-1-propanol : Eluted with hexane - EtOAc (1:1). Pale yellow solids. Mp 53-57 °C. ¹H-NMR (CDCl₃), δ : 0.92 (9H, s, ^tBu), 4.32 (2H, brs, CHOH), 7.06-7.35 (2H, m, C³ and C⁵-H), 7.61 (1H, ddd, J = 7.5 Hz, 7.5 Hz, 1.8 Hz, C⁴-H), 8.57 (1H, dd, J = 5.0 Hz, 1.8 Hz, C⁶-H).

3-Pivaloylpyridine : Eluted with hexane - EtOAc (5:1). Pale yellow liquids. ¹H-NMR (CDCl₃), δ : 1.36 (9H, s, ^tBu), 7.35 (1H, ddd, J = 8.1 Hz, 5.2 Hz, 0.5 Hz, C⁵-H), 8.00 (1H, ddd, J = 7.9 Hz, 2.0 Hz, 2.0 Hz, C⁴-H), 8.69 (1H, dd, J = 5.8 Hz, 1.5 Hz, C⁶-H), 8.97 (1H, d, J = 1.8 Hz, C²-H). MS 163 (M⁺).

2,2-Dimethyl-1-(3-pyridinyl)-1-propanol : Eluted with EtOAc. Pale yellow liquids. ¹H-NMR (CDCl₃), δ : 0.93 (9H, s, ^tBu), 4.42 (1H, s, CH), 7.24 (1H, dd, J = 7.8 Hz, 4.7 Hz, C⁵-H), 7.69 (1H, d, J = 10.0 Hz, C⁴-H), 8.38-8.63 (2H, m, C² and C⁶-H). MS 165 (M⁺).

REFERENCES

1. R. Yanada, S. Obika, Y. Kobayashi, T. Inokuma, M. Oyama, K. Yanada, and Y. Takemoto, *Adv. Synth. Catal.*, 2005, **347**, 1632.
2. C. J. Li and T. H. Chan, *Tetrahedron Lett.*, 1991, **32**, 7017.
3. B. C. Ranu, P. Dutta, and A. Sarkar, *Tetrahedron Lett.*, 1998, **39**, 9557.
4. Y. M. Chang, S. H. Lee, M. Y. Cho, B. W. Yoo, H. J. Rhee, S. H. Lee, and C. M. Yoon, *Synth. Commun.*, 2005, **35**, 1851.
5. M. R. Pitts, J. R. Harrison, and C. J. Moody, *J. Chem. Soc., Perkin Trans. 1*, 2001, 955.
6. We have reported the dehalogenation of haloheterocycles using indium in water as a preliminary report, see : N. Hirasawa, Y. Takahashi, E. Fukuda, O. Sugimoto, and K. Tanji, *Tetrahedron Lett.*, 2008, **49**, 1492.
7. I. Gomez, E. Alonso, D. J. Ramon, and M. Yus, *Tetrahedron*, 2000, **56**, 4043.
8. Y. Kondo, M. Shilai, M. Uchiyama, and T. Sakamoto, *J. Chem. Soc., Perkin Trans. 1*, 1996, 1781.
9. H. Gilman and S. M. Spats, *J. Am. Chem. Soc.*, 1940, **62**, 446.
10. P. G. Gassman and H. R. Drewes, *J. Am. Chem. Soc.*, 1978, **100**, 7600.
11. H. Nagano, Y. Nawata, and M. Hamana, *Chem. Pharm. Bull.*, 1987, **35**, 4068.
12. O. Sugimoto, M. Sudo, and K. Tanji, *Tetrahedron*, 2001, **57**, 2133.
13. H. S. Broadbent, E. L. Allred, L. Pendleton, and C. W. Whittle, *J. Am. Chem. Soc.*, 1960, **82**, 189.